The Vaccine Adverse Event Reporting System (VAERS) Results

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1108465-1	Heart was in A-fib, blood clot formed and had a Left Posterior Parietal Stroke
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1117313-1	Blood clot perpendicular to injection site
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1120494-1	Pulmonary embolism with acute Cor Pulmonale/hypoxia I had shortness of breath, dizziness, coughing. I had Covid test on 3/12/21 that was negative. Went to Urgent Care on 3/12/21. Gave me an albuterol inhaler which was not effective in relieving symptoms. I wasn't improving, so went to emergency room at the hospital on 03/18/2021. I was admitted to hospital and placed on oxygen and kept overnight. I was also given Xarelto. I am at home and on Xarelto.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1121937-1	Bloody nose on March 20th - threw up blood clot. Bloody nose again on March 21st, same clot formed. lasted about 15 minutes each time. Seeing my PC doctor today March 22nd.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1131671-1	Following the vaccine injection which was provided by the personnel, I have developed minor fainting spells and angina for the past 10 plus days. A large dried blood clot was noted on the first day of my period which occurred on March 16th. I continue to pass large blood clots while I was having my period. I also want to point out that a personnel who gave me an injection was incorrectly injected the vaccine outside of the area of the preferred injection zone. I have reported this event to the supervisor who was on site. I have not yet contacted my health care provider at this time but plan to contact her tomorrow as these symptoms have been persisting.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1132787-1	Within a week of getting the vaccine I developed a blood clot in the lining of my stomach
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1139939-1	PT CALLED ON 3-27 TO REPORT A BLOOD CLOT IN HER EYE AFTER RECEIVING THE VACCINE ON 3-13, STILL PRESENT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1146214-1	Was 5 weeks pregnant at time of vaccine with strong HCG levels, approximately 2 weeks after vaccination patient suffered a miscarriage. Vaginal bleeding, cramping and passed large clots including a sac like tissue.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1157533-1	DIZZINESS; FELL; HURT HIMSELF; SOME DAMAGE TO HIS KIDNEYS IN THE PROCESS (BLOOD CLOT); SOME DAMAGE TO HIS KIDNEYS IN THE PROCESS (BLOOD CLOT); CONFIRMED COVID-19 POSITIVE; This spontaneous report received from a patient via a company representative concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on 11-MAR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On 15-MAR-2021, the patient became ill with COVID like symptoms, and got tested for COVID on 22-Mar-2021 and received results that he was COVID positive on 25-Mar-2021 and was running high temperatures (fevers over 103), On 27-MAR-2021, the patient became dizzy, fell and hurt himself, was rushed to the hospital and was admitted same day. He did some damage to his kidneys in the process. On 25-MAR-2021, Laboratory data included: COVID-19 virus test positive (NR: not provided) Positive. On 25-MAR-2021, Laboratory data included: Body temperature (NR: not provided) Over 103. Laboratory data included: Oxygen consumption decreased (NR: not provided) 87. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the confirmed covid-19 positive, dizziness, fell, injury to kidney, hurt himself and blood clot was not reported. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition) This report was associated with product quality complaint : 90000174591.; Sender's Comments: V0: 20210355610- Covid-19 Vaccine Ad26.Cov2.S-Confirmed Covid-19 Positive. This event is considered Unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event. 20210355610- Covid-19 Vaccine Ad26.Cov2.S - Dizziness, Fell, Hurt himself, Some damage to his kidneys in the process (blood clot). This events are considered Unassessable. The events have a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1177000-1	Doctor determined superficial thrombophlebitis on left hand where IV 4 weeks before was taken out while at ER for kidney stone 2/18/2021. Hand was somewhat sore for weeks before vaccine. J & J vaccine on 3/12/2021. 5 days later 5/17/2021 left hand very painful and huge blood clot. Went to doctor and diagnosed blood clot and precribed clindamycin 300 mg 3X a day for 7 days. It wasn't until the 7th day the pain was gone and swelling of the vein back to normal

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THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1178023-1	Admitted with ischemic stroke and multiple thrombi in extremities
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1182945-1	Massive nose bleed with a huge blood clot to follow. Went to urgent care but it had stopped by then so they sent me home and told me if it happened again to go to ER. I haven?t had a nosebleed in years so it was scary to get it the day after my vaccine. The nose bleed lasted about 10 minutes and was very strong once the clot came through it slowed down and then stopped and did not return again.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1183089-1	Friday 3/19 Confusion, did not know her granddaughter, which grandchild she was kept confusing family names. That was very unusual for her. Saturday 3/20 didn't feel good that day, Sunday 3/21 around 3am got up to go to the restroom got very light headed and fell/fainted hitting her head on the night stand and falling to the floor. When EMS arrived she was in A-fib. That had never happened before. No known heart problems.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1183644-1	Shortness of breath, hypertension, hypoxia, tachycardia, lips and fingers turn blue, high fever, rash, hives, joint pain, whole body aches, left lower leg burning, artery blockages, infection, weakness, dizziness, tunnel vision, sweating, headache, new onset atrial fibrillation with rvr All of the above symptoms first occurred 30 minutes after injection of vaccine The atrial fibrillation occurred 6 days after injection Left lower leg started burning 15 days after injection The artial blockages and thrombus 18 days after injection by medical diagnosis due to instantaneous shut off of blood to left leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1185426-1	lymph node swelling with pain; blot clot, blue bruise like
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1192294-1	Fever 100 degrees for 36 hrs Fatigue for 48 hrs Headache for 1 week post Menstrual cycle started 6 days early Menstrual cycle was heavier and many clots noted
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1194975-1	Severe shaking. Admitted to ED for 180/90 blood pressure and rapid heartbeat (>120). Screened for potential pulmonary embolism and blood clots
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1195446-1	Blood Clots in legs and lungs
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1195850-1	Death by clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1199238-1	Approximately three weeks after receiving Johnson & Johnson vaccine, I began to get pains in legs. Calves were aching and throbbing. Didn't think anything of it at first....just thought achy muscles and such. The day before Easter (April 3rd 2021) decided since the pain was continuing and the internet will scare you to death - I went to a local Emergency Room in advance, I had an ultrasound on my left leg and it was noted I had DVT. (blood clot in left calf) To briefly explain: I had an achilles issue back in Nov. 2020 (slight micro tear - achilles tendonitis) I thought maybe this was related to that, as I had not been to the doctor to get it checked until early March 2021. and it bothered me off and on for a few months. I thought the calf pain may have been related to that? I explained to the emergency room doctor about my achilles and thought it was an issue related to that. They decided to perform an ultrasound on my left leg. Which noted the DVT/Blood Clot. They put me on a starter pack of Xarelto and I am to follow up with my normal doctor on April 22, 2021.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1199724-1	General muscle ache Dizziness Chills Very severe headache blew out my nostrils this morning and clotted blood came out
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200320-1	Flu-like symptoms. Muscle cramping in the left leg. Continuos period with heavy blood clotting.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200702-1	Had a huge blood clot come out of my nose after a nose bleed. I do not get nose bleeds so this was very strange. Took me 15 minutes to get it to stop bleeding.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200861-1	He had his vaccine in different state while visiting his daughter. Returned home and was found to have a blood clot behind his left knee. Also, he called the doctor last week and told him that he believes he has another one above the left ankle. He has an appointment on Thursday to see him again. He had two venous ablation procedure in both legs in late November/December of 2020. He was taken off of the baby aspirin and then put on Xarelto. The 2nd possible blood clot appeared about 10 days ago above the left ankle.

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THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200876-1	My mother received the Jansen vaccination on April 6, 2021. 2 days post vaccination she had 2 asthma attack at work. That following Friday she was not able to attend work due to shortness of breath. Saturday morning she cold not inhale and couldn?t stop coughing. She was then attempted to be taken to urgent care, but upon arrival she was coughing up blood and was instructed to go straight to the hospital. At the hospital upon test being done she was admitted to the covid unit and diagnosed with pneumonia due to covid 19 virus, bilateral pulmonary embolism in both lungs. She is now home in quarantine taking blood thinners for the next three months and unable to work.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200986-1	Findings in the great saphenous, common femoral, and proximal femoral veins suggesting nonocclusive thrombus.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1200992-1	"I developed leg pain in my right leg 2 weeks after vaccination. A week later, I went to the hospital and they confirmed ""Acute deep vein thrombosis) DVT of calf muscle vein of right lower extremity"" and at least one additional superficial clot. I was prescribed blood thinners apixaban (Eliquis) and was discharged. I'd previously had a clot in the same leg in 2009."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201057-1	1st night very out of breath. O2 at 93 on sensor at home. Felt better next day. 3/29/21 - went to ER for blood clot in left leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201200-1	Dermatitis on torso, hands and feet, skin peeling on left foot A few nosebleeds with small clots
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201222-1	small blood clot when tried to donate blood 2.5 weeks after shot, preventing me from finishing donation
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201310-1	I have a history of untreated fibroids that were diagnosed in December 2019. Because the coronavirus pandemic began in February 2020, elective surgeries were not allowed in my state. My bleeding and blood clots were caused by the fibroids during menstruation. On March 20, 2021, I received the Johnson & Johnson 1-shot COVID vaccine. By March 26th, the blood clots that were coming out of my body were enormous. They had NEVER been that large. Being concerned, I immediately saw my doctor as soon as my period ended. Today, I learned the J&J vaccine has been paused due to blood clots. I believe that my blood clots became so large because I was given this particular vaccine. WOMEN SHOULD BE ADVISED THAT IF THEY HAVE BLOOD CLOT CAUSING FIBROIDS, DO NOT TAKE THE JOHNSON & JOHNSON COVID-19 VACCINE.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201372-1	Resident c/o L arm pain on 4/7/2021 approx 2300. Resident transferred to ED for evaluation, CT abnormalities, transfer to higher level of care, clot manually removed on 4/8/2021, resident returned to nursing facility on 4/12/2021 without complications
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201378-1	Approximately three days after vaccine, I experienced a blood clot. This caused an acute ischemic stroke in my left frontal lobe, resulting in muscle weakness and partial loss of use in my right leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201386-1	developed a rare disorder involving blood clots within about two weeks of vaccination.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201389-1	Believe I may have passed a blood clot through urine or vagina early this morning. It appeared to be quarter sized. Non painful, one time event. Also have intense back pain and leg aching. Still feverish and have mild headache as well.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201418-1	Patient reports blood clots and longer bleeding time than usual with flossing.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201482-1	Patient had the vaccine on 3/5/21. She then had major abdominal surgery which included abdominal reconstruction, hernia repair and excess skin removal on 3/19/21. She was released from the hospital on 3/22/21. She was readmitted on 4/4/21 for blood clots.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201494-1	Supraficial blood clot in lower left leg. Pt contacted PCP, told to take 2 X 81mg aspirin for two weeks. Symptoms have resolved as of 4/13/21 when first reported to pharmacy.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201835-1	Low grade fever, aches & chills on 3/13/21 & 3/14/21. Felt better 3/15/21 through 3/18/21. Side effects (fever, aches, chills) returned on 3/19/21 & 3/20/21. Suffered stroke on 3/21/21 at 3:30 AM due to blood clots. Passed away on 3/22/21.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201961-1	10 after I received the Janssen Covid-19 vaccine, I admitted myself to the emergency room with severe chest pain and very difficult time breathing. They ran CT scans and found multiple blood clots in both lungs, pulmonary embolism. They put me on blood thinners and transferred me to another hospital. There they monitored me for a couple days and released me with a prescription for blood thinners.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202185-1	Blood clots running down the back of throat. Resolved by the following day.

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THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202402-1	Swollen of left leg and foot, took her to Delnor Hospital ER and they ran test.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202482-1	blood clots
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202568-1	Possible blood clot in L knee, appears to be superficial. I get these frequently (about once a year).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202579-1	On Saturday, 4/10/21, approximately 2 hours post vaccination, patient noted a knot on her left lower leg. No pain, redness or swelling noted to left lower leg. Patient took it easy the rest of the day. On Sunday, 4/11/21, patient began to have N/V, diarrhea and fatigue. No fever noted at this time or up to time of this report. As of 4/13/21, at 12:00pm the knot still remains present on left lower leg, along with fatigue and nausea. Patient was instructed to call PCP. Patient has an appointment with PCP on 4/14/21 at 1030am.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202608-1	started coughing on 03/27/2021 and was taking to hospital on 03/30/2021. There he was treated for blood clots with anticoagulants and a filter placed in the vena cava.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202639-1	Develop a blood clot lost vision in left eye due to stroke. I?m still in the hospital. I am 50 and had a stroke a week at getting that vaccine TPA shot. Given. Clot buster
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202661-1	I woke up 4am on April 6th 2021 to pee and I noticed blood after my pee. There was no blood in my pee or discoloration however I had to clean my penis tip with a tissue and noticed a blood cloth came out while I was wiping. This concerns me but that was the only time I saw blood, there was none afterward nor has it occur since then.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202689-1	PATIENT REPORTS COUGHING UP BLOOD CLOTS FOR THE PAST 3 DAYS AND SENT PICTURE OF 1 THAT WAS COUGHED UP
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202713-1	Slight fever 12 hours later. Headache the next day. Severe headache with nausea two weeks later that lasted more than a day. Notice a small clot in left forearm. No pain, though.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203268-1	3 blood clots along the left arm where the vaccine was administered.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203273-1	Patient reports that several hours after receiving the vaccine that he became short of breath, he was taken to the hospital via ems, Diagnosed with blood clots at hospital. Reports he is now on blood thinners. I spoke with the patient and told him I would be reporting this information and that he may be contacted.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203288-1	pt developed a blood clot in his ear and sought treatment 9 d after vaccination
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203308-1	Acute metabolic encephalopathy, BPH with urinary obstruction due to blood clot. Patient was vaccinated at a pharmacy with Janssen covid-19 vaccine and experienced headaches and dizziness. Then began to have urinary retention which was believed to be due to a large blood clot requiring 12x 100mL flushes. Patient went to ED at medical center on 3/30/21 and was admitted. He was discharged to nursing home on 4/11/21.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203443-1	The adverse symptoms I experienced were abnormally severe/heavy menstrual bleeding and abdominal pain. I was on Day 4 of my menstrual cycle on the day of my injection (April 3, 2021). My menstrual cycle usually starts to go away on Day 4 and concludes by Day 7. But the day after my injection, in addition to experiencing the common side effects of fever, chills, aches, and headaches, I also experienced an uptick in bleeding, which included severe/heavy menstrual bleeding with very frequent and large blood clots. That same day (April 4), I started my new birth control cycle as prescribed. By Day 8 (April 7, 2021), my menstrual bleeding continued to be so severe/heavy that I went to my OB-GYN for testing. The OB-GYN ruled out everything but the vaccine as an environmental factor. It was suggested that I take 600mg of Motrin every 6-hours for 48 hours. My menstrual bleeding continued for several more days until concluding on Day 12 (April 11, 2021) but I am still experiencing abdominal pain.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203460-1	Pt had Covid in November 2020. He was in the hospital and had blood clots in the left leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203902-1	Blood Clot which led to Pulmonary Embolism, Pneumonia, and death.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1203985-1	Pain, reddness, swelling in right leg, below the knee, when I got up in the morning. Symptoms continued over the weekend. Called Dr. on Monday, April 5, 2021 and got an appointment for the following day. Dr. diagnosed it as phlibitis and sent me for an ultr sound. Prescribed Xarelto 15 mg twice a

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				day for 21 days, then decrease to 20 mg daily.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204058-1	Patient started having abdominal pain on March 26, 14 days after vaccination. Went to ER on March 29 and was admitted. Diagnosed with blood clot in spleen. Discharged from hospital on March 31.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204231-1	Suffered from a stroke on 4/3/21 from a blood clot in his brain stem. Prior to vaccine he was healthy and 100% independent, still drove, did grocery shopping, yard work, etc. Now he cannot move his left side and was placed on hospice as he has lost his will to eat, drink, or live.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204459-1	blood clot in left ear
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204622-1	"Day of vaccine- Within hours, site soreness, raised red, approx size of a quarter, mildly painful, with a ""lump (that is still palpable) under the surface of the skin"". Within 5-6 hours following vaccine administration, she reports she developed fever, chills for approx 2 days with fatigue and hot flashes on and off. ** On 4/7/2021, patient states that in the morning she developed pain in her right calf. She saw her PCP."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204702-1	Patient seen in clinic on 04/07/21 reporting 2 weeks of right leg swelling and calf pain. Vascular ultrasound on 04/07/21 showed blood clots in legs. Lovenox injections started 04/07/21. Warfarin started 04/08/21.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1204744-1	Woke with: Blood Clots in my left arm at injection pt and lower arm, wrist, hand. Blood Clots in my left leg, mainly calf, lower leg, ankle, some in thigh. Larger, more numerically present than in past from antihistamines. None present before Vax, allergen-free diet prevents usually.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205012-1	Lower leg pain, confirmed clot about 4 inches long via an ultrasound on 4/9
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205079-1	Pt stated that approximately two weeks after vaccination she went to her physician office for an issue with her eye. Her eye was severely red. At the physician's office, the physician told the patient that there was a small blood clot in her eye. The patient stated she didn't think anything of it at the time, but now that the CDC came out with the statement about the Janssen vaccination, she decided to report to the pharmacy.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205086-1	Patient's husband called to report that his wife was in the hospital due to a blood clot that formed in her leg. He stated that she was having organ failure. Patient is currently being treated at a local hospital for blood clots that were moving and affecting her organs.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205385-1	Long period with heavy bleeding with blood clots. This has never happened before in my entire life.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205474-1	She fell twice. The second time was due to a blood clot on the brain that led to multiple bleeds under dura and the brain. She needed an emergency craniotomy and is now undergoing acute rehab for speech, PT and OT.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205729-1	My husband is still in hospital trying to recover. He had bad headaches that then led to seizures and confusion ,blood clots, and has been hospitalized since. He's literally fighting for his life and tomorrow Wednesday April 14 th he will be transferred to a rehabilitation center.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205766-1	I had rear end pain started March 20 in the morning, it gets worse and worse till 3/23 that I can not tolerate and had to call clinic to make appointment. They got me in on March 25 (Thursday). Dr inspected and told me it's thrombus that caused huge pain. They did the surgery right away, removed thrombus. I never had blood clots problem and I was healthy, which my blood test result can prove. Thrombus generated 12 days after my J&J vaccine shot, I am reporting this side effect. You may find all my medical record from Doctor's office.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205819-1	Had chest pain on and off for 5 days (started Thursday 3/25), which I thought was related to acid reflux. On Tuesday 3/30 when it became worse and was no longer managed with ibuprofen I went to patient 1st. I was given an ekg which was abnormal and then sent to ER. There it was determined that I had a blood clot and 100% blockage in my right coronary artery. That night I had surgery and they placed a stent.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1205996-1	Nearly a week after injection, injection site is still visibly swollen, slightly warm to touch. Blood clot passed vaginally (outside of normal menstruation schedule).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1206172-1	Severe blood clots,surgery sbc filter

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THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1206380-1	On 4/8/2021 I developed a headache after receiving the Johnson and Johnson vaccine. I took the next few days easy, and laid in bed for the majority of the weekend. On 4/10/2021 I left my residence to get dinner and noticed I was short of breath just walking from the building to my car. The shortness of breath worsened over the next few days. On 4/13/2021 I called a clinic and told them I had been short of breath and they directed me to the ER. I arrived to the ER around 2:30 in the afternoon. They ordered a chest x-ray and ran a test to determine if I could have a blood clot. The test came back with results indicating I could have a blood clot, so they did a chest CT and an MRI. The chest CT showed a blood clot in my left pulmonary artery. They prescribed a blood thinner, told me to stop taking oral birth control, and discharged me.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207088-1	The week of April 5, I believe it was April 5 & 6, I thought that I was coming down with a sinus infection. I was having some blood when blowing my nose and remember seeing some small, hard balls which I now realize were clots. I did not go to the doctor because my back was very bad and I was in a lot of pain and I had an upcoming appointment with my doctor. I believe that the symptoms resolved after 2 or 3 days and I did not even mention it to my doctor when I saw him. I felt a lot of pain in my leg, but this is probably due to my ongoing sciatica/radiculopathy /bursitis problems that I experience.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207377-1	Patient had vaccine on 03/15/2021. Started complaining of extreme pain in RLE by that evening. Patient contacted health agency and portable x-rays came out to view RLE. No fractures were seen. Pain continued and was unmanageable by 03/20/2021 and patient was transported to the ER. Patient was found to have blood clot in RLE and admitted to ICU. Patient further declined while admitted having multiple system failure and passed away on 03/25/2021.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207440-1	Massive heart attack due to blood clot. Heart Cath putting in a stint with over night stay in hospital
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207508-1	3/5/ 21 Received J & J at facility. 3/7/21 Client had troubles walking and went to Urgent Care. then he went home. He did not have good control (wife described that his legs did not have strength but was able to walk. Symptoms started getting better later that day. 3/15 He got up and he said he was not feeling to good. Chest hurt and shoulder hurt. Wife took him to the ER at Hospital. He was transported to Hospital. 3/16/21 Hospital did a Heart catherization. He had a complete blockage in the widow maker and 2 stents were put in. He had 2 blood clots in the Widow Maker. 3/18/21 He was discharged from the hospital. He is doing ok but he has no energy.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207534-1	Headache blood clots dizziness on and off and reduce of appetite
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207539-1	Chills/shaking began approximately 8 hours after injection. Other symptoms of an achy upper body, temperature of 103.1, and nausea came shortly after initial symptoms (within 1 hr.). In the middle of the night, I began heavily sweating (something I have never experienced). I was drenched with sweat and needed to change in the middle of the night. Absolutely no energy and a horrible headache. 24 hrs. after the injection I no longer had a high fever. However, consistently for a week, every day, I had at least one symptom (headache, nausea, chills, or lack of energy) at some point throughout the day. When I received the injection, I was on day 2 of my period. It ended one day later than normal. However, what was not normal was that on 4/3/21, I began to bleed again. I passed 2 clots and had spotting for an additional 5 days. This is in no way normal to my cycle.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207655-1	Day 1 & 2 after shot - left arm hurt around injection site. Day 2 I felt very tired and some body aches. By Day 5 or 6 - full body aches and my arms felt like there was pressure on them. I was feeling very tired. As more days passed; I was feeling tired, could not raise my arms, legs were weak and hurting. If I did anything (shower or carry a simple laundry basket) my arms would hurt so bad, I was out of breath and my heart was pounding so hard. By the week of March 29th - I was laying down in the middle of the day, no energy at all. Called the doctor and because of the shortness of breath & Body aches, they would only schedule a video visit for Monday, April 5th. After talking to me, they schedule to see me that day at 11:45 AM. I had weak muscle response in my arms, pale, and out of breath. They did blood work. Two hours after getting home; they called and said I needed to get to the emergency room. My red blood cell count was dangerously low, and they ordered a blood transfusion. I sent 6 days in the hospital.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1207994-1	Blood Clot on back of leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208026-1	"Patient received vaccine through pharmacy at employer vaccination event. Presented to clinic on 04/13/2021 and noted to have imaging confirmed ""There is occlusive superficial clot starting the upper lateral right thigh and extending distally through the mid to distal lateral calf. This appears to be within a superficial varicosity. Findings are consistent with extensive superficial thrombophlebitis of a varicose vein of the right lower extremity."""
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208107-1	Patient noticed blotches on both legs approximately 6 to 7 days after vaccine was administered. On 03/30/2021, patient fell to ground, unable to get up. Patient was taken by ambulance to Hospital emergency room. Scans revealed patient had multiple blood clots and was operated on. Later in the stay, patient developed widespread hematomas in groin, stomach, rear, and both legs. Patient was released from hospital on 4/8/2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208162-1	Blood Clot in right calf. deep arterial thrombosis
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208309-1	severe lethargy, shortness of breathe and fatigue (potential blood clot), went to PCP and prescribed Eliquis and Prednisone; however, written documentation said that he had no prior history of clotting prior to the vaccine.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208315-1	Patients power of attorney called to let us know patient had received Janssen vaccine on 3/12/21 and on 3/26/21 was hospitalized due to a clot and stroke. Patient had history of strokes in past.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208555-1	multiple large blood clots from her nares on Monday 4/12
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208568-1	Exact date and time of COVID vaccination unknown. Patient presented to Medical Center on 04/02/21 with left sided weakness and slurred speech. He reports receiving the COVID19 Janssen vaccine 15 days prior. Details of vaccination beyond patient report are unknown. Diagnosed with acute left cerebellar stroke, status post TPA on 04/02/2021. Extensive workup in hospital negative and unremarkable, including TEE, and hypercoagulable state so far negative. Follow up with hematologist as outpatient underway to evaluate for any possible coagulation disorders.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208701-1	Migraines, lung infection, blood clots, pain in lungs, shortness of breath.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1208750-1	Fell two weeks after immunization. Developed blood clot after procedure. Husband concerned immunization is cause of clot and not the surgery or down-time after the procedure. Transfered to new hospital from previous hospital and had significant stroke due to clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1209032-1	Persistent rectal bleeding and passing blood and blood clots for 7 days since vaccination. Bleeding occurs with stool and throughout the day, requiring the use of pads.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1209104-1	3 weeks post vaccination patient developed chest pain and shortness of breath. Was able to tolerate symptoms, then had radiation treatment and developed more shortness of breath. Was transported via EMS to hospital where an xray and MRI were completed and showed multiple blood clots in her lungs. Venous Doppler was also completed and blood clots were found in her legs. She was subsequently admitted to hospital.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1209183-1	Patient was vaccinated at 2PM (4/12/21) and caregiver said that at 4AM during the night(4/13/21) patient woke up very nauseated. She immediately vomiting and it contained blood. As she vomited more, numerous clots came out which resulted in caregiver calling 911. Patient was admitted immediately to hospital where she remains today (4/14/21). Doctors have performed an endoscopy and other tests to determine the source of the blood clots and treatment options. Caregiver is available if more details are needed.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1209679-1	Patient received his J&J vaccine for COVID-19 on 03/10/2021. On 3/17, he was at the grocery store when he felt a sting on his left lateral ankle. When he pulled up his pant leg, he saw the small little red spots that were exquisitely tender. He went immediately to his primary care physician's office, where he was prescribed Bactrim for suspected cellulitis. The rash steadily worsened with extension up his left leg and onto his right leg and blistering of some of the lesions most markedly in his groin. Pain associated with the rash increased and remained most severe on the lateral aspect of the left ankle. He presented back to his primary care physician's office on 3/19, where he was given methylprednisolone 80 milligrams IV, discharged with a prescription for prednisone 60 milligrams daily for the next few days. Pain worsened the morning of 3/20/21 to the point that he was unable to bear much weight on the left leg prompting his presentation to the emergency department.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1209797-1	vomiting, shortness of breath, hospitalized, chart shows she suffered from blood clot, hospital has extensive records from tests
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1210136-1	was found to have a blood clot in my left leg .
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1210630-1	"Mother called health department today, 4/14/21, to report reactions son experienced after receiving vaccine 3/4/21. She stated that on 3/6/21 he started having headaches, weakness and leg cramps. He went to the hospital two times with complaints. On 3/15/21 he was unable to walk/drive because headaches were severe. On 3/30/21 he passed out, squad was called. He was not responsive and was put on a vent at the hospital. A ""scan"" showed blood clots in brain and heart. This individual passed away on 4/4/21."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1210766-1	3/29/21 6 pm - Severe pain lower right leg, ankle, and foot, unable to weight bear on right leg About an hour later I was able to walk a little. Stayed off my feet. Right foot became very cold. 3/30/21 Saw my primary care physician at med center. She sent me for an ultrasound at medical arts. From there I was sent to the emergency room at hospital. I asked the attending physician assistant about a correlation between the blood clot and the Covid vaccine because I had read about it being a concern in other countries. He stated that was a bunch of baloney. Was told to have a repeat ultrasound in a few days and was prescribed a 30 day starter pack of Xarelto and released. 3/30 repeat penis ultrasound of right leg showed the same results. 4/1 appointment with hematologist. Prescribe Xarelto for three months and ordered a repeat ultrasound in three months and bloodwork at that time.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1210908-1	blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1211121-1	"Whether the adverse events were caused by the vaccine is to be determined. We are available to discuss more details if this information can be used to help others from experiencing these reactions. The morning after getting the vaccine, patient discovered that she could not swallow. As her condition worsened, she was taken to the emergency room. Over the past month, patient went to the emergency room 4 times. The last time resulted in extended hospitalization that continues to this day. In addition to the swallowing problem, two other major adverse events need to highlighted. First, on 27 March, while in the hospital, vaginal bleeding started. On 29 March, her doctor, performed a hysteriscopy/DNC finding "" ""an unexplained blood clot (hematoma)."" The second major event was when patient demonstrated significant confusion and lack of memory. On 7 April, she was diagnosed with Metabolic Encephalopathy. Neurologists at the hospital suspected that her MS was reacting to her UTI and once the infection is eliminated her memory should come back. But as of 10 April, she was still having issues including hallucinations. Although she is eating some as of 13 April, she is still very weak and will likely by moved from the Hospital to a rehab facility. She still does not remember much of the last 37 days."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212090-1	Systemic: Visual Changes/Disturbances-Severe, Additional Details: patients wife called to say that around 1 week after vaccine pt experienced vision issues. dr say he had a blood clot in eye which lead to partial blindess in one eye.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212127-1	Site: Pain at Injection Site-Mild, Systemic: Blood Disorder (diagnosed by MD)-Medium, Systemic: Dizziness / Lightheadness-Severe, Systemic: blood clots in stool-currently under care of primary care physician for clot formation investigation-Medium, Systemic: Flushed / Sweating-Medium, Systemic: Headache-Medium, Systemic: Nausea-Medium, Systemic: Shakiness-Medium, Systemic: Unable to Sleep-Medium, Systemic: Weakness-Medium, Additional Details: patient reported blood clots in stool on 4/11/21 and 4/12/21. Patient was flushed, dizzy and faint after clot was found. Patient has no significant history of hemorriods or colorectal conditions.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212435-1	He got the vaccine, was not able to breath real good and didn't think about it. It got to the point within 2 weeks that he was not able to get up from the chair due to the shortness of breath. By 4/1/21 he was not able to breath very well, but continued to stay home. Then by 4/12/21 he was not able to breath well at all and his wife took him to the ER. In the hospital they did x-rays and found out that his lungs are full of blood clots and both legs have blood clots in them. He was admitted and gave him Heparin drip for the 2 days to get the blood clots dissolved and was also on oxygen for 2 days. He was discharged home yesterday with Xarelto, and was told that he would be on that for the rest of his life, and is on oxygen for sleep and when he's up and around, but if he's resting and feels he's breathing okay he can remove it. They informed him that it would take approximately 2 months for all the blood clots to dissipate, and that he was full of blood clots, and had both legs and both lungs which are quite full.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212458-1	"2 weeks 8 days after vaccine, patient suffered which sent pt to emergency room. Husband states she had a ""blood clot"" to brain with skull fracture and traumatic brain injury. Transferred to Medical Center, then to long term facility. Has not recovered to date. Husband states unable to speak or move."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212529-1	I went on Sunday March, 14, 2021 to get the Johnson and Johnson shot offered through the school system. I noticed that my blurred vision intensified. I went to the eye doctor 3/16 and that is where they found blood clots behind my right eye close to my center. I was referred to a Retinol specialist 3/26/21 and sent me to get a physical to check for diabetes. It had started to heal a little. There was not clear diagnosis on what could have been causing the the blood behind my eye. 4/2/2021 Went for physical at PCP and results of physical was good with no Diabetes.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212673-1	Blood Clot and bruising on lower backside of leg, I noticed leg Pain (just uncomfortable) in the right leg at the location of clot. A time frame in relationship to the time of shot is unkown. Estimated within 24 hours. Bruising appeared morning of 04/14/21, PCP recommended in person visit. I went to urgent care, they had me drive to emergency Room at local hospital. Evaluation, blood testing and imaging concluded a blood clot in the lower right leg. Blod thinners was prescribed Eliquia 5 mg, discharged same day. It may be possible the hospital had submitted this form on my behalf.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212715-1	Developed pain in right calf and hamstring. No improvement over the next few days. Went to Vein specialist and they performed an ultrasound. A blood clot was found in my right calf.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212931-1	Stroke caused by blood clot - Vision Impairment and memory impairment
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212971-1	I had bloodwork showing increased D-Dimler and that I may have a blood clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1212978-1	Pt states that on Monday she noticed that she couldn't put pressure on her leg to walk. (04-12-21) Pt called and made an appointment on 04-13-21 and saw her PCP. Ultrasound was ordered and was dx with near occlusive vein thrombosis in the distal right femoral vein. Pt was prescribed Lovenox 100mg to take now and was also prescribed eliquis 5 mg twice daily, and acetaminophen-codeine to help with the pain.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1213131-1	PATIENT DIED FROM BLOOD CLOT 3/29/2021 - AUTOPSY PERFORMED AND CONFIRMED
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1213399-1	On April 11, 2021, patient passed a sizeable blood clot and has experienced labored breathing.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1213504-1	Deep Vein Blood Clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1213979-1	HOSPITALIZED FOR BLOOD CLOTS, UNRESPONSIVE AND TAKEN TO HOSPITAL
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214295-1	blood clot in arm
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214423-1	Charlie horse began 10 days after vaccine, leg felt full with cramping not allowing motion, went to ER diagnosed with blood clot in left calf
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214428-1	Patient has blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214606-1	4 hrs after onset of severe headache and body pain in lymph nodes under arm pit breast, on top of kidneys and in stomach fatigue low fever. After 4 days of that I got a really super heavy period with cramping and blood clots in my menustral flow. It was a horrible period, while the one after the surgery in Feb was very light. Dizzy and fatique 10 days after shot on the April 8 I took a fall from a chair outside, and from Thrusday to the Saturday 10th I ended up in the Emergency room with a concussion and head symptoms including dizziness and nose bleeds. This was from a simple fall, and i suspect the covid shot might have caused complications after the fall with additional swelling and brusing.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214739-1	I had the the vaccine on Friday then Monday i had a stroke from a blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1214815-1	Patient started not feeling well on 3/24. She was experiencing dizziness, headache, chills and fever. She contacted her doctor on 3/25 and he thought maybe she had an inner ear infection. Symptoms continued to worsen and eventually an ambulance was called on 3/27 because patient stated she was having severe SOB and headache along with dizziness and fever. Patient was transported to Hospital where they tested her for covid and pneumonia. Both came back negative. After several tests it was confirmed she had a blood clot in her lung and was also told she had clots in her leg and heart as well. She was in the hospital for 3 days where she was given heparin and several medications for the headache. Patient was discharged on lovenox and coumadin and eventually switched to xarelto by pcp who is managing her anticoagulation. She still has follow up appointments with a cardiologist and neurologist. She said she is still experiencing some dizziness and SOB but overall doing better
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1215019-1	Patient right leg became very swollen from knee down and was very painful to walk. Patient went to Urgent Care and was told that he would need to have ultrasound done to see what was going on. Patient went to Imaging and was told that he had several blood clots through out his right leg. They advised him to go to emergency room evaluation. Patient went to Medical Center and it was confirmed that patient had blood clots and was put on Xarelto for treatment.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1215158-1	Patient developed a blood clot

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1215435-1	Patient began experiencing back pain on March 14 and on or around March 18 he started coughing up blood clots according to a coworker. Patient lived alone and was unable to be reached on Sunday March 21. I went over to his house and found him deceased in his bed. Patient had a trashcan beside his bed and it appeared to have some blood in it. Patient had thick mucus coming out of his mouth and blood coming out of his nose. According to paramedics he had passed a few hours before finding him. Unfortunately Patient has been cremated so there is no way to say that this was related to the Covid 19 shot from Johnson and Johnson, however there are new reports that blood clots have been a side effect. I would like to speak with someone from the Department of Health to discuss this further. I feel this could be related to the vaccination and I would to know how long the Health Department knew about this possible side effect. If patient would have known sooner that the blood clots were a side effect I feel he would have gone to the Emergency Room. Patient had no health insurance and he was trying to prevent getting Covid. This has caused our family so much heartache and we are all very apprehensive about getting the Covid Vaccine ourselves. Please contact me as soon as possible, I am also patient's executor so I am able to speak to you on behalf of patient and our family. Thank you.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1215510-1	Woke up Sunday the day after and couldn't walk. Waited to see if it would wear off the next day was admitted into emergency with a minor stroke caused by a blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1215800-1	High fever for 3 days, lost of taste. Stomach pains and heavy menstrual cramps and bleeding. Blood cloths while menstrual cycle.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1216052-1	Approximately 5 hours after the vaccine I developed a very bad headache followed by chills, sweats, congestion, chest pain, body pain, muscle pain, fatigue and fever. The muscle pain and weakness was so bad I would loose balance when walking. I then started to menstruate clots, not my normal cycle timing, which lasted about 2 days. The clots subsided but I am still bleeding today. I was in bed for 4 days with severe immune response symptoms. After those symptoms subsided the fatigue was so extreme it was hard to function.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1216619-1	On 03/18/2021 at 03:30AM I had a lot of arm pain from my shoulder to my elbow in my left arm. This lasted until 03/31/2021. On 03/18/2021 I woke up with a headache that would not go away. Next, at about 03:15PM I was not able to walk for about 10 minutes. Then, at about 05:00PM I started menstruating. At 07:00PM I had severe pain in my pelvic area for an hour and a half. At approximately 08:30PM I passed a softball sized blood clot vaginally. I bled heavily through the night. On 03/19/2021 I still had a headache and no energy at all. I continued to bleed heavily all day as well. 03/20/2021 was the exact same as the days prior. 03/21/2021 I was still bleeding heavily throughout the day with no energy. I still had the headache as well. I started feeling dizzy while walking on this day. 03/22/2021 I went to work like normal. I was cold all day. I was still bleeding, headache and dizziness. It was hard to stand for 30 minutes. While climbing the stairs I got very light headed and dizzy. I could not go up the stairs at a normal pace. 03/23/2021 I had the same symptoms all day but I was getting short of breath walking short distances. I still had very little energy and a headache. I also lost all color in my face and hands. I was still cold all day. I noticed some slight swelling in my ankles. 03/24/2021 The symptoms were getting more intense. I was still cold. I finally took some aspirin for the headache because it was very severe. The medication didn't help with it. This routine went on for the next week and the symptoms intensified with each day. Throughout this time I was still passing clots that ranged in size from golf/ping pong ball to baseballs. They were very painful when they passed. The days from 03/25/2021 to 04/01/2021 were all the same as far as how I felt. On 04/02/2021 I woke up with the same symptoms but now I could not stay awake for anything. I slept on and off all day. I only ate one meal a day because I was so tired and I had no appetite. The headache was getting worse as the days went on. The bleeding was still happening. It was not as heavy as the weeks before, but enough to be painful still. I also started having abdominal pain. On 04/03/2021 I was seeing stars when I moved too quickly. My heart rate was elevated with shortness of breath along with the rest of the previous symptoms. My headache however was very intense. 04/04/2021 I still couldn't stay awake on top of all the other symptoms. The dizziness and stars were more frequent. I couldn't move my head without seeing stars. 04/05/2021 I couldn't get warm and I still felt the same way. I called and made a doctor?s appointment for Wednesday the 7th. I had all the same symptoms still and they were getting worse. I had to move in slow motion so I could try to function. 04/06/2021 I woke up drained of all my energy and color. I couldn't shower because of the pain in my body. I couldn't stand because I felt like I was going to fall down. I could barely stand up straight let alone walk very far. I was shaking from being so cold. I had all the same symptoms from the days prior. I was in severe pain. I couldn't sit because of the pain in my abdomen I couldn't stand because of the weakness in my legs. I could barely see anything because of the stars I was seeing. At this point I had no color and I was barely conscious. That is when my family took me to the emergency room.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1217619-1	Systemic: Blood Disorder (diagnosed by MD)-Severe, Systemic: blood clut in the lungs and legs-Severe, Additional Details: Per patient he went to the hospital on 3/17/21 and 3/23/21 and was diagnosed with a blood clut in his lungs as well as his legs. He was treated with Pradaxa but still continues to have difficulty walking and has overal body aches.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1217752-1	SHORTNESS OF BREATH; CHEST PAIN; SEVERE BLOOD CLOTS; This spontaneous report was received from a female patient of unspecified age. The patient's weight, height and medical history were not reported. The patient received COVID-19 vaccine Ad26.CoV2.S (suspension for injection, route of administration not reported), dose not reported, on 29-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, within a week of receiving the vaccine, the patient reported that she began experiencing shortness of breath and increasing chest pain. On 11-APR-2021, the patient was admitted to the hospital and a diagnosis of severe blood clots was made. After a full work-up, no etiology was determined. The action taken with COVID-19 vaccine Ad26.CoV2.S was not applicable. The outcomes of the shortness of breath, chest pain and severe blood clots were not reported. This report was serious (caused hospitalization).; Sender's Comments: V0:This female patient of unspecified age was hospitalized due to shortness of breath and increasing chest pain an unspecified time after blinded COVID-19 VACCINE Ad26.COV2.S was administered intramuscularly for prevention of symptomatic SAR-CoV-2 virus infection. No concomitant medications, past medical history has been reported. On an unspecified date, within a week of receiving the vaccine, the patient reported that she began experiencing shortness of breath and increasing chest pain. the patient was admitted to the hospital and a diagnosis of severe blood clots was made. After a full work-up, no etiology was determined. At the time of the report the outcome of the event is unknown. Based on the limited information the event is inconsistent with the causal association to immunization, per the WHO causality classification for adverse events following immunization. The event is considered not related to the blinded study vaccine. Additional information has been requested for further assessment.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1217874-1	"Per client self report: Experienced LLE extremity pain beginning on 4/11/2021, but thought nothing of it at the time. Client was previous scheduled for testing/procedure for asthma with pulmonologist on 4/15/2021, with pre-test/procedure lab work scheduled for Monday, 4/12/2021. Client presented for lab work on 4/12/2021 as scheduled and shortly thereafter received phone call from pulmonologist's office stating that the labs showed and elevated D-dimer and that she would be scheduled for a follow-up CT Scan of the chest and ultrasound of the lower extremities, which was scheduled on 4/14/2021. Client presented for CT scan of chest and ultrasound of lower extremities as directed and returned home on 4/14/2021. Client reported in conversation that she ""didn't feel right"" and noticed some SOB, and had planned to rest outside of going in for scheduled tests. Client received a phone call on 4/14 from pulmonologist's office advising her to go immediately to the emergency room due to a blood clot being found in the left lung as per CT scan report. Client presented to Emergency Room and was started on Eliquis 5mg tablets - 2 tablets by mouth twice daily X1 week and then 1 tablet by mouth twice daily and discharged to home. Client remains at home at the time of interview with no additional or worsening symptoms."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1218182-1	Asthma flair up, abdominal pain, blood clot, headache, and lack of sleep.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1218425-1	Patient received the COVID 19 vaccine on March 18, 2021 @6:00pm. On April 5, 2021 @8:00am the patient began to feel pain in L leg which progressed to area red and tender to touch and increased in pain with each day. Patient went to the ER on April 9, 2021 and an ultrasound revealed a 10 cm blood clot in left leg. Patient was not admitted but sent home on Zarelto 15 mg 2x daily. Patient did see physician as a follow up.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1218886-1	Bilateral arm swelling: 1. Occlusive thrombus is present within the right cephalic vein extending from the upper arm to the elbow. 2. Nonocclusive thrombus is present within the left basilic vein in the mid and distal arm. also developed acute appendicitis 04/06/2021 and underwent lap appendectomy.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1218973-1	"Per hospital chart notes, known timeline of events is pieced together as follows: 2021-03-23 - patient received J&J COVID-19 vaccine at Drug Store 2021-03-24 - patient developed myalgias, headache, shortness of breath, and chills - resolved within 24 hours 2021-04-02 - patient developed headache intermittent from right to left side of head for which patient was seen at an urgent care. Patient was prescribed Fioricet (butalbital/acetaminophen/caffeine). 2021-04-05 - patient developed right-sided neck pain 2021-04-06 - patient got tested for COVID-19, resulted negative on 2021-04-07 2021-04-08 - patient seen at Medical Center emergency department for headache and right-sided neck pain that shoots to (2021-04-08 cont.) the head and shoulder, for which patient self-medicated with ibuprofen. Patient reported to ED provider that using ibuprofen nearly relieves headache and makes her functional and had the most improvement the day she visited the ED prior to her visit. Patient received a CT scan of the head without contrast that was unremarkable. Patient was diagnosed with headache and neck sprain and was prescribed ibuprofen 600 mg, take 1 tablet by mouth every 8 hours, and diazepam 5 mg, take 1 tablet by mouth 3 times a day as needed for muscle spasm. 2021-04-12 - patient developed left lower extremity pain 2021-04-14 - patient followed up with Neurologist outpatient as directed by ED provider and was advised to get an MRV and left lower extremity doppler. Patient was prescribed tramadol 50 mg and butalbital/acetaminophen/caffeine (Fioricet) by the Neurologist. 2021-04-15 - patient presented to Medical Center ED before outpatient MRV and LLE doppler could be done due to worsening right-sided neck pain and inability to ambulate comfortably due to lower extremity pain that had been worsening over several days. Upon examination, there is subtle swelling to left lower extremity and doppler revealed multiple DVTs in left lower extremity. CT head without contrast was unremarkable, however, CT venogram of the brain reveals thrombi of the right transverse sinus and sigmoid sinuses, as well as IJ. Patient was initiated on argatroban and admitted to inpatient at the time of this report. ""Heparin induced platelet antibody"" and ""lupus anticoagulant evaluation"" tests are ordered and pending."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1219077-1	clotting in left leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1219798-1	Patient was in store and top of chest started hurting and down left arm. Patient states that they called 911 and ambulance took her to the hospital. She was tested and found to have mild heart attack with blood clot in vein of her heart. Had procedure to remove clot was put on blood thinner and is now wearing a heart monitor for the next 30 days. Is on Brelinta for the next 30 days as well. Patient went back to ER on Tuesday night (4/13/21) due to SOB spent 12 hours at ER before being transferred back to hospital. She spent another day and half before discharge diagnosed with fluid buildup.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1220641-1	Per pt report: Pt started having body aches within a couple hours of vaccine, developed severe 9/10 pain over the next several days. L shoulder pain radiating to ribs and neck w/ muscle spasms, causing HA, R hip and sacral pain, flank pain radiating down R leg. Pt reports that vision in L eye is blurry, R eye vision intact, numbness to L arm and hand, able to move. Pt reports 1in x 0.5 in blood clot when blowing nose Thursday morning, small clot subsequently about 1/2 the size, smaller clots later.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1220652-1	Blood clots (i started heavy menstrual cramps a day after the shot and I saw exceeding amount of blood clots which has never happened this excessively before) Fever, chills, headchae, nausea (all very severe level)
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1221632-1	Unusual Heavy menstrual bleeding with clots starting hours after receiving shot. Headache and fatigue day after shot. Menstrual heavy bleeding continued for 6 days outside of normal cycle. Day 7 having moderate leg pains and stomach pains and occasional headaches.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1222930-1	Patient woke up to get ready for work and noticed blood coming out of right ear. Clots were in the blood that came out of the ear.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223028-1	Seemed to very much effect my menstruation. My period came 10 days late, then this month?s period was 2 weeks early. Very bad clotting with menstrual bleeding. I am normally very regular so this was extremely unusual.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223054-1	It started with pain on the left side and shortness of breath. She was taken to Medical Center and there diagnosed with splenic infarction due to blood clotting. Was prescribed pain medication and blood thinners (Lovenox) to be injected daily for three months.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223301-1	"Patient reports tear in esophagus on 3/24/21 in the evening whenever she felt sick to her stomach. She did not feel well, drank one beer, and ate a packet of tuna. She continued to not feel well and felt the food did not go down correctly, so she reached into her throat and noticed blood. She called her doctor and the doctor recommended monitoring for increased bleeding. Later that night, she reports not feeling well again, made herself throw up, and noticed the vomit was all blood. The patient then called 9-1-1, and presents to the hospital. During the hospital visit she noticed that she saw 'blood clots about the size of a quarter' when she went to the bathroom. Patient reports the hospital gave her 4 pints of blood, 4 bags of 'yellow stuff' (patient assumes its plasma), protonix, antibiotics, and NS. Patient was in the ICU and reports being in the hospital for about 5 days total, whenever the doctor recommended being put on hospice. The patient did want to be put on hospice and wants to be treated, so she contacted her heart doctor and is waiting on an appointment with him on the 28th of April. Patient reports that doctor told her that she 'needs a new heart', but that she is not a good candidate, and that the Dr. told her this just a day or two ago. Patient reports that she did not know it was a tear in her throat (esophagus) until they ""stuck a scope down there""."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223502-1	received Janssen J &J shot 9:15 am on April 7th approx, 24 hours later had to leave work for home because I became incredibly weak, tired and without energy. struggled at driving home had to repeatedly slap my face to stay awake. checked my temp. when arrived home, had a low fever (100.1) went to bed slept for 4 hours, checked temp (100.1) went back to sleep. woke 9 am on 4/9/2021 checked temp. (normal) fever broke on 4/9/2021 used restroom at 11:01 am 4/9/2021 large amount of bleeding from rectum LARGE AMOUNTS of blood clots in stool (shaped like Candy) Contacted medical provider. Spoke to a triage nurse, explained my concerns about bleeding from rectum and blood clots in stool... she dismissed that my restroom issues were related to the vaccine,
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223667-1	Blood clots - minor stroke
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223789-1	My sister, received the J & J vaccine on March 5th. I now know that she was not feeling well for a few days afterwards. She then felt better until day nine after the vaccine. On the evening of the ninth day, she began having abdominal pain, vomiting & diarrhea, headache. This was Sunday evening. Unfortunately I wasn?t aware of how she was feeling and she didn?t go to the emergency rm. until Tuesday morning. Tests determined her platelet level was low and abdominal CT showed bleeding from her adrenal glands. She was given platelets, pain meds and admitted into the hospital. Later that night staff felt she wasn?t responsive and a head CT was performed. It was determined she had a brain hemorrhage and emergency surgery was performed. It is my understanding that there was brain damage at that point. During the early morning hours of the 10th day, post surgery, a repeat CT scan was done showing more hemorrhaging and blood clots. She also developed a DVT and PE and was kept on a respirator until the 13th day post vaccine to allow family to say their goodbyes. On the morning of the 13th day patient was removed from life support and she passed.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1223855-1	Chills 12 h after receipt of vaccine followed by low grade fever (101 F) and body ache with minor headache. Symptoms subsided by the end of the following day. 1st menstrual period post-vaccination was noticeably heavier with more clots than usual.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1224238-1	Left leg pain Mild Chest pain (resolved)
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1225799-1	4/11/2021 830PM I NOTICED MOISTURE IN MY UNDERWEAR. WENT INTO BATHROOM , PANTIES BLOODIED, BRIGHT RED AND A QUARTER SIZED CLOT WAS INTHE TOILET. I SUFFERED NO PAIN. HOWEVER I HAD A TOTAL HYSTERECTOMY IN 1997, AND SHOULD NOT HAVE ANY BLEEDING WHATSO EVER. TOOK A SHOWER, ADVISED MY HUSBAND OF THE INCIDENT. WENT TO BED , 230AM 4/12/2021 SAME THING HAPPENS, ONLY NOW I AM WIPING BRIGHT RED BLOOD AND I FELT A LARGER SIZED CLOT LEAVE MY BODY. I WENT TO THE ER IMMEDIATELY.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1226242-1	"Eight days after receiving the vaccine, my wife developed shortness of breath, headache, fatigue, and unusual vision problems. She was taken to the emergency room on 4/15 where she underwent a battery of tests (see details below). She was admitted to the hospital (where she currently is) and was diagnosed with blood clots in the lungs and two strokes in the area of the brain that affects vision. She then underwent ultrasound tests of her legs and carotid artery, which were negative for signs of blood clots. Although my wife's symptoms are different than the symptoms of others that led to the vaccine pause, her doctors strongly encouraged us to report it to the VAERS. My wife had no underlying or pre existing conditions that would lead to blood clots in her lungs, and stroke. After extensive testing there is no evidence that the blood clots in her lungs as well as the clot/clots that went to her brain initiated in her legs. They believe that some sort of ""event"" would have had to happen to cause these blood clots, and the only unusual thing in the time frame of this condition is the vaccine. As I mentioned earlier, my wife is still hospitalized."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1226722-1	Vomiting and Diarrhea day after vaccine. 6 days after shot blood clot and stroke, currently in ICU.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1226730-1	Big blood clots following Prostate Surgery 3 days after Vaccine applied Please see attached explanation and pictures
				THREE BLOOD CLOTS IN THE LUNG; TWO BLOOD CLOTS IN THE LEG; HEART ENLARGEMENT OF LOWER CHAMBERS; This spontaneous report received from a consumer concerned an 86 year old female patient. Initial information received on 13-APR-2021 and processed with additional information received on 15-APR-2021. The patient's weight was 161 pounds, and height was 67 inches. The patient's past medical history included hip fracture in 2015, and uncomplicated hip replacement surgery in 2015. Patient's concurrent conditions included non-smoker and no alcohol use. No recent travel or trauma was reported. No varicose veins or known cancer was reported. Patient saw her doctor for annual checkups and cancer screening. She is an active person who lives alone, still drives, and performs activities of daily living independently. The patient received vaccination with covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029) dose was not reported, administered on 08-MAR-2021 for prophylactic vaccination. Vaccination site was not reported. No concomitant medications were reported as it was reported that the patient denied HRT and chronic medications. Approximately 1 week later, on or about 15-MAR-2021, she began to feel fatigue, tiredness, and shortness of breath which she thought might be due to the start of a heavy pollen season. On 11-APR-2021, the patient's fatigue, shortness of breath, and tiredness did not improve, so her son took her to the Emergency Room (ER), where CT scan found 3 clots in her lungs and duplex scan found 2 clots in her legs (patient's son was not certain if bilateral or unilateral). Patient's oxygen saturation was in 80s. On 11-APR-2021, a test also revealed that the lower chambers of her heart were enlarged. On 12-APR-2021, patient underwent EKOS procedure to treat lung clots. Tissue Plasminogen Activator (TPA) had been put into the lungs via EKOS procedure 'with ultrasound inserted into blood clot in the lung'. Patient initially was treated with heparin but was stopped. Patient was treated with unspecified anticoagulant as patient's son did not know. Pulse rate was 139. On 12-APR-2021, patient was admitted to Intensive Care Unit (ICU) (no beds were available on 11-APR-2021). He had no information regarding lab results. On 15-APR-2021, patient was transferred out of ICU to a room on regular floor. Patient was still very tired. Oxygen saturation was in 90s. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from three blood clots in the lung, two blood clots in the leg, and heart enlargement of lower chambers. This report was serious (Hospitalization Caused / Prolonged, and Life Threatening).; Sender's Comments: V0: This 86 year-old active, independent female was hospitalized for 3 blood clots in her lungs and 2 blood clots in her legs 5 weeks after receiving Janssen Covid-19 vaccination. Medical history includes hip surgery after a hip fracture with uncomplicated course 6 years prior; she is a nonsmoker, does not have varicose veins, no known cancer, no history of recent trauma or travel and takes no chronic medications. Approximately 1 week after vaccination, the patient began to feel fatigue and shortness of breath. She initially thought her symptoms were due to the start of heavy pollen season. When symptoms did not improve 4 weeks later, she went to the emergency room where CT scan found 3 clots in her lungs and duplex scan found 2 clots in her legs. Oxygen saturation was in the 80s at that time. The following day she underwent EKOS procedure to treat the lung clots and was initially treated with heparin. The anticoagulant was subsequently changed after a public health announcement. She was admitted to the ICU and transferred to a regular room 3 days later. She was told the lower chambers of her heart were enlarged. She still feels very tired but her O2 saturation has improved to the 90's. No laboratory results were available to the reporter. Based on the limited information, the event is inconsistent with the causal association to immunization, per the WHO causality classification for adverse events following immunization.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227128-1	
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227129-1	"BLOOD CLOT; RIGHT LEG PAIN; HEADACHE; This spontaneous report received from a patient concerned a patient of unspecified age and sex. The patient's height, and weight were not reported. The patient's concurrent conditions included COVID-19. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805022, and expiry: UNKNOWN) dose was not reported, administered on 06-MAR-2021 at 02:19 for prophylactic vaccination. No concomitant medications were reported. The patient reported that she had a blood clot/pain in right leg 2-3 days after she received the vaccine. She reported that she used compression socks for a week. The patient also reported that she experienced headaches (date unspecified), which still come and go every other day. The patient stated that she had the same blood clot pain in the same leg as when she had COVID-19 infection in MAR-2020. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from headache, and the outcome of blood clot and right leg pain was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: A patient of unspecified age and sex reported a ""blood clot/pain"" in right leg 2-3 days after vaccine. The patient had a history of ""blood clot pain"" in the same leg when she had COVID infection one year ago. History of thrombosis would provide a plausible alternative explanation for the event, although there are insufficient details to make a meaningful medical assessment. The patient was contacted and could not be immediately reached. Additional information has been requested for further assessment."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227130-1	PNEUMONIA; BLOOD CLOTS; This spontaneous report received from a consumer concerned an 89 year old male. Initial information was received on 13-APR-2021 and processed with additional information received on 15-APR-2021. The patient's height, and weight were not reported. The patient's past medical history included congestive heart failure (ejection fraction 20%), large lower left groin hernia, atrial fibrillation, and dementia. No known drug allergies was reported. There was no history of blood clots. The patient received vaccination with covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown) dose was not reported, administered on 04-MAR-2021 for prophylactic vaccination. Vaccination site was not reported. Batch number was not reported, will be requested. Concomitant medications included apixaban twice a day for atrial fibrillation. On 11-MAR-2021, the patient was taken to Emergency Room (ER) and diagnosed with pneumonia. Patient was awake over 48 hours 'felt due to dementia and illness'. On 14-MAR-2021, the patient was taken back to hospital and X-ray and CAT scan showed saddle pulmonary embolism. On 15-MAR-2021, the patient was discharged to a home with hospice. On 23-MAR-2021, the patient deceased. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, Hospitalization Caused / Prolonged).; Sender's Comments: V0: An 89-year old man experienced fatal saddle pulmonary embolism 10 days after vaccine. Relevant medical history included congestive heart failure with ejection fraction 20%, atrial fibrillation (A Fib), and dementia. Relevant concomitant medication (others not reported) included Eliquis for A fib. The patient was diagnosed with pneumonia in the Emergency Department 7 days after vaccine, and 3 days later was brought back to the hospital and diagnosed with saddle pulmonary embolus. He was discharged home on hospice the next day and died 19 days after vaccine. There was no reported thrombocytopenia. The patient's age, concurrent pneumonia, and complicated past medical history are confounders. There is insufficient information to make a meaningful medical assessment. Additional information has been requested, including attempts to contact the patient's treating physicians.; Reported Cause(s) of Death: PNEUMONIA; BLOOD CLOTS
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227818-1	BLOOD CLOT; This spontaneous report received from a consumer via media by a company representative concerned a female of unspecified age. The patient's height, and weight were not reported. The patient was prone to blood clots her entire life, but managed it. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration and batch number were not reported) dose (1 total), start therapy date were not reported, for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient developed a blood clot. A day after the vaccination, the patient died from the blood clot. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender's Comments: This anecdotal report from media involves a female patient of unspecified age who was prone to blood clots her entire life and on an unspecified date developed a blood clot and died from the blood clot a day after received the Janssen COVID-19 Vaccine Ad26.COV2. Concomitant medications, and details of the event were not reported. This case has insufficient information to make a meaningful medical assessment. The case will be assessed further when additional information is received.; Reported Cause(s) of Death: BLOOD CLOT

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				BLOOD CLOT IN LEFT THIGH; SWELLING; FATIGUE; This spontaneous report received from a consumer concerned a 55 year old female. Initial information received on 13-APR-2021, processed along with information received via telephone communication on 15-APR-2021 The patient's weight was 100 pounds, and height was 152.4 centimeters. The patient's concurrent conditions included progressive multiple sclerosis, swelling of the left leg and foot, and non smoker, and other pre-existing medical conditions included the patient had no known drug allergies. the patient was not a pregnant at the time of reporting. The patient had no family history of thrombosis, no prior deep vein thrombosis (DVT). The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808609, expiry: UNKNOWN) dose was not reported, 1 total, administered on 22-MAR-2021 for prophylactic vaccination. Concomitant medications included baclofen, escitalopram oxalate, and risedronate sodium (received 2 doses and stopped after the event). The patient's husband reported that in MAR-2021, the patient had symptoms of fatigue. On 23-MAR-2021, the patient had swelling and fatigue. Her fatigue was resolved on 24-MAR-2021 and swelling was resolving. On 05-APR-2021, the patient developed left thigh swelling. On 06-APR-2021, the patient's swelling was not going down and she visited her primary care physician. The physician could not determine the cause of swelling (left thigh very noticeable clot swollen around where clot was located) and the physician suspected clot and performed an AD dimer (fibrin degradation) test to look for potential blood clotting. The patient also had a venous duplex ultrasound that showed left leg clot. At 19:00 on the same day, the patient's physician advised to visit emergency room and determined there was a clot on left thigh and she was prescribed Eliquis Starter Pack (apixaban) for 3 months treatment and recommended another scan in several months. On 06-APR-2021, the patient's vitals were reported as blood pressure 124/70 mmHg, heart rate 88 bpm and oxygen saturation of 99%. Laboratory data included prothrombin time (PT) 13.4 sec , International normalized ratio (INR) 1.2 and activated partial thromboplastin time (APTT) 34.8 sec. (Normal range not reported for all) d-dimer and platelet count results were not available and Covid-19 test was not performed. Body max index was 19.5 kg/m2. On an unspecified date the patient was feeling better but had not recovered from the event. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from fatigue on 24-MAR-2021, had not recovered from blood clot in left thigh, and recovering from swelling. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This case concerns a 55-year- old female subject, immobile due to progressive multiple sclerosis who developed a left thigh swelling 12 days after Janssen COVID-19 vaccine was administered intramuscularly for the prevention of symptomatic SARS-CoV-2 virus infection. The patient's concurrent conditions included body max index of 19.5 kg/m2, progressive multiple sclerosis and swelling of the left leg and foot. She is a non-smoker, no known drug allergies, no family history of thrombosis, or prior deep vein thrombosis (DVT). Symptoms started with swelling of the left thigh and fatigue. The fatigue resolved the next day. The patient's swelling in the left thigh was not going down and she visited her primary care physician who checked her D-dimer dimer (level not known to patient) and scheduled a venous duplex ultrasound that showed a clot in the left thigh. Her vital signs were blood pressure of 124/70 mmHg, heart rate 88 bpm and oxygen saturation of 99%. and she was prescribed Eliquis Starter Pack (apixaban) for 3 months treatment and recommended another scan in several months. Laboratory data included PT 13.4 sec, INR 1.2 and APTT 34.8 sec. D-dimer and platelet count results were not available and Covid-19 test was not performed. The patient is feeling better but had not recovered from blood clot in the left thigh, and is recovering from the swelling. Given underlying progressive multiple sclerosis with immobility and preexisting swelling of the left leg and foot on the same lower extremity, the clotting of the left thigh is considered not related to Janssen COVID-19 vaccine. Additional information has been requested.
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1227819-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227917-1	BLOOD CLOT IN LEG; This spontaneous report received from a patient concerned a 72 year old male. The patient's height, and weight were not reported. The patient was reported to be over 6 feet tall and not overweight. The patient's past medical history included cataract surgery (in his 60s); and concurrent conditions included former smoker (ex-smoker who quit 1 year ago). The patient had no chronic medical conditions and had no history of recent travel or trauma. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unk) dose was not reported, administered on 01-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. The patient was not taking any concomitant medications. On 04-APR-2021, the patient's leg felt abnormal and on 05-APR-2021, the patient went to urgent care center, where an ultrasound found a blood clot. The patient was prescribed Eliquis (apixaban) and was since feeling better, however, still had leg discomfort. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from blood clot in leg. This report was serious (Other Medically Important Condition). Additional information was received from consumer (daughter of patient) via telephone follow up on 16-APR-2021. The following information was updated and incorporated into the case narrative: patient details, medical history, concurrent conditions, medical story including treatment, and event details.; Sender's Comments: V1: Follow up from the patient's daughter provided additional clinical details. This case concerns a 72 year-old male without significant medical history who was diagnosed with a blood clot in his leg 4 days after receiving the Janssen Covid-19 vaccine. He is not overweight, is not taking chronic medication, and had no recent trauma or travel. He quit smoking one year ago. Three days after vaccination, his leg felt abnormal. The following day, he went to an urgent care center where an ultrasound revealed a lower extremity blood clot. He was sent home with apixaban. No information regarding blood testing was provided. As of the time of the report (i.e. 11 days after his diagnosis), he is feeling better but still has leg discomfort. Based on the limited information, the relationship with Janssen Covid-19 vaccine is considered inconsistent. Additional information is being sought.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227919-1	BLOOD CLOT IN LEG; BLOOD CLOT IN LUNG; This spontaneous self-report was received from a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029, expiry: unspecified) dose and therapy start date were not reported for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, 2 weeks after getting vaccination, the patient experienced blood clot in leg and blood clot in lung. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the thrombosis leg and thrombosis pulmonary was not reported. The patient also reported to VAERS (no reference number provided). This report was serious (Other Medically Important Condition).; Sender's Comments: A patient of unspecified age and sex experienced thrombosis leg and thrombosis pulmonary 2 weeks after vaccine. No concomitant medications were reported. No additional history, diagnostics, treatment, or other information was reported. There is insufficient information to make a meaningful medical assessment. Additional information has been requested.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227922-1	BLOOD CLOT; BROKE TAIL BONE; VACCINE EXPOSURE DURING PREGNANCY; This spontaneous pregnancy report was received from a pharmacist via a company representative, and concerned an approximately 40 year old female. The patient's height, weight, and medical history were not reported. The patient received Covid-19 vaccine Ad26.COV2.S (suspension for injection, route of administration not reported, batch number: unknown) dose and vaccination site were not reported, administered in 2021 for prophylactic vaccination. No concomitant medications were reported. In 2021, the patient experienced vaccine exposure during pregnancy. The date of the patient's last menstrual period and expected delivery date were not provided. In 2021, the patient experienced broke tail bone during labor and gave birth (live birth). On an unspecified date in 2021, the patient experienced a blood clot and died. It was noted that she was at high risk for clots because she was 4 weeks post partum (gravida 1, para 1). Action taken with Covid-19 vaccine Ad26.COV2.S was not applicable. The patient died of a blood clot and broke tail bone in 2021; the outcome of vaccine exposure during pregnancy was not reported. It was unspecified if an autopsy was performed. This report was serious (Death). This case, from the same reporter is linked to 20210430297.; Sender's Comments: V0: The case concerns a pregnant female subject around age of 40, who developed thrombosis, skeletal injury and exposure during pregnancy an unspecified time after Janssen COVID-19 vaccine was administered intramuscularly for prevention of symptomatic SARS-CoV-2 virus infection. The subject's past medial history, last menstrual period, estimated date of delivery and concomitant medications were not provided. Per the reporter (pharmacist) the patient was at a high risk for blood clots because she was 4 weeks post-partum. The patient broke her tail bone during the labor, gave a birth, and later died of a blood clot. No additional information was provided. It is not known whether the autopsy was performed. Given alternative explanation and risk factors of pregnancy, labor and skeletal injury (trauma) the event of thrombosis is considered inconsistent with the causal association to immunization, per the WHO causality classification for adverse events following immunization. Events of skeletal injury was result of an accident and therefore not considered related. Company causality for event of thrombosis is considered not related to Janssen COVID-19 vaccine (Level 4 - Insufficient information available to confirm a possible, probable or a definitive case of venous thrombosis, per the Brighton Collaboration case definition); Reported Cause(s) of Death: BLOOD CLOTS; BROKE TAIL BONE

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				COLD SYMPTOMS; DECREASED PLATELET COUNT; SADDLE PULMONARY EMBOLUS; BLOOD CLOT IN RIGHT LEG; This spontaneous report was received from a physician, and concerned a 58 year-old male patient. Initial report was processed along with additional information received on 16-APR-2021. The patient's height was 76 inches, weight was 266 pounds, body mass index 32.4. The patient's concurrent conditions included hypertension, and non-smoker. The patient had no personal or family history of clotting disorders, no recent trauma or travel. He had an active lifestyle (an outdoorsman who cut his own firewood). The patient received Covid-19 vaccine Ad26.COV2.S (suspension for injection, route of admin not reported, batch/lot number: 1808609) dose was not reported, administered on 02-APR-2021 vaccine anatomical site unknown, for prophylactic vaccination. Concomitant medications included hydrochlorothiazide for hypertension. On 03-APR-2021, the patient experienced cold symptoms. It was reported that some days later, he developed shortness of breath, dyspnea on exertion, decreased exercise tolerance and right calf swelling. On 13-APR-2021, the patient went to the doctor's office for his symptoms. An electrocardiogram (EKG) was normal, and the ultrasound of right leg revealed a blood clot. As the patient preferred to keep costs down, he was not hospitalized and was started on anticoagulant Eliquis (apixaban). He was not hypoxic. On 14-APR-2021, he called the doctor after a syncopal episode and was advised to go to the emergency room (ER) where a computerized tomography (CT) scan showed a saddle pulmonary embolus with evidence of right heart strain. His oxygen (O2) saturation was 97-98%. No treatment was given in the ER as he already had started apixaban (2 doses received by that time). Laboratory data included a normal comprehensive metabolic profile, normal troponin, white blood cell count 9, hemoglobin 16, platelet count 120 (lower limit normal: 130). No other tests were performed, and no prior complete blood count (CBC) was available. On 16-APR-2021, the doctor called the patient for follow up, and noted that he was feeling better. Action taken with Covid-19 vaccine Ad26.COV2.S was not applicable. The outcome of the blood clot in right leg, saddle pulmonary embolus, decreased platelet count, and cold symptoms was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This 58 year-old obese (BMI 32.4) male was diagnosed with a right leg blood clot and saddle pulmonary embolus 11 and 12 days, respectively after vaccination with the Janssen Covid-19 vaccine. Medical history included hypertension controlled with hydrochlorothiazide. He is a non-smoker who leads an active lifestyle; there was no recent trauma or travel and he had no personal or family history of clotting disorders. The day after the vaccination, he experienced cold symptoms. Then, an unspecified number of days later, he developed right calf swelling, shortness of breath, dyspnea on exertion, and decreased exercise tolerance. Eleven days after vaccination, he went to see his doctor who ordered an ultrasound which revealed a blood clot in his right leg. EKG was normal and he was not hypoxic. Apixaban was started as an outpatient. The following day, the patient called the doctor after a syncopal episode and was advised to go to the emergency room; CT scan revealed a saddle pulmonary embolus with evidence of right heart strain. Oxygen saturation was 97-98%. Labs included normal comprehensive metabolic profile, troponin, white blood cell count of 9, hemoglobin 16, platelet count 120 (lower limit normal: 130). No treatment was provided in the ER as he had already started apixaban. No other tests were performed, and the patient returned home. Two days later, when his physician called him in follow up, he was reportedly feeling better. Of note, platelet count was not checked on the day of leg thrombosis diagnosis prior to apixaban initiation and no other prior values were available for comparison; thrombocytopenia is a known adverse reaction with apixaban. Based on the available information, the relationship of the blood clot in the leg and pulmonary embolus with Janssen Covid-19 vaccine is considered indeterminant.
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1227923-1	
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1227926-1	BLOOD CLOT; This spontaneous report received from a consumer via a company representative and concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, a week or so after the Covid-19 vaccination the patient passed away in his sleep. The patient had no underlying condition. An autopsy was performed on an unspecified date and the patient was found to have blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender's Comments: V0: A male patient of unspecified age passed away in his sleep an unspecified time after receiving the Janssen COVID-19 vaccine for prevention of COVID-19 infection. It stated that the patient had no underlying condition. A blood clot was found by autopsy; no further details are provided. There is insufficient information to make a meaningful medical assessment. Additional information is being sought.; Reported Cause(s) of Death: BLOOD CLOT; Autopsy-determined Cause(s) of Death: BLOOD CLOT

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227928-1	"POTENTIAL DEEP VEIN THROMBOSIS; SMALL BLOOD CLOT; ANKLE CALF BOTH SWOLLEN; This spontaneous self-report was received from a patient and concerned a 64 year old female. The patient's height and weight were not reported. The patient's concurrent conditions included no known allergies and smoker. Other relevant history included no alcohol intake, nor any drug abuse/illicit drug use. The patient received COVID-19 VACCINE AD26.COV2.S (suspension for injection, route of administration and dose not reported, batch number: 1805029, and expiry: unknown) administered on 14-MAR-2021 on the left arm for prophylactic vaccination. No concomitant medications were reported. On 17-MAR-2021, the patient had ""ankle calf both swollen"" indicating potential deep vein thrombosis. She went to hospital emergency room, and ""HCP confirmed small clot"". The action taken with COVID-19 VACCINE AD26.COV2.S was not applicable. The patient was recovering from ""ankle calf both swollen"" and had not recovered from potential deep vein thrombosis and small blood clot. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a spontaneous report of a 64 year old female who developed a swollen calf and ankle swelling indicating a potential deep vein thrombosis 3 days after receipt of Janssen COVID 19 vaccine. Patient was seen in the emergency room by HCP, who confirmed a small clot. No other medical history was reported but patient was noted to be a smoker, reported as 2x a year. Age increases the risk of deep vein thrombosis, but the there is insufficient other details to make a meaningful medical assessment."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227930-1	BLOOD CLOT; ARM SORENESS; TIREDNESS; This spontaneous report received from a patient and concerned a 36 year old female. The patient's weight was 177 pounds and height was 165 centimeters. The patient's past medical history included fibroid surgery (fibroid was taken out 2 months), and concurrent conditions included alcohol drinker (once a week - occasionally), non-smoker, no drug abuse or illicit drug use, no known allergies. The patient received COVID-19 vaccine ad26.cov2.s (suspension for injection, intramuscular (right deltoid), batch number: 043A21A) dose was not reported, administered on 05-APR-2021 for prevention of covid-19 disease. No concomitant medications were reported. On 05-APR-2021, an unspecified amount of time after vaccination, the patient experienced arm soreness and tiredness that resolved within 3 days. On 15-APR-2021, 10 days after vaccination, the patient experienced calf pain and blood clot. It was reported the patient felt a very sharp pain on her right calf while walking, pain scale was 8/10, and it was so painful that she could not put pressure on it. Therefore, she decided to go to the hospital to have it checked. Her attending physician confirmed based on ultrasound that she has a blood clot on her right leg. She was prescribed blood thinner Eliquis (apixaban) however she was still undecided because her insurance denied to cover it and it was out of pocket. Once she returned home, she stated that the sharp pain was gone and it was just achy. After conversation with her physician, her physician postponed the medication for now because he wanted to make sure that her blood was not already thin before prescribing a blood thinner. She will go to the hospital again on 16-APR-2021 to see a hematologist for blood works and her physician advised her not to work until 20-APR-2021 as a preventive measure. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from arm soreness, and tiredness on 08-APR-2021 and had not recovered from blood clot. This report was serious (Other Medically Important Condition, and Life Threatening).; Sender's Comments: V0: a 36-year-old female experienced a blood clot in her right leg, 10 days after receiving the Janssen COVID-19 vaccine for prevention of COVID-19 infection. The patient is a non-smoker, her BMI is 34.5; she has no other relevant medical history and reported no concomitant medications. Ten days after receiving the vaccine, she felt a sharp pain, graded 8/10, in her right calf while walking. Ultrasound showed a blood clot on her right leg. She was prescribed Eliquis but did not fill the prescription and the pain subsided; she was to have further blood tests and see a physician the following day. Additional information is being sought.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1227933-1	"THROMBOSIS ARM; SHORTNESS OF BREATHE; ARM BURNING; ARM DISCOMFORT; SWOLLEN ARM; This spontaneous report received from a patient concerned a 59 year old of unspecified sex. The patient's height, and weight were not reported. The patient's past medical history included blood clots treated with Eliquis. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805020, and expiry: UNKNOWN) dose was not reported,1 total administered on 08-MAR-2021, left arm for prophylactic vaccination. No concomitant medications were reported. On 22-MAR-2021, two weeks after vaccination the patient experienced swollen arm and arm discomfort. On 29-MAR-2021, the patient experienced shortness of breathe and arm burning. On 30-MAR-2021, the patient experienced thrombosis arm. Laboratory data (dates unspecified) included: Scan (NR: not provided) Right Arm Occluded. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from swollen arm, and shortness of breathe, and had not recovered from thrombosis arm, arm discomfort, and arm burning. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This 59-year-old of unspecified gender, currently on Eliquis for history of blood clots noted swelling of arm with discomfort and burning (described as ""like from a match"") as well as shortness of breath 14 days after receiving COVID-19 VACCINE AD26.COV2.S on left arm. Scan showed right arm was occluded. Treatment for the events was not reported; the patient is recovering from swelling of arm and shortness of breath; the patient have not recovered from thrombosis of arm and arm discomfort/burning. The event is confounded by the underlying history of blood clots that is being treated medically. However, the events are assessed as indeterminate with a causal association to immunization, per the causality classification for adverse events following immunization based on a lack of a definitive plausible biological mechanism. Considering temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information has been requested for further assessment."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1228304-1	I was ill with diarrhea, sick to stomach, blood clots coming out of my rectum for 24 hrs., body aches.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1228668-1	My symptoms i had sever headache! Once April 1st Hit! Thats Was the day I Had a Heart stroke at store, i don?t remember anything and once i woke up i was at the hospital, and they told me that i had a heart stroke because i had lots of blood clots in my body. And they told me i vomit and peed myself durong the whole situation i went in store
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1229680-1	Received Janssen vaccine 4/6. Started menstrual cycle 12 hours after vaccine even though had just completed last cycle 1 week prior. Normally cycles are 28-30 days apart. They have not been quite as regular since Covid and has had some shorter periods between normal cycles. After the vaccine bleeding was heavier than normal with large clots but by end of last week had changed color and looked like it was ending. Today had no discharge upon awakening but again has a small amount of bright blood. Of course, there is concern for coagulopathy related to the vaccination. No leg pain, no shortness of breath. Has had HA off and on since had Covid but no worse now than in previous weeks. Left ear pain has been painful since Covid but now has pain in both ears when that occurs. Quit taking ibuprofen last week for discomfort and switched to Tylenol.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1229690-1	Janssen COVID-19 Vaccine EUA: patient presents to the emergency deaprtment (ED) reporting headache (10 out of 10 pain, difficult to keep eyes open) with neck pain, blurred vision, nausea, and photophobia for two weeks prior to arrival. Patient is four month post partum and breastfeeding. Patient diagnosed with cerebral venous sinus thrombosis and intraparenchymal hemorrhage. Admitted and underwent venous sinus thrombectomy. Currently still hospitalized.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1230134-1	Started to feel chest pain and shortness of breath 3 days after the vaccine. Ended up in the hospital 5 days after the vaccine and diagnosed with blot clots in the lungs.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1230341-1	According to patient, she woke up at 2:00 a.m. to use the restroom and noticed she had a very dry mouth and was thirsty. She passed out and when she came to she experienced decreased vision. She was taken to the emergency room via ambulance for tests. She was diagnosed with a blood clot to her lungs after CT scans. She also had aditional scans and blood work and stayed at Hospital in town (I am having trouble adding this to the shaded area in question 21).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1230405-1	Six days after receiving the Johnson & Johnson/Janssen vaccine, the patient complained about strong pain in the right arm (same as vaccination arm). She went to the ER on the 7th day after receiving the vaccination and after doing EKG, ultrasound, and CT scan, they found multiple blood clots in her right arm where she receive the vaccine.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1230877-1	patient came to clinic c/o right lower leg swelling on 4/6/2021, on 4/13/2021 she underwent a doppler us and was dx with right common femoral vein thrombus. her platelets are normal

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1231210-1	I received the vaccine on 3-12-21 and woke up 3-15-21 with pain in the bottom of my left foot. I scheduled an apt with my PCP on 3-16-21. My PCP scheduled an ultrasound on 3-24-21 where the nurse discovered blood clots in my lower left leg. I started taking Eloquis later that afternoon. I then had an apt with a blood dr. on 4-9-21 where she drew blood for tests. I was experiencing my heart racing and she scheduled me an apt with a cardiologist on 4-13-21 where the cardiologist suspected that the clots had moved into my lungs. The cardiologist did an EKG on the same day. I have a heart echo scheduled for 5-7-21. I am experiencing side effects from Eloquis so 4-19-21 I switched to Xeralto.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1231266-1	Around 5:30 pm in the evening following my vaccine I started having chills, headache, nausea, and fever of 101. Chills got worse, whole body was shaking and shivering. Nausea progressed into constant dry heaving. Head was pounding with the worst pain I have ever felt in my life, worse than natural child birth and worse than running marathons. The nausea was so bad and I could feel my fever burning and getting worse. I was about to vomit so got out of bed to go to the toilet to vomit and the whole room started spinning completely out of control. My heart was racing and I could hear it and feel it pounding rapidly through my ears. I couldn?t walk and was so disoriented I crawled to the bathroom to throw up in the toilet. My husband heard me and came and picked me up off the bathroom floor and carried me to our bed. As the night progressed fever got worse, head was in almost unbearable pain, severe body aches over my whole body, heart racing and horrible nausea and dry heaving. I made it through the night. For two days after I had a headache and fever and nausea and body aches and couldn?t get out of bed to care for my children. By the third and fourth day I was able to get out of bed at times and still felt fatigue, nausea, headache and body aches. Anytime I would stand up for 10 days after the vaccine it felt like I was rocking on a boat. I am almost at 2 weeks and still feel fatigue and dizziness at times. On April 15th, 9 days after my vaccination I had symptoms of a blood clot in my leg. I went to the Emergency Room and was diagnosed with a superficial blood clot of the leg as well as swelling in the vein. I am thankful to be alive!
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1231692-1	5 Blood clots coming from right nostril, headache, fatigue, stomach pains Nose bleed with clots lasted 15-20 minutes. Headache, fatigue, stomach pains lasted throughout the day.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1231985-1	pt developed symptoms on 4/19/2021 after receiving shot on 4/7/2021. Patient was admitted to hospital and discovered to have a clot in left leg. Patient is receiving Lovenox.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1231997-1	He developed a blood clot in his right arm.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1232464-1	pt says one of her toes on her left foot went completely number. Within a couple of days she started swelling between her left big toe and the toe next to it. Her entire left foot started swelling and then her leg started swelling. She went via ambulance to ER. They did an US of her leg and found 2 blood clots in the back of calve. She was given a RX for blood thinner. She went to 3 pharmacies and could not find it so she went home. She went home and then her right foot swelling as well. She went back via ambulance to ER. She was given blood thinner injection, and IV and took blood work. Pt was admitted where she stayed for 5 days. She was released and prescribed Eliquis. She has to take 20 mg a day for first week and then 10mg a day the following week for 3 months. She still has constant pain and her left foot continues to swell. Pt will have to have another US in 3 months.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1232940-1	"patient had a headache, weak arm, slept a lot more, experienced pain in arm as well, and a ""funny feeling"" in her chest. Patient went to the ER and said they found a blood clot."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233121-1	Received shot on 3/12. Starting having pain in chest on 3/22. Was diagnosed with blood clots in the lungs on 3/26 via blood test and CT scan.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233202-1	Patient states that while vacationing, 3 weeks and 1 day after receiving her Janssen vaccine, she woke up at 8:30am with numbness in her left foot. When attempting to stand she fell to floor. After checking her blood pressure and blood sugar she took 2 tylenol and sat down to rest. Around 10:30am she began to have numbness in her left hand and on the left side of her face. She was taken by ambulance at that time to Hospital. While there they ran a number of tests and scans diagnosing her with have a stroke caused by a small blood clot in right side of her brain. She was admitted to the hospital and was not discharged until 4/3/2021. Upon returning home, she has been seeing her PCP, APRN and a neurologist. She has since been released on 4/19/2021 to go back to work and has no residual side effects from stroke. She states she is now on blood pressure and cholesterol medication and is taking a daily baby Aspirin.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233450-1	A week and a half after taking the Janssen COVID shot I had a knot and pain come up in my lower back. After going to the chiropractor two times the knot and pain ended up in my right leg. I then went to Urgent Care where they diagnosed me with a large blood clot. They sent me to imaging and after scanning the area confirmed that there was a blood clot in my right leg. Urgent Care then set me up with a specialist.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233630-1	In the morning I noticed a pinching feeling in my lower right leg. I called my doctor to check on this. She sent me for an ultrasound of the leg on 04/15/2021. During a follow-up visit she informed me that it was a superficial blood clot and set up a second ultrasound on 04/28/2021 for confirmation that it had cleared.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233818-1	Purple , looking like blood clots on the back of her leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1233912-1	Intermediate Risk PE-diagnosed due to symptoms of pre-syncope, palpitations. Clinical history includes long car ride
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1234046-1	Received a Janssen vaccination on 4/7/21 from a home visiting nurse from the County Health Department. On approximately 4/13/21, the family noticed slight signs of an issue, with patient slumping towards the right, and showing some signs of weakness on right side of body. Family contacted the PCP, who advised to take her to the ED. Family was hesitant to do that because patient had been bedridden for past few years. She seemed to improve somewhat on 4/15/21. Then the morning of 4/16/21, the family found her on the floor of her bedroom. She appeared to have had a moderate to severe stroke. Right side of body paralyzed, cannot speak. Uncertain whether mental faculties further deteriorated. PCP ordered a hospice facility for care. Stroke likely caused by blood clot but unsure if related to JJ vaccine. She has not been evaluated in person by her health care providers.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1234108-1	Went to the hospital with chest pain. They performed a catheter to check for blockages and instead found an extremely rare blood clot that ultimately caused a mild heart attack.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1234132-1	Blood clot in left leg , Found out on April 15 2021
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1234170-1	We got a call from Doctor from ER to inform us that this patient came to the ER today & got swollen arm @ the injection site (left arm). They did the ultra sound & found there's a clot in brachial vein on the opposite arm (right arm). Doctor said they'll give pt Eliquist today.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1234257-1	"a 27 y.o. female with a history of Covid infection December 26, 2020, positive covid test December 29, 2020. She had approximately 2 weeks of symptoms including generalized body ache and generalized weakness. Fatigue and some headache. Had some persistence of symptoms after that time. Had some benefit from ibuprofen. Was seen at emergency room on January 27, 2021 stating that she felt that all her symptoms of Covid had resolved except for some chest pain which awoke her on January 27. She presented to the emergency room. Had sinus tachycardia at 113 but otherwise unremarkable physical exam, laboratory studies with white blood count elevated 16,004-20, hemoglobin 13.7 g hematocrit 41.8%, MCV 87.1, RDW 12.7%, platelet count 329,000, segs 72.8%, lymphs 20.6%, monocytes 4%, eosinophils 1%, basophil 0.6%. BUN 10.1, creatinine 0.8, AST 24, ALT elevated 56, alkaline phosphatase 68, total protein 7.6, total bilirubin 0.3, albumin 3.6, globulin by subtraction 4.0, troponin less than 0.02, D-dimer quantitative 0.22, urine pregnancy test normal. Chest x-ray normal. Chest pain appears to be musculoskeletal and was reproduced by palpation of her chest wall. No evidence of deep venous thrombosis or pulmonary embolism. D-dimer was normal. Heart rate returned to normal after some intravenous fluids. Patient discharged on Zithromax 500 mg day 1 and then to 50 mg days 2 through 5. Also continued on ibuprofen. Received Johnson and Johnson vaccine April 5, 2021. Over the next week patient started having increasing soreness. Her joints hurt more than usual and her headache was worse. She had a sensation that her legs were ""numb and tingly"" starting at her buttocks and extending down her leg. Because of reported incidence of cerebral sinus thrombosis or cerebral vein thrombosis patient appropriately presented for screening in the emergency room and was sent for appropriate imaging. April 16, 2021 CT scan of head with and without contrast no acute intracranial hemorrhage, no mass-effect or midline shift. On contrast-enhanced images there appears to be a lobular nonocclusive filling defect in the far lateral right transverse sinus. Lobular occlusive filling defect is also likely present in the mid to central right transverse sinus. Short segment filling defect in the medial left transverse sinus. Remainder of sinuses and internal cerebral veins are patent. Focal lobular filling defect within the confluence of the right and transverse sinus most likely related to arachnoid granulation. No edema hemorrhage of the cerebellum or cerebrum. No other significant findings. April 16, 2021 confirmatory MRI angiogram of head without contrast with no restricted diffusion to suggest acute or subacute infarct. Contrast void in the middle to central right transverse sinus consistent with occlusive thrombus. Nonocclusive thrombus in the far lateral right transverse thrombus. Narrowing of the medial left transverse sinus without complete occlusion. April 16, 2021 2345 hrs. inital hematology consultation. D-dimer and fibrinogen levels have been requested stat and are still pending. If fibrinogen level is low will replace with cryoprecipitate. If D-dimer is elevated barely confirms diagnosis of possible vaccine related thrombosis. The fact the patient is not thrombocytopenic at this time is encouraging, however despite the lack of thrombocytopenia she still has clearly documented symptomatic nonocclusive and occlusive thrombus in her cerebral sinus. From UPtoDate.com Ad26.COV2.S (Janssen COVID-19 vaccine, also referred to as

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	the Johnson & Johnson vaccine) ? On April 13, 2021, the US Food and Drug Adverse Event Description
				Administration (FDA) and Centers for Disease Control and Prevention (CDC) recommended pausing administration of AD26.COV2.S to further investigate rare cases of cerebral venous sinus thrombosis and thrombocytopenia [33]. As of that date, six cases had been reported, all in females aged 18 to 48 years with onset 6 to 13 days after vaccination; during this period, 6.8 million doses were administered (with 1.5 million doses in females of that age range) [121-123]. Since some vaccine recipients have not been followed for longer than the time frame over which these symptoms develop, the incidence may change with additional follow-up. Initial symptoms included headache, backache, chills, and fatigue, and progressed to focal neurologic deficits. Intracerebral hemorrhage and thromboses at other sites were also seen in some patients. All of the five patients who were tested for the anti-PF4 HIT antibody tested positive. Another case, in a young male who was a vaccine recipient in one of the pre-emergency use authorization efficacy trials, had been previously reported. These cases appear similar to those reported following ChAdOx1 nCoV-19/AZD1222, another adenovirus-vector vaccine. Although rare, the observed number of events exceeded the expected rate among females <50 years old. Given the extreme rarity of these events, the FDA and CDC acknowledge that they recommended the pause out of an abundance of caution and to ensure awareness of these rare events. ?Evaluation and management of possible thrombotic complications ? Recipients should be aware of the possible association and seek immediate care for signs and symptoms suggestive of thrombocytopenia (eg, new petechiae or bruising) or thrombotic complications (including shortness of breath, chest pain, lower extremity edema, persistent severe abdominal pain, unabating severe headache, severe backache, new focal neurologic symptoms, and seizures) [117]. In such cases, some experts suggest evaluation with complete blood count and differential (including the platelet count), quantitative D-dimer, HIT testing, and imaging of any suspected site of thrombosis [113,114,125]. Onset 4 to 20 days after vaccination, platelet count <150,000/microL, elevated D-dimer, and a positive anti-PF4 antibody (HIT antibody) suggest the diagnosis. Treatment with a non-heparin/non-warfarin anticoagulant (eg, argatroban or direct oral anticoagulant) and intravenous immune globulin has been suggested. The CDC recommends not using heparin in individuals with thromboses following receipt of Ad26.COV2.S unless HIT testing is negative [126]. (See ""Cerebral venous thrombosis: Etiology, clinical features, and diagnosis"" and ""Clinical presentation and diagnosis of heparin-induced thrombocytopenia"", section on 'Terminology and HIT variants' and ""Management of heparin-induced thrombocytopenia"", section on 'Role of IVIG'.) As stated above all the 6 cases were ""all in females aged 18 to 48 years with onset 6 to 13 days after vaccination"" this would certainly fit the timeframe that we are seeing in this case. Recommendations from national guidelines would be to start treatment with dexamethasone immediately while awaiting intravenous gammaglobulin. To pursue intravenous gammaglobulin treatment at 1 g/kg over 1 to 2 days. It is recommended to avoid heparin and instead use direct factor X inhibitor such as Eliquis, which this patient is already been started on. There is unfortunately considerable risk that even though patient's symptoms appear mild at this time, that she may deteriorate in the very near future and I would recommend treating her aggressively at this time with dexamethasone and intravenous gammaglobulin while we are awaiting further testing. Patient clearly has had unusual thrombosis, especially in a 27-year-old. With the fact that she had headache dating back to January, it is impossible to determine the acute versus chronic nature of her venous sinus thrombosis. Covid infection does cause increase in coagulation as well. However, given the small risk of tragic outcome with cerebral vein and sinus thrombosis in the setting of Covid vaccinations, we will proceed aggressively with treatment starting tonight with intravenous gammaglobulin and intravenous dexamethasone. Will require rapid fasting glucoses to evaluate possible hyperglycemia from high-dose dexamethasone. Will consider stopping dexamethasone at 48 hours. Current recommendation for gammaglobulin would be single 1 g/kg or 500 mg/kg over 2 days. No recommendations for treatment beyond this point. Current recommendations for gammaglobulin dosing would recommend using an adjusted dose for patients with greater than 125% of their ideal body weight. This patient at 122 kg on a 5 foot 2 inch frame does qualify for adjusted dose and her orders reflect that. I appreciate the opportunity see this patient this evening and consultation requested by Dr. We will follow patient in hospital. Hopefully she will have gradual improvement and avoid any serious or tragic complications of her cerebral sinus thrombosis. We will have pharmacy report possible adverse reaction to Johnson & Johnson vaccine to the FDA as required by law. April 17, 2021 follow up. No new events. Laboratory studies remained excellent. White blood count is further elevated, however, this may be secondary to high-dose dexamethasone with a white blood count of 16,060. Hemoglobin 14.0 g hematocrit 42.8%, MCV 86.8, RDW 12.9%. Platelet count is actually increased slightly at 374 as of 9:10 AM this morning. Differential shows left shift as expected with dexamethasone. Results from yesterday showed fibrinogen normal at 369 (180?415) pro time normal at 10.4 seconds with INR of 0.97, PTT 26.7. D-dimer 0.23 (0.19?0.50). With platelet count, fibrinogen and D-dimer normal I am inclined to believe that this is not an acute post vaccination HIT-like phenomena. If it were I would still expect to see some degree of thrombocytopenia elevated D-dimer and perhaps low fibrinogen. The other explanations for her cerebral venous sinus thrombosis are either idiopathic spontaneous and merely coincidentally associated at the exact same time as her Covid infection or more likely, cerebral venous sinus thrombosis secondary to hypercoagulable state secondary to her Covid infection from December 2020. Unfortunatly there is no way to accurately date the acuity of the venous thrombosis in her cerebral sinus. HIT Testing is still pending and most likely will not be available for 3-4 more days despite being ordered stat. Have called the

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				laboratory and asked them to check with our outside reference lab what the expected return time is on this test. (LabCorp says ""Tuesday April 20, 2021). Patient refused intravenous gammaglobulin infusion which has been ordered last night. Patient has been receiving dexamethasone which was a bridge until intravenous gammaglobulin could be administered. Current guidance from the expert hematology panel (EHP) on Covid vaccine induced thrombosis dated April 7, 2021 from the public health would state that a patient who has a reduced platelet count without thrombosis with a D-dimer normal and normal fibrinogen or thrombosis with a normal platelet count and D-dimer normal and normal fibrinogen are ""unlikely cases"". They recommend for probable cases who have D-dimers that are elevated to send HIT assay and then give immediate intravenous gammaglobulin while awaiting results. They recommend fibrinogen supplementation, if needed. They recommend a direct acting antithrombin agents such as the Eliquis this patient is receiving. They recommend steroids particularly if there is a delay in getting intravenous gammaglobulin delivered. With the fact that we now have 24 hours in the hospital with no evidence of thrombocytopenia, no evidence of increased fibrinolytic activity or active thrombosis, and no evidence of hypofibrinogenemia, I would argue that her presentation is subacute. In this setting, I do not feel intravenous gammaglobulin would be necessary and does involve a small risk as well as considerable cost. Dexamethasone also involves small risk and negligible cost. I have discontinued both of these medications. My overall impression would be that this was not related to her Johnson & Johnson vaccination which was recent, but rather her more distant Covid infection. It is impossible to definitively answer the question however. The next question is duration of anticoagulation. Risk of recurrent cerebral sinus venous thrombosis is relatively small but the morbidity can be high. Studies have been performed showing medications in the same class as Eliquis have been associated with a decrease in incidence of the recurrence. Most hematologist recommend 3 months of anticoagulation for provoked venous thrombosis including cerebral sinus venous thrombosis. I feel this patient's thrombosis is ""provoked"", with her association at the same time of her Covid infection (and possibly vaccination). I would recommend Eliquis (apixaban) 5 mg twice daily or Xarelto (rivaroxaban) 20 mg daily with food. I would feel comfortable in discharging patient for outpatient follow up IF Dr and other speciality services agree. Plan 3 months of full dose anticoagulation with direct factor X inhibitor. Circulation 2010 Jun 29;121(25):2740-6 ""Long-term evaluation of the risk of recurrence after cerebral sinus-venous thrombosis"", Background: The clinical course of cerebral sinus-venous thrombosis (CSVT) is largely unknown because prospective studies with a long follow-up and with the goal to assess thrombosis recurrence rate and predisposing factors for recurrence are lacking. Methods and results: One hundred forty-five patients with a first CSVT were followed up for a median of 6 years after discontinuation of anticoagulant treatment. End points were recurrent CSVT or other clinical manifestations of venous thromboembolism. CSVT recurred in 5 patients (3%) and other manifestations of venous thromboembolism (deep vein thrombosis of the lower limbs or pulmonary embolism) were seen in 10 additional patients (7%), for a recurrence rate of 2.03 per 100 person-years (95% confidence interval, 1.16 to 3.14) for all manifestations of venous thromboembolism and 0.53 per 100 person-years (95% confidence interval, 0.16 to 1.10) for CSVT. Nearly half of the recurrences occurred within the first year after discontinuation of anticoagulant therapy. Risk factors for recurrent venous thrombosis were male sex (adjusted hazard ratio, 9.66; 95% confidence interval, 2.86 to 32.7) and, for thromboses other than CSVT, severe thrombophilia resulting from antithrombin, protein C, protein S deficiency, anti-phospholipid antibodies, or combined abnormalities (adjusted hazard ratio, 4.71; 95% confidence interval, 1.34 to 16.5). Conclusions: The risk of recurrent CSVT is low and is higher in the first year after discontinuation of anticoagulant treatment and among men. Mild thrombophilia abnormalities are not associated with recurrent CSVT, but severe thrombophilia entails an increased risk of deep vein thrombosis of the lower limbs or pulmonary embolism. April 19, 2021 Follow up, no change in condition. Discussed case with Dr, will order Factor V Leiden, prothrombin gene mutation, anti phospholipid and anti cardiolipin antibodies, if she will allow draw. Expect Heparin Induced Thrombosis testing result tomorrow, but with no thrombocytopenia after given heparin, expect will be normal. Will await result. If discharged, would recommend 3 months direct thrombin inhibitor (Eliquis/Xarelto/Pradaxa) at full anticoagulation dose. Then stop, no taper. Overall still feel ""provoked"" thrombosis due to December Covid, unrelated to J&J Covid vaccine, but can not definitively prove that.."
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1234500-1	Blood clot leading to stroke
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1234526-1	Blood clot in left leg on 3/22/2021 and 3/29/2021
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1234907-1	On April 16 patient came home fatigue, pale and legs hurting. The next day complained of right knee pain, Fatigue and looked pale. He woke up in the middle of the night complaining of right calf tightness and pain. On Sunday 18th he started to limp and when we looked at his calf it was double in size, warm to touch, and redness. We then went to the ER and had an ultrasound and CT scan . We were told he has multiple blood clots in the right leg and clots in both lungs. He was then transported to a hospital and given blood thinner shots in stomach. He was released from the hospital 4/20 and given blood thinners and assigned a hematologist. He?ll be on blood thinners 6-12 months, out of work a month and tested regularly to check platelets.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1235447-1	"Patient presented to the ER on 4/20/21 with headache. On Saturday (4/17/2021), the patient had ""the worst headache I have ever had"" along with severe inner left ear pain. Patient took an ibuprofen which helped with the pain for about an hour then the pain came back. Pain got worse 2 hours later when she drove home from work. She called the RN line and was told to take ibuprofen and follow up in the AM. The next day 4/18/2021, patient went to urgent care and received Toradol and was told it was a migraine. Today 4/20/21, patient still had pain and noted feeling left sided facial numbness and blurry vision that correlated with when she got the headache. Patient went to an urgent care today and CT scan revealed a blood clot. She was then sent to the ED for evaluation. In the ED, she has no headache and facial numbness and has minimal left ear pain. Her headache is localized to the left side of her head. Patient denies fever, chills, and COVID symptoms. Patient was started on argatroban drip per hematology recommendation."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1235593-1	"I received the Janssen vaccine on 3/5/2021. My first menstrual cycle after the vaccine, which started on 3/21/21, was significantly different than any other period. I woke in the middle of the night to extreme pain in my abdomen and bled along the floor as I walked to the bathroom. There were clots in the blood in addition to extraordinarily heavy blood flow. The period was so significantly different from any other that I scheduled an appointment with a nurse practitioner that works with my primary care doctor. I asked her to ""run"" all tests that were available and approved by insurance, which she did. All came back normal. Due to the significant difference in my period this month, the ARNP requested an inner-uterine ultrasound which found the below:"
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1236116-1	"IN ICU WITH CLOTS; This spontaneous report was received from social media via a company representative and concerned a male patient of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown) dose, start therapy date were not reported, for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient was hospitalized in the intensive care unit (ICU) with clots (date unspecified). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome for the event of clots was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: v0 This report involves a male patient of unspecified age who received the Janssen COVID-19 Vaccine Ad26.COV2 and, after an unspecified period of time, was hospitalized in the intensive care unit (ICU) with ""clots"" (site unspecified). Medical history, concomitant medications, and details of the event were not reported. This case has insufficient information to make a meaningful medical assessment. The case will be assessed further when additional information is received."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1236650-1	I began having pain in my lower left leg later on Sunday (felt like a pulled muscle) but by Monday it was more painful. Since I had surgery (see above) my doctor sent me fir an ultrasound where a clot was found. Went to emergency room to further evaluate and get treated. Was given elequis and am taking for 3 months. Was told clot was likely related to immobility from injury, but wanted to report just for your consideration
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1236759-1	Foot and ankle swelled and went to doctor - found blood clot in right leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1237651-1	Possible cerebral venous sinus thrombosis
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1237936-1	Blood clot in right arm , SOB, CHEST PAIN, arm pain,
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1238061-1	Blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1238583-1	States she developed leg pain 6 days after shot and didn't think it was anything to worry about until she saw the J&J blood clot news and decided to get checked. Had appt and ultrasound on 4/19/2021 with NP. Found to have two non-occlusive, small blood clots in the right leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1238678-1	Adverse event: Ischemic Colitis most likely from a blood clot Symptoms: stomach pain -> severe diarrhea -> diarrhea of only blood, no stool for 24 hrs Symptoms began Saturday after vaccination (which was administered Thursday afternoon) Treatment: hospitalization, steroids, antibiotics
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1238925-1	Pt experienced pain/cramping in his right leg on or about 4-15-21. Thought this was due to new work shoes. Pain continued through the weekend and on Monday 19th visited his PCP where a scan was preformed confirming a clot in his right leg between his thigh and ankle. Pt started on therapy and has a follow up with his physician on Friday 23rd. Pt doing well and in good spirits.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1239051-1	2 weeks and 2 days after receiving the vaccination I had a blood clot blown from my nose no before issues with a bloody nose and not after it was a large blood clot there was not any continued bleeding before or after. Also where the shot was given is still sore in that area and does not seem to get better this is my right arm I am exercising and not any improvement.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1239726-1	Super sick 2-6 days after shot. Fever, chills, aches, headache, vomiting
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1240565-1	"BLOOD CLOT; This spontaneous report received from a male patient of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, frequency 1 total administered on MAR-2021(2weeks ago from date of reporting) for prophylactic vaccination.The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. The patient wanted to know about the blood clots with the JNJ shot and was thinking that he had a blood clot on 13-APR-2021. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: A consumer of unknown age or gender called company to inquire what is going on with blood clots after vaccine. The subject stated that he thinks he has a blood clot 2 weeks after vaccine, but did not provide details, did not see a doctor or have an any diagnosis, and refused to see a doctor despite encouragement. He did not share any contact information or other information stating he would ""wait for our lies to come out about clots."" This case is confounded by a lack of meaningful history, a lack of diagnosis, a lack of workup, and a lack of contact information for any follow up. A meaningful medical assessment can not be made."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1240708-1	Delayed menstrual cycle by 4 days, when started very painful and heavy. Much more clots than usual and lasting longer than usual. Uncomfortable enough to require rest. Still going heavy when it should be tapering off.?
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241054-1	BLOOD CLOT IN LEFT CALF; This spontaneous report received from a patient and concerned a female of unspecified age. The patient was called to obtain additional information on 21-APR-2021 and message was left on a voicemail. The patient's height, and weight were not reported. The patient has no family history of thrombosis. The patient does not have any underlying conditions. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, administered on 13-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On 14-APR-2021, the subject experienced blood clot in left calf. The patient noted that she did not have any underlying conditions or family history that may have made her more predisposed to this diagnosis. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot in left calf. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This spontaneous report concerns a female patient of unspecified age who experienced a blood clot one day after Janssen COVID-19 vaccine was administered for prevention of symptomatic SARS-CoV-2 virus infection. The patient's height, and weight were not reported. The patient has no family history of thrombosis and has not reported any underlying conditions. No concomitant medications were reported. The patient did not provide details related to the blood clot, except it was not resolved at the time of the report. Given lack of alternative explanation and temporal plausibility the event is considered possibly related to Janssen COVID-19 vaccine.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241143-1	BLOOD CLOT; This spontaneous report received from a consumer via a company representative concerned a female (consumer's wife) of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a spontaneous report of a female patient of unspecified age, who developed a blood clot an unspecified time after receiving the Janssen COVID-19 vaccine. No medical history or concomitant medications were provided. There is insufficient information to make a meaningful medical assessment. Additional information is being sought.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241185-1	BLOOD CLOT IN LEFT LEG; LEFT LEG CALF PAIN; This spontaneous report received from a patient concerned a 78 year old Asian, Not Hispanic or Latino male. The patient's weight was 220 pounds, and height was 67 inches. The patient's past medical history included stent (unknown where stent was placed), and concurrent conditions included non alcohol user, and non smoker. The patient had no known drug allergies and no any drug abuse / illicit drug use. The patient was previously treated with Clopidogrel bisulfate for drug used for unknown indication. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805031, and expiry: UNKNOWN) dose was not reported, 1 total administered on 18-MAR-2021 to left arm for prophylactic vaccination. Concomitant medications included amlodipine, metformin, metoprolol, and vitamin b complex all for drug used for unknown indication. On 28-MAR-2021, the subject experienced left leg calf pain. Patient reported that he experienced left leg pain below calf muscle and went to hospital and was further diagnosed with blood clot in left leg on 16-APR-2021. He was on Plavix and asked to stop Plavix on 16-APR-2021 by physician and placed on Eliquis 10 mg for the first 7 days twice a day and then 5 mg once a day for blood clot in left leg as treatment administered and with further concern to physician. He had appointment to hematologist for further follow up on 26-APR-2021. Laboratory data included: Diagnostic ultrasound (NR: not provided) Blot clot to left lower leg on 16-APR-2021. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from left leg calf pain, and blood clot in left leg. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a 78 year old male patient, who experienced left leg calf pain 10 days after receiving the Janssen COVID-19 vaccine on his left arm. 29 days after receiving the vaccine, patient sought consult for the left leg pain and was further diagnosed with blood clot on the left leg by diagnostic ultrasound. Patient's weight is 220 pounds and has a height of 67 inches. Patient is non smoker and does not take alcoholic beverages. Patient has no known history of drug allergies or drug abuse. Past medical history include having stent placed on him but he could not recall where the stent was placed. Patient was on Clopidogrel bisulfate but was asked to stop on the day he was diagnosed with the blood clot. Other concomitant medications include: amlopidpine, metformin, and vitamin B complex,. Indication for all medications being taken is unknown. Age and obesity (BMI = 34.5 per weight and height provided) are known risk factors for developing blood clot in the lower leg. Having a stent and concomitant medication suggest a cardiac problem. Based on the available information and considering the temporal relationship, the events of left leg calf pain and blood clot are assessed as indeterminate per WHO causality classification of adverse events following immunization.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241572-1	My period was supposed to start on 04/09 the day I got the vaccine. It was delayed until 04/13 when it began to bleed BLACK clotty sticky sludge. It had the appearance and consistency of vegemite or molasses. Black sludge. At first it was extremely heavy with very large black clots. Then it became tiny little black clots suspended in a brown sludge. This continued for 6 days until 04/18. Then on 04/19 my menstruation turned bright red with no clots, but has not abated at all for another 4 days. I am currently today on Thurday 04/22 still bleeding almost 2 weeks after taking the J&J shot. I cannot believe I am still bleeding after 10 full days!!! Never in 30 years of menstruating have I had a 10 day period. I am very scared and upset.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241772-1	I have severe swelling in both legs from knee to my feet, severe itching and hives, legs have blistered and burst open. I also have a Blood clot in my lower left leg. Have tried 2 kinds of antibiotics, I am on Xarelto 10 mg tab for the blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241867-1	Received the vaccine approximately 12:22 PM on Saturday, April 10, 2021. At approximately 6:55 pm that same day (Saturday, April 10, 2021) my husband collapsed in our living room. I called 911 and the operator talked me through how to do chest compression CPR on my husband until the paramedics arrived. My husband was then taken by ambulance to hospital where doctor performed an emergency procedure wherein he removed the blood clot in one of my husband's stints (he has three) that he had put in in 2017. Doctor was able to eventually stabilize my husband's heart. My husband spent 5 days in the hospital as a result of this.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241957-1	Thrombosed Hemorrhoid (blood clot). Rapid onset of initial pain. Still experiencing pain.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1241986-1	I had 2 adverse reactions to the Janssen Covid-19 Vaccine EUA: AE #1) Roughly 15 minutes after receiving the vaccination, the lower half of my face went numb for approximately 3 hours. The people who administered the vaccine made me stay at the site for about an hour and a half to make sure that the reaction did not get worse. AE #2) 10 days after receiving the vaccination, the lower part of my left leg began to swell significantly. I went to the emergency room on 4/13/2021. When I was there, they did tests and confirmed that I have a blood clot behind my knee. To treat the blood clot, the emergency room doctor gave me lovenox 110mg and warfarin 5mg. Since then, I have followed up with my primary doctor who has me taking warfarin 10mg (1x daily) and lovenox (as needed).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1242233-1	Blood clots in the Liver and Kidney also Pulmonary Embolism. went to ER was admitted and spent 6 days there. was put on Heparin as well as antibiotics. currently on Warfarin to maintain proper blood INR numbers.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1242493-1	On March 13, I received the Johnson & Johnson vaccine at the clinic The pharmacist monitored me for symptoms but there were none right away. I did have a fever and some body aches in the two or three days that followed but those were short-lived. While driving on March 16, I started experiencing some irregular breathing and periods of shortness of breath about three hours after leaving home. I stopped at the emergency room at the Health Center where Dr diagnosed me with an acute pulmonary embolism after discovering a small blood clot in one of my lungs. He prescribed me with Eliquis, which I am still taking today. Their phone number is provided if you have questions. I haven't had too many shortness of breath episodes lately but do not know if the blood clot is still there. I am scheduled to see my hematologist on April 27 to get further updates on this. My hematologist Dr is located in clinic number provided While it has not yet been proven that the vaccine caused the blood clot in my lung, I feel the need to report this in the wake of recent news about the Johnson & Johnson vaccine. My hematologist and primary care physician are both aware that I have received the vaccine but I have not received any further communication from them.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1242694-1	03/19/21 woke up with numbness on whole right side of body (head to toe). Referred to ER by Family Physician on 03/20/21. admitted to hospital for testing. MRI confirmed a blood clot caused a Lacuna Infarcts Event (type of mini stroke) The patient continues to have numbness and difficulty walking, driving and has affected her normal daily activities. The patient will be required to start Plavix and baby aspirin. Waiting for requirement for Prolong numbness which affects her daily activity.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1242748-1	I took the vaccine at 2:10pm on 4/8/21 and 20 minutes later at 2:30pm on 4/8/21 my cycle started a whole week early, which is very abnormal for me. I am bleeding clots of blood for 5 days now with weakness and pains. My normal cycle is very regular and lasts 3 days, the first day is heavy then moderate, then light on the third day i barely need a liner. This has been heavy bleeding with sizable blood clots.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1243646-1	On Sunday night, 04/11/2021 around 7:00pm, patient stated she started getting sick to her stomach. She complained of headaches and fatigue. At midnight she vomited, couldn't lay down and symptoms lasted all night. She complained of back pain too. she noticed blood in her urine with a clot of blood in it and had shortness of breath. She vomited twice. In contact by phone with her daughter at the time. Patient refused to go the the emergency room or call her physician. She has a cardiologist who is her primary care physician.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1244000-1	Swelling in ankle, then in right leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1244983-1	Experienced blood clots in my nose every night for about 10 nights after receiving vaccine injection. Didn't think anything of it until I mentioned it to my mom who received her vaccine at the same time & place from the SAME BOTTLE and she too was waking with bloody clots in her nose. I experienced a reoccurrence for 4 days in mid April. I wouldn't have said anything but friends nagged us after the J&J blood clotting issue. (We both also experienced severe piercing & throbbing pain in the bone beneath the injection for about 14 days.) I mentioned both events to my GP on 3/26/21
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1245227-1	My Mom received the J&J Vaccine. Apprx 10 days later fell, fractured hip, Surgery April 3rd - On April 5th doctor at medical center said she has a blood clot and low platelets. Was in Hospital from April 2 to April 12th
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1245290-1	Multiple pulmonary embolisms in bilateral lungs with lung infart, superficial blood clot in the right groin area, significant right abdominal pain, shortness of breath, low oxygen saturation levels admitted 3 days in the hospital. 36 hours of Lovanox injections followed by transition to Eliquis 10 mg twice per day, respiratory exercises, oxycodone given for pain Pain has diminished in lower right lung area, clotting appears to be under control
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1246172-1	PATIENT EXPERIENCE A BLOOD CLOT IN LEG AND HAD TO BE HOSPITALIZED
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1246956-1	Stroke and blood clot in right leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1246983-1	After vaccination, at 10pm I started feeling flu like symptoms, fever chills shakes feeling miserable and this lasted 3 days straight, I also had diarrhea, nausea and vomiting. I woke up with a swollen juggler vein that ended up being a blood clot, 10 days later my arm was swollen up twice the size of my leg, the juggler was still swollen. I call the urgent care nurse they said to go to ER. I went to hospital got a CAT scan and ultrasound, I was given a shot of warfarin that was supposed to last for 12 hours. I went home and the next morning instead of going to the hospital, I went to hospital and they found 2 blood clots in brain, arm and leg and neck. I have had no history of blood clots and this all happened after the J&J vaccine. They did a procedure where they tried to put a balloon to relieve the pressure. The blood clots in the brain cannot be removed because of their location and it would be dangerous to remove and I am on blood thinners for the rest of my life. Multiple blood clots in body brain, arm and shoulder.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1247921-1	4/15/2021 approximately 1030 am client found to be AMS, non-responsive, aphasic, aphagic, left-sided facial droop, teeth clenched, left eye closes, right eye with deviated gaze. 911 called, client to hospital ER, work up/evaluation, admission for stroke related to blood clots found left frontal and right temporal lobes, per spouse. Client discharged 4 days later to home s/p CVA, bed-bound, remains aphasic, aphagic, non-communicative, no response to verbal stimuli. PLEASE NOTE: hospital admission. Unable to complete section 21. Admit for 4 days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1248189-1	Blood clot in superficial vein in right heel of foot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1248235-1	Pain and swelling in Left Calf muscle. Treating the blood clot with Xeralto
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1248330-1	Patient has leg cramps starting on Monday, April 19th. She went to the hospital at later date. Doctor diagnosed her with leg clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249220-1	"ELEVATED CLOT LEVELS; DIZINESS/LIGHTHEADED; NAUSEA; BRUISE ON LEG; SWELLING; WISDOM TOOTH PULLED; This spontaneous report was received from a consumer and concerned a female of unspecified age and sex, White, Hispanic or Latino. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number and expiration date: Unknown) dose was not reported,1 total administered on 11-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The patient's father reported that on 06-Apr-2021, in urgent care patient's wisdom tooth was pulled. On 08-APR-2021, the subject experienced diziness/lightheaded, nausea, bruise on leg, swelling 4x4 inches on lower right calf. The next day, the patient's bruise/swelling move to inside of right knee, she had new bruise on leg and went to urgent care; the doctor told her she hit it. On 09-APR-2021, the bruise moved to inside of knee, and the patient was taken to emergency room (ER) where ""clot test"" showed elevated clot levels and rivaroxaban (Xarelto) was prescribed for treatment. On 10-APR-2021 (Saturday morning), ultrasound was done; result not reported. At the time of report there was decreased in swelling. The patient's father stated that, the event's might be related to wisdom tooth pulled. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from elevated clot levels, diziness/lightheaded, nausea, bruise on leg, and swelling. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a female patient, unspecified age, unspecified ethnicity, who experienced elevated clot levels 29 days after receiving the Janssen COVID-19 vaccine and 3 days after the patient's wisdom tooth was pulled out. The patient had her wisdom tooth pulled out 26th day post vaccination. 2 days later (28 days after the vaccination), the patient experienced dizziness and nausea and was noted to have bruise and swelling (around 4x4 inches) on the right lower leg. The next day, bruise/swelling was noted to reach the area of the right knee, probably the medial side, which prompted consult at the ER. The ""clot test"" done as was noted to be elevated (no actual results provided). An ultrasound was also done but the result was not reported. Long periods of immobility (sitting in the dental chair) and the temporal plausibility from the dental procedure to the event of elevated clot levels confounds the temporal plausibility of the elevation of clot level and the vaccine; hence this is assessed as indeterminate per WHO classification of adverse event following immunization. Additional information requested."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249239-1	BLOOD CLOTS; This spontaneous report received from a company representative via social media post and concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, one total dose is administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested. No concomitant medications were reported. On an unspecified date, the subject experienced blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This patient reported via social media to have blood clots and to have received the COVID-19 vaccine ad26.cov2.s. No other information provided. There is insufficient information to make a meaningful assessment. Additional information will be requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249328-1	BLOOD CLOT; This spontaneous report was received from a female patient of an unspecified age. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine Ad26.COV2.S (suspension for injection, route of administration, and batch number were not reported) dose, vaccination site, and start therapy date were not provided, for prophylactic vaccination; and was treated with XARELTO (rivaroxaban; film-coated tablet, oral, batch number was not reported) dose, frequency, and therapy dates were not provided, for an unknown indication. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced a blood clot. The patient reported she got a blood clot after receiving her vaccine, and that she had just started taking rivaroxaban. Action taken with Covid-19 vaccine Ad26.COV2.S was not applicable, and action taken with rivaroxaban was not reported. The patient outcome for the event of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: v0 This report involves a female patient of unspecified age who experienced a blood clot on an unspecified date after receiving the Janssen COVID-19 Vaccine Ad26.COV2. Concomitant medications included rivaroxaban (recently started). The patient's past medical history and details of the event were not reported. The recent starting of rivaroxaban infers a recent clot or a condition that would predispose the patient to develop a clot, however this case has insufficient information to make a meaningful medical assessment.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249336-1	"THROMBOSIS IN MY RECTAL AREA; ARM VERY SENSITIVE TO THE TOUCH; SEVERE HEADACHE; CHILLS; SORE ARM; This spontaneous self-report was received from a patient of unspecified sex, age, race and ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration, dose, vaccination site, and batch number were not reported,) administered on 16-MAR-2021 at 13:00 for prophylactic vaccination. The batch number will be requested in follow up. No concomitant medications were reported. On 16-Apr-2021 in the evening post vaccination, the patient had severe headache and chills. The headache continued the whole next day, but was not as bad as the first evening. Her arm was very sore, and it hurt to move it the first day, and then very sensitive to the touch the next day, and continued for a whole week. On 29-Apr-2021 (13 days post vaccination), "" thrombosis (according to my doctor) appeared in my rectal area that still hasn't gone away"". The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from arm very sensitive to the touch, severe headache, and sore arm, had not recovered from thrombosis in my rectal area, and the outcome of chills was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This patient of unspecified age, gender, and ethnicity reported to have ""thrombosis (according to my doctor) appeared in my rectal area that still hasn't gone away"" 13 days after receiving Janssen COVID-19 vaccine for the prevention of symptomatic SARS-CoV-2 virus infection. The patient also reported severe headache and chills on the day of vaccination that persisted until the following day, as well as arm soreness that persisted for a whole week. No other details was reported. Based on the information that is available, the event is assessed as indeterminate with the causal association to immunization, per WHO causality classification of adverse events following immunization based on a lack of a definitive plausible biological mechanism. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information is requested"
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249343-1	"CLOT OF ABOUT 3-4 INCHES; LOWER LEG PAIN; This spontaneous report received from a patient via a company representative concerned a 35 year old male. The patient's weight, height, and medical history were not reported. The patient had no history of blood clots and is active. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown) dose was not reported, 1 total administered on 23-MAR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date in 2021, the patient experienced lower leg pain. About a week after that, on an unspecified date in 2021, the patient experienced clot of about 3-4 inches confirmed by ultrasound. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the lower leg pain and clot of about 3-4 inches was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This 35-year-old male patient of unknown ethnicity was reported in social media to have ""clot of about 3-4 inches confirmed by ultrasound"" after an unspecified duration from receiving Janssen COVID-19 vaccine for the prevention of symptomatic SARS-CoV-2 virus infection. This was preceded by symptom of lower leg pain 1 week prior. No other details was reported. The information available precludes a complete and meaningful assessment. The case will be re-assessed once additional information is received."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1249414-1	"CLOTS ON RIGHT LEG; HEADACHE; PRESSURE ON BOTH LEGS; PRESSURE ON HANDS; This spontaneous report received from a patient via a company representative concerned a 64 year old Hispanic or Latino female. The patient's height, and weight were not reported. The patient's concurrent conditions included high blood pressure and diabetes. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029 expiry: UNKNOWN) dose was not reported, 1 total administered on 17-MAR-2021 at left arm for prophylactic vaccination. No concomitant medications were reported. On 17-MAR-2021, the patient had experienced headache after vaccination. On 27-MAR-2021, the patient began to experience intermittent pain on right leg. On 30-MAR-2021, she experienced headache again and was still having leg pain; she visited the Emergency Room (ER) and was admitted on 30-MAR-2021. On 02-APR-2021, the patient experienced clots on right leg. On APR-2021, a catheter was placed in catheterization procedure and her right leg vein was unclogged of a clot. A second clot did not require removal because the vein had already unclogged itself after medication. The patient was administered unspecified anticoagulants. She was discharged on 5-APR-2021 and was prescribed ticagrelor which was then changed to clopidogrel. On 18-APR-2021, the patient experienced a headache again and feeling of pressure on legs and hands. Patient was advised to visit ER. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from clots on right leg and intermittent pain on right leg on 02-APR-2021, recovered with sequelae from headache on 19-MAR-2021, and had not recovered from pressure on both legs and pressure on hands. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0: This 64 year-old Hispanic or Latino female with history of high blood pressure and diabetes was hospitalized for clots in her right leg 13 days after receiving the Janssen Covid-19 vaccine. Concomitant medications were not reported. On the day of her vaccination, she experienced a headache. Ten days later, she began to experience intermittent leg pain. Three days later (ie, 13 days post vaccination), she went to the emergency room (ER) and was admitted to the hospital. During the hospitalization, a clot was removed via catheterization and she was administered unspecified anticoagulants. A second clot resolved on its own without surgical intervention. After 6 days in the hospital, she was discharged with ticagrelor which was later changed to clopidogrel. She recovered from the clots. Thirteen days after discharge, she began to experience headache and ""pressure"" in both legs and hands. She was advised to go to the ER; no further information was provided. Based on the limited information, the relationship of the serious events with Janssen Covid-19 vaccine is considered indeterminant. Additional information is being sought."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251068-1	PT STATED HE WENT TO THE CLINIC AND THEN THE ER WITH A BUMP ON THE BACK OF HIS LEG. PT WAS TOLD IT WAS A BLOOD CLOT.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251202-1	Events: Vaccine was given on 3/19/21; Hysterectomy on 3/25/21; blood clots diagnosed on 3/30/21 Left forearm was swollen, with a light reddish tint to it, and tender. After getting an ultra sound on my forearm a DVT was found and I was sent to the emergency room. I then had a CT scan done of my chest and it was discovered I had several pulmonary emboli. I was admitted to the hospital and put on a heparin drip. I am currently taking Eliquis and will be following up with a pulmonologist and a hematologist in a week or two.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251240-1	13th of April. I had a blood clot in my hand. My index finger on my left hand was numb and black and blue all around the knuckle area and into my hand area. I went to my primary care - I went to the acute center there. She said the hand was cold and could see the black and blue. They put me on Plavix for 30 days and I haven't had any reoccurrence of that. I think it was the 21st, I couldn't sleep all night, I had a kidney stone - the pain was so bad I got nauseous - around my back and side and in my front. I didn't have a fever. I went to the doctor office and they checked urine - and said there was blood in the urine. I was prescribed - Tamsulosin HCL. They thought I had passed most of it by the time I got there.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251445-1	Feeling quite ill. Did not sleep night before trip to hospital. Terrible taste in mouth. Taken to hospital emergency room morning of April 18, 2021. Admitted to hospital by ER physician. Reason: blood c lots in both lungs.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251532-1	Patient stated she needed to go to hospital for adverse reaction of pain in leg. After asking MD to check her leg, they discovered a blood clot per patient. She is now doing well per her.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1251620-1	Severe pain in left thigh groin & knee area. Went to ER, confirmed blood clot. Sent home w/blood thinners - Xarelto
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1252018-1	"On 4/16/2021 went to the emergency room with high blood pressure that was noted as a ""hypertensive emergency"". Was admitted into the hospital the next day where a CT scan confirmed that a blood clot in the brain occurred which led to a basal ganglia stroke. After a 5 day stay in the hospital, patient is now in a full time rehab facility to attempt to recover her cognitive and speech skills, while also attempting to regain use of her right hand which is currently not functioning due to the stroke."
THROMBOSIS	COVID19 (COVID19 (JANSSEN))	JANSSEN	1256511-1	Left lower medial aspect swelling , warmth and thrombosis noted. Placed on Naproxen 500mg BID Compression stockings Heat and Cold compresses

Symptoms	Vaccine (1203)	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1256715-1	During my menstrual cycle I usually have no symptoms other than tender breasts. So these symptoms were very unusual. Leg pains, pain and pressure in pelvis, clitoral pain, pain around vaginal opening, stinging like acid during urination that sent a shock through my entire body, uncontrollable bladder, nausea, large blood clots, lower stomach pain, flank pain, hot/cold chills, if I touch my belly button the pain radiated to my vagina, ovaries hurt, cervix pain, dryness, unable to get comfortable, crawling out of my skin, and constant urge to urinate, depression. I landed myself in the ER where they told me to take Tylenol and Motrin and sent me on my way. These medications were not able to control the amount of pain I was in. I have now reached out to a couple specialists.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1256798-1	Patient's daughter came to the pharmacy on 04/24/2021 and stated that her mother, the patient, got the Janssen vaccine here and within a week, she had a blood clot and has been in the hospital ever since. This is all the information we received.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1257910-1	My first menstrual period following the vaccine was exceedingly heavy in flow for the first two days, which is atypical for me. Heavy clots of menstrual discharge were observed. It is not yet known whether these side effects will continue for my next menstrual cycle. Lesser effects followed two days after the vaccine with fatigue, sore arm, armpit, and glands in the injection arm. These side effects dissipated by day 3 status post vaccine administration.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1258387-1	3 Blood clots formed in the bottom of my left leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1258576-1	Heart went into Atrial fibrillation causing blood clotting. He has no prior heart condition. A blood clot traveled to his brain causing a Posterior Cerebral Artery Stroke. He has been in ICU since 04/22/2021.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1258629-1	Blood clots in the left leg and in the lungs.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1259560-1	ultrasound the left lower extremity shows extensive occlusive lower extremity DVT involving multiple veins. Labs otherwise unremarkable, patient appears well, denies chest pain or shortness of breath. He had the J+J vaccine less than 3 weeks ago.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1259704-1	Went to er on April 11 th for cough and shortness of breathe they said nothing was wrong and sent me home. Did a virtual apt on 21st because my leg hurt and shortness of breathe. They told me to go see doctor in person . Went to Er April 23rd and was diagnosed with a massive blood clot in my lower right leg.they prescribed xarelto.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1259807-1	I woke up from almost like a pop in my brain and fluid started pouring down my throat and I started choking. I woke up and was able to spit some of it out. It was clear and also tinged with blood. It stopped once I stood up. I immediately thought my brain was leaking. I told my daughter about this the next day. My balance was off and I had a weird flapping in my head for the next few days. As well as a headache. A few days after I heard about the blood clots. I am not 100% sure if it was from the shot but it was literally hours from when I received it. It was very scary.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1260272-1	Soreness & lump in left lower thigh size of tic tac Swollen and sore left foot Diagnosed with ultrasound as superficial blood clot Heat applied and aspirin Still swollen
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1260342-1	Patient came in this evening 4/26 after emergency room visit for blood clot in leg/shin. She received Janssen IMZ 4/10. Her symptoms of painful leg began several days after her IMZ and continued to worsen with her leg shin swollen, painful, bruised with a knot until she sought medical help in the ER. Ultrasound was done and ruled out deep vein thrombosis, determined to be peripheral blood clot. She was sent home on meloxicam for pain and inflammation.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1260781-1	posterior tibial vein non-occlusive thrombus. RLE pain starting 4/23, diagnosed 4/26

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261192-1	BLOOD CLOT FROM NOSE; TEMPORARILY UNABLE TO WALK, RICKETY KNEES; BURNING LEG PAIN; JOINT PAIN/ SHOULDER PAIN; SHOULDER FEEL TIGHT; This spontaneous report received from a patient concerned a 30 year old male. The patient's height, and weight were not reported. The patient's concurrent conditions included cigarette smoker (occasional pot; rarely). The patient's grandfather died from blot clot under knee (thrombosis). The patient had no history of bloody nose. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808609, expiry: UNKNOWN) dose was not reported, administered on left arm on 28-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On 30-MAR-2021 (two days after vaccination) the patient experienced really bad burning leg pain for few days and his joints in shoulder and knees were affected two days after vaccination and he could not walk (temporarily unable to walk). The patient also experienced rickety knees. On the same day, the patient blew out a huge blood clot from his nose. On 2021, the patient's shoulder felt tight. At the time of this report, the patient's knees and shoulders were still feeling tight like there was hardly any cartilage. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from blood clot from nose, and temporarily unable to walk, rickety knees, was recovering from joint pain/ shoulder pain, and the outcome of burning leg pain and shoulder feel tight was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-Covid-19 vaccine ad26.cov2.s-blood clot from nose. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261249-1	"BLOOD CLOT; LEG CRAMPS; ACHE; HEADACHE; FEVER; This spontaneous report received from a patient concerned a 50 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included overweight, high blood pressure, and high cholesterol. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 207A21A expiry: unknown) dose was not reported, 1 total administered on 06-APR-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. On 07-APR-2021, the patient experienced a mild fever. On 08-APR-2021, the patient was achy and had a mild headache. The headache continued for several day and became pretty bad for about two days. On an unspecified date in Apr-2021 the patient was treated with Tylenol for headache. On 12-APR-2021, the patient experienced leg cramps in her right leg. On 15-APR-2021, the patient had an ultrasound and a blood clot was found in her right leg. The patient started on Apixaban (Eliquis). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from headache on an unknown date in APR-2021, and the outcome of blood clot, ache, leg cramps and fever was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This 50-year-old female was found to have blood clot in her right leg 9 days after receiving Janssen COVID-19 vaccine for the prevention of symptomatic SARS-CoV-2 virus infection. There was reported mild fever, headache, and ""aches"" 1 to 2 days after vaccination that was treated with Tylenol. The symptom reported was leg cramps on right leg 5 days after vaccination, and ultrasound showed blood clot in her right leg 3 days after the symptom of cramps was noted. No other laboratory/diagnostic test reported. The patient was treated with apixaban. The outcome of blood clot in leg was not reported. Based on the information that is available, the event is assessed as plausible with the causal association to immunization, per causality classification of adverse events following immunization based on a lack of a definitive plausible biological mechanism. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information is requested"
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261282-1	LARGE BLOOD CLOT; SWOLLEN BALL ON SIDE OF KNEE; This spontaneous report received from a female patient of unspecified age reporting on self. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry: Unknown) dose was not reported, 1 total administered on 05-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced large blood clot and swollen ball on side of knee. The swollen ball on side of knee was going on for a couple days now. The action taken with covid-19 vaccine ad26.cov2.s was not reported. The patient had not recovered from large blood clot, and swollen ball on side of knee. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a report of a female patient, unspecified age, unknown ethnicity, who experienced a large blood clot and swelling on the side of the knee (laterality not reported) on an unspecified number of days after receiving the cobid-19 vaccine ad26.cov.2. The information provided precludes a meaningful medical assessment. Additional information requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261284-1	BLOOD CLOT IN FOOT; This spontaneous report was received from a patient via a company representative and concerned a male patient of unspecified age. The patient's weight, height, and medical history were not reported. The patient received COVID-19 vaccine Ad26.COV2.S (suspension for injection, route of admin not reported) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, two days after vaccination, the patient experienced a blood clot in his foot. The action taken with COVID-19 vaccine Ad26.COV2.S was not applicable. The outcome of blood clot in foot was not reported. This report was serious (other medically important condition).; Sender's Comments: V0: This male of unspecified age and ethnicity reported to have experience blood clot in foot 2 days after receiving Janssen COVID-19 vaccine for the prevention of symptomatic SARS-CoV-2 virus infection. No other information was reported. Based on the information that is available, the event is assessed as plausible with the causal association to immunization, per WHO causality classification of adverse events following immunization based on a lack of a definitive plausible biological mechanism. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information is requested
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261322-1	CLOT BLOOD; ACUTE ISCHEMIC STROKE IN LEFT FRONTAL LOBE; PARTIAL LOSS OF USE IN RIGHT LEG; MUSCLE WEAKNESS; This spontaneous report received from a patient concerned a 46 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included amoxicillin and flexeril allergy, and migraine. The patient experienced drug allergy when treated with cyclobenzaprine hydrochloride. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 042A21A, and batch number: 042A21A expiry: UNKNOWN) dose was not reported, administered on 08-APR-2021 11:30 on left arm for prophylactic vaccination. Concomitant medications included ascorbic acid, ascorbic acid/ergocalciferol/folic acid/nicotinamide/panthenol/retinol/riboflavin/thiamine hydrochloride, ergocalciferol, iron, and withania somnifera. Approximately three days after the vaccine, on 11-APR-2021 08:20, the patient experienced clot blood. Which caused an acute ischemic stroke in left frontal lobe resulting in partial loss of use in right leg and muscle weakness. The patient had a blood test, CT scan, Carotid artery ultrasound, diagnostic ultrasound, electrocardiogram (EKG) and an MRI, results are unknown. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from clot blood, acute ischemic stroke in left frontal lobe, muscle weakness, and partial loss of use in right leg. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This is a report of a 46 year old, female patient who experienced a blood clot that caused an acute ischemic stroke in the left fontal lobe 3 days after receiving the covid-19 vaccine ad26.cov.2. Patient's height and weight were not reported. Patient has migraine and allergies to amoxicillin, flexeril and cyclobenzapine hydrochloride. Concomitant medications included ascorbic acid, ascorbic acid/ ergocalciferol/folic acid/ nicotinamide/ panthenol/retinol/ riboflavin/ thiamine hydrochloride, ergocalciferol, iron, and withania somnifera. Smoking history, drug abuse and alcohol intake were not reported. Three (3) days after receiving the vaccine, patient was noted to have a blood clot (unspecified area) that caused an acute ischemic stroke in the left frontal lobe, resulting in muscle weakness and partial loss of use of the right leg. Patient had blood tests done as well as CT Scan, carotid artery ultrasound, electrocardiogram and MRI but the results were not reported. The information provided precludes a meaningful medical assessment. Additional information requested.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261329-1	THROMBOSIS; Dizziness; Pass Out; Weakness in the all side of the body; This spontaneous report received from a patient. The patient's height, and weight were not reported. The patient's concurrent conditions included sea food allergy, and other pre-existing medical conditions included the patient did not had any other illness at the time of vaccination and to one month prior and chronic or long standing health condition. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose was not reported, administered on 02-APR-2021 11:25 for prophylactic vaccination. The batch number was not reported. As per procedure, no follow-up will be requested for this case. Concomitant medications included metformin for drug used for unknown indication. On an unspecified date, the subject experienced thrombosis, dizziness, pass out, and weakness in the all side of the body, and was hospitalized (date unspecified). Laboratory data (dates unspecified) included: X-ray (NR: not provided) heart, head and lungs. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from thrombosis, dizziness, pass out, and weakness in the all side of the body. This report was serious (Hospitalization Caused / Prolonged, and Life Threatening).; Sender's Comments: V0: This is a report of a patient, who was noted to have thrombosis on unspecified time after receiving the covid-19 vaccine ad26.cov.2. Patient's height and weight were not reported. Patient medical history was not reported. Concomitant medications included metformin for unknown indication. Patient is allergic to seafoods. On an unspecified date, the subject experienced dizziness and passed out. She also had weakness on the left side of the body. Patient was noted to have thrombosis. Patient was hospitalized, where laboratory and diagnostic tests were done. Results were not reported. The information provided precludes a meaningful medical assessment. Additional information requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261384-1	"SWOLLEN NECK; SWOLLEN RIGHT ARM; 6 BLOOD CLOTS; SICK; This spontaneous report received from a patient concerned a male of unspecified age. The patient's height, and weight were not reported. The patient's past medical history included surgery. There was no history or family history of blood clots. Prior to vaccination, the patient had several physicals performed due to other health concerns and CAT-scans and MRIs were performed that showed his body was totally without any blood clots. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic indication. Batch number was not reported and has been requested. No concomitant medications were reported. After getting the vaccine patient was sick for 3 days. On an unspecified date, the patient woke with a with swollen neck, swollen right arm that was twice its normal size. The patient went to the hospital and they were concerned it was a blood clot further so he was then sent to hospital where they performed an ultrasound which revealed 6 blood clots. The patient was hospitalized and underwent surgery on his arm to ""open up the vein that was causing the swelling"". The patient reported he was scheduled for several more surgeries of this kind. Corrective treatment included intravenous blood thinners. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from swollen neck, swollen right arm, 6 blood clots, and sick. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0: This male patient of unspecified age noted swelling of righ arm and neck that the ultrasound showed 6 blood clots after an unspecified duration from receiving Janssen COVID-19 vaccine for the prevention of symptomatic SARS-CoV-2 virus infection. The patient was hospitalized and underwent corrective surgery and was started on unspecified blood thinners. No other details was reported. The information available precludes a complete and meaningful assessment. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information was requested."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261867-1	Blood clots, heart attack, died, cpr 5 times survived Blood is very thick blood thinners unable to thin
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262401-1	Throbbing pain at injection site and entire arm. Unable to move up without assistance. Pain grew more and more intense. Went to urgent care on unspecified date. NP examined me and discovered a lump in my arm. Both the lump and the vaccination site hurt when she pressed on them. She prescribed presdone and gave me a work order for a vascular ultrasound to be performed on my left arm. I went on unspecified date for the scan. It was discovered that a DVT blood clot had formed in my brachial vein in the left arm. I was then prescribed eliquis, 20 mg a day for 7 days and told not to take the predesone. NP called my pcp, his office immediately called me and asked me to come in as they are concerned about the high dosage of the eliquis.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262422-1	RESIDENT COMPAINED OF PAIN TO BILATERAL THIGHS AND LOWER BACK SENT TO ER. NOTIFIED BY ER, PATIENT HAS BLOOD CLOTS IN ABDOMIN
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262540-1	blood clot in my left eye, swelling of the eye and under the eye. headaches at the base of my skull so bad nothing helped. Joint pain so bad I couldn't walk for 2 days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262556-1	Clotting was present
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262566-1	Blood clot in my left forearm. I first noticed it on 4/20 around 7pm. There was a bump (the size of a dime) and small bruise. Overnight the bump got a bit smaller and the bruise expanded. Over the next several days the bump kept diminishing. The bruise remained about the same size and kept getting darker. It?s been one week now and the bump is very tiny (size of a small pea), and the bruise is incredibly dark, but showing more of a healing process color.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1262688-1	Patient was driving on 3/24 when her vision became blurry in her left eye. She saw her eye doctor on 4/21 and was sent to a specialist. On 4/23 she was diagnosed with a blood clot in her left eye. She is to begin eye injections for treatment.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263011-1	Fever over 105, chills, cough, soar throat. aches and pain for 2 days. Led patient to go to the ER on monday 4/12. MD found blood clot at that point. No treatment was given patient was told to follow up with her PCP. Did not go to PCP as yet.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263180-1	Increase short of breadth, chest pain, & blood clog
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263254-1	Blood clot in left calf muscle

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263472-1	patient had a blood clot 2 days after receiving the J & J vaccine; she was hospitalized and released. We just learned of this when the patient was in to pick up a Rx--we are unsure if this was already recorded by the hospital that treated her--we were advised by our market manager to submit just in case they didn't submit at the hospital.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263497-1	Blood clot, L arm (where pic line was located)
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1263525-1	"Left leg pain started about 4/16/21. Reports evaluated by nurse practitioner at the medical office of primary care provider on 4/23/21. Ultrasound ordered a hospital and diagnosed with a blood clot in the left leg on 4/23/2021. Placed on a ""blood thinner"" to take orally BID for one month then once daily for undetermined duration. States MD is not considering the vaccination as the reason for the blood clot because of the duration of time between when the vaccine was given and the onset of the blood clot. When asked, pt reports a platelet count was done."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1264000-1	My sister suffered a massive stroke with blood clots. Totally paralyzed on right side and limited speech. Unable to function on her own, receiving care at a Nursing Home
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1264082-1	The pt said she developed a blood clot on her posterior hand. I asked her if it was a bruise on the back of her hand and she said it was a blood clot. She said she slapped the bumb/clot and it subsequently went down/away.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1264375-1	Headache, leg swelling Sent to ED for evaluation blood clot in leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1265163-1	Blood clot in each lung, chest pain, labored breathing, leg pain and upper back pain.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1265873-1	STROKE; BLOOD CLOT IN LEG; This spontaneous report received from a consumer concerned a 51 year old male. The patient's height, and weight were not reported. The patient's pre-existing medical conditions included the patient was healthy with no health issues. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. It was reported that on an unspecified date in 2021, the patient started with symptoms after 13 days of vaccination. The patient woke with leg pain That morning, he got in the vehicle to go to work and pulled over due to heaviness in his left arm and slurred speech. An ambulance was called and they took him to the emergency room. From there he was life flighted to another hospital and was continued with laboratory tests. Laboratory data included: doppler ultrasound (NR: not provided) not reported, MRI (magnetic resonance imaging) (NR: not provided) not reported, scan (NR: not provided) not reported and other tests. It was due to the blood clot in his leg that caused him to suffer a stroke. He spent days in the hospital and had to wear a heart monitor for two weeks. He was on blood thinners until further notice to prevent this from happening again. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clot in leg and stroke was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: This case concerns a 51 year-old male with no prior medical issues who was hospitalized for a stroke and blood clot in his leg 13 days after receiving the Janssen Covid-19 vaccine. Concomitant medications, family history, and social history were not reported. Thirteen days post-vaccination, the patient awoke with leg pain. While driving to work, he experienced heaviness in his left arm and slurred speech and pulled his car over. An ambulance took him to a local hospital and from there, he was air-lifted to another hospital where doppler ultrasound, MRI, and other unspecified tests were performed. He spent an unspecified number of days in the hospital and wore a heart monitor for 2 weeks. He was treated with blood thinners. Per the reporter (patient's daughter), the blood clot caused the stroke. Outcome was not reported. Based on the available information, the relationship with Janssen Covid-19 vaccine is considered indeterminant. More information (e.g. platelet count, D-dimer, fibrinogen, anti-platelet factor 4 antibodies, medical records) is being sought.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1266095-1	As soon as I took the vaccine shot I had an immediate headache, after taking the vaccine I took Tylenol and slept in the rest of the day. The next day I woke up and my arm was red and swollen. I was running fever, still had the headache and now blurry vision. My entire body was hurting. My left side of my body felt numb and tingling. The next two days the pain became worse. I did not really have an appetite. When I did eat I could barely hold my food down. On Wednesday morning I could barely get out of bed, I also felt extreme pain in my lower legs, stabbing pain in stomach and right lower side. I also had ringing in my ears, more in the right ear. I then notice a bruise on my left leg, that it was not originally there, and was also tender to touch. I went to the emergency room on the same day and I was admitted. I was in there for 4 days, they tested my leg and stated it was not a blood clot, although it look and felt like one. They discharge me and advise me that it was a migraine that was causing all my pain. This was on Sunday, I had a follow up with my PCP on the Friday, however to being in so much pain and shortness of breath, and arm pain I was advise to go to the ER on the Thursday and was admitted again. I was told in the ER I had a blood clot in my right arm and it was most likely from the IV from the previous visit. My blood pressure and heart rate was so high that they put me on medication. No matter the pain medication they gave me, it would not completely take the pain away. They discharge me again on Monday and after me begging them to get run any and all test, they said they found nothing. I'm still in the same amount of pain and same symptoms, still having trouble keeping my food down. The only thing that changed is my left arm is not as sore, however I still have a huge knot and its still warm to touch. I don't know what else to do that is why im contacting the CDC for help. I'm in a extreme amount of pain and pounding headaches. Please help
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1266356-1	For the past 2 1/2 - 3 weeks I have had a pain in my left leg. Today it was extremely painful so I went into the dr. office and after an examination and ultra sound it was determined that I have a blood clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1266422-1	Patient stated that her doctor says she has blood clots and believe it could be due to the vaccine.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1266689-1	BLOOD CLOTS IN BOTH LUNGS. SHORTNESS OF BREATH. UPPER BACK PAIN. WEAKNESS. SWEATING AND COLD. NO COLOR IN FACE -- GREY. LOSS OF EYE SIGHT -- STARTED WITH BLURRED VISION.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1267199-1	"patient developed severe chest pain, ambulance was called and patient was taken to ER. Patient was transferred to higher level facility and was told he had a heart attack and that it was caused by a ""blood clot"""
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1267264-1	I have a major blood clot in my left leg. I had two ultrasounds and now starting blood thinners.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1267579-1	4/19/2021: Woke up in the middle of the night with an achy arm. Had this for a week. Went to chiropractor as she thought it was neck trouble. Chiropractor states arm is swollen and see MD. MD sent patient for ultrasound and blood clot confirmed in right arm pit. Patient started on eliquis.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1268089-1	Two blood clots in right leg below the knee, calf pain, swelling of lower leg and foot, painful to walk
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1268363-1	Patient received Johnson Johnson COVID vaccination on March 5th (Its not listed as an option for me to choose - Hence Janssen). Two weeks later patient and wife described symptoms of significant abdominal pain and headache, which improved after 24 hours, since then, abdominal pain on and off with worsening, nausea vomiting and diarrhea. Presented to Hospital with sepsis, negative evaluation, possibly GI etiology, given symptoms. MRI of the abdomen shows left portal vein thrombosis/which is an unusual site. Patient has no liver cirrhosis or metastatic cancer. Is being investigated for idiopathic or acquired thrombophilia
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1268669-1	Ended up with a blood clot in left leg, not a dvt clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1268686-1	Flu-like symptoms chills fever up and down shortness of breath itchy skin dry cough shaky hands restless urinating frequently urination blood clotting in calves

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1268728-1	HEADACHE/PAIN IN HEAD; HEAD FELT FLUIDY,FLOATING,FOGGY,/FREAKING OUT; FLUCTUATING LUCIDITY/FELT LUCID; PAIN IN NECK; WOKE UP AND STUMBLED; RED BLOOD (AND CLOT) WHEN BLEW NOSE; RED BLOOD (AND CLOT) WHEN BLEW NOSE; DREAMING DIZZY/DIZZY; FEELS JITTERY/NOT FEELING WELL; This spontaneous report received from a 49 year old female patient reporting on self. The patient's height, and weight were not reported. The patient's concurrent conditions included seasonal allergies. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808982,expiry: UNKNOWN) dose was not reported,1 total administered on 12-APR-2021 09:00 in right arm for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, in the evening she had headache. On 12-APR-2021 the patient was dizzy and not feeling well at night she woke up in sleep feeling dizzy as she was dreaming dizzy, got up from bed and stumbled. She told her husband not feeling well but he tried to brush it aside. up from bed and stumbled. She took Advil. The doctor on-call told her to calm down and to take 2 Advil and went to sleep and if it did not went away to ER. The consumer did so. On 13-APR-2021, (Tuesday morning) the consumer blew her nose and noticed red blood and clots (coming from further up higher than sinuses). She was concerned and talked to people (friends). On 14-APR-2021, around 3:00 Wednesday morning she was freaking out she called her doctor and was not available she was referred to another physician. The patient was lucid, head felt fluidy, floating and dizzy. She had pain in the head and the neck. On 15-APR-2021, Thursday morning the physician called her back and said to stop taking Advil and went to ER. Her head was heavy and fog, everything was slow and lucid. In the hospital as soon as they gave her saline IV fluid, the fog was half lifted. Laboratory data (dates unspecified) included they also did MRI and blood work which were normal. They discharged her the same day and suggested she could take Tylenol which she did Thursday night. On 16-APR-2021, Friday she felt jittery but good and will buy some Gatorade for hydration and patient also stated that both dizzy and heavy head was reduced to faint (coming down from 10 to 3.5 on a scale). The consumer suggested that the dose of the vaccine may be too much for certain individuals and suggested to reduce the dose to like half the dose or to give it in 2 shots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from red blood (and clot) when blew nose, and stumbling on 15-APR-2021, was recovering from dizziness, fluctuating lucidity, head felt fluidy, floating, foggy,/freaking out, pain in neck, and feels jittery/not feeling well, and the outcome of headache/pain in head was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: 20210436432- covid-19 vaccine ad26.cov2.s-head felt fluidy, floating, foggy,/freaking out, red blood (and clot) when blew nose. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1269209-1	On Jan. 26, 2021, I had an accident at home an suffered broken ankle, tibia and fibula broken from tarsus. I was rushed to hospital and had ORIF surgery. I later found out the surgeons failed to order blood thinners and about a month later, I was rushed to the hospital with 5 blood clots, one in each lobe or each lung and one forming in the calf muscle on the left leg. I have been taking Eliquis since late February and will be on it through May. I have had an echocardiogram which was normal but the doppler I had on my legs may have indicated the problem is still on-going. I don't know yet. I did contract JCCT to ask if I should get checked out due to recent clotting incidents and was told I did not. I decided I should report it anyway and my clotting situation probably has nothing to do with vaccine but I just want to make sure.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1269940-1	"BLOOD CLOT IN ARM; FEELING SICK; This spontaneous report received from a patient via a company representative concerned a 47 year old female. The patient's weight, height, and medical history were not reported. The patient was not pregnant at the time of report The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total, administered on 05-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date in APR-2021, the patient experienced burning sensation in arm, tenderness of arm, and pronounced veins on arm; also she was feeling sick. She saw the healthcare professional (HCP) after these symptoms. She was told about a blood clot (tenderness / burning sensation/ pronounced veins) in the arm. HCP recommended her taking aspirin and applying heat to the arm. Treatment medications (dates unspecified) included: acetylsalicylic acid. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot in arm, and the outcome of feeling sick was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: This 47-year-old female patient reported burning sensation and tenderness in arm, pronounced veins in arm, and feeling sick after an unspecified duration from receiving COVID-19 VACCINE AD26.COV2.S for the prevention of symptomatic SARS-CoV-2 virus infection. HCP was consulted and was ""told"" about a blood clot and prescribed aspirin with advice to apply her to the arm. No other details reported. The information available precludes a complete and meaningful assessment. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information was requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1269951-1	"BLOOD CLOT IN UPPER THIGH; NO PULSE FROM KNEE DOWN ON RIGHT LEG; RINGING AND BUZZING IN EARS; PURPLE TOE; COULD NOT MOVE; FEELING CRAPPY; TIRED; NAUSEOUS; This spontaneous report received from a patient concerned a 58 year old male. The patient's height, and weight were not reported. The patient's concurrent conditions included Barrett's esophagus, and controlled high blood pressure. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular batch number: 1805031, and expiry: unknown) dose was not reported, 1 total administered in left arm on 24-MAR-2021 for prophylactic vaccination. Concomitant medications included Nexium (esomeprazole sodium) for Barrett's esophagus, and lisinopril for high blood pressure. On 25-MAR-2021, the day after the vaccination, patient felt crappy and could not move, had tiredness and nauseous. After 10 to 12 days of vaccination on an unspecified date in APR-2021, the patient had ringing and buzzing in ears, pain in the bottom of feet on walking, and purple discoloration to front of the right toe. The patient had visited health care professional (HCP). The patient had undergone some tests and HCP identified no pulse from the knee down on his right leg and a blood clot in upper thigh. The patient had experienced pins and needles and buzzing on his leg. He has also has a ringing and a buzzing in his ears and the bottom of his feet hurt when he walked on them. The patient was scheduled for an appointment with cardiovascular surgeon on 28-APR-2021, prior to the visit the patient has to tested negative for COVID. The patient went to take test on 23-APR-2021 and results not yet received. Laboratory data included: Peripheral pulse absent (NR: not provided) no pulse from the knee down on right leg. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the feeling crappy, tired, nauseous, no pulse from knee down on right leg, blood clot in upper thigh, ringing and buzzing in ears and could not move was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to.This spontaneous report received from a patient concerned a 58 year old male.; Sender's Comments: V0: This 58-year-old male patient was found to have blood clot in upper thigh with no pulse from the knee down 10 to 12 days after receiving COVID-19 VACCINE AD26.COV2.S for the prevention of symptomatic SARS-CoV-2 virus infection. Concurrent conditions include Barret's esophagus and ""controlled"" high blood pressure; concomitant medications includes Nexium and lisinopril. The symptoms reported were ringing and buzzing in ears, pain in the bottom of feet on walking, pins and needles and buzzing on his leg, and purple discoloration to front of the right toe that prompted consult with HCP, who advised the patient that there was no pulse from the knee down on his right leg and a blood clot in upper thigh. Prior COVID test was negative. No other details reported. The information available precludes a complete and meaningful assessment. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information was requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"BLOOD CLOT IN LEG; SHORTNESS OF BREATH; HEART RATE INCREASED; GANGRENE IN TOE; RIGHT LEG SWOLLEN; RIGHT FOOT SWOLLEN; CHILLS; RUNNY NOSE; FEELING HOT; This spontaneous report received from a patient concerned a 50 year old male. The patient's height and weight were not reported. The patient's concurrent conditions included high blood pressure and diabetes. The patient received COVID-19 vaccine Ad26.CoV2.S (suspension for injection, route of administration not reported, batch number 1802070, expiry date unknown), dose not reported, 1 total administered on 19-APR-2021 for prophylactic vaccination on left arm. Concomitant medications included amlodipine besilate for drug used for unknown indication, empagliflozin for drug used for unknown indication, fenofibrate for drug used for unknown indication, insulin aspart for drug used for unknown indication, insulin glargine for drug used for unknown indication, losartan for drug used for unknown indication, and pioglitazone hydrochloride for drug used for unknown indication. On an unspecified date (reported as 22-MAR-2021; 28 days prior to vaccination), the patient experienced shortness of breath, heart rate increased, runny nose, chills and was hospitalized (date unspecified). He reported that on this past Saturday, 17-APR-2021 (2 days prior to vaccination), he went to the emergency room. He reported that his toenails were coming off, the skin on his toes was coming off, and he had swelling in his right foot and up his leg. He says there was no circulation going to his toe and he was diagnosed with having a blood clot in his right leg. He says he had surgery to open up the arteries in his leg, and also reported gangrene in his toe. The subject was discharged on 21-APR-2021. On 23-APR-2021, the patient experienced feeling hot. He also reported that he may need to have his right pinky (fifth) toe amputated because it has gangrene. He reported that he is going to go back to the hospital, to which he would seek medical attention if he was having issues. Treatment medications (dates unspecified) included paracetamol. When called back due to his vaccination dates being after event start dates, the patient said that his memory is not too good, so he 'had to take his time to remember the correct dates'. The action taken with COVID-19 vaccine Ad26.CoV2.S was not applicable. The patient was recovering from blood clot in leg, and had not recovered from runny nose, shortness of breath, heart rate increased, chills, feeling hot, gangrene in toe, right leg swollen, and right foot swollen. This report was serious (caused hospitalization).; Sender's Comments: V0: This 50-year-old hypertensive and diabetic male patient reported that he was feeling hot and may need to have his fifth toe amputated because it has gangrene 4 days after receiving COVID-19 VACCINE AD26.COV2.S for the prevention of symptomatic SARS-CoV-2 virus infection and 2 days after discharge form the hospital. The patient was hospitalized for shortness of breath, heart rate increased, runny nose, and chills 28 days prior to vaccination. The patient went to the emergency room 2 days prior to vaccination because his toenails/skin on his toes were coming off, swelling in his right foot and up his leg, gangrene in his toe; he reported that there was no circulation going to his toe and was diagnosed with having a blood clot in his right leg; the patient reported he had surgery to ""open up"" the arteries in his leg. Four days after vaccination, the patient felt hot and reported reported that he may need to have his right pinky (fifth) toe amputated because it has gangrene. The patient is taking paracetamol as treatment. No other details was reported. Based on the lack of temporal association, the causality is considered not related for the events blood clot in leg, shortness of breath, heart rate increased, right leg swelling, right foot swelling, chills, and runny nose. The information available regarding event gangrent of fifth toe and feeling hot precludes a complete and meaningful assessment. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded. Additional information was requested."
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1269952-1	
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1270153-1	Left leg start swelling And hurting
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1270278-1	Next day full body red rash, 103-104 degree fever, deep body aches, skin sensitivity, joint pain, headache, brain fog,lasted 4 days, One week later period started with large chicken-egg sized blood clots, heavy flow lasted 2 days longer than typical period for me. Second cycle after vaccine started 4.22.21 with same very large clots, intense cramping and bloating
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1270282-1	She got the vaccine, when they put the needle in she said it was far in. The pain was bad, but she could deal with it. Then 5 days later something hit her and hit her bad. She went to the ER and they gave her Bentyl, diarrhea medicine as she was vomiting, not eating anything, and they gave her some anti-vomiting medicine. They sent her home and said she would feel better. The diarrhea and vomiting stopped but she then started with a rash. it's all over her hands, arms, back, itching, and taking Claritin for it. it won't go away. Her symptoms continued to worsen, and woke up one morning and had so much pain that she could not move or breath and her husband called an ambulance and they took her hospital and diagnosed with a blood clot. She was in the hospital for 3 days and now discharged on Eliquis. She has bruises all over her coming out and very upset that she is still feeling bad and doesn't know how the rash is going to go away. She was also prescribed Oxycodone for pain and also Tessalon Perls to take for coughing.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1270837-1	a few days after the shot i was having a hard time breathing, made appointment with Pulmanologist he ordered a ct scan but no ultra sound. ct scan showed no clots in the lungs at that time. Woke up a day or two later with severe back and leg pain was transported to the emergency room. I was then transported to facillity because i could not move my left leg and was in severe pain. During my stay i coughed up a blood and requested the doctor to do blood work. The blood work showed high levels of clots. I was not taken to the hospital or treatment for the clots at that time. When i left the facility i saw my Doctor she ordered immediatly another ct scan of the chest and an ultra sound
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1271703-1	Blood clot in right wrist area: On the evening of 3/17/21, I felt sudden pain in the palm of my right hand, just above the wrist. Then the vein just below my wrist swelled up and turned blue. The next morning when I woke, the vein was no longer swollen, but the area below my wrist had a large bruise, which lasted several days. I contacted my physician that day (3/18) who said just to monitor/observe and come in if continuing issues (which there were not).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1271930-1	starting have shortness of breath/difficulty breathing and back and chest pain. light headed, pain unbearable. Saturday April 23rd Went to er was treated for kidney stones but they could only find one small one, Was sent home with pain medication. Shortness of breath persisted as did bloody cough and pain in chest and all over. Went back to the er on 4/28 as pain was unbearable (chest pain) once given a cat scan and diagnosed with blood clots in my lungs and then an ultra sound and found a blood clot in my leg (just above knee). It is thought that I was mis-diagnosed on the original trip to the er with kidney stones. I have been put on blood thinners and pain medication and will be following up with my doctor.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1272388-1	is at the er with blood clots in right leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1272492-1	Patient experienced a severe heart attack and was diagnosed with a blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1272700-1	"Patient received the Janssen COVID 19 Vaccine on 04/07/2021. No physical adverse assessment findings observed during the post-vaccine administration observation period. During the observation period, the patient verbalized that she was ""feeling fine and had taken an Alive(medication)"". Patient was later seen, after the observation period and engaged in a discussion stating that she ""continued to feel okay but felt a little queasy (confirmed as nauseated)"" and denied additional symptoms requiring immediate medical attention. On 04/29/2021, patient informed the clinical nursing team at the administering facility that she experienced adverse side effects 13 days after receiving the vaccine and was hospitalized on 04/20/2021 - 04/24/2021. Patients adverse side effects related to the vaccine are self-reported as blood clotting. Medical documentation not provided. The vaccine administrating facility's clinical staff is not unable to be verified by and confirm are deny adverse medical findings at the time of this dictation. Please contact patient for the selected healthcare provider where care was rendered for detail specific related adverse medical event."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1274347-1	On the 28th of April, 18 days from the shot, I went for a walk and came home around 6:30 p.m. to the outside of my right calf bright red and hot. I went to the ER where an ultrasound was performed and they found a DVT blood clot. I am now on an anticoagulant, Xarelto and have an appointment with my doctor on the 5th to schedule more imaging and blood work.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1274563-1	On April 16, 2021 (Day 13 after Vaccine) I was diagnosed with acute deep vein thrombosis (DVT) of my femoral vein in my left leg by Hospital. In other words, I was diagnosed with a blood clot in my upper left leg. Pain in both my legs had started on April 13, 2021 (Day 10 after Vaccine). On April 16, I had a virtual appointment with my doctor who prescribed an ultrasound to rule out DVT. I went to an ultrasound appointment at Radiology at 3:30pm where a blood clot was found in my left leg. I was admitted into the emergency room at Hospital and prescribed medication and follow ups from there...
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1274581-1	Patient states he had no soreness to his left arm. Patient states he got a blood clot. He went to the doctor and his doctor told him that the blood clot he had was not related to the vaccine.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1275188-1	Swelling in right leg following workout. Became painful as day went on. Swelling, pain continued. After three days, visited doctor.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276388-1	BLOOD CLOTS; SEVERE PAIN; This spontaneous report received from a consumer concerned a patient of unspecified sex and age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots and severe pain, and was hospitalized (date unspecified). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clots and severe pain was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: This is a report of a patient, unspecified age, unspecified gender, unknown ethnicity, who was hospitalized for pain and blood clots on an unspecified number of days after receiving the covid-19 vaccine ad26.cov.2.s. The information provided precludes a meaningful medical assessment. Additional information requested.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276394-1	LOSS OF BLOOD; VOMITING; BACK PAIN; ABDOMINAL PAIN; FEELING WEAK; HEADACHE; BLOOD CLOTTING; COULD NOT MOVE; ONLY COULD DO 5 MINUTES OF WORK; This spontaneous report received from a patient concerned a 45 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included fibroids. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1802070CCHD,expiry: UNKNOWN) dose was not reported,1 total administered on 20-MAR-2021 for prophylactic vaccination to left arm. No concomitant medications were reported. On 22-MAR-2021, the patient experienced blood clotting which lasted for three weeks and resolved as per physician on same day the patient experienced headache which got resolved on 22-Apr-2021. On 18-APR-2021, the patient experienced loss of blood, vomiting, back pain, abdominal pain and patient felt weak. On an unspecified date in 2021, the patient went to hospital, it was also reported that patient was not able to move and could do only 5 minutes of work. Patient got confused between clotting with menstruation. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from blood clotting on 12-APR-2021, and headache on 22-MAR-2021, was recovering from loss of blood, feeling weak, vomiting, back pain, and abdominal pain, and the outcome of could not move and only could do 5 minutes of work was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: In narrative of source document vaccination date is 20-Mar-2021 and in the structured field of source document vaccination date is 20-Apr-2021. So as per conservative approach, date given in narrative 20-Mar-2021 is taken as vaccination date. V0 20210440392-COVID-19 VACCINE AD26.COV2.S-Blood clotting and loss of blood. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276398-1	POSSIBLE BLOOD CLOT; INCREASED HEART RATE; CHEST TIGHTNESS; HIGH BLOOD PRESSURE; NECK PAIN; ARM PAIN; HEAD PAIN; BODY SHAKING; CHILLS; HIP JOINT PAIN; BACK PAIN; This spontaneous report received from a patient concerned a 48 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included high blood pressure, osteoarthritis, fibromyalgia, non smoker, and non alcohol user. The patient experienced drug allergy when treated with diphenhydramine hydrochloride. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 1805022) dose was not reported, 1 total, administered on right arm on 14-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On 14- MAR-2021, around 9 pm the patient experienced body shaking, and chills for which she took Theraflu. On 15-MAR-2021, morning the patient experienced high blood pressure, chest tightness and increased heart rate of 156 BPM for which she went to the ER. She also experienced neck pain, arm pain and head pain. lung and chest examination was normal. Laboratory data included: Blood pressure (NR: not provided) increased, Fibrin D dimer (NR: not provided) 780, and Heart rate (NR: not provided) 156 pm. As the- d dimer test result was high, the doctor said she has a blood clot. She was given a shot of Toradol at the ER. On 17-MAR-2021, Laboratory data included: Fibrin D dimer (NR: not provided) 650. On 30-MAR-2021, She had another episode, she was in a lot of pain. She went back to ER. Laboratory data included: CAT scan (NR: not provided) normal. They sent her home and prescribed her Meloxicam. The patient went to see her Cardiologist after that, She stated that her cardiologist didn't do any blood work or mention anything regarding a blood clot. Treatment medications (dates unspecified) included: ketorolac tromethamine, acetylsalicylic acid, meloxicam, and dextromethorphan. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from high blood pressure, chest tightness, increased heart rate, body shaking, chills, and head and neck pain and arm pain on MAR-2021, had not recovered from hip joint pain, back pain, and the outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:-covid-19 vaccine ad26.cov2.s-possible blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276413-1	COVID-19; BRONCHITIS; This spontaneous report received from a consumer (patient's husband) concerned a 43 year old female. The patient's height, and weight were not reported. The patient was not pregnant at the time of report. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: 042A21A, and expiry: 21-JUN-2021) dose was not reported, administered at Left Deltoid on 09-APR-2021 for prophylactic vaccination. No concomitant medications were reported. On 09-APR-2021, the patient was given a dose of the vaccine and she did not show up. On next day 10-APR-2021, she started shivering very badly and was feverish. On 11-APR-2021, the patient also had a headache. On 12-APR-2021, all symptoms gone. The next day on 13-APR-2021 when she got up she had chest tightness and went to doctor due to fever, chills, headache resumed. Since that day was the clot information in the news and they did testing for clot stuff which was all negative. Also the patient diagnosed with bronchitis. After the 13th she had 'on/off' days of experiencing symptoms. On 16-APR-2021, she tested for covid-19 via 'non-rapid' test and the results of test came on the 17th were positive for covid-19 (diarrhea, vomiting, chest tightness, shivering, chills, headache, feverish and nausea). On 17-APR-2021, she developed diarrhea and vomiting. By the 19-APR-2021, they went to the hospital where she received unspecified fluids and pain pills for the fever and to calm her down. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the bronchitis and covid-19 was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0-20210443606 - Covid-19 vaccine ad26.cov2.s-Covid 19. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276430-1	RARE BLOOD CLOT; LOW PLATELETS; This spontaneous report received from a consumer concerned about 50 years old female. The reporter obtained the information from news/media. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total administered on APR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unknown date in APR-2021 after vaccination, the patient experienced severe headache, abdominal pain, leg pain and shortness of breath. The patient developed a rare blood clot and low platelets on an unspecified date in APR-2021, within two weeks of receiving JANSSEN COVID-19 vaccine. The patient was hospitalized on an unspecified date in APR-2021. On APR-2021, the patient died from blood clot. The reporter was not sure whether the events was related to the vaccination. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of rare blood clot on APR-2021, and had not recovered from low platelets. This report was serious (Death, and Hospitalization Caused / Prolonged).; Sender's Comments: V0: This female patient in her 50s was reported to have developed a rare blood clot and low platelets within two weeks of receiving JANSSEN COVID-19 vaccine. On an unspecified date, symptoms reported were severe headache, abdominal pain, leg pain and shortness of breath. The patient was hospitalized on an unspecified date and subsequently died from blood clot. It was not known if autopsy was performed. No other details reported. The information available precludes a complete and meaningful assessment. However, considering the temporal relationship and recently evolving theories in the literature about COVID infections and vaccinations, potential vaccine contribution cannot be excluded.; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276432-1	BLOOD CLOTTING; This spontaneous report received via traditional media from a patient concerned an adult female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown and expire date: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced blood clotting and was hospitalized (date unspecified). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clotting was not reported. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0: 20210447114-Covid-19 vaccine ad26.cov2.s- BLOOD CLOTTING These events are considered unassessable. The events have a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276433-1	BLOOD CLOTS; This spontaneous report received via social media/news from a patient concerned an adult female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, 1 in total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient had clot blood and was hospitalized. The symptoms appeared were found to be consistent with the six cases reported elsewhere last week. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: -Covid-19 vaccine ad26.cov2.s-Blood clot. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276477-1	DIARRHEA; LETHARGIC; GENERALISED ACHING; HEADACHE; SWELLING AT INJECTION SITE; REDNESS AT INJECTION SITE; BLOOD CLOT FROM THIGH TO CALF; This spontaneous report received from a consumer concerned a 66 year old male. The patient's height, and weight were not reported. The patient's concurrent conditions included paralyzed on left side from a previous injury and not very active. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered 1 total administered on 06-APR-2021 on right arm for prophylactic vaccination. The batch number was not reported and it has been requested. No concomitant medications were reported. On 07-APR-2021, the patient had blow-out diarrhea, lethargic, body aches, headache. On APR-2021, the patient experienced leg pain (started week ago), swelling at injection site and redness at injection site. His left foot was swollen, and called physician to check for blood clot or break. The doctor said that the test showed a blood clot from his thigh all the way down to his calf that was a medical emergency. Reporter thinks there was a possibility that they were already exposed to COVID. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot from thigh to calf, diarrhea, lethargic, generalised aching, headache, swelling at injection site and redness at injection site was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0. 20210450241-COVID-19 VACCINE Ad26.COV2. S -Blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276478-1	BLOOD CLOT; This spontaneous report received from a consumer via news concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration was not reported, batch number: unknown) dose, start therapy date were not reported, 1 total, for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient developed a blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: -Blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1276487-1	"STROKE/HEADACHE; BLOOD CLOT; INJECTION SITE REACTION; PAIN IN ARM; This spontaneous report received from a patient concerned a 79 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805020, expiry: UNKNOWN) dose was not reported, administered to left arm on 11-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On 25-MAR-2021, 2 weeks after receiving the vaccination, the patient stated his injection site on his left arm ""felt horrible"" ""like someone was stabbing him in that joint"" (both events captured as injection site reaction). The patient reported his doctors did not know what was causing this and he could not get rid of it. On an unspecified date the patient experienced headaches with one being described as going from the back of his head to his left temple, he reported not usually having headaches. The patient also experienced severe pain when moving left arm with pain worsening at night when not moving. The patient was unable to reach up to grab things. On 24-APR-2021, the patient was taken to the hospital and was diagnosed as had a stroke. On an unspecified date results of Magnetic Resonance Imaging (MRI) were 'blood clots in the artery that goes up middle of spine and back of neck. The patient stated the results also had ""right side numbness, vertigo, slurred speech"". Results of Computerised Tomogram (CT) (date unspecified) were per the patient ""internal carotid arteries demonstrated calcified plaque, stenosis, arteries demonstrate localized three-four segment of left local artery"". The patient was discharged on an unspecified date and had not seen his regular physician yet. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the stroke, blood clot, injection site reaction, pain in arm and headache was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0 -covid-19 vaccine ad26.cov2.s-Stroke/Blood clot. This case concerns a patient of 79 year old male. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s)."
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1276504-1	TRANSIENT ISCHEMIC ATTACK MORE FREQUENT; POSSIBLE BLOOD CLOT; SHORTNESS OF BREATH; CHEST PAIN; SWELLING IN ONE LEG; PULSATING HEADACHE; This spontaneous report received from a patient via social media concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, 1 in total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, after took vaccine the patient had major health problems which was described as transient ischemic attack which was more frequent and lasted longer, headaches were a pulsating headache, shortness of breath(more than usual),chest pains, swelling in one leg and seems to be blood clots. The patient was enquire about the involvement of people in vaccine study with conditions like hepatitis C, strokes, chronic obstructive pulmonary disease, chronic headaches, liver problems, transient ischemic attack. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the transient ischemic attack more frequent, pulsating headache, shortness of breath, chest pain, swelling in one leg and possible blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-20210452291JANSSEN COVID-19 VACCINE Ad26.COV2.S This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1276515-1	CLOT IN LEGS; ESCALATING PAIN IN LOWER BACK; ESCALATING PAIN IN LEG; This spontaneous report received from a patient via news story concerned a 4 decade (early 40s) male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose was not reported, 1 total, administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. In APR-2021, the patient experienced clot in legs and was hospitalized. The patient was improving and scheduled to leave the hospital in few days. On 16-APR-2021, the Patient experienced escalating pain in lower back and leg. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from clot in legs, and the outcome of escalating pain in lower back and escalating pain in leg was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: - Clot in legs (PT-Thrombosis). This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276518-1	PROFOUND BLEEDING; CLOTTING; SUSPECT DYSFIBRINOGENEMIA; This spontaneous report received from a health care professional via a Regulatory Authority and concerned a 48 year old, female. The patient's height, weight and medical history were not reported. The patient was not pregnant at the time of reporting. The patient had no other illness at the time of vaccination, no chronic or long-standing health conditions, the patient was healthy without history of dysfibrinogenemia. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 1805020, and expiry: UNKNOWN) dose was not reported, administered on 09-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On 22-MAR-2021, the patient experienced profound bleeding, and clotting and was suspected to have dysfibrinogenemia. The patient was currently hospitalized for the events (date unspecified). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from profound bleeding, clotting, and suspected dysfibrinogenemia. This report was serious (Hospitalization Caused / Prolonged, Other Medically Important Condition, and Life Threatening). Sender's Comments: VO:-covid-19 vaccine ad26.cov2.s -profound bleeding, clotting, suspect dysfibrinogenemia. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276526-1	BLOOD CLOT WITH DYING PAIN IN RIGHT SHOULDER; This spontaneous report received from a patient concerned a 72 year old female. The patient's weight was 141 pounds, and height was 164 centimeters. The patient's concurrent conditions included back pain, two cysts on kidney, high blood pressure, anxiety, high cholesterol, non smoker, non alcoholic, and spine issue, and other pre-existing medical conditions included it was unknown that patient had drug abuse or illicit drug usage. she was health healthy and always doing exercises, she lifts 3kg dumbbells and walked and use indoor bicycle for exercise. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029, expiry: UNKNOWN) dose was not reported, frequency one total administered on 15-MAR-2021 for prophylactic vaccination. Concomitant medications included citalopram for anxiety, lorazepam for anxiety, oxycodone for back pain, and spine issue, and acetylsalicylic acid for drug used for unknown indication. Patient received vaccine on March 15, 2021 at a local vaccination center. she woke up with pain and had this for several days before. somebody suggested going to the Emergency room. Patient proceeded to ER on 26-March-2021 and was seen by Doctor. Doctor suspected a blood clot and told the if its blood clot it could moved to lung and cause stroke. The patient reports the following tests were performed X-ray, ultrasound and blood tests. Test results all showed blood clot that the patient describes as deep vein in the right shoulder between shoulder and neck. Patient reported that she was given a blood thinner in her tummy and then a prescription for 30 days of Eliquis 5mg. Patient reported that the pain was killing her and stated she was dying of pain. The pain was in the area of the blood clot (right shoulder). Patient was also seen by a specialist who prescribed Eliquis prescription for 1 year. she was taking baby aspirin which told to continued. she finished her 30 days prescription ,she planning to visit ER as pain had not go away. She has been told not to stop the blood thinner as it is dangerous. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot, and dying of pain s in the area of the blood clot (right shoulder). This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210453704-COVID-19 VACCINE AD26.COV2.S - BLOOD CLOT WITH DYING PAIN IN RIGHT SHOULDER. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276530-1	DEATH; BLOOD CLOT; This spontaneous report received from a consumer news/social media platform concerned a 5 decade old female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient developed a rare blood clot and died within two weeks of getting the Janssen covid vaccine. On an unspecified date, the patient died from unknown cause of death. It was unknown whether autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to.; Sender's Comments: V0-covid-19 vaccine ad26.cov2.s-This case concerns with 5 decade old female. Death, Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276539-1	BLOOD CLOT; This spontaneous report received from a patient concerned an adult female. This report was received from news/social media platform reported by a consumer/other non health care professional. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on 2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date in APR-2021, after vaccination the patient experienced blood clot, and was hospitalized (date unspecified). This report was notified through VAERS on 22-APR-2021. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Hospitalization Caused / Prolonged). This case, from the same reporter is linked to 20210453777.; Sender's Comments: V0: 20210454136-COVID-19 VACCINE AD26.COV2.S - BLOOD CLOT. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276551-1	BLOOD CLOTS; This spontaneous report received from a consumer (source: news report) concerned a female of age under 60. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case No concomitant medications were reported. It was reported from a news story that, on an unspecified date, the patient experienced blood clots which was in investigation by an agency. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210454646.; Sender's Comments: V0:20210454446-covid-19 vaccine ad26.cov2.s- Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276556-1	BLOOD CLOT; This spontaneous report received from a consumer concerned an adult male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot. The reporter stated that as per news report, the agency was investigating this case of blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210454446.; Sender's Comments: V0:20210454646-covid-19 vaccine ad26.cov2.s -blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276566-1	BLOOD CLOT; SEVERE HEADACHE/ FEEL LIKE A CAP ON HER HEAD; This spontaneous report received from a consumer concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot, and severe headache/ felt like a cap on her head. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the severe headache/ feel like a cap on her head and blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: AER 20210455412-Covid-19 vaccine ad26.cov2.s-BLOOD CLOT. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276571-1	THROMBOSIS (RED PATCHES ON HANDS, ARMS, LEGS AND FEET); SEVERE MIGRAINES; SEVERE VACCINATION REACTION; This spontaneous report received from a patient via a company representative concerned a 3 decade old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry date: Unknown) dose, 1 total administered, start therapy date was not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. The patient had a severe vaccination reaction, experienced red patches on hands, arms, legs and feet. The patient had visited the hospital and was diagnosed with thrombosis by the physician. Patient was monitored and later discharged. The patient had reattended hospital for severe migraines, also had contacted centers for disease control and prevention regarding vaccination reaction. The patient was unknown for company representative and had only this information. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the thrombosis (red patches on hands, arms, legs and feet), severe migraines and severe vaccination reaction was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0.20210455657- COVID-19 VACCINE Ad26.COV2. S - Thrombosis, severe migraines , severe vaccination reaction. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276632-1	CLOTS; This spontaneous report received from a physician concerned a male of unspecified age. The patient's weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The physician reported that on an unspecified date, the patient experienced clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of clots was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210456400.; Sender's Comments: V0.20210457363-COVID-19 VACCINE AD26.COV2.S -Thrombosis. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1276633-1	CLOTS; This spontaneous report received from a physician (source: news report) concerned 6 female patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry unknown) dose, start therapy date were not reported, frequency 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. The reporter called in regarding an event of clot experienced by 6 patients in the news on an unspecified date. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of clots was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210456400 and 20210457363.; Sender's Comments: 20210457370-covid-19 vaccine ad26.cov2.s-Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1277853-1	Pain on both legs, Tiredness, Fatigue.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1279696-1	patient reported to the pharmacy that they had to go to ER for blood clot on 04/19/2021
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1280115-1	See below
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1280655-1	First 2 days: fever 101.5, whole left side pain, abdominal pain, headache, lower right leg pain 3rd day: low grade fever 100.5, lower right leg pain, left arm pain 4th day: lower right leg pain increased along with redness, swelling, red lines, and eventually I couldn't apply weight on my right leg. After visiting Urgent Care hospital, blood clots were found in my small veins.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1280744-1	12 hours after injection, experienced normal side effects (e.g., muscle aches, chills, headache, fatigue) Within 1 week of receiving the vaccine, I began to experience blurriness in my left eye. After conferring with my primary care facility, it was recommended that I see an ophthalmologist. Confirmed that I had a Branch Retinal Vein Occlusion (BRVO) and recommended I meet with a retinal specialist. Dr. confirmed the BRVO was caused by a blood clot and commenced treatment via eye injection. I then met with my primary physician to discuss the situation. He recommend further tests to determine whether I had other clotting. In addition to blood work, he order MRIs of my brain, head and neck region and also recommended that I meet with a thrombosis specialist. I met with a Dr. who ordered additional blood work for additional testing. All tests performed to determine if I had additional clotting have come back negative.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1282231-1	I experienced blood clots, also had bilateral leg pain and lower back pain. I currently in the hospital.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1282273-1	Blood Clot in Right Leg
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1282638-1	Ultrasound done on 4-30-2021 for pain in calf of left leg and numbness of the toes for 7 days. Result came back positive for blood clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1283019-1	Patient present to ED on 5/3 with leg pain. found to have blood clot in left leg via ultrasound. started on rivaroxaban and discharged
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284648-1	PULMONARY EMBOLISM; THROMBOSIS; COR PULMONALE ACUTE/ DYSPNOEA; HYPOXIA; DIZZINESS; COUGHING; This spontaneous report received from a patient via a Regulatory Authority Vaccine Adverse Event Reporting System (VAERS) (VAER reference number 1120494) concerned a 68 year old female unknown ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 1805020, and expiry: UNKNOWN) dose was not reported, administered on 06-MAR-2021 14:15 for prophylactic vaccination. Concomitant medications included amitriptyline for sleep, and zolmitriptan for migraine. On 07-MAR-2021 08:30, the subject experienced pulmonary embolism, thrombosis, cor pulmonale acute/ dyspnoea, hypoxia, dizziness, coughing, and was hospitalized (date unspecified). On 12-MAR-2021, Laboratory data included: COVID-19 virus test negative (NR: not provided) Negative. Laboratory data (dates unspecified) included: Blood pressure (NR: not provided) normal, Blood test (NR: not provided) normal, CT scan (NR: not provided) Pulmonary Embolism, Chest X-ray (NR: not provided) normal, Diagnostic ultrasound (NR: not provided) No blood clots or vein thrombosis, Echocardiography (NR: not provided) Increased pressure on the right side of heart due to blood clots, and Heart rate (NR: not provided) elevated. Treatment medications (dates unspecified) included: oxygen, salbutamol, and rivaroxaban. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the pulmonary embolism, thrombosis, cor pulmonale acute/ dyspnoea, hypoxia, dizziness, coughing was not reported. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition); Sender's Comments: V2:Additional version created for MAC update. This updated information does not alter the causality of previously reported events. 20210441745-Covid-19 vaccine ad26.cov2.s-Thrombosis, pulmonary embolism ,Cor pulmonale acute, Hypoxia, dizziness, cough. These events are considered unassessable. These events has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284650-1	MIGRAINE; FEVER; HEADACHE; ARM PAIN; POTENTIAL BLOOD CLOT; This spontaneous report received from a patient concerned a 35 year old female. The patient's height, and weight were not reported. The patient's past medical history included stroke, and concurrent conditions included non-smoker, and sulfa allergy (anaphylactic). The patient experienced drug allergy when treated with nitrofurantoin. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 206A21A, expiry: UNKNOWN) dose was not reported, administered on 10-APR-2021 for prophylactic vaccination. No concomitant medications were reported. On 10-APR-2021, patients arm was hurting and on the 11-APR-2021, night patient started having a headache. On 11-APR-2021, the patient had a fever of 103 and a migraine on 12-APR-2021. Patient took 2 Tylenol and rested in bed for the whole day. On 12-APR-2021, patient saw the physician due to her migraine. The physician was concerned that the patient may have a potential blood clot since the patient has had a stroke in 2016. Physician wanted her to get a MRV (Magnetic Resonance Venography) with contrast of the brain to make sure patient does not have another stroke and also prescribed baby aspirin until she can get the MRV The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from arm pain, fever, and headache, had not recovered from migraine, and the outcome of potential blood clot was not reported.. This report was serious (Other Medically Important Condition).; Sender's Comments: 20210443400-Covid-19 vaccine ad26.cov2.s-Potential blood clot. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284656-1	ESCALATING PAIN IN THE LOWER BACK; PAIN IN LEG; BLOOD CLOT IN LEG; This spontaneous report received from a consumer (representative) concerned a 4 decade (early 30's) old adult male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, 1 total administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up would be requested for this case. No concomitant medications were reported. On 16-APR-2021, the patient experienced escalating pain in the lower back and in leg. The patient also experienced blood clot in leg, and was hospitalized in an unknown date in APR-2021. The patient had improvement and scheduled to leave the hospital in a few days at the time of this report. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from blood clot in leg, escalating pain in the lower back, and pain in leg. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210452829-COVID-19 VACCINE AD26.COV2.S-BLOOD CLOT IN LEG, ESCALATING PAIN IN THE LOWER BACK. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210452829-COVID-19 VACCINE AD26.COV2.S-PAIN IN LEG . This event(s) is labeled per RSI and is therefore considered potentially related.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284660-1	CLOTS; This spontaneous report received from a physician concerned a patient of unspecified age and sex. The patient's weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination.The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient died due to clot. It was unknown if autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died to clots on an unspecified date. This report was serious (Death). This case, from the same reporter is linked to 20210457363.; Sender's Comments: V0: 20210456400-Covid-19 vaccine ad26.cov2.s-Death. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: CLOTS

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284680-1	SIDE EFFECTS INVOLVING SANGUINEOUS CLOTS; This spontaneous report received from a consumer (source: article published) concerned 6 patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry: unknown) dose, start therapy date were not reported, frequency 1 total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. It was reported that the center for disease control and prevention (CDC) and arm of food and drug administration (FDA), the agency responsible for health surveillance, was about to discuss whether the pause in the use of covid-19 vaccine ad26.cov2.s is expected to continue after 6 people developed rare side effects involving sanguineous clots on an unspecified date. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of side effects involving sanguineous clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-20210460381-covid-19 vaccine ad26.cov2.s-side effects involving sanguineous clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1284969-1	I have APS and am on warfarin maintained at a therapeutic level of 2.5-3.5. I received the Johnson & Johnson Vaccine on April 7,2021. On April 9,2021, I noticed swelling and tightness in my left leg, on April 13,2021 I noticed severe swelling in my right leg too. I met with my primary care physician who sent me for imaging which confirmed clotting. I was referred to the Emergency Room since my INR was 3.7 but I was still clotting. I was sent home with out treatment from Medical Center because I was therapeutic on my warfarin. On Friday April 14, I went to the ER after speaking with a hematologist at the facility. I was admitted and was put on fondaparinux.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1285459-1	BLOOD CLOTS
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1287093-1	Right lower leg sudden onset severe swelling, pain, redness. Went to urgent care doctor who then performed assessment, blood work and ultrasound. Placed on Xarelto 15mg twice daily for 21 days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1287916-1	"ESCALATING PAIN IN LOWER BACK; ESCALATING PAIN IN LEG; BLOOD CLOT IN LEG; VACCINE-INDUCED IMMUNE THROMBOTIC THROMBOCYTOPENIA; This spontaneous report received from a patient via a company representative (media article) concerned a 4 decade old male (man in 30s). The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, frequency once total, administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date in APR-2021, within two weeks after receiving a dose of the J&J COVID-19 Vaccine, the patient experienced blood clot in leg. Patient began to experience escalating pain in the lower back and leg on 16-APR-2021. Patient was hospitalized on 21-APR-2021 with escalating pain in his lower back and leg, 13 days after taking the vaccine. It was reported that this was the first male patient with VITT (vaccine-induced immune thrombotic thrombocytopenia) syndrome. The patient was recuperating and making good progress and would be discharged within a few days. Patient's bloodwork showed signature low levels of platelets and fibrinogen, a blood-clotting factor made in the liver, which had been seen in other patients with vaccine-induced clots. Bloodwork showed the patient had the same syndrome as the other patients although initial imaging did not show a blood clot. Physicians later discovered a tiny clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from escalating pain in lower back, escalating pain in leg, blood clot in leg, and vaccine-induced immune thrombotic thrombocytopenia. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition). This case, from the same reporter is linked to 20210451418. Initial information was processed with additional information received on 28-APR-2021.; Sender's Comments: V0: This spontaneous report received from a media article concerned a man in 30s who experienced vaccine-induced thrombotic thrombocytopenia 13 days after vaccine. Medical history and concomitant medications were not reported. Eight days after vaccine, patient began to experience escalating pain in the lower back and leg. Initial imaging did not show a blood clot but physicians later discovered a ""tiny clot"" in the leg. Thirteen days after vaccine, patient was hospitalized with escalating pain in his lower back and leg. The patient was recuperating and making good progress and would be discharged within a few days. Patient's bloodwork showed ""signature low levels of platelets and fibrinogen."" Although the other risk factors (not reported) may have contributed, based on evolving knowledge of Thrombosis with Thrombocytopenia Syndrome (TTS, per definition from Brighton Collaboration - BC) and considering the low platelet count and temporal relationship to vaccination (BC Criteria level 1), the s events are assessed to have a plausible relationship with vaccination. Reporter contact information was not provided which precludes meaningful additional information in this version."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1288631-1	BLOOD CLOT IN LEFT LEG; DIFFICULTY SLEEPING; SOME LEG CRAMPS; SWELLING; This spontaneous report received from a patient concerned a 55 year old male. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown, expiry date: unknown) dose, 1 total on an unknown date in APR-2021 (reported as about three weeks ago from the date of reporting) on left arm for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. Since an unknown date in APR-2021 (reported as about a week ago), the patient experienced swelling and some leg cramps. It was reported that nothing (swelling and leg cramps) was dramatic initially, however, they worsened. On 28-APR-2021 (reported as last night), the patient had difficulty in sleeping. On 29-APR-2021, the patient was diagnosed with blood clot in left leg by his doctor. At the time of report (reported as currently), the patient was in transport to a hospital for further evaluation. The patient further mentioned that he did not have possession of the vaccination card, but his wife had it and she would be arriving at hospital later. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot in left leg, swelling, and some leg cramps, and the outcome of difficulty sleeping was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: -COVID-19 VACCINE AD26.COV2.S-Blood Clot in leg. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1288635-1	BLOOD CLOTS; This spontaneous report received from a consumer via social media through a company representative and concern a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry date: Unknown) dose, 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0- Covid-19 vaccine ad26.cov2.s-blood clots.This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1288959-1	Patient received vaccine on 4/10/21. Went to ED, got IV placed due to side effects. Came to clinic on 5/4/21 with right arm pain. Found to have occlusive cephalic vein clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1289927-1	Patient was discharged from the hospital after massive PE. She went to the hospital at 2am last Saturday. Patient described unbelievable pain from shoulder neck and arms. She could hardly walk to her car. Went to ER. There they ran tests and found many blood clots in lungs. Blood clots had stopped the blood in that area. They were not definitive but said it sounded like that. Got a breathing apparatus after because it's been hard to breathe and she has been shaky. Her shot was administered 04/12 during chemo. Hospital April 24. She was discharged and given a blood thinner and pain medication - Oxycodone & Eliquis.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1290307-1	I woke up at approximately 9:30 AM the morning of Sunday May 2nd 2021 the day after getting my covid vaccine and had to rush to the bathroom. I proceeded to have blood and blood clots come out my urethra very urgently. I had abdominal pain and pressure. I proceeded to need to urinate blood and blood clots for approximately 4 hours urgently. I could not go far from the bathroom without having to return quickly to the toilet. I wet myself several times. The clots stopped coming around 1:30 PM. I proceeded to have to have no control over urination urge and could not leave the bathroom. Around approximately 2:00 PM I could leave the bathroom without fear of wetting myself. I continued to have a small amount of a mix of blood and normal urine passing for another hour or so. I have a picture of the blood and clots that came out of my urethra. I continue to have dull pain and pressure in my abdomen.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1291581-1	POSSIBLE CLOT IN THE LEFT ARM; PAIN IN THE LEFT ARM/SIDE; This spontaneous report received from a patient via a company representative concerned a 6 decade old female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced possible clot in the left arm, and pain in the left arm/side, and was hospitalized (date unspecified). it was reported that the patient had reaction to the JnJ vaccine. She was being watched closely and taken care of. Pain on the left arm/side. They think it as clot in the left arm. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the pain in the left arm/side and possible clot in the left arm was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: v0;20210456346-Covid vaccine ad26.cov2.s-Possible clot in the left arm. Follow-up received regarding Clinical Details. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). v0;20210456346-Covid vaccine ad26.cov2.s-Pain in left arm. Follow-up received regarding Clinical Details. This event(s) is labeled per RSI and is therefore considered potentially related.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1291587-1	BLOOD CLOT; This spontaneous report received from a consumer via social media through a company representative and concerned a patient of unspecified age and sex The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unk) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot three weeks later without any underlying conditions. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210504574.; Sender's Comments: V0:20210503763-Covid-19 vaccine ad26.cov2.s-Blood clot. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1292143-1	lips and nail beds blue, 4/3/21 bumped left leg,,big bruise, felt clots, headache, metallic taste in mouth,
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1292685-1	Vaccine Induced Thrombotic Thrombocytopenia - confirmed with Lab testing Elisa PF4 highly positive 1.chills, generalized body aches, and noted that some glands in her neck were swollen. for 24-48 hours post vaccine 2. 4/27/2021, worsening headaches, bifrontal, along with some bruising in her right leg and left wrist area. Evaluated at ER. white count was 6.1, her hemoglobin was 12.5 and her platelet count was 91,000.(PLT previously 301,000). MR venogram of the head was performed with and without contrast. There was no dural, venous, sinus thrombosis or significant stenosis. The clinical impression at the time of the treating ER physician was that she had acute sinusitis 3. 4/30/2021 PCP visit Additional testing was obtained PLT 149,000. white count and hemoglobin were in the normal ranges. D-dimer was elevated at greater than 20. The fibrinogen was low at 143. Heparin platelet antibody test - lab - Negative 4. 5/3/2021 presents to Hospital. 12 to 16-hour history of back pain, bilaterally, left greater than right. It also was very painful to take a deep breath. She also noted that her fatigue was progressively worsening. CT Pulmonary Angiogram shows Thrombus within multiple right lower lobe, right middle lobe and posterior left upper lobe and medial left lower lobe pulmonary arteries. white count of 11.4, a hemoglobin of 13, and a platelet count of 254,000. Her ANC was 9. Her reticulocyte count was 1.3 and absolute reticulocytes were 0.06. Treated with Non heparin anticoagulant - argatroban IV infusion titrated to PTT lab results Discharged from Hospital with rivaroxaban (Xarelto) 5/6/2021
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1293356-1	4/10/2021 vaccination 4/11 or 4/12 at the latest, I started feeling a pain in my R calf. It was constant pain for a week and decided to go to the walk in clinic. Blood work which was high and they referred me to the ER and administered a sonogram. *positive for blood clot Eloquence for 30 day starter pack. PCM is going to extend till July in which he will administer another sonogram. *COVID +; 01/2021
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1293543-1	Feet turned black blood thinner 3 blood clots in lungs 1 clot in heart
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1293595-1	The evening of the shot I woke up with pain in my legs near my hips which lasted for a day. On 04/12/2021 my left leg was swollen and painful. I was already taking blood thinner. I went to the Emergency Room the next day and they found a blood clot in my leg. The Thrombosis was behind my left knee. My leg was swollen from the knee down and my ankle was very swollen. My Eliquis dosage has been increased from 2.5mg to 5mg twice daily. I did not have a repeat sonogram but my leg is no longer swollen.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1293612-1	Details of Hospital Stay History of Present Illness The patient is a 79-year-old woman with a past medical history of hypertension, hypothyroidism, lumbar compression fracture, hiatal hernia, metastatic adenocarcinoma primary lung to bone and question liver, on immunotherapy pembrolizumab started 12/3/2019, changed to osimertinib, right leg superficial thrombosis, on Xarelto presented to hospital on 3/10/2020 with complaints of 2 months duration shortness of breath, worsened over a 4-day period. She was admitted to the hospital with a diagnosis of acute respiratory failure with concerns of acute PE. For all details regarding the patient's initial presentation please refer to history and physical exam dated 3/10/2021.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1294490-1	blood clot in left leg calf
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1294816-1	BLOOD CLOTS; UNABLE TO WALK/TEMPORARY DISABLED; This spontaneous report received from a source from a consumer via a company representative concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. The consumer reported that the patient on an unspecified date experienced blood clots and was unable to walk. The patient was under doctor care and was temporary disabled. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clots, and unable to walk/temporary disabled. This report was serious (Other Medically Important Condition, and Disability Or Permanent Damage).; Sender's Comments: 20210502999-covid-19 vaccine ad26.cov2.s-Blood clots, unable to walk. This event(s) is considered un-assessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1296176-1	On 04/21 left knee pain, outside/back of knee 04/23 - left side tingling , went to urgent care and then referred to ED due to concern about stroke. Sent home after leg scan - was told it was normal and referred to neurologist 04/29 - back to ED, admitted overnight, scan from 04/23 showed blood clot behind the knee. Also diagnosed by neuro with TIA Also stated she had swelling, redness, and headaches for 4-5 days after vaccine
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1296456-1	Sharp pain on right leg, difficulty breathing.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1297529-1	pt told me she had the Johnson and Johnson one covid shot and slept for 3 days afterwards and then had to be hospitalized for a blood clot
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1300185-1	Blood clots in right arm and lungs. Swollen right arm.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1301256-1	Pain in right leg in knee area beginning on 4/6/2021. Pain worsened over the weekend. Went to the doctor on 4/14/2021. She sent me for a doppler on my legs where a blood clot was found behind my right knee. I started taking Xarelto that night - 15mg twice daily for 21 days. The pain increased considerable and by 4/20/2021 I could hardly walk. I went back to the doctor that day and she ordered a second doppler study. The blood clot was stable but there was no change. Continued on the Xarelto. Third visit to the doctor on 5/5/2021. Some lessening of the pain by then. Doctor changed the Xarelto dosage to 20 mg. one a day for the next two months. Then we'll schedule another doctor visit and probably another Doppler Study.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1301959-1	VARICOSE VEIN RED AND HARD; UPPER RIGHT THIGH PAIN; SUPERFICIAL CLOTS; This spontaneous report received from a patient concerned an adult female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose was not reported, 1 total administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On 09-APR-2021, the patient experienced varicose vein red and hard, and was hospitalized. She sought care at the Vein Institute in Voorhees, NJ, as she says she had been treated there in the past. On 09-APR-2021, the patient experienced upper right thigh pain. On Apr-2021, the patient experienced superficial clots, and was hospitalized. Laboratory data included: Diagnostic ultrasound (NR: not provided) confirmed Superficial clots. Treatment medications included Motrin and Heating pad for Varicose veins. She was told to return in 2 weeks for a follow up ultra sound. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the superficial clots, varicose vein red and hard and upper right thigh pain was not reported. This report was serious (Hospitalization Caused / Prolonged, Other Medically important condition). This case, from the same reporter is linked to 20210501531.; Sender's Comments: V0 : 20210500162 - Covid-19 vaccine ad26.cov2.s - Superficial clots and Varicose vein red and hard . This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1301969-1	BLOOD CLOT IN LEG; This spontaneous report received from a patient via a company representative concerned an adult male. The patient's weight, height, and medical history were not reported. The patient received ibrutinib (tablet, oral, batch number was not reported) dose, frequency, and therapy dates were not reported for an unspecified indication. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: UNKNOWN) dose, start therapy date were not reported, 1 in total for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot in leg, and was hospitalized (date unspecified) for two days. The action taken with covid-19 vaccine ad26.cov2.s was not applicable; and action taken with ibrutinib was not reported. The outcome of blood clot in leg was not reported. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0. 20210507935-covid-19 vaccine ad26.cov2.s -blood clot in leg. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1301971-1	BLOOD CLOT; This spontaneous report received from a consumer via a company representative concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received COVID-19 VACCINE AD26.COV2.S (suspension for injection, route of admin not reported, batch number: Unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The Batch number was not reported. The company is unable to perform follow up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the reporter stated that her husband had blood clot after vaccination. The action taken with COVID-19 VACCINE AD26.COV2.S was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210508168-COVID-19 VACCINE AD26.COV2.SI-BLOOD CLOT. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1304956-1	BLOOD CLOT; This spontaneous report received from a company representative concerned a 42 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry were unknown) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, between three to ten days later after vaccination the patient died due to blood clots. It was unknown, whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender's Comments: V0;20210510772-covid-19 vaccine ad26.cov2.s-Thrombosis. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1304971-1	BLOOD CLOT; This spontaneous report received from a traditional media article concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported, 1 total, administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient had blood clot after vaccination. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210514145 -Blood clot (PT-Thrombosis). This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1306313-1	THROMBOSIS; This spontaneous report received from a health care professional concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced thrombosis. Treatment medications (dates unspecified) included: apixaban for thrombosis. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The outcome of thrombosis was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210516823-Covid-19 vaccine ad26.cov2.s -Thrombosis. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1306915-1	"Client rec'd her vaccination on 04/09/2021. Approximately 2 weeks later, family states client started to not ""feel well."" Patient had leg pains and fatigue, as per family. On Saturday, May 8th, client drove her self to the ER, due to worsening symptoms of cough, leg pain, and shortness of breath. Client was evaluated in the ER and evaluation showed clots in bi-lateral lungs and lower legs. Surgery was performed to remove clots in lungs. Client was admitted. On 5/9/2021 client was intubated, and had an ECMO procedure. Client i remains in the hospital."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1307249-1	"Sudden onset of menstrual period, lasting 2 weeks with extremely heavy flow and pain. Saw my Dr. and she did a blood test to check for platelet count, which all labs done were normal. Period lasting from 4/7 to 4/21 all the while I take my birth control as normal with no missed doses. Now on 5/7 having severely heavy flow again with clotting. Dr. prescribed naproxen and an extra ""mini-pill"" hormone supplement on top of my regular birth control. Will monitor."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1307444-1	she?s had blood clots and has had surgery 3 times, she is unconscious at the moment. the whole left side of her brain has been removed and she is paralyzed on the left side. this is definitely from the shot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1308488-1	Patient called 5/11/21 stating today she blew a blood clot from her nose
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1309190-1	BLOOD CLOTS IN BOTH LOWER LEGS; LEG CRAMPS; This spontaneous report received from a patient concerned a 58 year old white non-Hispanic or Latino female. The patients weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 1805020, expiry: 25-MAY-2021) dose was not reported, frequency 1 total, administered on 18-MAR-2021 to left deltoid for prophylactic vaccination. No concomitant medications were reported. On 18-MAR-2021, the patient started experienced leg cramps. On 22-MAR-2021, the patient diagnosed with blood clots in both lower legs by ultrasound. Patient was put on unspecified blood thinner and started RX on 23-Mar-2021 Laboratory data included: Diagnostic ultrasound (NR: not provided) Positive for bilateral blood clots in lower legs. On 04-Apr-2021 patient went to ER for severe leg cramps, Patient got RX for Soma and started taking on 05-Apr-2021. Treatment medications (dates unspecified) included: carisoprodol. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clots, lower extremities, and leg cramps. This report was serious (Other Medically Important Condition).; Sender's Comments: V0- 20210506805 - Covid-19 vaccine ad26.cov2.s-Blood clots in both lower legs. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1309751-1	"PAIN IN UPPER BACK; CLOT IN LUNG; CLOT IN CALF; DIZZINESS; FLU LIKE SYMPTOMS; JOINT PAIN; STOMACH ACHES; SWELLING IN CALF; This spontaneous report was received from a patient and concerned a 55 year old male. The patient's height and weight were not reported. The patient's past medical history included fractured ribs and T6, and concurrent conditions included drug allergy to Vicodin (causes rash). The patient was healthy before the vaccine. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 042A21A expiry: 21-JUN-2021) dose was not reported, administered on and unknown date in APR-2021 (approximately ""middle of April"") in the left arm for prophylactic vaccination. There were no concomitant medications. The day after receiving the vaccine the patient started experiencing fever (no actual temp given), stomach aches, chills, joint pain, ""flu-like symptoms"" which lasted ""4 to 5 days after getting the vaccine"" (no actual dates given). He stated those symptoms went away. On the ""10th day after getting the vaccine"" he experienced pain in his calf and had a hard swollen spot which he assumed was cramps. This lasted until Sunday, 02-MAY-2021. On Monday, 03-MAY-2021 (also reported as 02-MAY-2021), the patient started experiencing pain in upper back which he felt ""in his lungs"". On Tuesday 04-MAY-2021 , the pain was still present. On Wednesday, 05-MAY-2021, he started experiencing difficulty breathing, dizziness, and shortness of breath and went to the emergency room (ER). This pain was unlike anything he had ever experienced. While at the hospital, they performed an ultrasound and found a clot in his calf. They also performed a CAT scan and found a clot in his lungs. The patient was treated with heparin, but after he expressed concern with heparin, the hematologist changed it to Eliquis (apixaban). The patient was discharged on 05-MAY-2021 with Eliquis 20 mg, one tablet by mouth twice a day for two weeks. He will be on it for the next 6 months. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from flu like symptoms, and stomach aches on APR-2021, and joint pain on an unspecified date, and the outcome of pain in upper back, dizziness, clot in calf, clot in lung and swelling in calf was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0:20210511212-covid-19 vaccine ad26.cov2.s-Pulmonary thrombosis, Thrombosis, Pain in upper back. This events are considered unassessable. The events has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1309770-1	On 4/15/21 came into office with bilateral edema in both lower legs as well as pain x 2 weeks. sent for doppler- Diagnosed with occlusive thrombus in R lower leg.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1311511-1	For thirty-six hours after the vaccine. I had body aches and chills and my eye sockets were sore and it hurt to look up. Ten days after the vaccine I was in the shower and a large blood clot came out of my nose. It was a large clot about the size of a quarter and there was no other blood with it. The clot came out of my left nostril but it was not like a nose bleed. I went to the Urgent Care but they did not do tests.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1312771-1	FEMALE ISSUES; BLOOD CLOTS; This spontaneous report received from a consumer concerned multiple patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported and has been requested. On an unspecified date, the consumer called and reported that he had read in newspaper that women had gotten blood clots and 3 passed away with female issues. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of female issues on an unspecified date, and the outcome of blood clots was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender's Comments: V0. 20210510294 -COVID-19 VACCINE AD26.COV2.S-Female issues, Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1313235-1	SUSPECTED URETHRAL MUCOSITIS; CLOT; PENIS TIP GOT RED; PENIS PAIN; SAW ANGELS AND DEMONS IN BED; TERRIBLE MALAISE; HEADACHE; FEVER 40 DEGREE; This spontaneous report received from a patient concerned a 26-year-old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 204A21A, and expiry: UNKNOWN) dose was not reported, 1 total, administered on 02-MAY-2021 for prophylactic vaccination. No concomitant medications were reported. On 02-MAY-2021 at night, the patient experienced some symptoms after vaccination including Fever of 40 degrees Celsius, headache, and a terrible malaise. The patient also reported seeing angels and demons in bed; Also, the patient noticed that the penis tip got red and experienced pain. On 05-MAY-2021, the patient continued with the symptoms, and it seemed as if a clot was generated. On 06-MAY-2021, the patient had an appointment with general physician who informed that he never saw an adverse reaction like this after vaccination and that it could be a urethral mucositis. On 10-May-2021, the patient would have an appointment with the urologist to inform the symptoms. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the fever 40 degree Celsius, saw angels and demons in bed, headache, terrible malaise, penis tip got red, penis pain, clot and urethral mucositis was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0.20210513944-covid-19 vaccine ad26.cov2.s - thrombosis. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1314221-1	Patient got small blood clot show on skin 4 days after shot, it went away lighter red in 1.2 week, blood pressure went up and down suddently. Sunday was fine, Monday the 26 was fine, went to bed at 9:00 PM. Was not able to wake up, went to ER around 8:30 AM on 4/27 AM by daughter call 911. In ER for 28 hour and found out she got stroke. Pnewnewmia. ER gave antibootic around 3:00 PM, transfer to intensive care, MRI around 7:00 PM 4/27/2021 due to CT scan shown stroke. Lost Vision on right side and parcial on left side due to stroke.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1314349-1	Blood clot developed in the leg 10 days after the injection
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1315704-1	BLOOD CLOTS BOTH LEGS; POSSIBLE HEARTBURN; SWELLING AT INJECTION SITE; This spontaneous report received from a patient concerned a 64 year old male. The patients weight, height, and medical history were not reported.The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805022, and expiry: UNKNOWN) dose was not reported, frequency 1 total, administered on 16-MAR-2021 at left arm for prophylactic vaccination. No concomitant medications were reported. On 16-MAR-2021, the patient experienced swelling at injection site. On 19-APR-2021, patient had blood clots (1 in each leg), Hematologist put him on xarelto 10mg, in hospital for few days for blood clots. Patient initially went to hospital ER on 19-APR-2021 because he thought having chest pains or heartburn. Emergency Room ran tests and heart was normal, but found blood clots in his both legs. Lab test (NR: not provided) Heart normal. Treatment medications (dates unspecified) included: rivaroxaban. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from thought having chest pains/heartburn on 22-APR-2021, had not recovered from blood clots both legs, and the outcome of swelling at injection site was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210504455-COVID-19 VACCINE AD26.COV2.S-Blood clot both legs. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1315710-1	BLOOD CLOT IN LEFT LEG; This spontaneous report received from a patient concerned a 48 year old, White and Hispanic or Latino female. The patient's height, weight and medical history were not reported. The patient had no blood clots in the past or no recent trauma and surgery. The patient was not on any medications. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1802072 and expiry: UNKNOWN) dose was not reported, 1 total, administered to the left arm, on 06-MAY-2021 for prophylactic vaccination. No concomitant medications were reported. On 09-MAY-2021, the patient came into the hospital with swelling in left leg, a lot of heat and a lot of pain. In emergency room (ER) they did scans and tests and it was found that the patient had a blood clot in left leg. The patient has to stay in the hospital for 24 hours until the blood clot was resolved. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot in left leg. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0;20210516127- covid-19 vaccine ad26.cov2.s-Blood Clot in left leg. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1315725-1	PASSED AWAY BECAUSE OF CLOT; This spontaneous report received from a consumer concerned a patient of Unspecified Race, ethnic origin, age and sex. The patient's weight, height and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry: unknown) dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported and has been requested. On an unspecified date, the patient passed away because of clot. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender's Comments: V0: 20210517449- covid-19 vaccine ad26.cov2.s-thrombosis. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1315726-1	BLOOD CLOT IN LEFT CALF; This spontaneous report received from a patient via a company representative through social media concerned a male of unspecified age and ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose, start therapy date were not reported, 1 total, administered for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, the subject experienced blood clot in left calf. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot in left calf was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0;0210517568-covid-19 vaccine ad26.cov2.-Blood clot in left calf. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1315751-1	BLOOD CLOT IN LEFT CALF; PAIN IN ANKLE; PAIN SLOWLY MOVING UP TO THE CALF; SLIGHT HEADACHE; This spontaneous report received from a patient concerned a 79 year old female of unknown race and ethnicity. The patient's height, and weight were not reported. The patient's concurrent conditions included high blood pressure, and penicillin allergy. The patient experienced drug allergy when treated with ciprofloxacin, and pethidine hydrochloride. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808980, and expiry: 21-JUN-2021) dose was not reported, 1 total administered on 02-APR-2021 to left arm for prophylactic vaccination. Concomitant medications included blood-pressure pills, water pills and Omega-3. On 02-APR-2021, the patient experienced a slight headache for about 2 days. On 29-APR-2021, the patient felt some pain in ankle, which was slowly moving up to the calf (left leg). The patient was brought to the emergency room, where an ultrasound and blood work was performed. Ultrasound detected a blood clot in her left calf and prescribed with Xarelto daily for 3 months. Patient stated that pain reached its most intense status on 01-MAY-2021 and is slowly getting better. Treatment medications (dates unspecified) included: rivaroxaban. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from headache on 04-APR-2021, was recovering from pain in ankle, and pain moving up to the calf, and had not recovered from blood clot in left calf. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210520686 -JANSSEN COVID-19 VACCINE Ad26.COV2.S-Thrombosis. The event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1316349-1	This spontaneous report received from a consumer via a company representative concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient died from blood clot. It was unspecified if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender's Comments: V0. 20210514012-COVID-19 VACCINE AD26.COV2.S-Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1316350-1	BLOOD CLOTS IN LEGS; This spontaneous report received from a patient via a company representative concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported 1 total for prophylactic vaccination. The batch number was not reported and had been requested. No concomitant medications were reported. On an unspecified date, on 2021, the patient experienced blood clots in legs. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clots in legs. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210515265-covid-19 vaccine ad26.cov2.s-thrombosis. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1316351-1	THROMBOSIS IN RIGHT HAND MIDDLE FINGER KNUCKLE; RIGHT ARM AND HAND COLD; RIGHT ARM AND HAND NUMB,LEFT HAND NUMB; WEAK LEFT HAND AND WEAK LEGS; CRAMP IN INNER MIDDLE FOOT LEFT FOOT; PAIN ALL OVER BODY; CHILLS; FEVER; HAND AND LEG STIFFNESS,STIFF NECK; RED/BLUE BRUISE; This spontaneous report received from a patient concerned a 38 year old white and Hispanic or Latino female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported and batch number: 205A21A expiry: UNKNOWN) dose was not reported, 1 total administered on 03-MAY-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. On MAY-2021, the patient experienced red/blue bruise. On 03-MAY-2021, the patient experienced pain all over body, hand and leg stiffness for 10 minutes, stiff neck, chills and fever. On 04-MAY-2021, the patient experienced cramp in inner middle foot left foot. On 09-MAY-2021, the patient experienced thrombosis in right hand middle finger knuckle, right arm and hand cold, right arm and hand numb, left hand numb and weak left hand and weak legs. Treatment medications (dates unspecified) included: acetylsalicylic acid. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from cramp in inner middle foot left foot on 04-MAY-2021, pain all over body on MAY-2021, and chills, and fever on 03-MAY-2021, was recovering from right arm and hand cold, right arm and hand numb, left hand numb, and weak left hand and weak legs, had not recovered from thrombosis in right hand middle finger knuckle, and hand and leg stiffness, stiff neck, and the outcome of red/blue bruise was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210516231-covid-19 vaccine ad26.cov2.s - thrombosis in right hand middle finger in Hispanic or Latino patient. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1316357-1	CLOTTING; MPV IS HIGH; This spontaneous report received from a patient via a company representative concerned to patient of unspecified age and gender with unknown race and ethnicity. The patient's weight, height, and medical history were not reported. Patient reported that last year Mean Platelet Volume MPV was in the normal range. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry date were unknown) dose was not reported,1 total administered on 20-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On 08-APR-2021, the patient performed yearly bloodwork and physical test which showed Mean Platelet Volume (MPV) was high and it lead to clotting. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the MPV was high and clotting was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0;20210517660-covid-19 vaccine ad26.cov2-MPV was high and clotting. This events are considered unassessable. The events have a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1316948-1	Patient's husband reported that his wife had gone to the ED with s/s of blood clot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1317095-1	Chills (no fever) within 24 hours of the Janssen vaccination, lethargic, no appetite. Very dizzy and lightheaded on day 5 and day 6 after the Janssen vaccination with continued lethargy and loss of appetite for approximately 3 weeks. Heavy nosebleed in left side of nostril with large blood clots for 1.5 hours approximately 3 weeks after the Janssen vaccination
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1317607-1	"Patient describes muscle pain and bruising/""blood clots"" on left side. The onset was 5/9/2021. Describes as severe pain that increased till seen at healthcare provider's office by healthcare provider."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1319774-1	BLOOD CLOT IN LEFT LEG/MUSCLE IN LEG TIGHTENED UP/CHARLEY HORSE LIKE/LEFT THIGH STILL HURT; FATTY LIVER; This spontaneous report received from a patient concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company was unable to perform follow-up to request batch/lot numbers. Concomitant medications included ustekinumab used for unknown indication. On an unspecified date, the patient experienced blood clot in left leg/muscle in leg tightened up/charley horse like/left thigh still hurt, and had a fatty liver a few weeks ago and was hospitalized. The patient stated that she took the vaccine of covid-19 three weeks ago and developed a left leg blood clot and had muscle in leg tightened up (real tight) like charley horse. Then, she went to the PCP (primary care physician) who told her for the valley imaging an ultrasound laboratory test which discovered the clot. On 30-APR-2021, the patient got admitted to the hospital and was discharged on 02-MAY-2021. The patient was hospitalized for two days. Treatment medications (dates unspecified) included: apixaban and she was doubled up on eliquis (apixaban) for blood clot in left leg. On the day of report, she stated that left thigh hurt still ongoing. (The patient have a colonoscopy laboratory test on 09-JUN-2021). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clot in left leg/muscle in leg tightened up/charley horse like/left thigh still hurt and fatty liver was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0 20210506567 -COVID-19 VACCINE AD26.COV2.S- Blood clot in left leg/Muscle in leg tightened up/Charley horse like/Left thigh still hurt, Fatty liver. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1319778-1	DEATH; BLOOD CLOT; This spontaneous report received from a health care professional concerned a 66 year old White and not Hispanic or Latino female. Initial information received from the health care professional on 05-MAY-2021 was processed with additional information obtained from live follow up with health care professional on 06-MAY-2021. The patient's height, and weight were not reported. The patient's concurrent conditions included high blood pressure, diabetes, high cholesterol, non smoker, non alcohol user and other pre-existing medical conditions included no known allergies, no drug abuse or illicit drug use. Lab work was done 3-4 months ago, the results of which were unavailable. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 202A21A, and expiry: Unknown) dose was not reported, 1 total, administered on 10-APR-2021 possibly left arm for prophylactic vaccination. Unspecified concomitant medications were reported. On 28-APR-2021, the patient developed right leg pain and right leg swelling also began around this time as well. On 04-MAY-2021, the patient died from blood clot. An autopsy was not performed. The reporter stated that the patient's death was related to Janssen covid-19 vaccination and blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of blood clot on 04-MAY-2021. This report was serious (Death).; Sender's Comments: V0:20210509157-JANSSEN COVID-19 VACCINE- Death, blood clot - These events are considered unassessable. The events have a compatible/suggestive temporal relationship, are unlabeled, and have unknown scientific plausibility. There is no information on any other factors potentially associated with the events.; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1319803-1	BLOOD CLOTS; HEART ATTACK; This spontaneous report received from a social media via a company representative concerned a patient of unspecified age and sex with unknown ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry were unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced heart attack due to blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the heart attack and blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210522101-Covid-19 vaccine ad26.cov2.s-Blood clots, Heart attack. This event(s) is considered un-assessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1320413-1	Heavy menstrual bleeding, large and excessive menstrual blood clots

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1321357-1	after getting the vaccine in morning, later that same evening patient felt sluggish, nauseaus, tired. Daughter finally convinced patient to see a doctor. Md prescribed her a cough medication. She still felt bad for another 2-3 weeks, so she finally went to the emergency room where she diagnosed with blood clots in the legs and the lungs. She staying in hospital approx. 4-5 days before being released.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1321418-1	On Saturday, April 10, 2021, I had started having pain and swelling in my right calf that lasted for several days on and Tuesday, April 13th, I started having pain on the left side of my chest. On Thursday, April 15th, I went to the ER due to pain and selling in my leg and it was discovered that I had 2 clots in my leg (on at my calf and a second one on the right side of my knee). Not realizing my chest pain was related, I discussed with my primary care at a follow up visit on Monday, April 19th, and a CT scan was performed confirming I have an pulmonary embolism (extensive pulmonary emboli in both lungs). I was immediately put on Xarelto blood thinner and sent home.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1321436-1	Is that I got blood lock? Help me please!
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322563-1	BLOOD CLOTS; This spontaneous report received from a consumer who reported hearing a news report concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number not reported, expiry not reported frequency one total, dose, therapy start date were not reported administered for prophylactic vaccination. The batch number was not reported and has been requested.No concomitant medications were reported. On an unspecified date, the patient had blood clots. On an unspecified date in 2021, the patient was died due to blood clots. it was not reported whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died due to blood clot on an unspecified date in 2021. This report was serious (Death). This case, from the same reporter is linked to 20210523500.; Sender's Comments: V0. 20210522533-COVID-19 VACCINE AD26.COV2.S-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOTS
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322564-1	BLOOD CLOT; This spontaneous report received from a pharmacist concerned a 26 year old female of unknown ethnicity. The patient's height, and weight were not reported. The patient had no known drug allergies. The patient received Covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, and batch number: 043A21A expiry: unknown) dose was not reported, 1 total dose, administered on 07-APR-2021 for prophylactic vaccination. No concomitant medications were reported. On an unspecified date (after vaccination), the patient experienced blood clot. The pharmacist thought, that she has been treated, and at the time of this report she was at home, but he was not 100% sure. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210522731-Covid-19 vaccine ad26.cov2.s-blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322565-1	BLOOD CLOTS IN LEFT LEG; BLOOD CLOTS IN BOTH LUNGS; This spontaneous report received from a patient via social media concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: UNKNOWN) dose was not reported, 1 total administered on 19-MAR-2021 for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. The patient reports that 3 weeks later(after vaccination) on an unspecified date APR-2021, the patient was in urgent care with blood clots in left leg and blood clots in both lungs. The patient also reports that she was literally headed to her 1 week follow up when it was flashed on the news about blood clots. And reports that since month she was on very expensive unspecified medication, injection and pills and also undergone unspecified labs and diagnostics. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clots in left leg and blood clots in both lungs was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0;20210523088-covid-19 vaccine ad26.cov2.s-Blood clots in left leg, blood clots in both lungs. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322566-1	BLOOD CLOTS; This spontaneous report received from a consumer via social media (internet) concerned a patient of unspecified age and sex. The patient's height, weight and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown expiry: UNKNOWN) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the consumer reported that he heard about issues of blood clot associated with the Janssen Covid 19 vaccine. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210523657-covid-19 vaccine ad26.cov2.s-Blood clots. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322586-1	LEG CLOT; SICK; This spontaneous report received from a consumer concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported ,1 in total administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the patient experienced leg clot and sick. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of leg clot and sick was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210526445 and 20210525676.; Sender's Comments: V0:20210526416 -Covid-19 vaccine ad26.cov2.s-Thrombosis - This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1323456-1	Within 24 hours pain in left side of left chest, in 48 hours pain on full left side of chest, within 72 hours pain through entire chest. Hospitalized within 72 hours, chest pains, catheterized within 120 hours, had a blood clotting issue, put into medically induced coma for three and a half days, followed by a pulmonary embolism and four (4) clogs, two (2) in either leg, within two weeks. Hospitalized twice, currently on three blood thinners, a beta blocker, pepcid, a statin, and one other. Expected to be on three blood thinners for at least three (3), most likely six (6) months. Start cardiac rehab on 5/19/21. Have chronic cough and chest discomfort. Expected recovery time: six (6) months.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1323725-1	The day of the vaccine I was told that my arm might be sore. I felt sore right away and as soon as I got in the car to drive home I was extremely tired and I slept for the entire six hour car ride home. I was very fatigued that night and then I got a bad headache, fever and it lasted for 48 hours. After that, the headache was less intense and they were intermittent but my arm still hurt and I couldn't lay on it. I was still fatigued for a week. On the fourth day I started to get a weird feeling of vibrations and twitching all over my body. It is especially worse in my temple. It has been a month but I still get the twitching in different parts of my body. My face, arms, legs and temples. It is in different places. It happens less often now, it is not all day but it is still daily and mostly in my head and my legs. I had it happen in my temples last night. I had my menstrual cycle since the vaccine and it was about three days later than expected. I had a lot of blood clots which was unusual for me. I have had flu like symptoms this week and I have a cough, runny nose and headaches but I am planning on trying to go to the doctor this week.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1323830-1	I developed a cough on or about beginning of May 1, 2021. After accidentally double dosing myself with my ACE inhibitor I went into the ER on 5/11 with light headedness/high heart rate and cough. After multiple tests were done it was discovered that I had a large satellite embolism. in my lung and another one in my left leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1323838-1	Suspected VITT treated with bivalirudin 0.08 mg/kg/hr, IVIG, hemoptysis, cough, chest pain

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1325834-1	"LOW PLATELET COUNT; INFLAMED LIVER; BLOOD CLOTS; HEADACHES; This spontaneous report received from a patient, via social media, via a company representative concerned a 33 year old male. The patient's weight, height, and medical history were not reported. The patient received COVID-19 VACCINE AD26.COV2.S (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, two weeks after vaccination, the patient experienced headaches, low platelet count, inflamed liver, and blood clots. It was stated the patient had an inflamed liver because of all the blood clots found; the patient is now on blood thinners for a minimum of 6 months. Laboratory data (dates unspecified) included: Low platelets (NR: not provided) 26K. The action taken with COVID-19 VACCINE AD26.COV2.S was not applicable. The outcome of the headaches, inflamed liver, blood clots and low platelet count was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: This spontaneous report received from a patient, via social media, via a company representative concerned a 33-year-old man who experienced blood clots and low platelet (26,000) count two weeks after vaccine. Medical history, concomitant medications, and other details were not reported. It was stated the patient had an ""inflamed liver because of all the blood clots found""; the patient is now on blood thinners for a minimum of 6 months. Information is limited in this case, and the occurrence of blood clots and low platelet count could represent background incidence of such events in the general population. However, a relationship with Janssen Covid-19 vaccine cannot be ruled out and thus the relationship is considered indeterminate (Brighton Collaboration Criteria level 5)."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"BLOOD CLOTS IN BOTH LEGS; PLATELET COUNT DECREASED; ABDOMINAL PAIN; LEG CRAMPS; SEVERE CHILLS; SEVERE HEADACHE; FEVER OF UP TO 102 DEGREES; SEEING LINES; DIFFICULTY IN WALKING; LEG PAIN; This spontaneous report received from a patient concerned a 40 year old female. The patient's weight was 155 pounds, and height was 67 inches. The patient's past medical history included back surgery, and kidney stone removal, and concurrent conditions included alcohol use (one drink once or twice a month), and non smoker, and other pre-existing medical conditions included patient had no drug abuse or illicit drug usage. January 2021, she had a procedure and her platelet count was 300 (unit unspecified). The patient experienced drug allergy when treated with amitriptyline hydrochloride, heparin, oxybutynin, tramadol hydrochloride, and hydrocodone bitartrate/paracetamol. The patient was not pregnant at the time of report. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808980, expiry: 20-JUN-2021) frequency 1 total, dose was not reported, administered to left deltoid on 07-APR-2021 for prophylactic vaccination. No concomitant medications were reported. One week later, on 14-APR-2021, patient began experiencing a severe headache, severe chills, abdominal pain, leg cramps, and a fever of up to 102 degrees. She reported no history of headache or migraines, yet along with the headache, she said she was seeing lines. She said that she called her doctor on 15-APR-2021 (the next day) who advised her to go to the hospital. On 16-APR-2021, patient went to the Hospital, where she was admitted for vaccine induced blood clots in both legs. After receiving the Janssen COVID vaccine, on 16-APR-2021, her platelet count was 27 (coded as platelet count decreased). While in the hospital, she was placed on a blood thinner drip, and her platelet count went as low as 9 (units unspecified). On 19-APR-2021, she was flown to another hospital. She remained in that hospital until 27-APR-2021 when she was discharged. Patient was hospitalized for a total of 11 days. She was currently taking Eliquis 5mg twice a day. She was still having leg pain, and could barely walk (coded as difficulty in walking). She was also concerned about her medical bills. She said that before getting the Janssen COVID vaccine, she was a healthy individual and had not taken any birth control or estrogen. She was continuing to follow up with her physician. Treatment medications (dates unspecified) included: apixaban. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from severe chills on 27-APR-2021, was recovering from severe headache, and fever of up to 102 degrees, had not recovered from abdominal pain, leg cramps, and leg pain, and the outcome of blood clots in both legs, seeing lines, difficulty in walking and platelet count decreased was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: This spontaneous report received from a patient concerned a 40-year-old female (BMI 24.3 kg/m2), who experienced ""blood clots in both legs"" and platelet count decreased 9 days after vaccine. Past medical history included back surgery and kidney stone removal. Her baseline platelet count was 300,000 last checked 4 months before vaccine. The patient was not pregnant at the time of report. No concomitant medications were reported but she denied taking oral contraceptives or other forms of estrogen. Seven days after vaccine, the patient began experiencing a severe headache, severe chills, abdominal pain, leg cramps, and a fever of up to 102 degrees. She said that she called her doctor on day 8 who advised her to go to the hospital. On day 9, patient went to the hospital, where she was admitted for vaccine induced blood clots in both legs. No specific imaging modality or results were reported. Nine days after vaccine, her platelet count was 27,000. While in the hospital, she was placed on an unspecified blood thinner drip, and her platelet count went as low as 9,000. On day 12, she was flown to another hospital. She remained in that hospital until day 20 when she was discharged. She was started on Eliquis 5mg twice a day. As of this report, she was still having leg pain and difficulty walking. She was continuing to follow up with her physician. Based on evolving knowledge of Thrombosis with Thrombocytopenia Syndrome (TTS, per definition from Brighton Collaboration-BC), the low platelet count and temporal relationship to vaccination (BC Criteria level 5), the events are assessed to have a plausible relationship with vaccination."
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1325835-1	
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1325857-1	BLOOD CLOT; MULTIPLE RED SPOTS ON NECK/RASH; This spontaneous report received from a patient concerned a 19 year old Asian and not Hispanic or Latino female. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 203A21A, and expiry: UNKNOWN) dose was not reported, 1 total administered on 11-MAY-2021 14:30 to left arm for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, the patient experienced symptoms of blood clot and multiple red spots on neck. Patient reported that the red spots on neck might be a rash. The patient's father wanted to know about treatment of rash. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from multiple red spots on neck/rash and the outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210524274- Covid-19 vaccine ad26.cov2.s- A 19- year old Asian and not Hispanic or Latino female presents with Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1325868-1	BLOOD CLOTS; This spontaneous report received from a consumer concerned multiple 6 female patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported), frequency once total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the consumer stated that 6 patients experienced blood clots from the vaccine. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-20210526046-Covid 19 vaccine- blood clots. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1326824-1	3 weeks plus 1 day following vaccine, i developed pain in my left calf. After having an ultrasound, two blood clots were discovered in my left leg (one DTV and one superficial). I was sent to the ER, a CT was performed, blood work was taken, and I was sent home with a prescription for Xarelto blood thinner. I have to be on this for 3 months, and we will evaluate to make sure I'm no longer at risk. My medical bills for the date of 5/3/21 were over $11,000, and my portion so far is over $3300 after insurance.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1326951-1	She been feels ill slightly ill ever since the shot. Always short of breathe. Now she died of a blood clot / heart attack while at night in her chair.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1327350-1	Blood Clot right leg after flight.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1327541-1	The next day of vaccine on 03/31/2021 I had a severe headache, nauseas, I had diarrhea. As the days passed I started getting worst and worst I was loosing mobility in my body my husband had to take me to the restroom. On 04/06/2021 is when I went to the Emergency Room. When I got to the hospital I could not breath my ribs hurt a lot & I had horrible back pain. They put me oxygen and gave me fluids but they said I had to get transfer to another hospital and they transferred me on an ambulance. In the other hospital us where they transferred me they kept poking my stomach when I asked them why they kept poking it they told me that it was to stop the blood cloths. I was hospitalized for 5 days. When I was sent home I was still not feeling good I was feeling sick still I've been trying to recover slowly but I am not the same. I'm still in a lot of pain I can't move my arms because of how much it hurts my back keeps hurting a lot as well. I want to go back to normal to how I was I can't do anything I can't even cook for my kids. I now have depression.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1327567-1	Day 12 post-injection hard a sharp pain in lower part of torso, then immediately passed a large clot from my vagina, bright blood accompanied. Then no post-period or anything with it. I'm post-menopausal and have no women health issues at all and haven't had a period in about one year.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1327968-1	5 days after dose had pain in left calf. On May 8th went to the hospital with pain under my left rib cage. It was determined by testing that I had a blood clot in left leg and one in each lung. There was also heart arrhythmia. I was given blood thinners for the clots which I am still taking. I was also given medication to help with high blood pressure and to help with heart beat.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1328996-1	abdominal pain. The pain was so severe that my knees would buckle and several times if I was holding anything, it caused me to drop it. After about a week, I had an appointment at 4:30 and was scheduled for the next morning at 8:30 to meet with a back and spine doctor for xrays. Xrays were negative. I was then schedule with a neck and hand specialist a few days later. This appointment could not provide a diagnosis. Back and abdominal pain increase and soreness had spread to the stomach and groin. After the news broke of the cerebral blood clots in 6 women. See attached Continuation Page.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329086-1	BILATERAL PULMONARY EMBOLISM; EXTENSIVE CLOTTING TO RIGHT LEFT AND ABDOMEN; DEEP VEIN THROMBOSIS; This spontaneous report received from a patient concerned a patient of unspecified age and sex. The patient's height, and weight were not reported. The patient's past medical history included four spine surgeries, and concurrent conditions included tendinosis, high blood pressure, type 2 diabetes, and platelet rich plasma therapy, and other pre-existing medical conditions included patient had no genetic conditions which would explain the clotting. patient had no prior history of blood clots. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, frequency 1 total, administered on 21-MAR-2021 for prophylactic vaccination. Batch number was not reported and has been requested. No concomitant medications were reported. On 25-MAR-2021 and 1-APR-2021 patient underwent a series of Platelet Rich Plasma (PRP) Injections for left elbow for tendinosis. On 02-APR-2021, the patient experienced deep vein thrombosis and extensive clotting to right left and abdomen, and was hospitalized. Laboratory data included: Diagnostic ultrasound (NR: not provided) DVT and clot blood confirmed. Treatment medications included: heparin non-company. The patient was discharged on 03-APR-2021 with instructions to take Eliquis. On 07-APR-2021, the patient experienced bilateral pulmonary embolism and the patient was readmitted to the hospital. Laboratory data included: computed tomography (CT) scan (NR: not provided) Bilateral pulmonary embolism. Discharged on 09-APR-2021. On 23-APR-2021, Laboratory data included: Diagnostic ultrasound (NR: not provided) not provided. On 26-APR-2021, the patient was readmitted and thrombectomy was performed. The patient was discharged on 27-APR-2021. On 30-APR-2021, the patient experienced rethrombosis. Laboratory data included: Diagnostic ultrasound (NR: not provided) Rethrombosis confirmed. Treatment medications included: bivalirudin, where doctor placed a catheter and administered an Angiomax IV for 72 hours. On 03-MAY-2021, thrombolysis and thrombectomy surgery was performed. Patient was discharged on an unspcified date with additional treatment medications (dates unspecified) included: clopidogrel bisulfate, acetylsalicylic acid, and levofloxacin. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the extensive clotting to right left and abdomen, deep vein thrombosis, and bilateral pulmonary embolism was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210525688-COVID-19 VACCINE AD26.COV2.S-extensive clotting to right left and abdomen, deep vein thrombosis, bilateral pulmonary embolism. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY, UNDERLYING DISEASE
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329659-1	LEFT ANKLE STARTED SWELLING, LEFT ANKLE SWELLING INCREASED AND REACHED LEFT THIGH, WENT ULTRASOUND WHICH RESULT BLOOD CLOT; This spontaneous report received from a patient concerned a 47 year old female. The patient's weight was 95.70 kilogram, and height was 167 centimeter. The patient's concurrent condition included Cholesterol. Patient was non alcohol user and non smoker.. Patient had no known allergies and no history of drug abuse or illicit drug use. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry date were unknown) dose was not reported, administered on 11-MAR-2021 in right arm for prophylactic vaccination. The batch number was not reported and has been requested. Concomitant medications included pravastatin for cholesterol. Around 19-MAR-2021 (approximately 1 week after the vaccine), the patient experienced left ankle swelling (as per patient, she did not think much of it, and just elevated her leg and applied hot-cold compress). Around 26-Apr-2021 the patient experienced increased left ankle swelling and had reached her left thigh. On 26-April-2021 patient went to the hospital and was hospitalized, was given an ultrasound procedure which result stated that she had blood clot. Patient was given Eliquis right away, while in the hospital. On an unspecified date patient was discharged and upon discharge she was prescribed Eliquis 5 mg for 2 times per day. Patient stated that she will go back to the doctor will be doing another ultrasound. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot. Patient stated her left leg swelling was not yet completely resolved, but not as worse, whenever she elevates her leg, the pain and swelling gets better, but once she gets up and move around, the swelling starts again. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0:20210529680-covid-19 vaccine ad26.cov2.s ,LEFT ANKLE STARTED SWELLING, LEFT ANKLE SWELLING INCREASED AND REACHED LEFT THIGH, WENT ULTRASOUND WHICH RESULT BLOOD CLOT. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329661-1	BLOOD CLOT IN LEFT LEG; This spontaneous report received from a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced blood clot in left leg from the vaccine. Patient stated that never in life had a blood clot before The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot in left leg was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210529802-Covid-19 vaccine ad26.cov2.s-Blood clot in left leg. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329662-1	BLOOD CLOTS; This spontaneous report received from a consumer concerned 3 women with unknown race and ethnicity . The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number and expiry was unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date,3 women died from blood clots after getting (Janssen) covid 19 vaccine. It was unknown whether autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death).; Sender's Comments: V0 : 20210530454-COVID-19 VACCINE AD26.COV2.S-Blood clots . This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329664-1	BLOOD CLOTS; This spontaneous report received from a patient concerned multiple patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose 1 total ,start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On 2021, the subject experienced blood clots, and was hospitalized (date unspecified). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots leading to hospitalization was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: v0-20210530971-covid-19 vaccine ad26.cov2.s-Blood Clots. This event(s) is considered Un assessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329666-1	BLOOD CLOT; This spontaneous report received from a patient concerned a patient of unspecified ethnicity, age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown and expiry: Unknown) dose was not reported, 1 total, administered on 06-APR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On 2021, the patient developed a blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210531990-COVID-19 VACCINE AD26.COV2.S-Blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1329869-1	Bloody noses lasting more than one hour with clots
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1330508-1	patient stated he ended up in the hospital with blood clot in the lungs
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1330554-1	Upon speaking with patient here at the hospital she reports having vaccine on March 31, 2021 and reports experiencing clot that had been previously reported to CDC and she asked that this hospitalization be reported to hospitalization as concerns that it may be related. Patient admitted for possible stroke/TIA symptoms resolving
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1331063-1	I woke up Wednesday and I felt like I was hit by a train. I felt like a wet dish rag hanging on the sink. Everything hurt, even taking a step, I just felt bad, tired like the flu but no respiratory issues. By the 14th I started getting short of breath, couldn't walk without SOB, my legs started to swell, joints were sore, I felt tired. Went to the doctor and my blood pressure was high, I was prescribed medication for my blood pressure and fluid pills. It hurt just to brush against the skin on my calf. An ultrasound revealed 4 blood clots in my left calf. Went to the ER at . Hospital in and sent to hematologist Dr. and there were no findings.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1332964-1	"BLOOD CLOTS GALORE; FEELING SICK (NOT FEELING WELL); FEELING OF WOULD NOT MAKE IT AND SAYING GOODBYE TOMORROW; MUSCLE ACHES; TIRED; LETHARGIC; BODY ACHES; This spontaneous report received from a consumer concerned a 55 year old female of unknown race and ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, expiry date: UNKNOWN) dose was not reported, 1 total, administered on 16-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On 16-APR-2021 after vaccination, the patient got tired, lethargic, was not ""feeling well"" and body and muscles started to ache and progressed further. The consumer reported that on 08-MAY-2021, the patient was feeling sick and drove herself into the hospital. On 11-MAY-2021, the patient had blood clots galore and was on dialysis. On 12-MAY-2021, the consumer had one note stating the patient was holding steady and during night the consumer stated there's no change in patient's condition. On 13-MAY-2021, patient was going through with computerised tomography scan and told later that patient would not make it and would be saying goodbye tomorrow. On 13-MAY-2021, Laboratory data included: CT scan (NR: not provided) unknown. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clots galore, and the outcome of tired, lethargic, body aches, feeling sick (not feeling well) and muscle aches was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0. 20210529809 - Covid-19 vaccine ad26.cov2.s-blood clots galore. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s)."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1332971-1	BLOOD CLOTS; This spontaneous report received from a consumer concerned multiple patients (8 patients) Patients weight, height, and medical history were not reported. Patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, 8 patients got blood clots so they stopped using JANSSEN COVID VACCINE. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210533746-COVID-19 VACCINE AD26.COV2.S-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1334549-1	On early morning of 5-20-2021. Shortness of breath, felt weird, coughing up blood.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1334558-1	had redness and tightening of the legs, went to ER, found very small blood clot in right calf. Gave rx blood thinner eloquis along with current plavix and asa

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1334756-1	POSSIBLE CVA; HIGH BLOOD PRESSURE; HEADACHE; POTENTIAL BLOOD CLOT; STOMACH PAIN; LEG PAIN; This spontaneous report received from a patient concerned a 56 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included polycythemia, and Meniere's disease, and other pre-existing medical conditions included no known allergies. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 1805022, expiry: 25-MAY-2021) dose was not reported,1 total administered on 10-MAR-2021 to right arm for prophylactic vaccination. No concomitant medications were reported. On 10-MAR-2021, the patient had no side effects right after the vaccination. On 25-MAR-2021, the patient felt a really bad leg pain in left leg behind the knee which had lasted for five days until 31-MAR-2021 or 01-APR-2021. On 31-MAR-2021, the patient had very bad stomach pain and felt if somebody was stabbing her until 11-APR-2021. On 23-APR-2021, the patient had pain in head, it woke the patient up in her sleep; the patient had no control over the right side of body and was not able to see, walk or talk. The patient did not know how long it lasted. On 24-APR-2021, the patient was lying in bed and felt that somebody taken the bottom of the bed and flipped her out; she also had blurred vision. On 26-APR-2021, patient went to the hospital. The patient underwent a CT (computed tomography) scan and found no bleeding in brain but a possible CVA (cerebrovascular accident-stroke). The patient was asked to admit in the hospital but the patient wanted to go back home. The patient's blood pressure was really high although she had a perfect blood pressure and never took medication before. The patient was scared to take the pill and left the hospital. On an unspecified date, after coming back home, the patient suddenly lost control over left side of the body while taking bath and had blurred vision. The patient was taken to the hospital and found no bleeding in her brain. On an unspecified date in MAY-2021, a week prior to this report, the patient undergone phlebotomy and had to draw 500 ml of blood because patient had polycythemia and her blood was super thick. The patient also had a CT (computed tomography) scan and the results were unknown. On 13-May-2021, the patient had MRI (magnetic resonance imaging) and all were normal. On 14-MAY-2021, patient would be taking CT (computed tomography) angiography for finding out blood clot in brain. The patient was concerned about all events and was scared to find out if there would be blood clots in her brain. The PCP (primary care physician) of two hospitals she visited told her that she should not have taken the vaccine shot because of her polycythemia. On 14-MAY-2021, the patient's blood pressure was 175/112 (unit unspecified) and in the morning it was 156/117(unit unspecified). The patient was started with unspecified blood pressure medication and also acetylsalicylic acid (Aspirin) to thin her blood. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from stomach pain on 11-APR-2021, and leg pain on 31-MAR-2021, had not recovered from high blood pressure, and the outcome of headache, possible cva and potential blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0;20210530382-COVID-19 VACCINE AD26.COV2.S-Cerebrovascular accident, Thrombosis, Hypertension. This event(s) is considered not related. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1334771-1	BLOOD CLOT IN LEG; SWEATING; This spontaneous report received from a consumer concerned a 70 year old male of unspecified ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date in 2021, the patient experienced sweating which lasted about one day. It was stated that patient received vaccine around 18-APR-2021. It was also reported that actual date of vaccination may have been prior to 18-APR-2021 but definitely in April. On 17-MAY-2021, patient visited hospital and got diagnosed with blood clot in leg. The patient was on unspecified medications to dissolve clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from sweating on 2021, and had not recovered from blood clot in leg. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0- 20210534558-Covid 19 Vaccine- This case concerns a 70 yr old male. Blood clot in leg, Sweating. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1337394-1	On 5/10/21 got up with sever pain in my left side neck, shoulder and arm pain. Initially thought it was a muscle pull. Pain was out of control on Tuesday 5/11/21 Night. Took Advil, used equate cool & heat pain relieving liquid. On Wednesday 5/12 discovered I have bruises and blood clots on my back and left side. Took appointment with PCP on 5/14. Prescribed Methylprednisolone 4 MG tables for 6 days. Pain continued with blood clots and bruise marks. Developed left side ear pain, Shivering and mild fever. Visited PCP again on 5/21/21. Prescribed Cyclobenzaprine. Over the past 1 months have lost weight, had a sever pain on left leg, blurry vision.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1337662-1	Patient developed a nose bleed lasting from 7:10pm to 7:40pm on the evening that he received his vaccine. He coughed up a silver dollar sized blood clot around 7:35pm. He was feeling fine the following day, but had another nose bleed that started at bedtime the night after his vaccine. Patient's wife called on call doctor and was told to pinch his nose, which they'd been doing for 35 minutes and to go to ER if needed. Patient saw doctor, who told him to stop using Flonase for a few days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1337708-1	The day before she got her vaccine she had started her menstruation cycle. She is not on any medications, everything was normal, went to work and went to get her vaccine. She got her vaccine and around 4:15 she started gushing, which is abnormal for her and having clots the size of a half dollar and covered in blood. This is not normal for her periods, and she is now in the bathroom every hour and a half and looks like a murder scene. Her normal menstrual cycle is heavy for one day and then gradually lessens and only lasts a few days. She does occasionally have small clots with hers, but this are large and bleeding profusely. The only side effect she has is that she is extremely tired. She slept from 4:30 yesterday to 2:00 AM and feels like she could sleep the rest of the day. Her arm is a little bit sore, but otherwise fine from the injection site. Is concerned due to the heavy volume of bleeding, which is extremely abnormal for her. She has taken Tylenol and drinking a lot of fluids.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1340202-1	BLOOD CLOT; This spontaneous report received from a consumer concerned a 76 year old male. The patient's weight was 82 kilograms, and height was not reported. The patient's past medical history included angina pectoris, peptic ulcer disease, and tension headache, and concurrent conditions included asthma, atherosclerotic heart disease of native coronary artery, benign hypertension, stage 3 chronic kidney disease, chronic obstructive pulmonary disease with acute exacerbation, cirrhosis of liver, diabetes mellitus type 2, generalized osteoarthritis, morbid obesity, mixed hyperlipidemia, stenosis of bilateral carotid artery, peripheral vascular disease, malignant neoplasm large intestine, carotid artery occlusion, and hepatocellular carcinoma. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, administered on 31-MAR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. Non-company suspect drugs included: cabozantinib s-malate (tablet, oral, batch number was not reported) 40 mg, 4 times every 1 day, from 04-MAR-2021 for hepatocellular/liver cancer. Concomitant medications included apixaban, atorvastatin calcium, carvedilol, ergocalciferol, hydrocodone bitartrate/paracetamol, levothyroxine sodium, losartan potassium, mecobalamin, metformin hydrochloride, salbutamol sulfate, tadalafil, tamsulosin hydrochloride, and vitamin b complex. On 03-APR-2021, the patient experienced blood clot. On an unspecified date in Apr-2021 the patient was hospitalized for one day. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0:20210530464-covid-19 vaccine ad26.cov2.s-Blood clot. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: CONMEDS-OTHER SUSPECT DRUGS, MEDICAL HISTORY
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1340229-1	CLOTS IN LUNG; BLOOD CLOTS IN BACK; SEVERE ABDOMINAL PAIN; GROIN PAIN; This spontaneous report received from a patient concerned a 52 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown), dose was not reported, 1 total, administered on 16-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The patient reported that on an unspecified date in MAR-2021, within 5 days of the shot, the adverse reaction began with severe back pain (subsumed under blood clots in back) and abdominal pains. After two additional weeks and four doctor's appointments, the pain was so severe, patient went to the emergency room on 14-APR-2021. A computerised tomogram (CT) scan with contrast revealed clots in lung and back. Patient consulted with his doctor and decided to go to the emergency room. Patient was still suffering from severe abdominal and back pain, especially when sitting or laying down. On 2021, pain had moved to the groin area as well. Laboratory data included: CT scan (NR: not provided) revealed clots in lung and back. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from severe abdominal pain, and groin pain, and the outcome of blood clots in back and clots in lung was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210536709-COVID-19 VACCINE AD26.COV2.S-Clots in lung, Blood clots in back . This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1340597-1	I had a brain tia-stroke. Blood clots. Paralyzation.loss of vision. Loss of voice

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1340812-1	Johnson and Johnson Vaccine, shortness of breath, elevated blood pressure, elevated temperature, increased heart rate, decreased kidney functions, blood clots in both arms and jugular vein, slurred speech, severe pain in right side, fluid in right lung and around heart. Admitted to MICU from emergency room. Intubated for 12 days. Chest tubes both sides, central line, Hickman Catheter. Discharged and admitted to hospital. Currently home receiving outpatient therapy.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1341471-1	The same day I took vaccineon April 6, 2021, I had pelvic pain and cramping that increase over the next two. Weeks. I made a appointment with my physician. On 4/29/2021 I was examined by my physician and was told to go to the ER. I had my appendix removed ans the pathology report showed that I had a bllod clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1341674-1	Could not get air into lungs, made sugar get out of control, had a seizure and turned purple, passing out , ambulance had to come get me from vehicle and take me to emergency room in the hospital, kept me there for about 6 hours then released me to go home, went to my Doctor next day and he immediately sent me to hospital. I was checked for Blood Cloth due to the signs being there along with other tests I am home however I still do not feel well.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1343040-1	I went in for a scheduled MRI of my foot and the MRI found that I have a 7mm blood clot on my foot. It was found on my left foot plantar region (bottom of my foot). A few days after the vaccine I had swelling on the top and bottom of my left foot. I got two vascular images completed and the MRI was never sent to the vascular department. My podiatrist confirmed the blood clot on an ultrasound. I'm still experiencing swelling.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1344020-1	had bakers cyst burst and had 3 blood clots
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1344199-1	Blood clot in left thigh
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345688-1	BLOOD CLOT; This spontaneous report received from a consumer concerned a patient of unspecified age, sex, race and ethnic origin. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, 1 total administered for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient got blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210534929-covid-19 vaccine ad26.cov2.s-blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345702-1	BLOOD CLOT; This spontaneous report received from a patient via a company representative concerned a female of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The Company is unable to perform follow-up to request batch/lot numbers.No concomitant medications were reported. On an unspecified date, the subject experienced blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-20210537368-covid-19 vaccine ad26.cov2.s -Blood clot-his event is considered Un assessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345734-1	FEELING HORRIBLE; BLOOD CLOT; BODY ACHES; JOINT PAIN; SPAMS ALL OVER BODY; TOE SWELLING; KNEE SWELLING/ ANKLE SWELLING; This spontaneous report received from a patient concerned a 54 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 042A21A expiry: unknown) dose was not reported, 1 in total administered on 03-APR-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. After getting the vaccine, the patient had immediate joint pain and body aches. He had spasms since day one. After two weeks, he developed swollen toe looked like a bunyon, then it moved up to the ankle then it moved up to the knee being swollen. The patient was sent to the imagining center. He had following imaging tests done, bilateral Lower extremity Doppler Exam. On 30-APR-2021, he was recommended to go to the emergency room (ER) and was admitted. He was diagnosed with blood clot and stayed overnight. He was prescribed on Xarelto (rivaroxaban) 15 mg twice a day (BID), titrating to 20 mg once a day. The patient was hospitalised for 2 days. He got another follow up on 14-MAY-2021, gave him another prescription for blood thinner to be on for two months. On 18-MAY-2021, he was given tramadol 7 pills for spams which had helped with his pain and currently patient was feeling horrible. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from joint pain, body aches, toe swelling, knee swelling/ ankle swelling, spams all over body, and blood clot, and the outcome of feeling horrible was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210540030- Covid-19 vaccine ad26.cov2.s- A 54 yearold male presents with Blood clot, Body aches, Toe swelling, Knee swellng /Ankle swelling, Spasms all over body, feeling horrible. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210540030- Covid-19 vaccine ad26.cov2.s-Joint pain. This event(s) is labeled per RSI and is therefore considered potentially related.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345749-1	BLOOD CLOT; HEADACHES; This spontaneous report received from a consumer via social media/news concerned a male of unspecified age. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown exp: unknown) frequency 1 total, dose, start therapy date were not reported for prophylactic vaccination.The batch number was not reported and has been requested. No concomitant medications were reported. Reporter talked about how he saw report in the news about male who got blood clot after receiving Janssen Covid-19 Vaccine. On an unspecified date, the patient experienced blood clot, and headaches. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the headaches and blood clot was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210540327.; Sender's Comments: V0- 20210540314- Covid-19 vaccine ad26.cov2.s-Blood clot.This event is considered unassessable. The event has an unknown temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345936-1	BLOOD CLOTS IN BOTH ARMS; NUMBNESS IN BILATERAL ARMS; REDNESS ON BOTH LEGS; SWELLING OF BOTH LEGS; PAIN IN BOTH LEGS; This spontaneous report received from a patient concerned a 71 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included breast cancer. The patient experienced drug allergy when treated with hydrocodone, and oxycodone for drug used for unknown indication. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029, and expiry: unknown) dose was not reported, 1 total administered on 20-MAR-2021 on left arm for prophylactic vaccination. No concomitant medications were reported. On an unspecified date in 2021, the patient felt numbness on both arms. Numbness got worse and patient went to emergency room. On an unspecified date in 2021, Laboratory data included: MRI brain results of which were pending. On an unspecified date in MAR-2021, a week after the vaccination patient went to emergency room due to redness, swelling, and pain on both legs. The doctor prescribed antibiotics which relieved those symptoms and was encouraged to get blood circulation assessed once patient returned to home. On 17-MAY-2021, patient got hospitalized, and on same day, ultrasound test showed a blood clot in each arm. Patient underwent chest X-ray and a test with and without contrast for the lungs. It was reported that patient was started on anticoagulants and physical therapy. On 18-MAY-2021, ultrasound of legs was negative for blood clots. On 19-MAY-2021, test in a machine with and without contrast to look at lungs (Lung scan) was within normal limits. On 19-MAY-2021, the patient got discharged from hospital. Patient stated that, consulted two doctors if it may have been the Janssen covid-19 vaccine that caused this and doctors said possibly and recommended that patient should report the information. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from redness on both legs, swelling of both legs, and numbness in bilateral arms, and had not recovered from pain in both legs, and blood clots in both arms. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210540231-COVID-19 VACCINE AD26.COV2.S-BLOOD CLOTS IN BOTH ARMS, NUMBNESS IN BILATERAL ARMS. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1345947-1	BLOOD CLOTS; This spontaneous report received from a consumer by a other manufacturer company (Pfizer Inc.) received on14-MAY-2021 and concerned multiple (few) patients. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) frequency 1 total, dose, start therapy date were not reported for prophylactic vaccination. No concomitant medications were reported. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. Reporter stated that Johnson and Johnson vaccine caused blood clots and few people died. It was not reported whether autopsy was performed. On an unspecified date, the patients experienced blood clots. On an unspecified date, the patients died from blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death, and Other Medically Important Condition).; Sender's Comments: V0; 20210545044-covid-19 vaccine ad26.cov2. s Blood clots. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.; Reported Cause(s) of Death: BLOOD CLOTS
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1346470-1	Three weeks after J& J vaccine abnormally heavy vaginal bleeding with large clots.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1346853-1	2 weeks after receiving johnson and johnson vaccine I began a second period in one month. I am very regular with my menstrual cycle so it was weird. And what is very strange it has large clots and not alot of bleeding. I have never had an period like that since I am 11 years old.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1347829-1	Janssen COVID-19 Vaccine EUA I got the vaccine the evening of 03/23/21. Overnight I noticed my body starting to hurt and felt chills. The morning of 03/24/21 I woke up with full body aches, chills, and fever over 100 F. These symptoms lasted all day. Fever went up to 102 F. Recently I have noticed a change in my menses since having the vaccine and I wonder if it is linked. I have been on birth control since last year to control heavy bleeding. It was helping very well. The period I had in April and May were like it was before I started the pill. It was heavy and I had numerous large clots. I will be scheduling an appt with my ob/gyn regarding this change.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1349017-1	MULTIPLE PULMONARY EMBOLISM; MULTIPLE BLOOD CLOTS; SUSPECTED CLINICAL VACCINATION FAILURE; SUSPECTED COVID-19 INFECTION; This spontaneous report received from a health care professional concerned a 55 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry: Unknown) dose not reported, 1 total, on APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient developed multiple pulmonary embolism included multiple blood clots. On 19-MAY-2021, after 4 week of vaccination, the patient was diagnosed with suspected covid-19 infection (suspected clinical vaccination failure) and was not doing well. The patient got hospitalized. Number of hospitalization days was not reported. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the suspected covid-19 infection, suspected clinical vaccination failure, multiple pulmonary embolism and multiple blood clots was not reported. This report was serious (Hospitalization Caused / Prolonged and Life Threatening). This report was associated with product quality complaint number: 90000180239. The suspected product quality complaint has been confirmed to be not voided based on the PQC evaluation/investigation performed.; Sender's Comments: V0:20210539965-covid-19 vaccine ad26.cov2.s - multiple pulmonary embolism,multiple blood clots,suspected covid 19 infection. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). V0:20210539965 covid-19 vaccine ad26.cov2.s - suspected clinical vaccination failure This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically:SPECIAL SITUATIONS

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1349025-1	THROMBOSIS/MORE MILD CLOT ONE IN THE LEG/BLOOD CLOT DISORDER; THROMBOCYTOPENIA; INCREASING LOWER BACK; LEG PAIN; This spontaneous report received from a company representative Via social media concerned a 30 year old male. The patient's weight, height, and medical history were not reported. The patient received Covid-19 vaccine (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, 1 total dose was administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported. The Company was unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On unknown date in APR-2021, the patient experienced increasing low back pain and leg pain (around one week) after vaccination. The patient was hospitalized (date unspecified) and was treated for vaccine-induced thrombotic thrombocytopenia (VITT) (coded as thrombosis and thrombocytopenia). The patient received treatment, which includes intravenous immune globulin, the anticoagulant argatroban and prednisone. Doctors stated, patient was responding well to treatment and should be released soon (discharge details unspecified). It was reported that, patient had more mild clot, and one in the leg. But certainly it was a real clot, and as patient was in his early 30's, so it was likely and could be linked to the vaccine. The action taken with covid-19 vaccine was not applicable. The patient was recovering from thrombocytopenia, thrombosis, low back pain and leg pain. This report was serious (Hospitalization Caused / Prolonged). This is a spontaneous report concerning a 30 year old male, unknown ethnicity, patient who experienced increasing low back pain and leg pain around one week after receiving the Covid-19 vaccine. The patient was hospitalized on an unspecified date and was treated for vaccine-induced thrombotic thrombocytopenia (VITT). The patient was responding well to treatment, which included intravenous immune globulin, an anticoagulant (argatroban) and prednisone. The patient's weight, height, and medical history were not reported. There is mention of patient having vaccine-induced thrombotic thrombocytopenia but information is limited in this case. A relationship with Janssen Covid-19 vaccine cannot be ruled out; thus the relationship is considered indeterminate. Additional information has been requested.
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1349042-1	DROPPING DEAD; HEART ATTACKS; BLOOD CLOTS; STROKES; BRAIN DAMAGE; HEART CONDITIONS; This spontaneous report received from a consumer who reported reading from many personal social media accounts which concerned multiple patients of unspecified age and sex. No past medical history or concurrent conditions were reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. It was reported that, the patients were suffering from blood clots, dropping dead, had strokes, heart attacks, heart conditions and brain damage after vaccination. It was also reported that, the patients were perfectly healthy before and now they would never be the same. On an unspecified date, the patient died from unknown cause of death. It was unknown if an autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patients died of unknown cause on an unspecified date, and the outcome of blood clots, strokes, heart attacks, heart conditions and brain damage was not reported. This report was serious (Death, and Other Medically Important Condition). This case, from the same reporter is linked to 20210534943.; Sender's Comments: V0: 20210544901-covid-19 vaccine ad26.cov2.s -Dropping dead, brain damage, blood clots, heart attacks and strokes. This event(s) are considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH
THROMBOSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1349749-1	BLOOD CLOTS; HEADACHES; This spontaneous report received from a consumer who reported by seeing a news report concerned a female of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. It was reported that, on an unspecified date, the patient had blood clots, and headaches. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the headaches and blood clots was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210540314, 20210539987 and 20210547453.; Sender's Comments: V0- 20210540327- Covid-19 vaccine ad26.cov2.s-This case concerns with female of unspecified age.-Blood clot. This event is considered unassessable. The event has an unknown temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1349750-1	GOT A BLOOD CLOT IN LEG; This spontaneous report received from a patient via a company representative concerned a patient of unspecified age and sex. The patient's height and weight were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient got a blood clot in their leg and our vaccine is not good to compare to mRNA Vaccine. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of got a blood clot in leg was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210541027-COVID-19 VACCINE AD26.COV2.S-Got a Blood Clot in Leg. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1350574-1	Both sides of my neck were very swollen. There was extreme pain on my right arm. The pain was severe and numbness in my finger. I went to the ER and they said I have radial thrombus, a clot in the radial artery. Before the shot, my heart was checked multiple times and I was fine. After the shot, my coronary arteries are now very small. There are now luminal changes inside my coronary arteries which are still now very small. I now have been diagnosed with CAD. I had a heart cath on March 29. My arm is still in a lot of pain. My neck is still swollen.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1353015-1	BLOOD CLOTS; This spontaneous report received from a consumer via other company (Pfizer, report number: 2021503212) concerned multiple patients of unspecified age and sex. Patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unk) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The batch number was not reported. Per procedure, no follow up will be requested for this case. No concomitant medications were reported. On an unspecified date, reporter stated that patients experienced blood clots that scared people all the way around (no further information provided or obtained). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0 : 20210547337 - COVID-19 VACCINE AD26.COV2.S-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1353017-1	BLOOD CLOTS; HEADACHE; This spontaneous report received from a consumer who reported by seeing a news report concerned a 14 female of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patients received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. It was reported that, on an unspecified date, 14 patients had blood clots, and headaches The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the headache and blood clots was not reported. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210540327.; Sender's Comments: V0:20210547453-covid-19 vaccine ad26.cov2.s-This case concerns to 14 female of unspecified age. Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1353762-1	BLOOD CLOT; DISTURBING CHANGES IN MENSTRUAL CYCLES; EARLY PERIODS; UNUSUALLY HEAVY PERIODS; OTHER IRREGULARITIES; This spontaneous report via social media received from a patient concerned multiple patients. The patient's height and weight was not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. The reporter shared from a news article stating that blood clots were not the only vaccine side effects and that women experienced several weird side effects including disturbing changes in menstrual cycles, early and unusually heavy periods and other irregularities. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the other irregularities, early periods, unusually heavy periods, blood clot and disturbing changes in menstrual cycles was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210545709-COVID-19 VACCINE AD26.COV2.S-blood clots. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1354724-1	bi-lateral leg pain within 1 week of shot. then severe back pain after 2 weeks. difficulty breathing week 3. Admitted to hospital for multiple blood clots in legs & lungs.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1354733-1	on 5/17/2021 pt developed a fever, pain in the left side of his ribs and had SOB. He was brought to Hospital ER. Pt had Chest x-ray, blood work, Doppler on both legs, and CT. His DX was pneumonia, blood clot in left lung and bilateral blood clot in legs. He was admitted and stayed for 5 days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1354760-1	It was really bizarre, I remember feeling like all my senses were heightened , I got my period, it didn't stop for 3 weeks. Called my OBGYN, they said let it run it's coarse and they had they had many similar phone calls. I take a Nuva ring for birth control, I was concerned about estrogen and the blood clots. I did not feel well at all for about a week after my vaccine, everyday around 4pm I would get really bad headaches for about 3-4 weeks. I was nauseated and sweating.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1354801-1	About 1 week after vaccination. I had pain in my legs and swelling in my feet, it made it difficult to walk. It was like I was carrying around a dead leg. I did have blood clots in leg. I ended up up being hospitalized 3 days to receive blood transfusions due to anemia after taking blood thinners.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1354808-1	2 days after applying the vaccine I felt body pain, then a week later I began to have pressure in my chest that reached my throat so I was for a full week the third week I decided to go to the hospital and they discovered that I had blood clots in an artery in my heart. Then they installed 2 stents in my heart 2 days later they sent me home and 1 week later I had to return to the hospital after the studies they detected that I had blood clots in both legs, I had no problem before the vaccine and I'm almost 100 percent sure this is what caused these problems.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1355551-1	"April 13 (I had been traveling before it and came back) and I was preparing for work and my foot hurt. I hobbled around that day and noticed my foot was swollen. The next morning, I went to Piedmont Hospital ER(on the 14th) - they gave me a starter dose of blood thinners and I was able to schedule appt with Dr. Shamm -my hematologist - Zeralto - was what she prescribed. Everything is ""back to normal."" But based on my health history - the doctor care will be ongoing and I'll continue to see this doctor for treatment."
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1357423-1	BLOOD CLOT IN RIGHT ARM; ARM SWOLLEN; This spontaneous report received from a patient concerned a 56 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included high blood pressure, alcohol use (beer at home), and smoker. The patient experienced drug allergy when treated with acetylsalicylic acid. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown and expiry: unknown) dose was not reported, 1 total, administered on 05-APR-2021 to right arm for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The patient did not feel sick after vaccination. On an unspecified date in 2021, the patient believed that, she had a blood clot in right arm. In past 2-3 weeks, it had been getting bigger and was moving up in her arm. She probably had the clot for longer but started noticing it more 2-3 weeks ago. Clot was moving from her elbow to the shoulder blade and arm was getting big. Patient did not have any pain but her arm was swollen and due to dark skinned she could not tell if her arm was red. Patient called emergency medical service and told them that she had a blood clot, she could feel and see the blood clot and it was told that she needs to go to the doctor but she did not bother. The blood clot kept moving up. Patient had a regular doctor but she did not inform her doctor. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clot in right arm and arm swollen was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210547027-covid-19 vaccine ad26.cov2.s- blood clot in right arm. This event is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1357434-1	DEATHS; BLOOD CLOTS; This spontaneous report received from a consumer who reported reading and seeing on the news concerned a patient of unspecified age, race, ethnic origin and sex. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number not reported, expiry not reported) frequency one total, dose, therapy start date were not reported administered for prophylactic vaccination. The batch number was not reported. Per procedure, no follow-up will be requested for this case. No concomitant medications were reported. On an unspecified date, the consumer stated that he read and saw on the news that this vaccine causes patient's deaths, and blood clots. On an unspecified date, the patient died from unknown cause of death. It was not reported whether autopsy was performed or not. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient died of deaths on an unspecified date, and the outcome of blood clots was not reported. This report was serious (Death, and Other Medically Important Condition).; Sender's Comments: V0: 20210550011-covid-19 vaccine ad26.cov2.s-deaths, blood clots. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: UNKNOWN CAUSE OF DEATH
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1361603-1	Stroke .. blood clot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1362194-1	Sudden numbness in leg 5:20pm. Went to emergency room via ambulance. Did a CT scan, found blood clot in upper leg. Sent via ambulance to Hospital had Surgery at 10pm to remove clot and save leg.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363777-1	BLOOD CLOT IN BRAIN; BLOOD CLOT IN RIGHT LUNG; BLOOD CLOT IN STOMACH AND NECK; This spontaneous report received from a patient concerned a 64-year-old, female. The patient's height, and weight were not reported. The patient's past medical history included tendonitis in right shoulder, surgeries on right shoulder and concurrent conditions included arthritis in right shoulder, non-smoker, abstains from alcohol, and seasonal allergies, and other pre-existing medical conditions included the normal wear and tear on her body at the time of this report. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, one total, administered on 04-MAY-2021 at the right arm for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. The Patient reported that on 04-MAY-2021, after vaccination, she began feeling unwell with a low grade fever, her highest body temperature was 102F. She experienced headache that she initially felt on the right side of her head but could not determine the source for sure. On 14-MAY-2021, she coughed up blood, but she didn't think anything of it and continued her normal routine then on 15-MAY-2021, she coughed up more blood but it had some phlegm in it, so she went to the ER (emergency room) on 18-MAY-2021. She received a Computerized axial tomography (CAT) scan of her chest which showed blood clots in her right lung. She got a secondary CAT scan and blood clots were found in her stomach, brain and in her neck. She was admitted to hospital on 19-MAY-2021. The Patient was reportedly hospitalized for 6 days and was supposedly discharged on 24/May/2021. For treatment, she was receiving an Intravenous (IV) blood thinner but later changed to an oral blood thinner. She was also receiving Tramadol for pain. She was receiving other medications when first admitted but could not recall specific names. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot in brain, blood clot in right lung, and blood clot in stomach and neck. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210546778-covid-19 vaccine ad26.cov2.s- blood clot in brain, blood clot in right lung, and blood clot in stomach and neck. This events are considered unassessable. The events has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363787-1	BLOOD CLOT IN LEG; This spontaneous report received from a consumer concerned a 73 year old female unspecified race and ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: Unknown, expiry: unknown) dose was not reported, 1 total, administered on 16-MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On 13- MAY-2021, the patient was taken to the emergency room (ER). On 14-MAY-2021, she was hospitalized with a clot/clots in her leg(s). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot in leg was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0-20210551429 - Covid-19 vaccine ad26.cov2.s-Blood clots in leg. This case concerns with 73 years old female. This event is considered unassessable. The event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363797-1	BLOOD CLOT; This spontaneous report received from a consumer concerned a 24 year old female. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, therapy start date were not reported, with frequency 1 total administered for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date in APR-2021, the patient experienced blood clot and was hospitalized (date unspecified) and it was reported caller said his brother would called to provide more information regarding the patient's case. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210552268-COVID-19 VACCINE AD26.COV2.S-BLOOD CLOT. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363809-1	BLOOD CLOT FROM GROIN AREA ALL THE WAY TO THE LEG; This spontaneous report received from a patient concerned a 63-year-old female. The patient's height, and weight were not reported. The patient's concurrent conditions included osteoporosis, osteoarthritis, bursitis, rheumatoid arthritis (RA), migraine headache, anxiety, and cholesterol, and other pre-existing medical conditions included patient had no history of blood clots and no prior heart conditions. On 16-Mar-2021 around 09:00, the patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805029, expiry: Unknown) dose was not reported, frequency 1 total, administered on left arm for prophylactic vaccination. Concomitant medications included pantoprazole for acid reflux, citalopram for anxiety, simvastatin for cholesterol, and prn 1371 for migraine headache. On an unspecified date in Feb-2021, the patient mentioned (heart doctor) had done some tests on veins making sure the patient's heart was open and does not have any blood clots. Laboratory data included: Diagnostic ultrasound (NR: not provided) Normal, and Doppler ultrasound (NR: not provided) No blood clots. 7 days later the patient was informed everything was fine and she got vaccinated on 16-Mar-2021. On 07-May-2021, the patient reported that her left leg started hurting. On 09-May-2021, the leg started swelling. On 11-May-2021, the patient was admitted to the hospital with a massive blood clot from groin area all the way to the leg. On 14-May-2021, she was discharged from hospital. The patient was hospitalized for 3 days. The patient stated her doctor will let her know if she can move around more next week, the patient is currently in lot of pain. Treatment medications (dates unspecified) included: apixaban, paracetamol, tramadol hydrochloride, gabapentin, and baclofen. The action taken with covid-19 vaccine ad26.cov2. s was not applicable. The patient had not recovered from blood clot from groin area all the way to the leg. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0-20210500586-COVID-19 VACCINE AD26.COV2.S-This case concerns with 63 year old female û blood clot from groin area all the way to the leg -This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363813-1	BLOOD CLOTS IN BOTH LUNGS; LEFT SIDE OF BODY HURTING; GENERALIZED WEAKNESS; TIREDNESS; BLOOD CLOT IN BOTH LEGS; TROUBLE WALKING; This spontaneous report received from a consumer concerned a 77- year old White Not Hispanic or Latino male. The patient's height and weight were not reported. The patient's past medical history included polyp removal on the nose, and concurrent conditions included prostate cancer, asthma, chronic obstructive pulmonary disease, allergic to sulfa drugs, alcohol user (drinks wine 1-2 times a week), and non-smoker, and other pre-existing medical conditions included no drug abuse or illicit drug usage. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 205A21A expiry: UNKNOWN) 1 total, the dose was not reported, administered on right arm on 09-MAY-2021 for prophylactic vaccination. Concomitant medications included prednisone for loss sense of taste, and ascorbic acid/biotin/cyanocobalamin/folic acid/nicotinamide/pantothenic acid/pyridoxine hydrochloride/riboflavin/thiamine hydrochloride, and vitamins and minerals. On 14-MAY-2021, the patient began experiencing symptoms of both legs hurting from hips to knees and generalized body weakness as well as tiredness. a Patient complained about legs that whole week stating was in a lot of pain, having trouble walking. Consumer stated pain level was around at a 7 or 8 and that patient had a lot of difficulties getting out of the car and not being able to walk from kitchen to the bedroom. They were concerned. On, Monday 24-May-2021, the patient stated whole left side of the body was hurting and could not put any pressure or sit on the left side of the body. the Patient also had shortness of breath and chest pain on Monday 24-May-2021. Patient was hospitalized on 25-May- 2021 and found multiple clots in bilateral lungs and both legs. The patient was hospitalized for 2 days. Laboratory data included: Blood test (NR: not provided) blood clots bilateral lungs and both legs, Computerised tomogram scan (NR: not provided) blood clots bilateral lungs and both legs, Venous Doppler (NR: not provided) blood clots in legs, and X-ray (NR: not provided) blood clots bilateral in lungs. On 26-MAY-2021, Laboratory data included: Echocardiogram (NR: not provided) clear. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clots in both lungs, blood clot in both legs, trouble walking, left side of body hurting, generalized weakness, and tiredness. This report was serious (Hospitalization Caused / Prolonged); Sender's Comments: V0:20210554302- blood clots in both lungs, blood clot in both legs, trouble walking, left side of body hurting -This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY, UNDERLYING DISEASE V0:20210554302- generalized weakness, and tiredness.This event(s) is labeled per RSI and is therefore considered potentially related.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363851-1	BLOOD CLOT; STROKE; This spontaneous report received from a patient concerned a 46 year old male. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported and batch number: 202A21A expiry: 23-JUN-2021) dose was not reported, 1 total administered on 07-MAY-2021 for prophylactic vaccination. No concomitant medications were reported. On 19-May-2021, the patient begun to experiencing confusion and could not think straight. On 23-MAY-2021, the symptoms progressed when the patient went to the hospital. The patient was diagnosed with blood clot and had stroke. The patient was hospitalized on 23-MAY-2021 and was discharged on 26-MAY-2021. The patient was hospitalized for 3 days. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clot and stroke was not reported. This report was serious (Hospitalization Caused/Prolonged).; Sender's Comments: V0: 20210554839-covid-19 vaccine ad26.cov2.s-blood clot and stroke. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363896-1	PAIN; PITTING EDEMA; SWELLING; BLOOD CLOT LEFT AND RIGHT LEG; This spontaneous report received from a patient concerned a 66 year old female. The patient's height, and weight were not reported. The patient's past medical history included factor 5 positive.The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805022, expiry: UNKNOWN) dose was not reported, administered on 08-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, the patient experienced pain, pitting edema, and swelling. The patient had an Ultrasound done on 02-MAY-2021 at the emergency room that showed a blood clot in her left leg. On 27-MAY-2021 she went to her doctor and had another ultrasound on her right leg that indicated a blood clot. She was started on Zorelto as a treatment for blood clot on 02-May-2021 15 mg BID Three times daily and is now taking 20mg 1 times daily, but that she may not be able to stay on Zorelto because it was not working. Also stated this vaccine had ruined her life and was very sorry that she ever got the vaccine. She was extremely frustrated and unhappy and had pain and swelling that has not allowed her to play tennis and experience inactivity. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from blood clot left and right leg, and the outcome of pain, pitting edema and swelling was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: 20210556299-covid-19 vaccine ad26.cov2.s-Blood clot left and right leg. This event(s) is considered not related. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There are other factors more likely to be associated with the event(s) than the drug. Specifically: MEDICAL HISTORY
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1363921-1	BLOOD CLOTS OF LUNGS; BLOOD CLOTS OF LEGS; This spontaneous report received from a patient via a company representative concerned a patient of unspecified age and sex. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown, Expiry: Unknown) dose, start therapy date were not reported, 1 total for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient ended up getting blood clots of lungs and legs and was wanting to know more information. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the blood clots of lungs and blood clots of legs was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210556674-COVID-19 VACCINE AD26.COV2.S-Blood clots of lungs, blood clots of legs. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1366784-1	BELL'S PALSY; BLOOD CLOT; GUILLAIN BARRE SYNDROME; ANAPHYLAXIS; This spontaneous report received from a patient via a company representative (other manufacturer Pfizer) concerned multiple patients. The patient height and weight was not reported .No past medical history or concurrent conditions were reported.The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin, and batch number were not reported)frequency one total , dose, start therapy date were not reported for prophylactic vaccination.The batch no was not reported ,The Company is unable to performed follow up to request batch /Lot numbers. No concomitant medications were reported. On an unspecified date, the subject experienced bell's palsy, blood clot, guillain barre syndrome, and anaphylaxis. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the bell's palsy, blood clot, guillain barre syndrome and anaphylaxis was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0. 20210556206-COVID-19 VACCINE AD26.COV2.S. Bell's palsy, Blood clot, Guillain Barre syndrome. This events is considered unassessable. The events has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events. 20210556206-COVID-19 VACCINE AD26.COV2.S. Anaphylaxis. This event is labeled per RSI and is therefore considered potentially related.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1366787-1	BLOOD CLOT; BELLY BUTTON PAIN; LOWER LEG PAIN; STOMACH PROBLEMS; This spontaneous report received from a patient concerned a 57 year old white(Not Hispanic or Latino) female. The patient's height, and weight were not reported. No past medical history was reported. The patient had no known allergies. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1805025,expiry: UNKNOWN) dose was not reported, administered on 09-MAR-2021 for prophylactic vaccination. No concomitant medications were reported. On 21-MAY-2021, the patient experienced stomach problems(pain, cramping, bloating). On an unspecified date, the patient experienced blood clot, belly button pain, and lower leg pain. The patient did not made any changes to diet and no new medications took since getting the vaccine. The patient had tried yogurt and probiotic. Treatment medications (dates unspecified) included: bismuth subsalicylate for unknown indication. On 28-May-2021, she started taking turmeric, but so far nothing had resolved the issues. The patient had a tele visit with family doctor of medicine and would go next week wednesday. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from blood clot, and lower leg pain, and had not recovered from stomach problems, and belly button pain. This report was serious (Other Medically Important Condition). This case, from the same reporter is linked to 20210556734.; Sender's Comments: V0 20210556510- COVID-19 VACCINE AD26.COV2.S- blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1367531-1	BLOOD CLOTS; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age, race and ethnic origin. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210553711-COVID-19 VACCINE AD26.COV2.S-Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1367532-1	BLOOD CLOTS INTO THE LUNGS; BLOOD CLOTS ON RIGHT ARM AND SHOULDER; FLU LIKE SYMPTOMS; DISCOLORATION ON THE RIGHT ARM AND IT TURNS BLUE; SWOLLEN RIGHT ARM; This spontaneous report received from a patient concerned a 26 year old. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 207A21A, and expiry: UNKNOWN) dose was not reported, 1 total dose administered on 07-APR-2021 in left arm for prophylactic vaccination. No concomitant medications were reported. On 09-APR-2021, two days later after getting the vaccine patient had experienced flu like symptoms. After 3 weeks, patient had discoloration on the right arm and it turned blue and swollen. On 03-MAY-2021, the patient was hospitalized. Patient was administered through the emergency care on 05-MAY-2021. Then the patient got transferred to the floor. Blood clots were formed on arm, shoulder, and into the lungs. At that moment patient had shortness of breath. Patient was hospitalized for 5 days and discharged on 07-MAY-2021. Treatment medications (dates unspecified) included: apixaban. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from swollen right arm, had not recovered from discoloration on the right arm and it turns blue, blood clots into the lungs, and blood clots on right arm and shoulder, and the outcome of flu like symptoms was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0:20210554390-covid-19 vaccine ad26.cov2.s- blood clots into the lungs, and blood clots on right arm and shoulder. These events are considered unassessable. The events have a compatible/suggestive temporal relationship, are unlabeled, and have unknown scientific plausibility. There is no information on any other factors potentially associated with the events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1367535-1	BLOOD CLOTS IN HEART; BLOOD CLOTS IN LUNGS; BLOOD CLOTS IN LEGS; CHILLS; SORE ARM; This spontaneous report received from a patient concerned a 55 year old male. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, and batch number: 1808982 expiry: unknown) dose was not reported, administered on 06-APR-2021 for prophylactic vaccination. No concomitant medications were reported. It was reported by patient that on 06-APR-2021 patient experienced sore arm and on 07-APR-2021 patient had experienced chill. The patient stated that on 27-MAY-2021, the patient experienced chest pain which led to him hospitalization on an unspecified date. While hospitalized they discovered that he had blood clots in the heart and lungs (also reported as blood clots in the legs). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from sore arm, and chills, and the outcome of blood clots in heart, blood clots in lungs and blood clots in legs was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0:20210557848-covid-19 vaccine ad26.cov2.s-Blood clots in heart, Blood clots in lungs, Blood clots in legs. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1369341-1	Review of ED notes show she received IV infusion of normal saline, and IV push of benadryl and compazine after going to the ER for severe headache on 04/17/2021. She first noticed pain when streching the arm shortly after the IV infusion but didn't pay attention to it, then 5 day later she noticed localized swelling and redness. Patient was sent for US doppler which left cephalic vein thrombus detected on duplex US. Was treated with aspirin and had follow up doppler of UE for follow up on LUE cephalic vein thrombosis-negative 5/13/21.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1370292-1	BLOOD CLOTS; This spontaneous report received from a patient concerned a male of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine (suspension for injection, route of admin not reported, batch number: unknown) dose was not reported, with frequency 1 total administered on 08-APR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots. It was reported that the patient experienced blood clots after receiving the vaccination for which the prison has failed to provide the medical treatment. The action taken with covid-19 vaccine was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: covid-19 vaccine Blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1370997-1	Systemic: Blood Disorder (diagnosed by MD)-Severe, Additional Details: Patient was diagonised with blood clot few days after J&J covid vaccine. Patient was treated in the hospital and doing better.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1371273-1	am facing stomach pain and some blood clots and gas problem at stomach
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1371683-1	The day after, I woke up with a bad headache and I had the headache for a good 19 days. The doctors thought I was over-reacting. It was a bad headache on left side of head. I was just not feeling well. On the 19th day, it went away. I still from time to time I do still feel pain in the head on the left side. I had a lot of tightness in my left leg. In previous years, I have had blood clot in my leg and after birth in my stomachs - so with the pain in calf I called a doctor and went. They did a visual. After a lump appearing in my leg, an ultrasound was scheduled. I went Urgent care on May 10 - she ordered an ultrasound but her office didn't ever follow up with me. On 17th, I saw my Primary care and she sent me straight to get an Ultrasound - on the 18th, it revealed I had a blood clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1371791-1	THE MORNING AFTER MY VACCINE MY TOES AND LEGS WERE NUMB AND TINGLY AND FELT HEAVY AND FELT WEIRD WALKING. I HAVE NEVER FELT THIS FEELING BEFORE, IT WENT AWAY PRETTY MUCH AFTER A COUPLE DAYS AND THEN CAME BACK A WEEK LATER E VEN WORSE THEN BEFORE AND STAYED FOR AT LEAST 3 DAYS SO I WENT TO URGENT CARE AND THEY SAID IT WAS NOT A BLOOD CLOT AND THEY DIDNT KNOW MUCH ABOUT THE VACCINE. NOW OVER 3 WEEKS LATER I STILL HAVE NUMBNESS IN MY TOES AND LEGS. IT COMES AND GOES, SOME DAYS ARE WORSE THAN OTHERS.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1372605-1	i had a blood clot which caused a stroke

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1373240-1	TWO BLOOD CLOTS; LEFT CALF PAIN FOR A COUPLE OF DAYS/FELT LIKE MUSCLE STRAIN; This spontaneous report received from a patient concerned a male of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 042A21A, expiry: unknown) dose was not reported, 1 total, administered on 03-APR-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. On 25-MAY-2021, the patient experienced left calf pain for a couple of days which the patient believed that it was a muscle strain and was increasingly painful. On 28-MAY-2021, the patient was sent to the hospital to do an ultrasound and the HCP (health care professional) suggested for the patient to go to the emergency room visit because the patient had two blood clots. On 28-MAY-2021, the patient was hospitalized and was discharged on 28-MAY-2021. The patient was hospitalized for 1 day. Laboratory data included: Diagnostic ultrasound (NR: not provided) 2 blood clots. On the day of reporting, it was informed that the patient was taking treatment medications (dates unspecified) which included: Eliquis (apixaban) for blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from two blood clot, and the outcome of left calf pain for a couple of days/felt like muscle strain was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210601553-covid-19 vaccine ad26.cov2.s - two blood clots in patient. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1374698-1	Swelling of the right foot and leg. Went to ER and they performed a doppler ultrasound and found a blood clot in the lower right leg. Prescribed blood thinner. So far, the swelling has gone down
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1375949-1	My menstrual cycle has been incredibly irregular. I have been having breakthrough bleeding and extremely heavy clotting which I have never had before. I have been bleeding for 25 days straight at the time of reporting this, and my clots are getting bigger and the bleeding is getting heavier.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1376342-1	Patient called today, He stated that 18 hours after his dose of Janssen he had a nose bleed that lasted 8 hours and expeled 2 large blood clots from his nostils. I asked if he had seen a physician and he had only called his friend who was a PA. I encouraged him to contact his physician and inform him/her of the situation.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1376703-1	Blindness in left eye, blood clot in left eye- no treatment available. Permanently blind in left eye now.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1380631-1	BLOOD CLOTS; This spontaneous report received from a consumer via social media concerned a male of unspecified age with unknown race and ethnicity. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, once total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0 : 20210602521-COVID-19 VACCINE AD26.COV2.S - Blood clots. This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1380635-1	SEVERE BLOOD CLOTTING; OTHER ISSUES; This spontaneous report received from a consumer via a company representative concerned a female of unspecified age and unknown race and ethnicity. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: not reported) dose, 1 total, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient experienced severe blood clotting and other issues (coded to unspecified adverse event) after receiving the one-dose COVID-19 vaccine. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the severe blood clotting and other issues was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210604196-covid-19 vaccine ad26.cov2.s-Severe blood clotting. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1380641-1	BROKEN HIP; HAIRLINE FRACTURE IN ARM; FELL; PAIN IN BACK OF CALF; BLOOD CLOT; MILDLY SORE ARM AROUND INJECTION SITE; This spontaneous report received from a consumer concerned a 67 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included shoulder pain. The patient did not have a history of blood clots, and was not pregnant at the time of vaccination. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1816022 expiry: Unknown) dose was not reported, 1 total, administered on 09-MAY-2021, about 17:00 pm, to left arm for prophylactic vaccination. Concomitant medications included ibuprofen for shoulder pain. On an unspecified date in MAY-2021, the patient had mildly sore arm around injection site which had resolved (Unspecified date). On 11-MAY-2021, two days after receiving the vaccine, about 8:00 am in the morning, the patient fell and broke her hip and arm. The hip needed a rod and the patient had surgery to repair the hip by 17:00-18:00pm, the arm was a hair-line fracture and the patient was hospitalized. About a week and a half after the surgery, the patient experienced pain in back of calf, had an ultrasound, and found blood clot. She was put on blood thinners (unknown name). The blood clot was resolving, breaking up and going away. The reporter did not think that blood clot had anything to do with suspect company. On 02-JUN-2021, the patient was discharged from the rehab within the hospital, where she had her surgery and other care. The patient was hospitalized for 23 days. Its marked 3 weeks since she fell, she had very health exercises and does boot camps. The patient could now take 2 steps with walker, could stand up and pivot, go to bathroom by herself, physical therapy would be given at the patient's house, and it is expected that she will make full recovery. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient recovered from mildly sore arm around injection site, was recovering from blood clot, and the outcome of fell, broken hip, pain in back of calf and hairline fracture in arm was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210605202-Covid-19 vaccine ad26.cov2.-Blood clot ,Broken hip, Hairline fracture in arm, Fell. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210605202-Covid-19 vaccine ad26.cov2.-Pain in back of calf. This event(s) is labeled per RSI and is therefore considered potentially related.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1380658-1	BLOOD CLOT IN RIGHT CALF; BLOOD CLOT IN LEFT LUNG; COULDN'T WALK; FATIGUED WHEN WALKING; MUSCLE PAIN; WALKING SLOW / STUMBLING; LEFT CALF PAIN; This spontaneous report received from a consumer concerned a 40-year-old female of unknown race and ethnicity. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number:1808980 00040604, expiry: not reported) dose was not reported,1 total, administered on 09-APR-2021 for prophylactic vaccination. No concomitant medications were reported. In APR-2021, about a week after vaccination, the patient experienced right calf pain/right lower leg pain and muscle pain. Patient was unable to walk, experienced fatigue while walking and was out of breath/unable to breath and on 20-APR-2021, the patient went to the urgent care (Emergency room) and then to the hospital. At hospital, patient had done Ultrasound Test and found blood clot in right calf (right leg) and in left lung. Patient was unable to breath. At hospital, patient was given with medication for blood clotting. After taking medication, patient was able to take breath. On 21-APR-2021, patient was discharged from hospital with medication to prevent blood clotting. Patient was hospitalized for 2 days. Patient was taking the medication one each in morning and evening. On 02-JUN-2021, patient was still walking slowly, tumbling and was not able to walk. On same day, patient also had a pain in left calf and fatigued. On 03-JUN-2021, patient went to the Emergency room(ER). Laboratory data included: Ultrasound scan (NR: not provided) which showed a blood clot in the right leg and left lung. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from fatigued when walking, had not recovered from couldn't walk and left calf pain, and the outcome of walking slow / stumbling, muscle pain, blood clot in right calf and blood clot in left lung was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0: 20210607546-covid-19 vaccine ad26.cov2.s -Blood clot in right calf, Blood clot in left lung, Couldn't walk. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s). 20210607546-covid-19 vaccine ad26.cov2.s -Fatigue when walking, muscle pain. This event(s) is labeled per RSI and is therefore considered potentially related.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1383640-1	Shortly after vaccine, upper respiratory issue started. Fatigue, falling, soiling bed sheets and pants regularly. Then went to hospital on june 7th with abdominal pain. Low bloid pressure and hight white blood count. Had blood clot. Urinary Infection Possible sepsis. Still in hospital.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1385289-1	5/21/21 10:00pm shortness breath, chest pain, shoulder pain, jaw pain, metallic taste, fatigue ?mini heart attack? symptoms 5/26-5/27 trouble breathing, fatigue, light headed 5/29 pain in left arm 5/30 night - second ? mini heart attack? symptoms General fatigue, hard to breath, overall feeling bad 6/2 - bruise on arm and lump Extreme headache, fatigue, chest pain, arm pain Emergency room visit = blood clots

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1387017-1	Four days after receiving her Covid vaccination patient started bleeding vaginally. Had blood clots and large amount of blood and it lasted 2 weeks. Patient has been post-menopausal for 10+ years.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1387475-1	LOSS OF APPETITE; REALLY ILL; LACK OF TASTE; COULD NOT WALK DUE TO PAIN; LEG FELT LIKE IT WAS BURNING; BLOOD CLOT; This spontaneous report received from a patient concerned a female of unspecified age, unknown race and ethnicity. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported and batch number: 180890 expiry: 20-JUN-2021) dose was not reported, 1 total, administered on 06-APR-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. On 06-APR-2021, about 30 minutes after the vaccination, the patient noticed her ankle bruising that started to hurt real bad. The entire ankle and part of her leg was severely bruised due to the blood clot in her leg as diagnosed by her doctor. The bruised changed her life for the worse and was extremely embarrassing. It still hurts when the site was touched. The doctor told the patient that her bruise was going to be permanent. On an unspecified date, the patient experienced loss of appetite, as if she got the virus and was really ill, lack of taste (could not taste anymore), could not walk due to pain The doctor was trying to break up the dry blood. Her leg felt at one point like it was burning. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from loss of appetite and lack of taste, and the outcome of blood clot, really ill, could not walk due to pain and leg felt like it was burning was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210605693-covid-19 vaccine ad26.cov2.s-blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1388886-1	pt started with arm pain that would not go away. On May 6, 2021 her arm swelled up to 3 times its size. She went to an urgent care and was diagnosed with a blood clot. She was given a prescription for Eliquis. She was not hospitalized and is doing OK now. She reported this to us on 06/10/2021
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1389031-1	I developed a blood clot. I went to an urgent care and was directed to the emergency room.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1391499-1	Received vaccine on 6/9/2021, the next day she started feeling cramp in left leg and swelling. Went to see her physician. Diagnosed with blood clots to left leg. Given 1 lovenox shot at time of visit and put on 81 mg ASA twice a day for 2 days.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1391726-1	Blood clots in right leg and both lungs
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1391789-1	BLOOD CLOTS IN EYE
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1391958-1	shortly after (few weeks) receiving the vaccine, I experienced a large amount of pain in my right forearm which resulted in a painful lump later identified as a thrombosis which had to be removed. It was removed on 5/27/21.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1394032-1	THROMBOSIS OF THE CLOT; INCREASED FREQUENCY OF MIGRAINES/VERTIGO; STROKE; This spontaneous report received from a consumer concerned a female of unspecified age. The patient's height, and weight were not reported. The patient's past medical history included vertigo, and concurrent conditions included lupus antiphospholipid syndrome, and migraine. The patient had no known allergies. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose was not reported, administered on APR-2021, 1 total dose for prophylactic vaccination. Concomitant medications included Xarelto (rivaroxaban) for lupus antiphospholipid syndrome. On APR-2021, the patient developed thrombosis of the clot and increased frequency of migraines/vertigo (over the past month or two). On 04-JUN-2021 (3 night ago), she was diagnosed with stroke. Later, she underwent procedure for thrombosis of the clot and she was stable. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from stroke, and thrombosis of the clot, and the outcome of increased frequency of migraines/vertigo was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210617520-covid-19 vaccine ad26.cov2.s-Stroke, Thrombosis Of The Clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1395994-1	I had no regular symptoms, but a few days later I sat down in my chair and it felt like someone shot me in the stomach, I took oxycodone, it kept getting worse and I dealt with it for a week. I started getting pain in my kidney and went to the ER because of the pain and the pain level on a scale of 1-10 was a 8-9 at all times. I couldn't' breathe and was rushed to the hospital. I have portal vein thrombosis and have three blood clots in my stomach. I tried to figure out what to do to relieve the pain, I couldn't eat or during and was on IV's. I was in so much pain to even eat a cracker, they kept me at the hospital. While in the hospital they did a CAT scan to see if it was diverticulitis and found out that I had 3 blood clots, they also did the dye or contrast. I spent days in the hospital and was transferred to the hospitals skilled nursing facility so that they can watch me because of the clots. I was told that the blood clots could burst in the vein.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1396453-1	Blood clot in right leg - 6.5?
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1398437-1	BLOOD CLOT IN ARM; HARD TIME BREATHING/COULDN'T BREATH; HIGH FEVER; This spontaneous report received from a a 63-year-old female patient of unspecified of race and ethnic origin and her daughter via a traditional media and via a company representative. The patient's height, and weight were not reported. The patient's concurrent condition included diabetes. The patient never tested positive for coronavirus disease (COVID). The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration was not reported, batch number: unknown, and expiry date: unknown) dose was not reported, 1 total, administered on APR-2021 for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, in APR-2021, shortly after the vaccination, the patient experienced a high fever. A day later, the patient had a hard time breathing/could not breath and was rushed to the hospital (date unspecified). The patient also developed a blood clot in her arm. The doctors told the patient's kids that the patient might not recover. In a hospital bed, the patient spent 21 days on a ventilator. The patient was recovering in the hospital in the last month of MAY-2021 and was expected to leave the rehabilitation facility by the end of JUN-2021. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient was recovering from high fever, hard time breathing/couldn't breath, and blood clot in arm. This report was serious (Hospitalization Caused / Prolonged, and Life Threatening).; Sender's Comments: V0: 20210619186-COVID-19 VACCINE AD26.COV2.S-Blood clot in arm, Hard time breathing/couldn't breath, High fever. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1398450-1	BLOOD CLOT; This spontaneous report received from a patient concerned a male of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine (suspension for injection, route of admin not reported) 1 total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient had blood clot. The action taken with covid-19 vaccine was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210621271- covid-19 vaccine Blood clot. This event(s) is considered unassessable. The event(s) has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1398645-1	BLOOD CLOTS; This spontaneous report received from a patient concerned a female of unspecified age, race and ethnic origin. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported. The company is unable to perform follow-up to request batch/lot no. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots, and was hospitalized (date unspecified) for 3 to 4 weeks. Treatment medications (dates unspecified) included: Xarelto (Rivaroxaban). The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clots was not reported. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: V0 20210614970-COVID-19 VACCINE AD26.COV2.S-blood clots. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1399289-1	Pain, fever
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1401117-1	Shortness of breath chest pain severe headache. Multiple blood clots stent placed in one and taking medication now to help with the other blockages.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1401290-1	Blood clot in ankle
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1401676-1	BLOOD CLOT; This spontaneous report received from a consumer concerned a male of unspecified age. The patient's height, and weight were not reported. No past medical history or concurrent conditions were reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of administration not reported, batch number: Unknown) frequency one total, dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. It was reported that on an unspecified date, the patient experienced a blood clot. There was no additional information provided. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of blood clot was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:20210626844-COVID-19 VACCINE AD26.COV2.S-blood clot. This event is considered unassessable. The event has an unknown/unclear temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1402224-1	I have had pain in my calves so I stopped the birth control mini pill. My period had also been incredibly irregular, super heavy and clotted bleeding. I?ve felt dizzy and weak on the heaviest days I?ve been bleeding as well. I?m passing clots - similar to when I?ve miscarried in the past.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1402920-1	Ankle swelled up beginning of May.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1403469-1	Pt developed a blood clot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1403654-1	PT states after her J&J vaccine of 05-05-2021, she experienced phlebitis in the vein, left leg that resulted in a blood clot as per PCP. She states her PCP told her that her phlebitis would go away in a few weeks. PT states she still continues with the phlebitis in her left leg but now is less painful but still present.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1404270-1	Had her Johnson and Johnson covid vaccine on 4/8/21 LMP 4/14/21 Pt began spotting 5/9/21 which was day 25 of her cycle Day 2-5 very heavy bleeding Day 5 passed clot 11 days of bleeding Pregnancy test positive on 5/19/21 and 5/20/21 Presented to ED w/ intermittent RLQ abdominal pain and vaginal bleeding and spotting since 5/9 with positive hCG, 52 on 5/20 and 209 on 5/27 with OB ultrasound on May 28 that showed no IUP and small hyperechoic nodule adjacent to the right ovary that could represent an ectopic pregnancy and patient opted to avoid methotrexate treatment at the time. Returned to the ED on 5/30 with ongoing symptoms of RLQ cramping but no vaginal bleeding. OB ultrasound shows right adnexal structure measuring 2.6 cm, larger than structure 2 days ago. Pt diagnosed w/ Right ectopic pregnancy - successfully treated with methotrexate
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1406251-1	BLOOD CLOT; This spontaneous report received from a physician concerned a 21 year old male of unspecified race and ethnicity. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: unknown) 1 total dose, start therapy date were not reported for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the patient had blood clot at some point following vaccination and died due to it. It was not reported, if the autopsy was performed. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. This report was serious (Death). This case is linked to 20210456400 and 20210457370 (same reporter). .; Sender's Comments: V0: 20210626581-covid-19 vaccine ad26.cov2.s-This case concerns a 21 year old male, Blood Clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).; Reported Cause(s) of Death: BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1406337-1	That day, light headed ness and heart palpitations. Heart palpitations for short time only but felt periodically over the next few days. June 11, lower left led swelled and painful and shortness of breath. June 14, went to ER. They found multiple blood clots in upper thigh, behind knee and lower left leg. They also found approximately 20 clots in the lungs and administered Heparin drip and I was impatient. They put me on Eliquis and released me the next day. Now convalescing at my daughter?s house. Diagnosis. Acute Deep Vein Thromboses and Pulmonary Embolism bilateral.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1406519-1	Right leg was swollen up and doctor confirmed that she has blood clots. Swelling started two or three weeks after the shot was given.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1407106-1	On 04-24-2021 I had four days of extreme headaches that would not go away so I went to the ER where I got checked out. Also on 03-28-2021, I woke up with extreme left leg pain. I had difficulty holding things with my arms and I almost collapsed to the floor due to my left leg pain and weakness. I also had heavy menstrual bleeding with giant clots for the Month of March 2021. I can say today I have no symptoms of my left leg or headaches today.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1407133-1	ON APRIL 16, 2021 I WOKE UP AND I COULDN'T SEE OUT OF MY LEFT EYE, I WAITED A WHILE TOO SEE IF MY SIGHT WILL COME BACK AND IT DIDN'T SO I CALLED MY EYE DOCTOR AND MADE APPOINTMENT AND THEY SCHEDULED ME FOR APRIL 23, 2021. A WEEK LATER I WENT BACK TO THE DOCTOR AND HE TOLD ME TOO GO TO HOSPITAL A.S.A.P. THEY TOOK A CAT SCAN AND SAW THE OPHTAQLMOLOGIST IN HOSPITAL THE DIAGNOSIS WAS A BLOOD CLOT IN MY LEFT EYE AND I TOOK A STROKE.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1407730-1	Received the J&J vaccine in May 20. Started experiencing leg muscle and back pain June 1, 2021. Progressively got worse over the following weeks. On June 15 I woke up with a new pain in my leg. Went to urgent care and had a D-Dimer test that came out very high. I was sent to ER where they discovered one clot in each leg and two in my lungs via ultra sound and ct scan. I was immediately hooked up to herapin blood thinner through an IV. I was released the following afternoon with Eliquis blood thinner tablets.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1408240-1	patient stated he went to the hospital for blood clots. Says he went in last Tuesday or Wednesday for heavy breathing. Normal activity was making him very tired. Had COVID infection in March. The doctors cannot determine what other cause could have led to him having blood clots. Patient was released yesterday from being in since last Tuesday or Wednesday. Patient says he's having nose bleeds starting today. He was on the blood thinner shots but not sure of the name. Now he is on the pills (Eliquis) from pharmacy instead of the shot.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1409479-1	BLOOD CLOTS; This solicited report received from a consumer via PPSCVM000808: Janssen CarePath for XARELTO concerned a male of unspecified age. The patient's height, and weight were not reported. The patient's past medical history included covid-19 infection. The patient was treated with rivaroxaban (film-coated tablet, route of admin, and batch number were not reported) dose, frequency, and therapy dates were not reported for an unspecified indication. The patient initiated treatment with covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: Unknown) dose, start therapy date were not reported, 1 total, administered for prophylactic vaccination. The batch number was not reported. The Company is unable to perform follow-up to request batch/lot numbers. No concomitant medications were reported. On an unspecified date, the patient experienced blood clots and not sure what was causing it. The action taken with covid-19 vaccine ad26.cov2.s was not applicable; and action taken with rivaroxaban was not reported. The outcome of blood clots was not reported. The reporter provided no causality assessment. Company causality between covid-19 vaccine ad26.cov2.s, and blood clots was not related; and between rivaroxaban, and blood clots was not related. This report was serious (Other Medically Important Condition).; Sender's Comments: V0-20210627971- covid-19 vaccine ad26.cov2.s-Blood Clots-This event is considered un-assessable. This event has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the events.
THROMBOSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1410216-1	Blood clot in lower left leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0908846-1	Very mild, but unexpected nose bloody nose from left nostril. Only noted when blowing nose. With 1 clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0919376-1	Patient received Moderna vaccine, Wednesday 12/30. On Saturday 1/3/2021 patient felt pressure/tightness in lower extremity. When patient touched area, a noticeable ball was felt under the skin, tender to the touch and warm. Patient went into urgent care on Monday 1/4/2021 with a confirmed dx of a superficial blood clot. Unknown etiology of whether this is from current birth control or the COVID19 Moderna vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0937518-1	LEFT FACIAL SWELLING WITH LARGE AMOUNT OF NASAL BLEEDING AND CLOTTING. SHE SAW HER PCP AND HAD LAB TESTING AND GIVEN BENEDRYL. SHE WILL F/U WITH PCP AGAIN TODAY.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0938147-1	I am not completely convinced that this is related to the vaccine but thought I should report it just in case. I thought I had a spider bite on my left foot, and then I thought it was shingles and then cellulitis. I was then diagnosed with a blood clot today. I have never had blood clots before.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0941080-1	5 days after Moderna vaccine, developed severe abd pain, mid epigastrium. No Nausea or vomiting. No fever. Mild diarrhea. after 48 hrs with no improvement went to ED
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0942005-1	swelling in neck and down chest blood clots enlarged lymph nodes

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0946141-1	Moderna COVID-19 Vaccine At 2 PM I went blind in my left eye. Went to emergency room at Hospital Was told I have Blood clot in my eye causing the blindness and Ophthamologist says it will probably be permanent
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0954804-1	Started with severe chills, body aches and feverish. The. Slight leg pain which worsened with time , swelling on the right leg calf, warm to touch and difficulty breathing. Got hospitalized on 1/16 21 with multiple clots in my right leg and clot in the lung. Still in the hospital now.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0969230-1	Developed blood clot in left leg (DVT)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0978002-1	The morning of Jan 20, 2021, Patient was disoriented, could not communicate well, and fell when he tried to get out of bed. He was taken to The Hospital by ambulance about 11:30 am. He is still hospitalized. Emerging symptoms include: passing a blood clot from his mouth, slight pneumonia, high white count, low kidney values, high fever, all of which fluctuated. A blood transfusion was given because of blood in the urine. He has been confused and disoriented until this morning.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0979630-1	blood clot in lungs, PE,
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0988302-1	Very heavy period. Passed multiple golf ball sized clots for several hours.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0989737-1	Received vaccine on 1/4. Started menstrual cycle on 1/16 which was 1 week early. 3 days into the cycle, heavy vaginal bleeding with large blot clots (golf ball size) starting on 1/20/201 and continuing until now 1/31, bleeding still going on.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0990361-1	blindness in left eye; stroke in back of the eye; blood clot; A spontaneous report was received from a consumer concerning an 83-year-old, female patient who received Moderna's COVID-19 Vaccine (mRNA-1273) and experienced blindness in left eye, blood clot, and stroke in back of the eye. The patient's medical history was not provided. Concomitant medications reported included comerdol, losartan, hydrocortisone, meloxicam, and unspecified stomach pills. On 14 Jan 2021 at 10:15 am, approximately 3 hours and 45 minutes prior to the onset of the events, the patient received a dose of mRNA-1273 (Lot number: 013L20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 14 Jan 2021 around 2:00 pm, the patient lost sight in her left eye as a result of a blood clot and stroke in the back of her eye. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events, blindness in left eye, blood clot, and stroke in back of eye, was unknown.; Reporter's Comments: This case concerns an 83-years-old female patient, who experienced a serious unexpected event of blindness unilateral, retinal artery occlusion, and thrombosis. The event of blindness unilateral and retinal artery occlusion occurred 3 hrs. after first dose of mRNA-1273, lot # 013L20A. The event of thrombosis occurred on an unspecified date after first dose of mRNA-1273, lot # 013L20A. Treatment included details were not provided. Concomitant medications included Comerdol, Losartan for blood pressure, Hydrocortisone, Meloxicam and stomach pills. Very limited information regarding this event has been provided at this time. Based on the current available information and temporal association between the use of the product and onset of the event a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0994135-1	I am 38 years old with no history of medical problems. I do NOT have a history of miscarriages and have one healthy child who is 22 months old. On 1/13/21, I took a home pregnancy test which came back positive. At that time, I had a missed period but also had several common pregnancy symptoms such as bloating, acne, fatigue and tender breast. later that week, I called OB/Gyn and spoke to an RN to schedule my 8/9 week ultrasound and to inquiry about the vaccine during pregnancy since I had no clue whether it was recommended/safe or not. the RN, very confident and without any disclaimer, stated that hospital is recommending all of their pregnant patients to receive the vaccine. Obviously, I decided to trust this medical professional who was so confident in her response. My normal pregnancy symptoms continued. On 1/19/21, I was 5 weeks pregnant and received my first dosage of the vaccine. felt fine other than a sore left arm. on 1/20/21, I woke up with a lot of abdominal cramping and pain. It was new to me but assumed it was normal. My cramping and pain continued until 1/21/21. On 1/21/21, I woke up without the cramping and pain. But, I also noticed that my breast were no longer tender and my skin had completely cleared up. I became concerned but prayed everything was fine since my home pregnancy test was still positive. On 1/22/21, by cramps continued once again but more mild. My pregnancy symptoms seemed as if they were no longer present but remained hopeful. On 1/23/21, I woke up with light spotting that only lasted through the morning. Soon after, I started having extreme abdominal pains. I prayed everything was fine. The pain continued and became worse. That night, the pain was so bad that I just went to bed. Right before going to bed, I noticed I had started spotting again. A little heavier than in the morning. I made sure to lay on my left side, hoping it was normal in pregnancy. On 1/24/21, I woke up with heavy bleeding and clotting. I went to the doctor and got an ultrasound and blood test. I was told by the doctor at Hospital that I had a miscarriage.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0995017-1	On 1/28/2021 at 0545 hours as I was on my way into work when I started to have severe right side chest pain, so much so that I had to pull over. Shortly thereafter ambulance arrived on scene at which time it was determined that I would be transported to the hospital. Once there, a series of tests were conducted, one of which was a D-Dimer blood test to rule out blood clots. That resulted in an elevated count to which a CT scan was ordered. The result of the CT scan revealed that I had 2 Pulmonary Embolisms, one in the upper lobe of my right lung and one in the middle lobe of my right lung. Due to me being what the doctors said was a healthy 45 year old with no underlying conditions or signs of DVT (deep vein thrombosis) they deemed me low risk at that time and advised I was at low risk of dying in the next 30days. However, they advised that at any time, the clot could possibly become dislodged and cause worse problems, possibly death if time is not given to the body to dissolve the clots. Ultimately it was determined that I be discharged and prescribed blood thinning medications to help thin the blood to prevent further clotting should I have a clotting issue (undetermined at this time).
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1008808-1	I started my menstrual cycle about one week early, experienced dark, brown tinged spotting for 2 days followed by heavy flow with more clotting than I have ever experienced before. Cycle lasted about 8 days, versus my normal 4 days. These symptoms concern me due to several reports of miscarriages while receiving vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1012962-1	On Wednesday 1/27 the day after my second dose I felt a minor pinch/discomfort on my lower leg. I did not think much of that is reason I did not report it right away. Day by day the discomfort and pain got worse. Exactly one week after my second dose I was not able to sleep due to the pain. On Wednesday 2/3 I decided to go urgent care. From there I was referred for an ultra sound a few hours after. After the ultra sound I was contacted by my doctor, letting me know that the results showed I had a blood clot on my left leg. I was put on Eliquis medicine right away. On 2/5 I was able to physically see my doctor who told me that this was caused by the second dose of the moderne vaccine that I took on 1/26.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1014720-1	Very heavy menstrual cycle. Numerous clots. Severe cramping. All unusual for me
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1016123-1	Day after (2/8/21): Severe chills, tiredness, body aches, headache, loss of appetite, fever (101.4), bloody nose w/blood clots (left nostril only). 2 days after (2/9/21): Intermittent headache, aches in joints, bloody nose w/blood clots (left nostril only), hive around injection site.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1019785-1	1 week after the first dose patient started to have her period. The period consisted of heavy clots and would go through 6-8 pads /day. Period lasted 3 weeks and went to see GYN and they prescribed Provera, which helped stop the bleeding.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1034409-1	Very early menstruation, occurring a couple of hours after the vaccine. It started as spotting that night, (I never spot) then a gradually heavier flow, working up to an unusually heavy & painful period over the course of the next 2 days. Also some very light clotting, which rarely happens for me. I was not due for my period for 2.5 weeks. My cycle is extremely regular and has never deviated from 28-31 days in the last 6+ years. I have a copper IUD (for 6 years), and have never been pregnant.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1036912-1	Initial injection site pain for two days. Around day five, both armpits were discolored significantly. On day nine I was in the ER with left side abdominal cramps that was diagnosed after a CT Scan as a splenic infarction caused by a blot clot. I have not had a blot clot previously and have always been healthy with no surgeries or prescription medications except propranolol for anxiety (since the pandemic started).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				On January 21, 2021 I started feeling pains in my right shoulder (1st shot was left shoulder) and the following day it had spread across my body, shoulders, chest, arms, legs and calves. I also had a severe headache along with the muscle pains and woke up each day since in a pool of sweat with the sheet soaked to this day. Due to the severe muscle pains, I started taking Tylenol and all the symptoms pointed to the virus itself. I was tested for the virus on Monday, January 25th with negative results on the morning of the 26th. I continued to self-medicate with Advil every four hours to remove the severe muscle pains noted above. To back track slightly, I had a colonoscopy on January 19, 2021 and one large polyp was removed without issue. Due to the full body muscles pains and taking the Advil, it likely caused bleeding from the polyp site of the colonoscopy and went to the ER and they transferred me back the hospital on January 28, 2021, the location of the colonoscopy, and hospital found nothing wrong with the colonoscopy during my 2.5 day stay there. The did an x-ray of my chest to test for the flu and related items and found everything to be clear. They also a CT of my abdomen to look for any abnormalities and found none. Through all this the muscle pains continued when I would move, the muscles would fire up and intense pain persisted until I could calm them down after about 30 mins of intense pain. Hospital also tested me twice for the COVID virus and each turned out negative. Overall, I?ve been tested for the virus nearly 12 times and all were negative. From the visit to Hospital, with the colonoscopy and no additional bleeding occurring, Hospital gave me a steroid called Prednisone to mask the pain on Saturday, January 30, 2021 at about 1pm and by 4pm they fully released me with no idea what was causing the severe muscle pain across my body when I moved. I literally just walked out of the hospital since the steroid masked the muscle pain issue. The steroids worked but I was still walking up in a pool of sweat and very clammy and wet each morning from 1am until I woke up. On February 2, 2021 I had 3 bowel movements that were all dark purple and full of blood clots which led me back to Hospital and the ER could not get in touch with the Hospital Doctors for transfer so they admitted me to the Hospital, thank God. On February 4, 2021 Methodist performed an emergency colonoscopy to clamp the polyp site, took two additional polyps out and did an endoscopy to ensure my upper and lower GI track were clear, and it was and the two additional polyps were benign. Through all this the serve muscle pains persisted and Hospital moved me to a patient room and out of the ER. Over the next several days, February 4 thru the 11th Hospital cleared me of everything they could test for, over 85 different tests were ran based on MyChart. All my blood counts were all over the place, WBC were 19,000, Platelets were nearly 700, my sedimentation rate reached 64+ and the server muscle pains persisted and I was basically incapacitated during my entire stay at Hospital. Hospital had several specialists seem me from Internal Medicine, Infectious Disease, Neurology to Rheumatoid ologist and none of them found anything wrong with me other than my COVID anti-bodies were enormously high with not signs of slowing down. The conclusion by each of the specialist is that my symptoms all point to an Adverse Level 3 Inflammatory Reaction to the first COVID Vaccine shot. The Infectious Disease specialists emphasized that I DO NOT get the second vaccine shot since it would like have killed me. (We had cancelled my 2nd shot, scheduled for February 3, 2021 via Cancer Center the week of January 30, 2021.) Hospital concluded that they could not do anything further for me since all their extensive testing all came up negative other then the COVID antibody levels and started me on the same steroid via IV on February 10, 2021 and on February 11, 2021 I was released from Hospital with a 28-day gradual reduction of the steroid over that timeframe to see if the COVID antibody production would simply stop. They had no other recourse or follow up with for any of the specialists. I?m on my own at this point and scared. To help with the inflammation the only thing my wife and I saw we could do is see a health and well Doctor to try and purge myself of the inflammation and assist my bowel and liver health with trying to rid me of the vaccine and hopefully the antibody production. I?m on a strict diet and taking supplements to assists in the reduction of inflammation and cleaning my system out. Starting today, February 18, 2021 I start stepping back on the steroids prescribed by Hospital and praying that the symptoms and pain simply go away but I?m still having night sweats, weak and shaky.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1038305-1	I started bleeding on the 4th, doctor stated threaten miscarriage. I had blood clots and vomiting'. I was taking to emergency room. I was giving a DNC because I had a miscarriage.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1041297-1	Patient admitted with acute stroke, no prior history of same. Mild hyperlipidemia, but otherwise no clear risk factors for stroke. Unclear if anything to do with COVID vaccine.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1042458-1	Heart palpitations. Fast heart beat. 99-120. 95 when at rest. Up to 130 when stabding or moving from laying position. Chest pain heart area.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1043738-1	diarrhea low gas pain chills / sweating dry mouth sleepy no apatite cough with phlegm dizzy loss of bladder control blood clots in nose mucus
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1047576-1	4 days after receiving the vaccine in his left arm, my father developed a blood clot in his left arm that required emergency hospitalization and large doses of blood thinners.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1049991-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1053191-1	Vaccine administered 02/08/2021 , by Thursday 02/11/2021 patient almost nonverbal, by Monday 02/15/2021 patient went to the hospital with bruising, sores on her stomach and clots reported as thrombocytopenia, deceased by Friday 02/19/2021.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1056011-1	"My grandpa had a stroke on the 15th of February. He claimed he had been feeling ""off"" for a few days, but didn't say anything. A blood clot had formed in his brain. He was doing better and about to go to rehab to strength his right side of his body. On the 22nd he took a turn for the worst. He was having trouble breathing and they sedated and partially paralyzed him to put a tube in his mouth. I believe another blood clot had formed and oxygen wasn't properly going through his body. They could not stabilize him, and he passed away the same day."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1056972-1	5-6 days after receiving first Moderna covid vaccine pt. began not feeling well. On 02/10/2021 she saw a provider in an office for eval of abdominal pain and diarrhea and sent home. On 02/15/2021 she presented to a local ED with continuing symptoms, transferred to Medical Center, She is currently an inpatient there with a diagnosis of multiple blood clots in abdomen and brain and antiphospholipid syndrome.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1057786-1	Within 12 hours of receiving my second vaccine, My left leg felt like it internally rotated and started to hurt and feel like lead. On January 28, 2021, I went to the ER and an extensive blood clot (DVT) was found.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1059017-1	MODERNA COVID-19 VACCINE EUA Patient's arm was sore after injection and later that evening pain spread down to her elbow. In the following days she states that it spread to her fingers and included temporary numbness which resolved in a short period of time. Her arm remained sore with warmth and redness about the size of a quarter at the injection site. The warmth, redness and pain did not resolve so she went to the emergency room today.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1059327-1	Moderna COVID-19 Vaccine EUA. PT - Multiple blood clots both lungs (no other presentable causes i.e. diagnosed as unprovoked).
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1060219-1	Trouble breathing, severe fatigue, blood clots in lung and leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1064361-1	Four blood clots in left leg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1069054-1	Anxious; Tired; Blood clots in left leg, right leg and brain; A spontaneous report was received from a consumer who was also a 66-year-old female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and developed blood clots in the left leg, right leg, and brain, anxious and tired. The patient's medical history was not provided. Concomitant medications reported included vitamin D, magnesium, lisinopril, and vitamin B12. On 14 Jan 2021, approximately 3 days prior to the onset of symptoms, the patient received their first of two planned doses of mRNA-1273 (Batch number not provided) intramuscularly in the left deltoid for prophylaxis of COVID-19 infection. On 16 Jan 2021, the patient experienced immense pain in the middle of the night, and subsequently saw her primary physician. An ultrasound revealed blood clots in her left leg, right leg and brain. A hematologist and vascular surgeon were consulted. Patient was treated with apixaban while they are doing blood work. The patient also became tired and anxious. She noted that she never had comorbidities before and was upset that her life has completely changed. A repeat ultrasound was scheduled for 19 Feb 2021. The mRNA-1273 dose was discontinued in response to the event of blood clots in the left leg, right leg, and brain, anxious and tiered. The outcome of events, blood clots in the left leg, right leg, and brain was considered unknown at the time of this report. The outcome of the events, tired and anxious were considered not resolved at the time of this report.; Reporter's Comments: Very limited information regarding the events has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1072189-1	Started dry cough for three day, pain across the lower back began two days after cough. Began intense pain with nausea and cold and clammy sweats. Called the nurse on my insurance who advised to get to the ER.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1073412-1	Initial flulike symptoms with fever Tues -Friday. Had a right side stroke 6 AM Saturday morning . Rushed to hospital and had a procedure involving a catheter through the body to the clot in the brain. Came out of that with a weekend left side and was medicated through Sunday night. At 10:30 PM Sunday night had a second stroke on the right side. Did not wake from that. Now I?m not expected to survive.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1074082-1	My stomach was hurting while urinating and there was blood in my urine. At another instance, there was a small blood clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1075784-1	About a week after receiving my second Moderna Covid shot I felt shortness of breath. 9 days after the shot I had a mild fever (100.5f), severe shortness of breath and chest pain (level 5+) and went to the emergency room. They found D/Dima was elevated and sent me for a CT scan. This showed a significant blood clot in the right lung and a lesser clot in the left. I am usually extremly health, walking 2 miles at least 5 days a week and bicycing about 50 miles per week

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1079224-1	3 DAYS AFTER THE 2ND INJECTION, I DEVELOPED 2 BLOOD CLOTS AT THE BASE OF MY LUNGS - 1 ON EACH LUNG. IN CONNECTION WITH THAT, I DEVELOPED SEVER PAIN IN MY LOWER RIGHT FLANK AREA THAT RADIATED TO MY SHOULDER AS WELL AS HIGH BLOOD PRESSURE. THERE IS NO HISTORY IN MY FAMILY OF ANY CLOTTING ISSUES AND I MYSELF HAD NEVER HAD SUCH A THING OCCUR. THE PAIN BECAME SO BAD IT FORCED ME TO GO TO A HOSPITAL EMERGENCY DEPT. AFTER CT SCANS CONFIRMED THE PRESENCE OF THE CLOTS, I WAS SUBSEQUENTLY ADMITTED FOR OBSERVATION ON A REGIMEN OF DRUGS, BLOOD THINNER, PAIN MED, AND A CLOT BUSTER (NOT SURE WHICH MANUFACTURER.) I AM STILL EXPERIENCING PAIN IN MY RIGHT FLANK AREA - SOMEWHAT RADIATING TO MY RIGHT SHOULDER DAY 1 POST DISCHARGE AND AM HAVING TO RELY ON OPIOID PAIN MED AS WELL AS THE CLOT BUSTING MED AND A HIGH BLOOD PRESSURE MED. TO ME, IT SEEMS THAT THIS MAY NOT BE JUST COINCIDENTAL. I WAS FEELING FINE PRIOR TO THE S 2ND DOSE OF THE MODERNA VAC AND THIS SEEMS TO ME THAT IT MIGHT BE RELATED TO THE 2 DOSE.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1080646-1	- Headache: starting about 12 hours after the vaccine and continuing for 2 days - Fatigue: starting about 12 hours after the vaccine and continuing for about 2 days - Chills: starting about 16 hours after the vaccine and intensifying until 24 hours after the vaccine administration - Fever: starting about 16 hours after the vaccine and intensifying until 24 hours after the vaccine administration - Blood clots: expulsion of clots during the same period as the fever and chills. Had completed menstruation 2 weeks prior, so not related to that.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1081909-1	DVT blood clot and surface blood clot in lower left leg diagnosed at the ER on 2/12/21. Levonox injections and warfarin have been started. Twice weekly INR checks. No longer need the injections; warfarin is continued. Next INR check is scheduled for Thursday March 11, 2021. Warfarin may need to continue life-long.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1082086-1	Patient was vaccinated at pharmacy on 2/9/2021, first dose of Moderna COVID-19 vaccine. Per medical records from hospital: patient developed fever, diarrhea, nausea and abdominal pain on 2/25/2021 and presented to the hospital E.R. on 3/1/2021. Patient was diagnosed with Sepsis and Pneumonia. Cardiac arrest on 3/6/21, renal failure, seizures. Patient tested negative for COVID-19 on 3/1/2021 and 3/8/2021. Patient has declined, was placed intubated and placed on a ventilator. Patient admitted to hospice services on 3/8/2021 and plan is for compassionate removal of life support at hospice. Prognosis is poor.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1086033-1	Blood clots in arm; Arm was sore; A spontaneous report was received from a healthcare professional concerning a 67-year-old, female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced sore arm/pain in arm and blood clot in her arm/thrombosis. The patient's medical history, as provided by the reporter included diabetes. Concomitant medications were not included. On 15 Feb 2021, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (Lot number: 031M20A) in the left arm for prophylaxis of COVID-19 infection. On 15 Feb 2021, post vaccination the patient experienced sore arm and her condition got worse. She was then taken to emergency room with blood clots in her arm. Action taken with mRNA-1273 in response to the events were not reported. The outcome of the events, sore arm and blood clot in her arm, were not known.; Reporter's Comments: This case concerns a 67-year-old, female patient, who experienced thrombosis and pain in arm. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. The patient's medical history of diabetes is a risk factor. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1086384-1	On the following day after receiving my first dose of the vaccination, my arm was throbbing, completely swollen, decreased ROM and skin was RED and severely tender to the touch. Took OTC Tylenol (2) po and had no signs of relief. Pain , swelling, and skin redness lasted for 3 days. On February 10th, 2021 I had a positive pregnancy test (confirmed by bloodwork and urine) which indicated I was 6-7 weeks pregnant and had a miscarriage on March 2021, after giving live births 3 prior times. The cause of the miscarriage is unknown. I also developed a blood clot on the arm that was inserted for an IV. I have no family history or personal history of needing blood thinners.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1086484-1	I got the Covid vaccine on 2/13/21, I lost a small clot and started to bleed on 2/19/21. I was due around Oct 4th, at the time of the vaccine I was 6 weeks pregnant. I miscarried at home on 2/25/21- I lost a very large clot and was bleeding heavily for a week. Doctor confirmed with HCG test that I miscarried. This was my second pregnancy, I had my first child 1/10/20.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1092490-1	Patient became lethargic, extremely tired, unable to care well for self. Patient had no appetite, refused most food and spent many hours simply sitting in her chair. She felt that she was having a heart attack and called 911 using her emergency response button on 2/21/2021. Doctors indicated that the patient's life was in serious danger, but because of age they were hesitant to try to remove the clot pressing on her heart. She was only allowed to go back home because we utilized Hospice. When asked about her life expectancy, we were told she could die any minute or might live a few more months, but that she was not expected to live much longer. SHE WAS IN GOOD HEALTH PRIOR TO THE IMMUNIZATION!

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1092640-1	atrial flutter pain in right arm , pain kept going up arm but only if you touched it, got on xartorl (blood thinner) . right forearm was really swollen , tingling and turned blue, went to hospital they did a ultrasound to check pressure in veins. if she had hand at 90 degree angel it wouldnt turn blue but if I drop it it would turn blue. next day ran more test. MRI on arm. taped vitamin c to spot where pain was in arm. developed blood clot. told her to put heat on arm and take pain meds. felt better until she stopped blood thinner feb22, 2021. two days later had more blood clots on arms, veins were showing huge in breast and foot, pants got tighter. feb 27veins in lower leg and neck started appearing more. feb28 my left hand was cold and the veins were showing more and u could see veins. right back veins started appearing , got period but after 12 hours it stopped. march 3 veins started appearing again. march 4 right arm swelled and tingling in fingers and right right arm was so painful. Nerve pain. put back on blood thinner. march 4 in evening was put on new meds. I'm still dealing with pain and different issues. march 9 had a Val moment but had tingling from butt to brain . next morning felt miserable, nauseous , throwing up and couldnt control it. 3/11 neck is swollen. referring her to university because no doctor can figure out what it is.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1092762-1	I experienced a runny nose with mucus, blood running from nose with blood clots four to five times a day from March 8th 2021 up to March 10th 2021. It is unknown rather I have blood clots within my body.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1093487-1	Thrombosis of the vein branch of the left eye.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1094443-1	Sweating, low-grade fever, chills, body aches, exhaustion, headache. I took Tylenol for 24 hours starting when I woke up at 6am to help with symptoms. It took about 36 hours for my symptoms to be gone after they started. I was also on my period when I received the vaccine. My period seemed to slow down almost completely while my symptoms were happening, and then my period resumed with abnormal cramping and clotting after my symptoms were gone.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1094625-1	Thrombosis in popliteal vein
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1095030-1	After 1st injection (1/11/21) experienced severe shortness of breath with minimal activity with accelerated heart rate and drop in O2 stat. within 2 days of injection then developed wheezing cough. Primary MD prescribed Augmentin nd prednisone with eventual success. With normal blood work, chest x-ray and negative Covid test all done by 1/15. On 3/9/21 (28) days after second dose (2/11/21) experienced burning and swelling on right leg. Went to Hospital, ER and found extensive blood clots in right leg and saddle pulmonary embolism . Also had 50% Platelet blood count decrese since the 1/15 bloodwork Admitted for 3 nights on heparin dip. Discharge on home Lovenox injections for 30days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1095915-1	I got a blood clot in my left leg, as well as numerous blood clots in both my lungs. I have never had a blood clot before, and my family does not have a history of them.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1096672-1	About 16 hours later I had a heart attack, 20% plaque and the rest was a blood clot that acute my artery. I was in the Hospital for two days. I ended up having angioplasty and a stent. When I went home I had post heart attack issues and a sore arm for about 3 weeks.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1096774-1	superficial blood clots in left leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1099050-1	On day 5 after injection, patient developed right inner thigh discomfort, leg swelling and accentuation of existing varicose veins.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1101157-1	blood clot in left forearm; Sore arm; A spontaneous report was received from a consumer concerning a78, year, old, female patient who developed a blood clot in the left forearm. The patient's medical history included hypertension, hiigh cholesterol and hypothyroidism. On 19-JAN-2021 the patient had two stents placed in her heart. Products known to have been used by the patient, within two weeks prior to the event, included clopidogrel bisulfate (One 5 milligram AM and PM) and apxaban (One 5 milligram AM and PM) and Asprin. On 10-FEB-2021, approximately six days prior to the onset of the symptoms, the patient received their first of two planned doses of mRNA-1273 (Batch number not provided) intramuscularly in the left arm for prophylaxis of COVID-19 infection. The patient had a sore arm for about a day after the vaccination. Six days later the left arm was red and swollen, inside forearm four inches down below the elbow it was stinging, slightly swollen, just to a soft touch she felt lumps of swelling. The patient went to the emergency room and had blood work and an ultrasound. The emergency room doctor told her that she had a blood clot in the left forearm four inches down below the elbow. The lab test done at the emergency were blood work and ultrasound. Treatment for the event included changing her current medication to one clopidogrel bisulfate 5 milligrams AM and PM and apxaban two tablets of 5 milligrams AM and PM and to discontinue the Asprin for one week. There was no change planned to the dosing schedule of mRNA-1273 in response to the event(s) and is scheduled to get her second vaccination 10-MAR-2021. The outcome of the events were considered as unknown. Follow up: No follow up information received.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1103001-1	Development of blood clots in lower right leg causing swelling and pain. Pt. visited doctor who prescribed an ultrasound which revealed the clots. Doctor has now put pt. on blood thinners indefinitely.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1104499-1	Developed numerous blood clots in left saphenous vein and likely pulomary embolism (did not do test for embolism b/c I am allergic to the contrast but I was very short of breath for several weeks). Not sure if this is related to the vaccine since I was post bone surgery and also on estrogen therapy (ovaries removed 5 years ago) but I have not ever developed a problem before after surgery and while on estrogen.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1104841-1	My husband received the first injection on Feb 3, 2021 he developed a cough afterwards, cough would come and go. He was scheduled to receive 2nd dose on March 11 and I was concerned about having a cough at the time of 2nd vaccine. He went to walk-in clinic and got a covid test was negative, he said his exam was normal, they gave him prescription of Benzonatate 100mg he took one and it did not help so he never took any more. On March 5 while working in the yard he collapsed, paramedics arrived and he was in cardiac arrest, they started CPR, they used AED was in vfib, he was taken to Hospital then transferred to second hospital. I was told he had blood clots in both lungs, one in his leg, and possible clots in the mesenteric area of abdomen. His heart stopped due to the blood clots in his lungs. He remains in the hospital in the Intermediate Coronary Care Unit. He has not yet gained full consciousness.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1105092-1	PT WAS VACCINATED WITH SECOND MODERNA DOSE ON 02/16/2021 . DR CALLED ME TO INFORM OF PT ADVERSE EVENT. PT PRESENTED TO DR WITH SWELLING IN LEFT LOWER LEG. DR ORDERED VENUS DOPLAR SHOWING A LARGE BLOOD CLOT FROM PT'S GROIN TO HER CALF MUSCLE. DUE TO NO PREVIOUS HX OF CLOTTING OR BLOOD DISORDERS NOR FAMILY HX AND PROXIMITY TO VACCINATION, DR BELIEVES VACCINE CAUSED THE CLOT. DR PLACED PATIENT ON ELIQUIS. DR WISHES TO BE CONTACTED IF FURTHER DETAIL IS NEEDED.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1105441-1	Pt developed RUE erythema and edema. RUE US demonstrated clot in 1 of 3 brachial veins in mid-portion of RUE. Pt prescribed Eliquis x 6 weeks (5 mg PO BID once. 10 mg PO BID x 7 days. Then 5 mg PO BID x 5 weeks) followed by repeat RUE US.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1105496-1	you got second Moderna vaccine on March 10, Wednesday. Saturday night began with left leg swelling and pain. Thought it was a side effect of the vaccine. Tuesday, march 16 the pain got so bad he went to ER. Has bleed clots in main artery in leg femoral artery. Pt. was transferred to larger hospital.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1105535-1	Multiple blood clots in right leg (DVT) and multiple blood clots in lungs (PE)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1106403-1	Blood clots in the left leg, same side as the vaccine was given. Woke up approximately two days afterward (2/11/21) with muscle pain in the left thigh. The pain did not lessen over the week, assumed it was just muscle. The pain migrated down to my calf and then was too painful to stand on or walk on 2/18/21.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1106505-1	Ocular stroke. A black curtain dropped down over right eye. Went to ER on the 28th, and was admitted. Had a blood clot in the right eye. Carotid artery right side did not have stenosis, they do not know why she had a stroke. Doctor can't say that it was caused by the vaccine. She now has a permanent blind spot in right eye.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1106515-1	Severe middle back pain beginning February 21. Initially thought it was a kidney stone, but a visit to a urologist on February 25, determined it wasn't a kidney stone. The urologist ordered some blood work and that indicated that I most likely had blood clots. Went to the ER on the 25th and had further tests that determined that I indeed did have blood clots in my lungs

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				further tests that determined that I indeed did have blood clots in my lungs.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1106802-1	One week after 2nd Moderna, started coughing periodically in day and night chest conjestion started to build at night. After three days of cough periodic and each night became more conjested with problem being to not inhale deeply, and lower right side adobonem pain. Sought medical appointment March 20 but did not get appointment until March 9 with medical facility due to increasing pain and not sleeping from conjestion. March 10 doctor visit ordered blood and urine tests and CT scan. Early Mar11 test B type natriuretic protein was high (CHF), Doctor ordered chest xray, ekg and echogram. CT scan found clot in right lower lung. Told to go to ER Mar 10 evening, ER ordered Ultrasound images of the legs.BILATERAL LOWER EXTRE BILATERAL LOWER EXTREMITY DEEP VEINS: Nonocclusive thrombus in the distal right femoral vein and popliteal vein extending into the tibioperoneal trunk. Thrombus is hypoechoic and expansile consistent with acute thrombus.Left lower extremity veins are patent and compressible. Left common femoral vein demonstrates normal waveform without loss of respiratory phasicity.Positive for right lower extremity DVT of the distal femoral vein, popliteal vein, and tibioperoneal trunk. CT;LUNG BASES: Small right pleural effusion with overlying compressive atelectasis and consolidation. There are addition, there are apparent filling defects within the subsegmental pulmonary arterial branches suggestive of emboli. Many /multiple blood tests March 9, 10 ,11 , 12 and 16 available to examine details..one perhaps low Platelets count 151 K/uL (range) 140 - 400 K/uL. I can give permission for these tests results to study to correlate lung and leg clots . This is unique because hospital data available to diagnose in detail my reaction after 2nd Moderna shot. Please look into it with experts..Did Moderna cause my clots I never had before?
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1107202-1	Migraines have became more frequent and more intense to the point of were she could not function; Developed a welt at the injection site that she describes as COVID arm which was a 3-4inch raised circle underneath the injection site; Pain under lymph nodes; Developed thrombosis on her head, but it went away; A spontaneous report was received from a consumer on 04 Mar 2021 concerning a 45-years-old, female patient who received Moderna's COVID-19 Vaccine (mRNA-1273) and developed thrombosis on her head, but it went away, migraines have become more frequent and more intense to the point of were she could not function, developed a welt at the injection site that she describes as COVID arm which was a 3-4inch raised circle underneath the injection site, lymph nodes pain, which she still has. The patient's medical history was not reported. Current conditions included migraine. The patinet had allergy to latex and pineapple. No relevant concomitant medications were reported On 14 Feb 2021, the patient received their first of two planned doses of mRNA-1273 (lot: not provided) intramuscularly in right arm for prophylaxis of COVID-19 infection. Since 14 Feb 2021, patient's migraines had became more frequent and more intense to the point of were she could not function. She also developed thrombosis on her head, but it went away. The patient also developed a welt at the injection site that she described as COVID arm which was a 3-4 inch raised circle underneath the injection site. She also developed pain under lymph nodes, which she still had. No treatment information was provided. Action taken with mRNA-1273 in response to the events was unknown. The event of migraines have became more frequent and more intense to the point of were she could not function and pain under lymph nodes which she still has were unresolved, the event of developed thrombosis on her head but it went away was resolved, while the outcome of the event of developed a welt at the injection site that she describes as COVID arm which was a 3-4inch raised circle underneath the injection site was unknown. The reporter did not provide an assessment for the events thrombosis on her head, but it went away, migraines have became more frequent and more intense to the point of were she could not function, developed a welt at the injection site that she describes as COVID arm which was a 3-4inch raised circle underneath the injection site, Pain under lymph nodes, which she still has.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1107342-1	Blod clot, stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1107356-1	Blood clot in blood vessel in the left hand at the base of the middle finger causing darkening in the lower part of the finger and numbness in the whole finger.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1108629-1	Patient developed shortness of breath upon exertion on 2/16/21 after receiving his second COVID vaccine on 2/12/21. He had an office visit on 3/3/21 with his PCP where he was found to have an elevated d-dimer. He was sent to the ED for a CT-scan that confirmed an acute bilateral pulmonary embolism with large burden of thrombus within the right lung. He was admitted from 3/3 to 3/11. He underwent a pulmonary arterial embolism on 3/5. He was also on a heparin drip while inpatient. He was discharged on Eliquis.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1108656-1	The next day, I had short term memory loss. I went to the hospital, and they said I had some type of heart episode. I was taken by ambulance to the hospital and was told I had a blood clot in my brain and had suffered a mini stroke. I was hospitalized for 4 nights. I was given blood thinner and I improved.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1110391-1	Received vaccine, and no reactions. Nine days after the vaccine slowly had drooping on the right side of his face, numbness on the right side of his face, mainly around the mouth, a little on the eye, hard to close the right eye. General soreness around his mouth when he tries to eat or talk a lot. Limited movement of his face do to swelling, difficulty closing his eye, and headache. Went to the ER on 3/7/21 and admitted him as they believed he had a blood clot, diagnosed him with Bell's palsy on Monday 3/8/21. Was given corticosteroids for 9 days Prednisone 60 mg. The symptoms are slowly improving but still has the Bell's palsy.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1111503-1	I had the shot on a Tuesday and developed blood tests by Friday.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1112628-1	I received my vaccine on Sunday March 7 and in the morning of March 12, I woke up to a large bruise with a hard clot on the inside of my right forearm measuring 7cm x 4cm with a large clot in the bruise. I also have a smaller spot which has a firm bump under the skin. I did not hit my arm or injure it during my sleep or before going to bed.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1115043-1	Blood Clot in Leg; Tenderness; A spontaneous report was received from a consumer concerning a 68-year-old, male patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced blood clot in leg and tenderness. The patient's medical history was not provided by the reporter. Concomitant medications reported included diltiazem hydrochloride, atorvastatin calcium and acetyl salicylic acid. On 13-Feb-2021, prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (Lot number:031M2QA) through intramuscularly in the right arm for prophylaxis of COVID-19 infection. On 13-Feb-2021, following administration of the vaccine, the patient had tenderness in his leg that continued until he sought out medical treatment. After medical treatment is was concluded that the patient has a blood clot in his leg and has to receive medical treatment to resolve that condition. The treatment information included rivaroxaban. Action taken with mRNA-1273 in response to the events was not applicable. The events, blood clot in leg and tenderness were considered recovered.; Reporter's Comments: There is not enough information to assess the causa association between the reported event of blood clot in leg and the administration if the mRNA-1273 vaccine. Critical details such as the patient's medical history is lacking. Additional information has been requested. Howe ever, based on temporal association between product administration and the event of tenderness, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1115432-1	Jan 27th Began to run fever from the shot Jan 28th Continued to experience fever, uncontrollable chills, upset stomach, diarrhea through the day around 6PM I began to sweat, my stomach became upset again, diarrhea grew worse. Chest pain began and call to 911 was placed. Taken to hospital by flight
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1115659-1	"Patient states he had a fever ""a day or two"" after he received the 2nd Moderna vaccine. The fever reached 101.7 degree F and resolved within 24 hours. A few days after that the patient states he had a ""minor stroke."" Patient states the doctor told him that there was a small clot on the left side of his brain. Patient has resulting numbness of the right side of his body. Per the patient the doctor says the clot had nothing to do with the stroke. When asked if the doctor had thought that the vaccine had anything to do with either stroke or clot the patient said the doctor said he didn't think so but also kind of ignored the question."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1116566-1	Pt was admitted for right lower extremity swelling. has had DVT and pulmonary embolism. Pt found to have thrombosis of right popliteal vein. pt following up with hematologist.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1117090-1	I HAD HEADACHES, CHILLS, DIARRHEA, ACHES AND TIREDNESS BUT MY BIGGEST CONCERN IS THE FACT THAT I WOUND UP IN THE HOSPITAL ON FEB 8 WITH BLOOD CLOTS DOWN MY ENTIRE RIGHT LEG AND IN MY LUNGS. I NEED TO KNOW IF THE MODERNA CAUSED THESE OR MADE THEM WORSE. I HAD NO IDEA I HAD BLOOD CLOTS BUT MY FOOT, LEG WERE SWOLLEN AND I ACHED FROM MY HIP DOWN THAT LEG. I ACTUALLY THOUGHT IT WAS SIADACA.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1117927-1	Blood clot running from ankle to groin; entire leg swollen; still swollen after 6 weeks; received shot of heparin and put on Xarelto; blood clot was just short of where it could break off and go to heart or lungs.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1117950-1	A few minutes after the shot, I became dizzy, watery eyes, blurry vision, resolved in a few minutes and I drove home, woke up to severe pain in my right eye, foggy brain, sharp/shooting pain in left side of chest and left breast, headache (very rare for me to have a headache), toes began to hurt and brown spots with excessive dry peeling skin appeared with pain (more on right toes than left) and was unable to wear closed toe shoes, hit my thumb and a blood clot appeared (never had a blood clot in my life). Both the toes and clot is just now beginning to get better (and is just now starting to get better). I continue to have blurry vision and occssional foggy brain,

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1118197-1	2 days after receiving the second moderna vaccine my father experienced a stroke. He was transported to the emergency room medical center where he went thru a procedure to remove a clot in his left side brain. Prior to this vaccine my father was in good health and was very active and still works and owns and operates a restaurant. He has never had any problems like this before the vaccine. One day after the vaccine he was complaining about a pain on the left side of his neck area. He was doing paperwork at approx 7:40pm when he experienced the stroke. I had to call 911. My father is currently at the medical center recovering from the incident.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1118872-1	Pt started having muscle pain in left leg on 3/18/2021 went to hospital on 3/19/2021 had doppler done and confirmed blood clot in left leg. Pt has been started on Eliquis. Currently out of hospital and at home.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1120842-1	this is all per family, 4 to 5 days after 2nd COVID vaccine he was acting unusual and was taken to the hospital. He had a clot in his brain and underwent brain surgery. He experienced seizures after the surgery, but it was ultimately reported the surgery went well. He remained intubated and on a ventilator after surgery. He developed complications of his lungs and kidneys while on the ventilator. Ventilator was removed 3/16/2021 and he passed away that day. The hospital providers thought the clot in the brain may have been from hitting his head over a month ago. From my understanding he was A&O, independent with ADLs, and lived in his private residence prior to these complications.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1122044-1	"Patient stopped at the health department on 3/18/21 and spoke with the secretary stating that he had a blood clot on 2/26/21 after receiving the covid vaccine. She took his information and RN followed up with him on 3/19/21. Patient states he had a blood clot in his left leg a few years ago and on the 26th he experienced the same pain as with his original blood clot. He said it was 10/10 pain in his left leg from just above the knee to right above the ankle around the whole leg. He states it felt warm. He did not speak with his doctor or seek any medical attention. He states he ""suffered through it"" and it went away after a couple days."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1122253-1	Stroke (blood clot in vein in left side of brain), occurred Wednesday March 17 at 5:15 pm. Immediately admitted to emergency room/stroke unit and given blood thinner IV and anti-seizure medicine
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1124590-1	Swollen right lower leg. Doppler study revealed multiple blood clots. Stared on Eliquis. Continues on Eliquis at this time and leg still swollen
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1124727-1	Pulmonary Embolism Multiple blood clots in legs and lungs
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1126577-1	Patient reported the following: I went to the ER last Thursday because my left leg was red and swollen. They did a sonogram and the results came back as tendentious and a superficial blood clot. The ER physician at Health suggested that I report the blood clot because it is of unknown origin and I did receive both vaccines.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1126609-1	CARDIOPULMONARY ARREST 2 DAYS AFTER RECIEVING SECOND MODERNA DOSE
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1127730-1	A stroke occurred approximately 3 hrs after receiving vaccine. He was flown by helicopter to Medical Center. He had dysphasia which did resolve by the time he was admitted into the ER. He has since had CHF and Afib and is now on anticoagulation and diuretics. He continues to have problems with CHF, SOB, weakness and is undergoing treatment as outpatient. He is now followed by Cardiology, Neurology.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1128838-1	lost appetite; so tired, thought could not get out of bed; Big/ huge clots, came out in a spit; had 3 nose bleeds; A spontaneous report was received from a 77-year old, female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced 3 nose bleeds/ Epistaxis, big/huge clots, clot came out in a spit/ Thrombosis, tiredness/ Fatigue, and loss of appetite/ Decreased appetite. The patient's medical history was not provided. No concomitant product use was reported. On 12 Feb 2021, the patient received their first of two planned doses of mRNA-1273 (Lot number: unknown) intramuscularly for prophylaxis of COVID-19 infection. On 13-Feb-2021, the patient experienced 3 nose bleeds with huge blood clots. The bleeding was persistent throughout the day, stopping at the intervals and later on the same day it was reported the patient had a huge clot come out in spit. On 14 Feb 2021, the patient was tired, could not get out of bed, and had lost appetite. On 18-Feb-2021, the patient woke up feeling normal. Treatment information was unknown. Action taken with mRNA-1273 in response to the events was not provided. The outcomes of the events were not provided.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1129475-1	Early onset of menstrual period-14 days early-lasted 11 days versus normal 5 and heavy bleeding including large clots for 10 days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1130191-1	Developed extreme pain in my left leg. Went to emergency room Ultrasound showed I have a blood clot in left calf. Doctor prescribed a blood thinner, told me to elevate my leg and apply moist heat.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1130407-1	Reported side effects of fever, chills and headache that began within 24 hours, also reports burning in throat. Was evaluated by a specialist after 1 week due to continued burning and pain in throat. Reports was diagnosed with edema around vocal chords. Approximately 2 weeks after vaccination developed swelling in left arm. She was seen by her physician who sent her for an ultrasound and they found a blood clot. Patient called to report these symptoms.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131196-1	blood clot; Vertigo; Nausea; Vomiting; A spontaneous report, was received from a consumer concerning a 53 year old female patient who received Moderna's COVID-19 Vaccine (mRNA-1273), and experienced vertigo , nausea, vomiting and also developed a blood clot (thrombosis) during hospitalization. Patient had no medical history. The concomitant medication included were zolpidem tartrate 10mg nightly, allopurinol 100mg daily, duloxetine 50mg daily, multivitamin, vitamin d and acetylsalicylic acid 81mg. On 05 Feb 2021, prior to the onset of events, the patient received the first of two planned doses of mRNA-1273 (Batch no- 010MZ0A), intramuscularly in the left upper arm for prophylaxis of COVID-19 infection. On 17 Feb 2021, few days after administering the vaccine, the consumer experienced vertigo related nausea and vomiting. The patient was hospitalized for two days and experienced a blood clot during hospitalization. Patient was treated with only diazepam on first two days of treatment. In hospital, patient was given meclizine, diazepam and other anti-nausea medication whereas at home, patient was treated with ondansetron 8 mg, meclizine and diazepam. On 05 Mar 2021, the patient received the second of two planned doses of mRNA-1273 (Batch no- 036A21A), intramuscularly in the left upper arm for prophylaxis of COVID-19 infection. The action taken with the second dose of mRNA-1273 in response to the event was not applicable. The outcome of the events, vertigo, nausea, vomiting was not recovered whereas the outcome of blood clot was unknown at the time of the report.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131197-1	blood clot in his left leg; A spontaneous report was received from a consumer concerning a 70-year-old, male patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and developed a blood clot in his left leg/thrombosis. The patient's medical history was not provided. No relevant concomitant medications were provided. On 10 Feb 2021, approximately two weeks prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (Lot number: 030M20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. Two weeks after the vaccination, the patient developed a blood clot in his left leg. He got his second vaccine on 10 Mar 2021. Treatment information was not provided. There was no change planned to the dosing schedule of mRNA-1273 in response to the event. The outcome of the event blood clot in his left leg was not reported.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the onset date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131365-1	On March 6, I woke up feeling like I had a Charlie horse in my left calf. I tried to walk and it was sore but I felt it would go away in a day or two. By Wednesday, March 10, I called my doctor and after telling them my left calf hurt , they had me come right in. After seeing dr I was sent to get a sonogram immediately and was told I had a blood clot inside my left knee. I was sent back to Dr and was given Xerelto 15 mg twice a day for 21 days and the 20 mg once a day until gone. Today,, it is still sore. I am asking if I should take my second shot on March 29 or not.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131417-1	Fever, headache, chills, night sweats. All on same day. 3 days later swollen knees. 10 days later blood clot in right leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131574-1	Weird headache at top of head which would come on quickly for a minute or two then vanish and repeat every 10-15 minutes or so. Very tired. Had started menstrual cycle on March 17, but post vaccine it was lighter than normal and then on the 22nd it became very heavy and there were lots of large blood clots. This has lasted an additional three days. My cycle usually lasts 5-7 days but day 2/3 are the heaviest and it?s usually very light by day 6/7. I have not had blood clots in several years. These are very large.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1131644-1	2 large Blood Clots in the lungs
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1134671-1	Patient went to have labs done for her leukemia checkup 3/24. Patient has had leukemia for many years and has had many blood draws. Patient had blood taken at her appointment and was waiting when the nurse came back out and said her blood had clotted so they needed to redraw her blood. She states this has never happened before and wondered if her blood clotting would have been due to her covid vaccine the week prior. She states the first girl who drew her blood was slower than the second time around when a different nurse drew her blood.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1135298-1	blood clot; Fever; sick feeling; chills; nausea; night sweats; A spontaneous report was received from a healthcare professional concerning a 31-year-old, male patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced sick feeling/illness, chills, nausea, night sweats, blood clot/thrombosis, and fever/pyrexia. The patient's medical history included Cystic Fibrosis and only has half of their liver. No relevant concomitant medications were reported. On 26-JAN-2021, prior to the onset of the events the patient received their first of two planned doses of mRNA-1273 (lot/batch: unknown) intramuscularly for prophylaxis of COVID-19 infection. On 26JAN2021, the patient began experiencing symptoms including sick feeing, chills, nausea, and night sweats. The patient had blood clot and fever on 21-FEB-2021. Patient was admitted into the hospital on 22-FEB-2021 and discharged on 26-FEB-2021. He received the following tests while in the hospital: MRI, Bone Density, and LABS. Treatment of the events included Eliquis. Action taken with mRNA-1273 in response to the events was not reported. On 25 Feb 2021 the outcome of event, fever was resolved. The outcome of events, sick feeling, chills, nausea, night sweats, blood clot were unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1135376-1	I am writing this to let you know that two weeks after receiving my first COVID 19 vaccination I developed a severe blood clot from my right ankle to my right groin. I am Factor 5 deficient, and believe that the vaccine caused the clot. I had no symptoms at all until I noticed my right calf was swollen (no pain). My wife insisted I go to emergency because I am Factor 5 deficient and inclined to thickening of the blood. I have not been on a plane or taken a long trip by car. I walk 3 miles a day, play golf and am very active. I strongly believe the vaccine caused the clot. I will be on blood thinners the rest of my life..
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1135692-1	Pneumonia; Thrombus; Incoherent; dehydrated; threw up; Fever; A Spontaneous report was received from Consumer concerning 73-year-old of male patient who received Moderna's COVID-19 vaccine (mRNA-1273) and fever, incoherent, threw up, dehydrated, blood clot and he was diagnosed with Pneumonia. The patients relevant medical history included. The concomitant medication was not reported. On 28 Jan 2021, prior to the onset of the event, the patient received their first dose mRNA-1273 (Lot number: 012M20A, Expiration date: not provided) via unknown route in the left arm for prophylaxis of COVID-19 infection. On same day, after vaccination patient experienced fever, incoherent, threw up, dehydrated. Few days later he went to the hospital because of shortness of breath, the hospital diagnosed him with a blood clot (on 31 Jan 2021), Then on 09 Feb 2021, he went to see his family doctor because he still did not feel well and he was diagnosed with Pneumonia. Treatment was included as Eliquis and Antibiotic. Action taken with the mRNA-1273 in response to the events was not provided. On 29 Jan 2021 the outcome were recovered for fever, incoherent, threw up, dehydrated. The outcome were unknown for pneumonia and shortness of breath. On 01 Feb 2021 the outcome blood clot was recovered.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events of pyrexia, vomiting, dehydration , incoherent and thrombosis, a causal relationship cannot be excluded. Based on the current available information and the mechanism of action of mRNA-1237 vaccine, the event of pneumonia is assessed as unlikely related. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1135707-1	Blacked out; Multiple blood clots; inflammation in spine; A spontaneous report was received from a consumer, concerning a female patient of unspecified age, who received Moderna's COVID-19 vaccine and experienced multiple blood clots/ thrombosis, inflammation in spine/inflammation, blacked out/loss of consciousness. The patient's medical history included blood clots. Concomitant product use was not provided by the reporter. On an unspecified date, prior to the onset of events, the patient received their first of two planned doses of mRNA-1273 via unknown route for COVID-19 infection prophylaxis. On 11 Mar 2021, the reporter stated that her sister received the Moderna vaccine and she had multiple blood clots, inflammation in spine and blacked out. The events multiple blood clots and blacked out were assessed as serious based on IME list. Treatment information was not reported. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events multiple blood clots, inflammation in spine and blacked out was reported to be unknown at the time of this report.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1135720-1	blood clots in upper thigh down to foot; swelling of the foot; tingling; clot was burning and hot; A spontaneous report was received from a consumer concerning a 62-years-old patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced events blood clots in upper thigh down to foot, swelling of the foot, tingling, and clot was burning and hot. The patient's medical history included insertion of a blood clot filter. Concomitant medications reported were Xarelto for drug use for unknown indication. She is also taking 17 other medications (not provided) and has other medical issues (not provided). She also has a blood clot filter inside her body (location not provided). She also wears compression and diabetic socks. On 25 Feb 2021, prior to the onset of the events the patient received the first of two planned doses of mRNA-1273 (lot/batch: L014M20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On an unknown date, the patient experienced the event(s) blood clots in upper thigh down to foot, swelling of the foot, tingling, clot was burning and hot. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not reported. The outcome of event(s), blood clots in upper thigh down to foot, swelling of the foot, tingling, clot was burning and hot was unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1137221-1	Blood clots in both legs; Cramps and spasms on her right and left legs; Painful nodules under both arms and shoulders; Itching down to her toes and left side of the body; Injection site itching; A spontaneous report was received from a consumer concerning a 79-year-old female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced Cramps and spasms on her right and left legs, injection site itching, itching down to her toes and left side of the body/pruritus, painful nodules under both arms and shoulders/nodules and blood clot in both legs/Thrombosis. Medical history was not reported. Concomitant medications included losartan, cetirizine hydrochloride and pantoprazole. On 30 Jan 2021, the patient received her first planned dose of mRNA-1273 (batch number: 007MZ0A) intramuscularly for prophylaxis of COVID-19 infection. On 27 Feb 2021, she had her second planned dose of mRNA-1273 (batch number: 01021A) intramuscularly on left arm. Soon, she developed itching at injection site down to her toes and left side of the body. She developed painful nodules on shoulders as well as both under arms and it was mentioned that the one on left side was severe. On 09 Mar 2021, she began experiencing cramps and spasms on her right and left legs. She was diagnosed with blood clots in both legs while in the emergency room (ER). She was treated with hydroxyzine and apixaban. The event blood clot in both legs was considered to be medically significant. The action taken with mRNA-1273 in response to the events injection site itching, itching down to her toes and left side of the body, painful nodules under both arms and shoulders and blood clot in both legs was unknown. The outcome of the events Cramps and spasms on her right and left legs, injection site itching, itching down to her toes and left side of the body and blood clot in both legs was unknown whereas painful nodules under both arms were still sore but better.. The reporter did not provide any causal relationship between mRNA-1273 and the events injection site itching, itching down to her toes and left side of the body, painful nodules under both arms and shoulders and blood clot in both legs.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1137224-1	Develop a blood clot on their left knee; Left leg kept swelling up; A spontaneous report was received from a physician concerning a 83-year-old, female patient, who received Moderna's COVID-19 Vaccine (mRNA-1273) and experienced a blood clot on her left knee and left leg kept swelling up. The patient's medical history included cholesterol unspecified. Products known to have been used by the patient, within two weeks prior to the event, included vitamin D and medications for cholesterol. On 26 Feb 2021, prior to the onset of the symptoms, the patient received her first of two planned doses of mRNA-1273 (Batch number: 038K20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On an unspecified date after the vaccination patient developed blood clot on left leg, leg also kept swelling. Relevant treatment for the event included blood thinners and aspirin. Action taken with mRNA-1273 in response to the events was not provided. The outcome of the events, blood clot on their left knee and left leg kept swelling up, was unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1137739-1	2 DAYS AFTER SM RECIEVED HIS MODERNA VACCINE IN HIS LEFT ARM HIS RIGHT ARM TURNED BLUE. HE WENT TO THE EMERGENCY ROOM TO FIND OUT HE RECIEVED A BLOOD CLOT WHICH THEY THINK WAS CAUSED BY THE VACCINE. HE WAS PUT ON BLOOD THINNERS THAT DAY AND SENT HOME 6 HOURS LATER. HE WILL BE ON BLOOD THINNERS FOR UP TO 6 MONTHS.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1137940-1	About 11:30 PM on 03/15/2021, started gasping for breath when laying down, sat up in lazy boy for a hour and half, beathing returned to normal. About 3:30 PM on 3/16/2021, started gasping for breath again. Called my daughter and she transported me to Hospital. They tested for covid - results negative. They did a C-Scan of my chest and saw a blood clot in each lung. While hooked to the monitors, they found I had Afib. They also did an ultra-sound of my legs and found a small clot in each leg below the knees in a small vein. I have never had any record of clots or Afib. They transferred me to Medical Center early on 3/17/2021. They started a heparin drip to soften the clots. In the afternoon of 3/17/2021, they performed a Suction Thrombectomy to remove the clots from my lungs. On 3/19/2021, they put me on Eliquis Tab. I was released from the hospital in the evening of 3/20/2021.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1139147-1	I went into DKA, my left eye bled, my left arm has a blood clot I was in hospital 9 days and a rehabilitation place for 11 days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1142665-1	Working full time during this post injecting time but 2days later sarted with nausea and vomiting and then itching and then next day asthma started to flare up and by the next day it developed into full blown exacerbation. As everything worsened and couldn't control anymore with inhaler and even tried nebulizer, I went to hospital. I went to ER that day on 2/17. They did breathing treatments, several Blood tests, chest xrays, Epi injection, IV steroids and several other meds, oxygen, and was admitted. As it worsened I had to be intubated 3 different times, where I would get to point they thought could be weaned off and for a day or two and would be a little better and then would worsen again requiring me to be reintubated for a total of about 10 days on the ventilator. During this time on vent second time I developed pneumonia, staph/MRSA, UTI. After being put on vent 3rd time I had to be life flighted to bigger hospital ICU. Then after a few days they were able to wean of ventilator but during which time I developed laryngospasms. I continued to improve with breathing but mentioned I had leg pain and the said probably due to low sodium and lack of mobility. I was finally released on 3/5 and discharged home with home health but sent home in supplemental oxygen due to says dropping and poore tolerance without sitting O2 stay at hospital prior to DC home. After getting home the next day or two right leg pain worsened along with breathing worsenining and HH sent me to PCP who discovered blood clots and sent me to local er, that day which was 3/8. That hospital evaluated breathing issues and took blood work and chest xrays, started me on heparin drip and sepsis protocol and then called in life flight to send me to bigger ICU hospital within a few hours after arrival which they debated on ventilator or bipap but ended up on bipap and flown to hospital. There I was treated for breathing issues, infection, clot, vocal cord dysfunction from the multiple intubations then sent home when stable again with home health RN, PT, PT and ST. Also during all this hospital stay I was treated 4x for covid and all negative. Then after coming home this last time on 3/13 i had 15-20lb with gain in a little over a week and in 3 of those last days almost 9lbs with abdominal distension and chest pain/ nausea/ dizziness/low BP but fluctuating high at times / high Hr, so referred back to PCP 3/23 who sent me to ER that day concerned of abd blessed, kidney and or liver issues/sepsis/etc. PCP sent to ER. They did blood work and Abd CT and xray not showing anything and blood levels of liver slightly elevated, with blood in urine and so they said nothing of concern even though I went from looking normal to 6/7 months pregnant in less than a week. Followed up with pulmonologist who said bases of lungs hypoventilated, PFT very poor, wheezing and tight and WBC still up and gave me more steroids and asthma meds. Went to urologist due to blood in urine and also having urinary retention. They advised me to self cath and I will follow up in 1 month to determine if they need to do more tests to look into it more. PCP still concerned about these new abd. And other symptoms, and referring to specialists (cardiologist, Gastrointerologist, and Nephrologist) along with others already seeing too find the root of what are causing these new problems and if other organs are being affected with this whole immune response. I work full time, had 3 kids and works out with no limitations in my daily activities or ability to function, however since all this and being the hospital for pretty much most of laar month, the ongoing breathing issues (which I'm still requiring O2 at home) and all the other medical issues and the physical weakness/ lack of cardiopulmonary endurance I am unable to work (on short term disability/FMLA) and limited to home with ability to do much at all. Now I'm left with decreased income and tons of medical bills to come which is very concerning in itself how I will even pay since i can't work right now, but the most concerning is I'm still in this waiting game to figure out why things are not improving that much and why these other symptoms are arising. I have worked my whole life and love my work and my life and was trying to protect from this horrible virus by doing my part and getting vaccinated. But it seems that my life went from nothing wrong to one crazy medical reaction to another after this 2nd covid-19 19 moderna injection.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1142915-1	Blue/black blood clot on middle finger right hand; Burning in middle finger on right hand; Itchiness on middle finger on right hand; Pain in middle finger on right hand; A spontaneous report was received from a Consumer concerning a 55-years-old female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced events burning in middle finger on right hand/burning sensation, itchiness on middle finger on right hand/pruritus, pain in middle finger on right hand/pain in extremity, blue black blood clot on middle finger right hand/thrombosis. The patient's medical history was not provided. No relevant concomitant medications were reported. On 2 Jan 2021, prior to the onset of the events the patient received their first of two planned doses of mRNA-1273 (lot batch: 037K20A) intramuscularly for prophylaxis of COVID-19 infection. On 20 JAN 2021 1.00 pm, after taking mRNA-1273, the patient felt weird sensation in right hand middle finger. She felt burning, itchiness, pain and had a dark blue/black blood clot on inside of finger. 2 hours later it started to fade down (purple in color). In the evening it was considerably lighter. On 21 Jan 2021 there was a light imprint of the blood clot. Blood clot went away later that day. Patient did not received treatment. Action taken with mRNA-1273 in response to the events was not reported. On 20 Jan 2021 the outcome of events burning in middle finger, itchiness on middle finger on right hand, pain in middle finger on right hand was resolving. On 21 Jan 2021 the outcome of event blue black blood clot on middle finger right hand was resolved.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1143465-1	Superficial blood clot in the leg/area is sore to touch; A spontaneous report was received from a consumer (patient), concerning a 82-years-old female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced superficial blood clot in the leg/area is sore to touch (thrombosis). No medical history was reported. No concomitant medications were reported. On 19 Jan 2021, the patient received the first of two planned doses of mRNA-1273 (Lot number: not provided), on 15 Feb 2021, the patient received the second of two planned doses of mRNA-1273 (Lot number: 027L20A) intramuscularly for prophylaxis of COVID-19 infection. On an unknown date, the patient experienced a superficial blood clot in her leg (medically significant). She also reported that the area was sore to touch. The doctor prescribed medication to treat her problem, but they said to wait a week to see if it solved on its own before taking it. She was also applying ice pack on the affected area. No further information as reported. The patient received both scheduled doses of mRNA-1273 prior to the event, therefore action taken with the drug in response to the event is not applicable. The event of superficial blood clot in the leg/area is sore to touch (thrombosis) was unresolved. The reporter did not provide assessment for the event of superficial blood clot in the leg/area is sore to touch (thrombosis).; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1144314-1	patient complained of shortness of breath of 3/22/2021 to primary care physician approximately 1 week post covid vaccine. CT Scan was ordered. Confirmed on 3/26/2021 as Left Lower lobe occlusion and diagnosed with Pulmonary embolism. On 3/26/2021 at Emergency department, he also showed bilateral PE with large clot burden. Patient is hospitalized a this point since 3/26/2021 and has been treated for PE since. He was discharged on 3/28/2021 and continue treatment as appropriate with Rivaroxaban.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1144392-1	Last Moderna vaccine March 3, 2021. On March 17 began not feeling well. Never felt this way. Then on March 25 had severe low abdominal pain that sent her to ER.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1145194-1	Patient had Moderna covid vaccine #1 on 3/21/21. Patient presented to the emergency dept on 3/23/21 complaining of a discolored right arm (turning purplish) and swollen. Patient had a blood clot in Sept 2020 and was on anticoagulation until Jan 2021. Ultrasound of the RUE found an occlusive thrombus, R subclavian and right axillary vein as well as a nonocclusvie thrombus within the inferior R internal jugular. Patient was initiated on apixaban anticoagulation. Patient was seen back in the office and had a repeat ultrasound due to lack of improvement on 3/26, thrombus was unchanged. Patient continued on apixaban. Patient called the office on 3/29 with no improvement in his arm. Patient to be re-evaluated
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1146300-1	Systemic: blood clots-Medium, Systemic: Nausea-Medium, Systemic: Neurological Disorder (diagnosed by MD)-Medium, Systemic: Stroke-Severe

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1147138-1	Very hard to breath for few minutes; Swollen arm; Nausea; Headaches; Blood clogging; A spontaneous report was received from a consumer concerning a 63 years old female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced events very hard to breath for few minutes/dyspnoea, blood clogging/thrombosis, headaches, nausea, swollen arm/peripheral swelling. The patient's medical history included high blood pressure. Concomitant product use was not provided by the reporter. On 15 Jan 2021, the patient received their first of two planned doses of mRNA-1273 (lot: 027L20A) for prophylaxis of COVID-19 infection. On 08 Feb 2021, the patient received their second of two planned doses of mRNA-1273 (lot: 0101M20A) for prophylaxis of COVID-19 infection. The first dose of vaccine was taken in non dominant arm and second dose was taken in right dominant arm. On an unknown date, the patient experienced swollen arm. In between the two doses she felt very hard to breath for few minutes and when she took a deep breath it was fine. She had tylenol for the headaches after the second dose on 08 February 2021. She also had nausea along with headaches. She had blood clogging when she was in a blood donation camp. The event very hard to breath for few minutes and blood clogging were considered as medically significant. Thiamine was provided as medication The patient received both scheduled doses of mRNA-1273 prior to the events; therefore, action taken with the drug in response to the events is not applicable. Follow-up received on 17 Mar 2021 included updated date for first dose of vaccine, and the events very hard to breath for few minutes/dyspnoea, blood clogging/thrombosis.; Reporter's Comments: Based on the current available information which shows a strong temporal association between the use of the mRNA-1273 and the reported events and excluding all other etiology, a causal relationship cannot be excluded. Nausea and headache are consistent with the safety profile of the product
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1148421-1	At 2AM on Saturday, March 13, 2021, I began passing blood into the toilet. That evening I went to the Emergency Department and was admitted as an inpatient. I was treated there and discharged to home on Tuesday, March 16. Over the next two days I passed substantial amounts of coagulated blood. My diagnosis was colitis.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1148710-1	blood clot in lower leg; achy; Pain in the back of leg that would not go away; body ache/ache; headache; A spontaneous report was received from a consumer concerning a patient age ,57 years old male patient who developed Blood clot in lower leg/Thrombosis, Body ache/Myalgia, Headache/Headache and pain in the back of the leg that would not go away/pain in extremity. The patient's medical history included Central Retinal vein occlusion (CRVO). Concomitant product use was not provided/unknown by the reporter. The patient received their first of two planned doses of mRNA-1273 (Batch number not provided) on unknown date. The patient received their second of two planned doses of mRNA-1273 (Batch number not provided) on 03 Feb 2021, intramuscularly in the (unknown injection site) for prophylaxis of COVID-19 infection. On the same day as receiving the vaccine 03 Feb 2021 patient reported as having pain in back of the leg that would not go away. Patient also had body aches, felt achy and had headache. Patient went to Emergency room on 03 Feb 2021 and was diagnosed a blood clot in lower leg. The event blood clot in lower leg was assessed as medically significant based on IME list. Treatment information was not provided/unknown. Patient is following up with a hematologist. The patient received both scheduled doses of mRNA-1273 prior to the event(s); therefore, action taken with the drug in response to the event(s) is not applicable. The outcome of the event(s), Blood clot in lower leg/Thrombosis, Body ache/Myalgia, Headache/Headache and pain the was unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events (pain, myalgia, and headache) a causal relationship cannot be excluded. A very limited information regarding these event (Thrombosis) has been provided at this time. Noting the history of central retinal vein occlusion may remain as a risk factor for thrombosis. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1148711-1	Her leg started bothering her, the pain was quite heavy and consistent; pain under her left arm; blood cloth in her leg; a little nauseous; headache; tired; pins; A spontaneous report was received from a consumer concerning a female patient of 70-years-old, who received Moderna's COVID-19 vaccine(mRNA-1273) and experienced Pins/ Paraesthesia,nausea,tiredness,headache,pain in leg was quite heavy and consistent, blood cloth in leg, pain under left arm. The patients medical history was not provided .Concomitant medications reported included Eliquis starter pack twice a day,Latanoprost for Glaucoma,Omeprazole for heart burn. On 15 Jan 2021,prior to the onset of events, the Patient received their first of two planned dose of mRNA-1273(Lot number: 037K20A) vaccine via unknown route for prophylaxis of COVID-19 infection. On 12 Feb 2021,prior to the onset of events, the Patient received their second of two planned dose of mRNA-1273(Lot number: 024M20A) vaccine via unknown route for prophylaxis of COVID-19 infection On 12 feb 2021,That evening she got pins again.The next day she was a little nauseous, tired and had a headache. Her leg started bothering her 3 days to a week after the second dose.And the pain in her leg was quite heavy and consistent, she thought it might be her sciatica. She went to the doctor and ended up getting a doppler and they discovered a blood clott there. She had never had a blood clott before in her life.she was feeling pain under her left arm as well. No Treatment for the event was provided. Action taken with the mRNA-1273 in response to the event was not applicable. the event pins were considered resolved on 13 Feb 2021. The events nauseous,tiredness,headache were considered resolved on 14 Feb 2021. The outcome of the events pain in leg was quite heavy and consistent, blood cloth in leg, pain under left arm was unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of these events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1150203-1	pain in left arm, numbness. Then developed a itchy rash and a lump in the cubital fossa and swollen lymph nodes. Initially took benadryl, then went to the ED. An ultra sound showed a clot located in her left cubital fossa. Transferred to Med center started on eliquis and released from the hospital. Now complaining of pain in the left leg. Adv caller to follow up with her pcp about the pain.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1151065-1	Stroke; Blood clot on the brain; I Just sort of zoned out on them; Arm soreness; I Just kept getting tireder; A spontaneous report was received from a consumer, concerning a 69-years old male patient, who received Moderna's COVID-19 vaccine and experienced stroke, blood clot on brain, arm soreness without redness and swelling, he just kept getting tireder, he just sort of zoned out on them. The patient's medical history was not provided. Concomitant product use was not provided by the reporter. On 05 Jan 2021, prior to the onset of events, the patient received the first of two planned doses of mRNA-1273 (lot number 037K20A) via unknown route for COVID-19 infection prophylaxis. On 05-FEB-2021, prior to the onset of events, the patient received the second of two planned doses of mRNA-1273 (lot number 032L20A) via unknown route for COVID-19 infection prophylaxis. On 05 Feb 2021, after the second dose of vaccine, the patient experienced arm soreness and fatigue, He did not get any medication or treatments for his arm soreness without redness and swelling and and just kept getting tireder events. On10-MAR-2021 patient experienced just sort of zoned out of them, stroke, blood clot on brain. An ambulance was called for him and he was taken to the hospital where he received MRIs, and CT scans. He states they think it was a stroke. No treatment medication was reported. Action taken with mRNA-1273 in response to the event was not applicable. The outcome of the events (arm soreness without redness and swelling and he was just kept getting tireder) was recovered but the events (stroke, blood clot on brain, he just sort of zoned out on them) not resolved at the time of this report.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1152023-1	developed a blood clot in right leg DVT
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1152349-1	developed a blood clot in left calf
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1153133-1	Blood clot; Swelling in lower leg; A spontaneous report was received from a consumer concerning a male patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced swelling in lower leg and blood clot . The patient's medical history was not provided.Concomitant medications reported included blood thinners Xarelto. On 21 Feb 2021 prior to the onset of the events, the patient received his first of two planned doses of mRNA-1273 (Lot number: unknown) for prophylaxis of COVID-19 infection. Treatment for the events was unknown. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events, swelling in lower leg and blood clot was unknown.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1153172-1	Ischemic stroke; Right side of brain and blood vessels affected badly and found blood clots because of ischemic stroke; A spontaneous report was received from a consumer concerning an 89-years-old male patient, who received Moderna's COVID-19 vaccine (mRNA-1273) experienced ischemic stroke and right side of brain and blood vessels affected badly and found blood clots because of ischemic stroke/thrombosis. The patient's medical history included hearing issue. Concomitant medication included losartan. On 24 Feb 2021, approximately 12 days prior to onset of the events, the patient received their first of two planned doses of mRNA-1273 (batch number: 001A21A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 08 Mar 2021, the patient got ischemic stroke. The patient was in hospital for 3 days and doctor put him on blood thinner. MRI (magnetic resonance imaging) of his upper body revealed his right side of brain and blood vessels affected badly and found blood clots because of ischemic stroke. Treatment for the event included blood thinner. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events ischemic stroke and right side of brain and blood vessels affected badly and found blood clots because of ischemic stroke was not reported.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1155160-1	"Patient received Moderna vaccine dose #1 on 3/3/21 at approximately 0900. In the afternoon of 3/4/21, patient had sudden onset shortness of breath while at home, as well as fatigue and low energy. Patient knew that a potential side effect of vaccine was fatigue, so he assumed this was just a normal side effect to be expected. States that his difficulty breathing and fatigue continued for about a week. After continued urging from wife to seek medical care, patient visited urgent care center on 3/15/21. The urgent care did a D-Dimer lab, and an x-ray of chest. X-ray ""showed something abnormal"" so they did a CT scan as well, which showed a ""huge pulmonary embolism."" D-Dimer lab was elevated. Patient was transported to hospital by ambulance and had immediate surgery to remove blood clot from lungs at approximately 2100 on 3/15/21. Per patient, the surgeon said the clot was about the size of his palm. Surgeon reported that there was ""100% blockage to left lung from the aortic artery and 90% blockage to right lung from aortic artery."" A DVT in popliteal vein was identified, and the surgeon assumes the clot started in the popliteal vein and broke off and traveled to lungs. Patient was admitted and hospitalized until he was discharged home on 3/18/21. During course of hospitalization, genetic testing was done and it was determined that the ""genetic marker for clots was negative, so they think the clot is from a one-time event--patient's family does not need to be concerned that they have clotting issues."" Patient states that he was started on heparin in the hospital, and was discharged home on Eliquis blood thinner, which is being tapered. Patient states that the doctors think he will be on Eliquis for about 6 months and will not need to be on it for life. Doctors say the DVT in leg will dissolve on its own. During hospitalization, an echogram and cardiac cath was performed, which showed ""an old heart attack on the left side, but that collateral vessels have built up already."" Right side of heart was weaker from having to work so hard due to blood clots in lung, but prior to discharge from hospital, the right side of heart was already showing improvement. A heart cath was done to check vessels and it was determined no stents were needed, although there was a small clot in left anterior descending artery. Patient was also newly diagnosed during course of hospitalization with Type II Diabetes in a non-obese person, HTN (which patient had never had before), Coronary Artery Disease, Hyperlipidemia, 1st degree AV block. Was discharged home with metformin, losartan, chlorthalidone, atorvastatin, and Aspirin. The chlorthalidone was discontinued, due to ""blood pressure dropping too low--they think his high blood pressure was situational"" per wife of patient. Medical doctors advised against receiving the 2nd dose of Moderna vaccine. Patient being followed by Internist at hospital. Patient's shortness of breath resolved ""immediately after clot was removed"" but is continuing to be followed while on new medications and blood thinner dosing. The above information was provided during telephone interview of patient and his wife."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1155524-1	Pt. states she started to spit blood approximately 12 hours after Covid 19 vaccines. It happened both times. 1st dose given on 2/25/21, reporting second dose from 3/25/21 because pt. sought ER care at . She states the blood was coming from throat area, not vomiting. She just spit out blood and blood clots. Dr. will send her to ENT to rule out lesion.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1159805-1	Patient received first dose of Moderna vaccine. Later that day, patient experienced heavy vaginal bleeding that included blood clots and required the use of a sanitary pad. She had her period the previous week and it had since stopped so she does not believe it's her period. At the time of the call to the pharmacy, 8:30am the following day, patient was still actively bleeding. Advised patient to monitor for other signs and symptoms of bleeding such as excessive bruising, nose bleeds, gum bleeds, and to check her stool for the appearance of coffee grounds. Patient was calling to try to find a doctor nearby who could see her. Advised patient that if she could not find a doctor to go to urgent care if the bleeding continued or worsened.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1159908-1	She got her vaccine and had no reactions at all. On 3/12/21 she was in her car and started having stabbing pain in the left side of her chest under the rib. She thought it was gas pains and dismissed it, and then it started radiating towards the front and the stomach. On 3/16/21 she went to the ER and they did a CT scan that showed that she had a blood clot on her spleen. With that they gave her the diagnosis of idiopathic splenic infarction. She was then sent by ambulance to another hospital to be admitted in case she was needing surgery. She was given heparin and on that for 2 1/2 days. They determined that 60% of her spleen is dead, blood supply cut off to it, and the other 40% is nonfunctional. On 3/19/21 they gave her vaccines of Prevnar 13, HIB, meningococcal, meningococcal B, and a flu vaccine. She was discharged on the 3/20/21 with Plavix. She saw her regular doctor yesterday and he diagnosed her with functional asplenia and to continue the Plavix. She is no longer having any symptoms after dissolving the blood clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1160353-1	Sore arm at injection site, extreme fatigue, sudden onset uncontrolable menstrual bleeding & blood clots, felt faint & had some blurred vision. Heavy bleeding & heavy clotting continued until next morning (I have not had any issues with my cycle since my surgery last year so this was not normal). Woke up with fever of 100.7 Saturday morning at 4:57am; fever did not go away until Saturday 8:45pm. Had flu-like symptoms, fatigue, fever, chills, intense all over muscle soreness. Felt better & symptoms were gone by Tuesday 03/02/2021. Menstrual cycles went back to normal following month after 2nd vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1163787-1	Patient presented to our ED on 3/30/21 @ 1149 with complaints of SOB and chest pain. D-Dimer was elevated > 3 times upper normal limit so CTA was performed. CTA showed bilateral lower lobe PE with moderate thrombus burden. Patient was given Lovenox 1mg/kg (90 mg) x 1, pain medication, and home medications he had not yet taken at home on 3/30. He was then transferred to a higher level of care. Course at transfer hospital unknown at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1163967-1	Felt crummy on day one and two better on day three and on day four developed four blood clots on Coumadin with an INR of 4.5 and died the next day on 3/31/2021.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1169237-1	Blood Clot, lower leg (left calf muscle). MD. recommed Xeralto 20mg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1171092-1	Thrombosis on right foot
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1172377-1	I had an extensive oral surgery about 36 hours after getting the vaccination. I had non profuse bleeding after the surgery. I experience a condition called Liver Clot (currant jelly clot). This condition and the bleeding continued for over 5 days following the surgery or 6.5 days after the vaccine dose 1.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1172529-1	Blood clot that caused total vision loss in right eye. It occurred on 04/03/2021. Was treated by ophthalmologist at eye Center, and at Hospital,
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1173053-1	Patient states that he woke up the next day extremely fatigued. About 3pm, he could not move his arm/right hand and had trouble walking. He called his doctor's office who told him to call 911. At the hospital, he was given emergency medication for a blood clot. He states the ER told him his stoke was from the covid vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1173594-1	Blood clot on his left leg, from his groin to his ankle/still having the blood clot; A spontaneous report was received from a consumer concerning an 80-years-old male patient, who received Moderna's COVID-19 vaccine (mRNA -1273) and had blood clot on his left leg, from his groin to his ankle. The patient's medical history was not provided. No relevant concomitant medications were reported. On 02 Jan 2021, prior to the onset of the symptoms, the patient received their first of two planned doses of mRNA-1273 (Batch number: 039KZ014) intramuscularly for prophylaxis of COVID-19 infection. On 30 Jan 2021, approximately two weeks prior to the onset of the symptoms, the patient received their second of two planned doses of mRNA-1273 (Batch number: unknown) via unknown route in the left arm for prophylaxis of COVID-19 infection. On an unknown date in Feb 2021, the patient reported having a medically significant event, blood clot on his left leg, from his groin to his ankle and is still having it. The patient taking medication for it. The patient received both scheduled doses of mRNA-1273 prior to the events, therefore action taken with the drug in response to the events was not applicable. At the time of this report, the outcome of the event had blood clot on his left leg, from his groin to his ankle was not resolved.; Reporter's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1174091-1	Very heavy period with clotting. I was on day 3 of my period at the time of vaccination. Immediately upon receiving the vaccine I had uterine cramps and heavy bleeding with large clots continued throughout the day and into the next day. Had some dark bleeding/discharge about 4 days after my period ended for 2 days. This also happened during my first dose of Moderna, although my first dose was on the first day of my period so I didn? t quite register it since bleeding is normally heavy at the beginning of my period. First period was also substantially heavier with clots and had some dark bleeding/discharge 4 days after my period ended

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1175650-1	After the first Dose I exp tingling on right side of my face. I woke up next day exp headache and fatigue realized right eye lid wasn't closing same time as left eyelid. I went to walk in clinic next morning confirmed Bell's Palsy started Prednisone. I do have a genetic blood clotting disorder for 18 years.Since the vaccine on 3/31 I woke up with pain in my foot on lower right leg applied ice and took Tylenol. I scheduled a doppler at the Imaging center on 2/7 confirmed blood clots and prescribed Xarelto.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1176103-1	Pelvic blood clot in right common iliac artery
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1176319-1	Started with stomach pain that continued to increase. Went to E.R. After variouse test found a blood clot near stomach in vein.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1177465-1	The patient started to feel swelling, redness on the interior of upper right leg.It was a swollen vein. She went to the physician on 2-15-2021. She was diagnosed with having a blood clot and had to have it removed. She is wearing compression stockings now 24/7.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1177623-1	Woke up on 4/4 with a golf ball sized, raised welt at the injection site. It was hot and very red. The next day, it was even larger, by 4/7, it had completely gone away. Everyone thinks I should also mention that I was admitted to the hospital with chest pain, high BP/Pulse on 3/31, they did a D-Dimer on me and it was elevated, suggesting a clot has formed and is in the process of breaking down. I had a CT of my chest and no PE was found, but I had thickening of my left lobar pulmonary artery wall, further supporting the suggestion of a possible blood clot. After an echo stress test, I was discharged the next day on HTN meds. I was experiencing chest discomfort prior to my vaccine, but again, EVERYONE is telling me I need to report this, so I am. :)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1178258-1	Blood clot in Left Arm; itchy rash from elbow to shoulder; lump on the underside of her biceps; nauseous; couldn't lift arm up; arm sore at the injection site; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot in Left Arm) in a 63-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No reported medical history.). Concomitant products included CLONAZEPAM for Anxiety, STEROIDS for Back disorder, CITALOPRAM and MORPHINE SULFATE for an unknown indication. On 23-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 23-Mar-2021, the patient experienced VACCINATION SITE PAIN (arm sore at the injection site). On 24-Mar-2021, the patient experienced MOVEMENT DISORDER (couldn't lift arm up). On 25-Mar-2021, the patient experienced RASH PRURITIC (itchy rash from elbow to shoulder), MASS (lump on the underside of her biceps) and NAUSEA (nauseous). On 27-Mar-2021, the patient experienced THROMBOSIS (Blood clot in Left Arm) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clot in Left Arm), RASH PRURITIC (itchy rash from elbow to shoulder), MASS (lump on the underside of her biceps), MOVEMENT DISORDER (couldn't lift arm up), NAUSEA (nauseous) and VACCINATION SITE PAIN (arm sore at the injection site) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Treatment included Benadryl for symptoms and Eliquis for blood clot.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1178589-1	Significant swelling at base of skull with severe headache starting 11 hrs after vaccination. Superficial swelling of vein on left forearm started 46 hrs after vaccination in left upper arm. This vein had a clot that proceeded to swell and burst causing significance pooling of blood with tracking on inside of L forearm. Left arm remains warm and throbbing.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1178615-1	Pain in right leg, visited doctor, had ultrasound, found blood clot, prescribed aspirin.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1179009-1	Blood clot (brain)/ stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1179370-1	blood clots in right leg and left lung
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1180000-1	Pain in left calf began on April 2 during a long road trip. Pain continued for several days. On April 5, I visited Urgent Care for a suspected muscle strain. They referred me to the Hospital emergency care for an ultrasound. The ultrasound detected deep vein thrombosis (DVT) (blood clot) in the popliteal vein in my left leg, above and below my knee. I am currently taking anticoagulant (Eliquis) medication to treat the DVT.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1180842-1	Received 2nd Moderna Covid shot on 03/29/2021 and on 03/31/2021, felt what I thought was a muscle cramp that never went away. Went to urgent care where the shot was given on unspecified date and they diagnosed what I thought was a muscle cramp area, as a blood clot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1181614-1	"Moderna EUA Pt started having left leg pain, discoloration (blue-purple) and swelling on 3/9/2021. Pt went to the hospital and was later called to come back to the hospital because she had a blood clot. Pt was sent to clinic and had a procedure by to remove clots. Pt states that the clot was ""2/3 size of thigh."" Pt states that she is scheduled to have a second procedure in May to remove blockages from iliacs."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1181756-1	blood clot in left arm causing enormous swelling
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1182802-1	pain in legs on 3/22/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1184699-1	Heart attack (blood clot in major artery) approximately 6 weeks post 2nd Moderna dose. Surgery done in cath lab to address blockage. Released from hospital several days later.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1184977-1	3/14/2021 Tightness in chest, attributed to physical work outside, 3/15 still felt tightness, 3/16 getting winded and went to the ER around 5:30pm, they took blood and ran test EKG, X-ray and ECHO complete with TTCV, ultrasound of chest, and test showed multiple blood clots in both lungs; by this time he was having trouble breathing and was admitted because of the blood clots. He was given Heparin and Vicodin through IV, on 3/17 they ran more test to see where blood clots came from and where they came from but nothing was found in legs. he was in a lot of pain Wednesday and he was sent home on 3/19.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1185321-1	I started to feel pain in both knees going down both legs around 4/1/21. An ultrasound was performed and a blood clot was found in my left lower leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1186196-1	fever and associate shivering day after the injection. Sever incapacitating muscle spasm in left leg two days after ijection . Three weeks later second muscle spasm but not as sever. Discomfort in leg prsisted and diagnosed through ultra sound blood clot in left leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1186830-1	Patient reported to the pharmacy that she was admitted to the hospital for blood clots shortly after receiving the vaccination and was there for at least 10 days
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1187825-1	blood clot in superior mesenteric vein found with Abd CT scan placed on lovenox then xarelto with relief of sxs - abd pain
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1188223-1	3/16/2021 Received 2nd Moderna Covid-19 vaccine 3/25/2021 Trouble breathing when exercising so stopped after 5 min and usually do 20 mim; temp elevated for me normal 97.6 and was between 98.6 and 99.3, hacking up yellowish phelgm. Phelgm had been coming up for longer period of time. 3/26/2021 went to clinic and put on treatment for possible bronchitus. Z-Pak and Prednisone 5 days and asked to schedule x-ray to see if something more showed. 3/30/2021 Completed meds and x ray scheduled. No improvement in breathing 4/1/2021 was finally able to get chest x-ray 4/2/2021 x-ray showed nodule on lung & calcified grandular - told to schedule ct to check out nodule - told clinic still no change in condition 4/5/2021 called clinic again due to pain overnight in left lung. Trouble getting ct scheduled so clinic sent to me emergency room rather than schedule ct 4/5/2021 - At ER blood work, x-ray and ct scan completed. CT chest was significant for large bilateral pulomanry emboli w r ventricular strain. Also r middle lobe lung nodule and r adrenal nodule. Started on heparin drip and admitted. Hospitalized 4/5 and sent home 4/8. Sent home on Xarelto to continue at least 3 months and consult w family doctor to determine cause of submassive clots and further treatment.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1188275-1	Pulmonary Embolism with Acute Core Pulmonale -no sign of DVT. Multiple blood clots in major arteries of both lungs. Event began with tiredness over about. 2 week period and ended in acute event with trouble breathing. Taken to hospital via ambulance. Tests also showed right heart damage due to stress from clots.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1188842-1	Blood clot; Fatigue extreme; muscle soreness; loss of appetite; Unable to get out of bed for four days so then they took her to the ER; Lack of circulation on legs due to blood clot produced a wound to develop in her leg; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot) and FATIGUE (Fatigue extreme) in a 74-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 025A21A and 007M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event. On 28-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 26-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criteria hospitalization and medically significant), FATIGUE (Fatigue extreme) (seriousness criterion hospitalization), MYALGIA (muscle soreness), DECREASED APPETITE (loss of appetite), MOBILITY DECREASED (Unable to get out of bed for four days so then they took her to the ER) and LIMB INJURY (Lack of circulation on legs due to blood clot produced a wound to develop in her leg). The patient was hospitalized on 22-Mar-2021 due to FATIGUE and THROMBOSIS. At the time of the report, THROMBOSIS (Blood clot), FATIGUE (Fatigue extreme), MYALGIA (muscle soreness), DECREASED APPETITE (loss of appetite), MOBILITY DECREASED (Unable to get out of bed for four days so then they took her to the ER) and LIMB INJURY (Lack of circulation on legs due to blood clot produced a wound to develop in her leg) had not resolved. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown Route), the reporter did not provide any causality assessments.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1188857-1	experienced blood clot; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (experienced blood clot) in a 55-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 038A21A) for COVID-19 vaccination. Concurrent medical conditions included Factor V Leiden mutation. Concomitant products included RIVAROXABAN (XARELTO) for Factor V Leiden mutation. On 16-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 23-Mar-2021, the patient experienced THROMBOSIS (experienced blood clot) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (experienced blood clot) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. The patient was treated with blood thinner Xarelto (rivaroxaban) on a daily basis for factor V mutation and was treated with the same for blood clot.; Sender's Comments: Limited information regarding the event has been provided at this time and a causal relationship cannot be excluded
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1189629-1	Blood clot on her eye after vaccination; Her HCP told her that she could wait even 3 months to get the 2nd dose; This spontaneous case was reported by a patient (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clot on her eye after vaccination) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Enucleation of eyeball. In March 2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot on her eye after vaccination) (seriousness criterion hospitalization) and OFF LABEL USE (Her HCP told her that she could wait even 3 months to get the 2nd dose). At the time of the report, THROMBOSIS (Blood clot on her eye after vaccination) and OFF LABEL USE (Her HCP told her that she could wait even 3 months to get the 2nd dose) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was unknown. Treatment information not provided. Her HCP told her that she could wait even 3 months to get the 2nd dose Patient decided not to give more information as she believed it wouldn¦t help her. Reporter did not allow further contact; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1189636-1	Blood clot in leg; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clot in leg) in an 81-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 039KZ014) for COVID-19 vaccination. The patient's past medical history included No adverse reaction (No reported medical history.). On 02-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 16-Jan-2021, the patient experienced THROMBOSIS (Blood clot in leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clot in leg) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Treatment information was not provided. Company Comment Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1191982-1	Swelling in lower right leg caused by blod clot in thigh

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1192235-1	New onset DVT/PE blood clots. Diagnosed after patient had new onset right calf pain and cramping about 9 days after receiving second Moderna COVID vaccine, came to hospital had an ultrasound which showed clot, then had CTA chest which confirmed pulmonary embolism (both done on 4/7/2021). She has no other known risk factors for thromboembolism, however a hypercoagulability work-up is also underway and will not be completed for several months.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1192464-1	Fatigue, achy all over, pain under right rib cage that hurt threw to my back. Went to med center and was seen by Dr. He was concerned so that he insisted that I go to the hospital to be admitted on Tuesday 4/6/21. Was admitted and blood work was done and ct abdomen and car which was negative. More lab work was done on 4/7/21 which showed a elevated d-diner. At that time a ultrasound of legs done which showed a clot in my right leg. Then a ct chest was scheduled and done 4/8/21 which showed a clot in my right lung.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1192802-1	Blood Clots, Stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1193450-1	Extensive arterial thrombus of the left lower extremity. Partial right lower extremity thrombus. Interventions include heparin therapy and evaluation by cardiovascular surgery and interventional radiology
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1194118-1	Blood clots; Felt extremely ill; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Cancer and Autoimmune disorder. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant) and MALAISE (Felt extremely ill). At the time of the report, THROMBOSIS (Blood clots) and MALAISE (Felt extremely ill) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1194136-1	was hospitalized with multiple blood clots; This spontaneous case was reported by a physician and describes the occurrence of THROMBOSIS (was hospitalized with multiple blood clots) in a 61-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 017B21A) for COVID-19 vaccination. The patient's past medical history included Surgery (on his shoulder.) on 12-Mar-2021. On 26-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 31-Mar-2021, the patient experienced THROMBOSIS (was hospitalized with multiple blood clots) (seriousness criterion hospitalization). At the time of the report, THROMBOSIS (was hospitalized with multiple blood clots) had not resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. Concomitant drugs were not reported. Treatment information was not provided.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the onset date of the reported event of thrombosis, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1194141-1	very much like a blood clot; R leg swelling; very hematoma; big red patch; warm sensation in the the leg; pain that he never had; normal pain in the injection site; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (very much like a blood clot) in a 49-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No reported medical history). On 01-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 01-Apr-2021, the patient experienced VACCINATION SITE PAIN (normal pain in the injection site). On 04-Apr-2021, the patient experienced THROMBOSIS (very much like a blood clot) (seriousness criterion medically significant), PERIPHERAL SWELLING (R leg swelling), HAEMATOMA (very hematoma), ERYTHEMA (big red patch), FEELING HOT (warm sensation in the the leg) and PAIN IN EXTREMITY (pain that he never had). At the time of the report, THROMBOSIS (very much like a blood clot), PERIPHERAL SWELLING (R leg swelling), HAEMATOMA (very hematoma), ERYTHEMA (big red patch), FEELING HOT (warm sensation in the the leg), PAIN IN EXTREMITY (pain that he never had) and VACCINATION SITE PAIN (normal pain in the injection site) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown Route) was unknown.; Sender's Comments: Very limited information regarding these events has been provided at this time. The events are probably related to the patient's comorbidities
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1194371-1	Blood clots both times in left leg after each dose. Emergency room visit required

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1194716-1	Abnormal menses. Huge clots. Velocity and volume of blood extreme. (Different from previous experiences of menorrhagia). The only other time I experienced this type of dramatic and concerning bleeding was after I took the Hepatitis vaccine (first shot) about 2 years ago. I think that's the only other vaccine I've had as an adult. There is mention of heavy bleeding as a rare side effect for Hepatitis vaccine, and I surely hope you will study this effect from Covid vaccine as it took nearly a year to regulate my cycle after the Hepatitis vaccine reaction. I've seen some reports suggesting that the excessive bleeding that women are experiencing after the Covid vaccine is due to stress and that is just plain insulting. Stress does affect the menstrual cycle; that is well-known, but this is something entirely different. I have also had small clots in the nose during the past several days, which again, is highly abnormal for me. Thank you for looking into the bleeding side effects that hundreds (if not more) women are struggling with at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1196028-1	Abnormal uterine bleeding. Started morning after an evening 2nd dose. Normally 28 day cycle very routine. Started a week early and has not stopped since, 8 and a half weeks so far. Large clots at times. Is a change from baseline, not blamed on stress please. As still ongoing and labs pending, uncertain outcome at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1196171-1	Blood clots (DVT in right leg and three pulmonary embolisms in right lung), symptoms appeared around 4/5/2021 and diagnosed on 4/9 and 4/10. Also started experiencing diarrhea during the same time period. Being treated with Xarelto and Azithromycin.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1197104-1	I had diarrhea about 20-40 minutes after receiving my first dose. I then got my period a few hours later. It was on time, however, it was extremely heavy. On the 3rd day, I began to have very large blood clots. I spoke to my gynecologist and she started me on birth control that sane day in order to stop the bleeding.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1197857-1	PATIENT BEGAN SLURRED SPEACH AND WEAKNESS IN LEFT SIDE ONE WEEK AFTER VACCINATION. DAUGHTER, WHO IS A NURSE, NOTICED AND MADE PATIENT GO TO ER. THERE, PATIENT HAD A SCAN DONE THAT REVEALED TWO BLOOD CLOTS, WHICH HAD CAUSED A STROKE, IN THE BACK OF THE BRAIN. PATIENT WAS KEPT FOR OBSERVATION OVER NIGHT IN THE HOSPITAL AND THEN RELEASED.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1198032-1	Patient has had L leg blood clot and multiple mental co-morbity-Anemia, hypertension, heart disease, hypokalemia, liver enzyme off per MD, poor nutrition, tremors, hypothyrodism, COPD, small vessel disease of the brain
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1198041-1	Menstrual period began two days after 1st Moderna vaccine with spotting on 3/27/21. One week after vaccine, heavy period began with several fist-size clots the evening of 4/1/21. Today is 4/12/21 and I am still having my period. It has lightened up but not stopped after more than two weeks. My age is 50 and I am in peri-menopause.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1198613-1	Eight days following the first vaccine I was admitted with massive blood clots to my lets and lungs. I was hospitalized for 8 days. Four days were in ICU.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1198615-1	Received Moderna COVID vaccine on 2/26/2021 and 3/26/2021. After initial dose she was fatigued. had also taken Plan B 3/1/2021. Fatigue lasted 1 month. 5-6 days after second dose she developed shortness of breath and chest pain. and 9-10 days lateraled noticed a prominent vein in the leg. Called physician office 4/8/2021 and sent to urgent care then ER, found to have superficial thrombosis in leg and multiple bilateral segmental and subsegmental pulmonary emboli.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1199088-1	My last period prior to my first vaccine shot had been in June 2020. All signs, including blood work, pointed to me having entered menopause. My mom also went through menopause at age 45 when her periods suddenly stopped, so I figured I was following in her footsteps. On Saturday, February 13, 2021, I received my first dose of the Moderna vaccine, and by Thursday morning, I was spotting. This was very surprising since I had not had a period in several months. Over the course of a couple of days, it turned into a full-blown period with the most severe cramping I've ever had in my life. By Saturday, they felt like labor contractions, and I was releasing sizable blood clots. I made an appointment at our health clinic, and all bloodwork again pointed to menopause. After my 2nd Moderna vaccine, I again experienced just a tiny bit of brown spotting a few days after the vaccine. However, it did not progress into a full-blown period that time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1200509-1	Pain started on inside of right calf on 3/23/2021, but continued to spread up and down from just above ankle to groin area. Three areas were extremely painful to touch. Diagnosed as blood clots in the saphenous vein. Put on ibuprofen, keflex and famotidine on 3/29 and advised to elevate and use moist heat to help clots resolve. Pain started subsiding on 4/3 but is still there, just not as bad. Referred to vascular surgeon, on 4/6/2021. He recommended following advice from since they seem to be resolving and are not in the deep femoral vein. As of today, 4/13/2021, pain is still there although somewhat improved.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1201270-1	Blood clot on middle joint of right hand pinky finger. Pressure bandage, heat, Advil. Decreasing in size after 18 days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1201747-1	Father started complaining about his right leg after first dose of Vaccine. After second dose on Feb 12, 2021, Father started complaining much more about his leg. On Feb 23, 2021 it was diagnosed Father had a Blood Clot in his leg.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1201948-1	X4 BLOOD CLOTS WITH 1 MONTH
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1201955-1	Shortness of breath , extreme pain 3 hospital visits until blood clot was detected,now on blood thinner Eliquis
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1202135-1	Reporting for a friend she is in the hospital, all events may not be filled in, but I thought it was important to report sooner than later due to the news on other vaccines. Patient had her second shot around 3/7/2021 u sure as she is in a rehab facility time of day was 1630. On the 17 th of March she started having back pain then wasn?t able to get herself to the hospital called 911 and was transported to the hospital and that night had emergency surgery for a blood clot in her neck. She was in rehab and got rushed back to the hospital and had a pacemaker placed. She is still in a rehab facility.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1202196-1	LOWER LIMB EMBOLISM-BLOOD CLOT, TIGHTNESS, HEAT AND PAIN WHEN WALKING, TREATING PHYSICIAN COULD NOT CONTRIBUTE TO ANY OTHER HEALTH CONDITIONS OR REASONS
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1202613-1	Seven days after Moderna vaccine I got up and started feeling a warmth in my leg that started in foot and went up and it started hurting really bad, called an ambulance and went to hospital and because of blood clots, I had to have emergency surgery, it was very huge arterial clot, and developed shingles as well.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1202649-1	Blood clots then death
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1202749-1	Pt was seen by me for full arm thrombosis and CNS thrombosis which occurred approximately 1 week following second vaccine
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1203350-1	Severe headaches began 3/19/21; Urgent Care treated with meds and did CT Scan; sent to ER on 3/25/21. Admitted to hospital. MRI and CT scans done; Lumbar Puncture done--no bacteria; diagnosed as Cerebral Vascular Sinus Thrombosis; treated with Lovonox for 5 days then switched to Pradaxa when released from hospital; given oxycodon, dilaudid, and tylenol for pain. Starting 4/1/21, numbness in right arm down to fingers. Sent to ER again on 4/8/21 as headaches and numbness worsened. MRI and CT done. Clot dissolving but pain continued at 10/10 level. Released and continues on Oxycodon, Tylenol, Elavil, without relief as of today, 4/13/21. Still incapacitated by pain and unable to walk at length, work, or any normal activities.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1203471-1	I had a blood clot in my right leg . It started to bother me on 3/19/21. I called my doctor on 3/22/21 and she advised that I be seen in the ER. The ER doctor confirmed that I had a blood clot in my leg and I was treated with Lovenex. In addition to the clot in my leg, I have multiple bruises on my arms and legs where blood vessels broke in the couple days before and after the ER visit. I also felt very out of shape, with trouble getting enough breath. Within 2 days of getting the Lovenex, I felt like I could breath normally and my symptoms improved. Not sure if it is connected to the vaccine but thought it was important to report. I am an active, otherwise healthy person who walk at least 5 miles daily. The blood clot was very unexpected.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204047-1	Same day:Tingly arm. Metal taste in mouth. That night: very sore arm. Nausea started and dizzy spell at night. Nausea lasted 2 days 3 weeks after shot: spotting about 6 days before period. Heavy period with clots that came a few days early.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204419-1	Swelling in lower right leg, ankle, foot; Venous Doppler ordered; blood clot detected; doctor examination; Eliquis prescribed.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204779-1	Clot blood; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Clot blood) in an 80-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 031M20A) for COVID-19 vaccination. The patient's past medical history included No adverse event (No medically reported history). On 20-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 17-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 31-Mar-2021, the patient experienced THROMBOSIS (Clot blood) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Clot blood) outcome was unknown. The patient was diagnosed with blood clots in legs and lung. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204782-1	Blood clot; COVID-19; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot) in a 62-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history reported.). Concomitant products included PARACETAMOL (TYLENOL) for an unknown indication. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion medically significant) and COVID-19 (COVID-19). At the time of the report, THROMBOSIS (Blood clot) and COVID-19 (COVID-19) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In 2021, SARS-CoV-2 test: positive (Positive) postive. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Products known to have been used by the patient, within two weeks prior to the event, included blood thinner medication. Treatment information was not provided. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204868-1	I peed a blood clot and had to go to the ER on 03/30/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1205036-1	On March 30th started with headaches then progressed to shortness of breath while moving. Then on April 4th transport to hospital via EMT was tested for Covid 19 results was negative. It was determined that blood clots where found in Right leg & Left leg and both lungs and now blood clot was found in heart. PT has low platelets, low blood pressure. While in the hospital pt was given herapin . Patient has remained in ICU since the 4th to present time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1205392-1	blood clots in left leg and both lungs diagnosed on 2/22, died 2/24
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1205860-1	Fatigue, dizziness, weakness. Issues with breathing caused hospitalization 3 days after shot. Blood clot located in lung and 2 in right leg. Hospital stay 11 days, step down facility roughly 2 weeks, PT/OT upon arrival home.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1205903-1	What appeared to be blood clots on both forearms.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1206430-1	I have endometriosis and my periods are heavy with painful cramps first 2 days and my period is done by the 4th day. I take aleve a day before my period which helps with my cramps. My last period started April 9th and I am still on my period today April 15th. I had heavy bleeding and blood clots the size of quarters for the first 5 days. The aleve didn?t seem to help and my pain level was at a 10, I would pass out from the pain. My period usually last 4 days and I am on day 7 and still bleeding, but not experiencing any cramps. I?ve read that Covid effects the blood clotting process and also read other women are experiencing heavier menstrual cycles after their covid shot. I also read that moderna didn?t study any side effects related to the menstrual cycle. I have an appointment today (April 15th) to get my second shot. I am not sure if I should get the second shot, due to the amount of clots I had.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1206902-1	her left leg was swollen; two red clots on left leg; right feet was hurting; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (two red clots on left leg) in a 65-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event. On 03-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced PERIPHERAL SWELLING (her left leg was swollen), THROMBOSIS (two red clots on left leg) (seriousness criterion medically significant) and PAIN IN EXTREMITY (right feet was hurting). At the time of the report, PERIPHERAL SWELLING (her left leg was swollen), THROMBOSIS (two red clots on left leg) and PAIN IN EXTREMITY (right feet was hurting) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1206927-1	"burning sensation in the lung; had labored breathing,they found several clots in right leg and lungs; It was shiny and hard and calf blewup twice the size,they found several clots in right leg; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PULMONARY EMBOLISM (had labored breathing,they found several clots in right leg and lungs) and THROMBOSIS (It was shiny and hard and calf blewup twice the size,they found several clots in right leg) in a 63-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 002B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Family history included Clot blood (Family history of clots; her father and his sister had clots after surgery). Concurrent medical conditions included Burning sensation. On 12-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 04-Apr-2021, the patient experienced PULMONARY EMBOLISM (had labored breathing,they found several clots in right leg and lungs) (seriousness criterion medically significant) and THROMBOSIS (It was shiny and hard and calf blewup twice the size,they found several clots in right leg) (seriousness criterion medically significant). On an unknown date, the patient experienced BURNING SENSATION (burning sensation in the lung). At the time of the report, PULMONARY EMBOLISM (had labored breathing,they found several clots in right leg and lungs), THROMBOSIS (It was shiny and hard and calf blewup twice the size,they found several clots in right leg) and BURNING SENSATION (burning sensation in the lung) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Ultrasound Doppler: several clots in right leg and lungs (abnormal) several clots in right leg and lungs. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant products were reported. Patient was taken to the ER (emergency room). The physician suspected blood clots and treatment included was blood thinner. Patient reported that ""patient got ultrasound done on Monday, and they found several clots in right leg and lungs"".; Sender's Comments: Based on the current available information which includes a strong temporal association between the use of the product and onset of the reported events, and excluding other etiologies, a causal relationship cannot be excluded"
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1207099-1	My menstrual cycle came on 5 days early but when it came on I had large blood clots. I?ve never had my cycle come on early. I take birth control tablets the correct way and the Saturday when I started bleed, I had taken my tablet at my regular time that day. I also usually don?t have heavy periods at all so the amount of blood so early was unusual for me.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1207493-1	moderate pain in arm; moderate headache then sudden bloody nose 2 days later from right nostril with two large blood clots. Headache remains.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1207844-1	He got the vaccine, and didn't have any real reaction, anything unpleasant, he had a numb lip and 2 fingers were numb. He called his doctor and she told him to go to the ER, which he did at Hospital. They did a CAT scan of his brain. They tried to do an MRI and they pushed his head against the machine as his back is curved and he stopped breathing. They had given him 2 narcotics for anxiety before the scan. He was then transferred to Hospital in . He saw Cardiophysiologist/IM and he found that he had a blood clot (? where), and they inserted a pacemaker through his groin as his pulse dropped to 40 BPM. He was hospitalized for 7 days, and discharged him with instructions on what he is able to do. They started him on new blood pressure medicines for his kidney's. On 3/26/21, lot# 028A21A. He subsequently had the 2nd vaccine, and all liquids emptied from his body (vomiting and diarrhea), at the same time, and then he was fine once everything settled down which lasted for about 3-4 minutes and then he was fine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1208115-1	on 03/22/2021 I had a red inflamed area on left Achilles tendon area, off work for two days, negative US for DVT I had twisted my knee on 03/29/2021, negative DVT MRI meniscus tear I got a blood clot behind my left knee on 04/04/2021 left calf started hurting on 04/01/2021, got intense on 04/04/2021 went to UC US was positive for DVT I have never had one, no health issues, was advised to report blood clot
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1208289-1	Approximately one week after receiving the second dose of the vaccine my wife felt a pain in her left leg. The pain did not feel like a muscle pain and continued for a few days. She went to an urgent care and an MRI of her leg was conducted. She was diagnosed with a blood clot and has begun treatment. My wife is healthy, young and has no family history of blood clots prior to this incident.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1208579-1	Developed blood clot in heart -- > went into Atrial Fibrilation with no prior history of A FIb -- > Large clot was thrown from the heart into the brain splitting into two clots, one blocking brain stem and one blocking right hemisphere -- > Death on 2/10/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1208878-1	Two blood clots in right lung two days after second vaccine. I was hospitalized and I am now on oxygen 24/7 and I?m taking blood thinner Eliquis.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1209527-1	Blood clot in left leg

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1210261-1	patient admitted to hospital with blood clot on 4.8.21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1210394-1	Woke up with pain in both arms
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1210412-1	I had my second shot and the same night developed a clot in my arm. I had blood coagulation labs drawn and my C protein and cardiolipin have altered and show I may get clots now. I had labs previous to this because of my gallbladder surgery and all my blood was normal.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1210977-1	Lower right leg blood clot. Currently on Xarelto blood thinner.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1211176-1	walked a mile a day and has always taken care of himself. He ran marathons. He never was bothered by walking the daily miles. Nothing happened while he walked even though he has a sarcoma. One night, after he got into bed, his hip and leg started hurting and have continuously hurt since then. His pain is debilitating and his leg is very heavy. He can not lift his leg to get into bed. His wife has to lift it with two hands. He cannot lift his leg to get into a car. He has been given Gabapentin, Norco, Tramadol, and Tylenol for pain. He needs ice packs during the day and night. He cannot sleep in his bed all night. He has gotten up to sleep in a recliner.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1212634-1	Patient presented to the ER with complaint of swelling in right lower extremity for 3 weeks.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1213492-1	"on Saturday, March 27th, patient noticed swelling in left lower leg with Shortness of breath, and heart racing. on 3/39, the patient presented to the ED and was admitted and diagnosed with a blood clot from left groin to left ankle, with ""clot in heart and lungs"". heparinized and discharged home on Coumadin currently taking 5 mg on Sunday, Monday, Wednesday, and Saturday and 2.5 mg on Tuesday and Friday each week."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1214422-1	Blood clots in left leg, hip, bilateral lungs; treated with blood thinners; was admitted to hospital from 4/1-4/3; discharged to home, stable
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1214597-1	2nd dose Moderna administered on Feb 3rd, fatigue/ shortness of breath week of Feb 23rd, admitted to Hospital on March 1st, spent 18 days in ICU, developed Blood CLOTS which one attached to lung, blood clots In left leg as well, no family history of blood clots and no injuries to cause blood clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1216137-1	The patient called to make us aware that he was diagnosed with a blood clot on Saturday and he wasn't sure if there was a connection or not but wanted us to report it.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1216324-1	Exactly 1 week from vaccination, menstruation began approximately 2 weeks before schedule and roughly 10-12 days after date of start of previous cycle. Heavy with many clots, extreme fatigue, headache, and heavier than usual cramping. Completely unexpected and NOT accompanying any usual, predictable signs of PMS or anticipated cycle. Women need to know this is happening.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1218318-1	I have a history of heavy periods which resulted in the need for an ablation in October 2018. Since the ablation my periods have been much more manageable. I got my period 13 days after receiving the shot and it started out much heavier than normal, by the second day I was bleeding extremely heavily, soaking an overnight pad in less than 2 hours with bright red blood and passing clots the size of dimes and pennies. Day 3 has started out the same as day 2. Prior to my period starting my back hurt and I had much worse than normal abdominal cramps. Nothing in my life has changed besides the vaccine, I am certain there is a connection between the two. I am not sure how long this will last, I was instructed to go ahead an report the adverse event.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1219004-1	Menstrual cycle started 9 days early and pretty heavy. Usually on schedule and not as heavy bleeding or clotting/spotting.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1219326-1	Body aches night of injection. Extreme bloody nose with clots on 3rd day after injection. Swollen and very itchy right wrist 3rd day after injection. Was able to stop bloody nose after 1 hour, did not need medical help. Have not experienced a bloody nose in over 50 years so this was quite unusual.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1219450-1	pain, swelling, tightness and increased temperature in lower right leg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1219828-1	Patient received his first Moderna vaccine 3-19-21. He was due for his 2nd dose 4-16-21. We followed up with his sister due to missed appointment and was informed that he passed away 4-15-21 due to a blood clot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1220899-1	blood clots in both legs; severe pain in both his legs; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots in both legs) in a 67-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 008D21A and 011A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history was reported.). On 25-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 23-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 01-Apr-2021, the patient experienced THROMBOSIS (blood clots in both legs) (seriousness criterion medically significant) and PAIN IN EXTREMITY (severe pain in both his legs). At the time of the report, THROMBOSIS (blood clots in both legs) and PAIN IN EXTREMITY (severe pain in both his legs) outcome was unknown. Not Provided For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant product use was not provided. Treatment included Eliquis (apixaban) for three months. Reportedly, the patient was put on apixaban 2 tablets every morning and evening for 14 days and will then take 1 tablet every morning and evening for a total of 3 months. He does think that the medicine is helping. Patient stated that he never had blood clots before and suspects that it was from the shot.; Sender's Comments: Very limited information has been provided at this time. Further information has been requested. Based on the current information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1220922-1	Blood clot; Pain; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot) in a 30-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event. On 11-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion medically significant) and PAIN (Pain). At the time of the report, THROMBOSIS (Blood clot) and PAIN (Pain) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Company comment: Limited information regarding the events has been provided at this time and a causal relationship cannot be excluded
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1220961-1	"Thrombosis on left arm and chest; ""you can see all of the veins""; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Thrombosis on left arm and chest) in a 56-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 037B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Neuralgia. On 08-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 08-Apr-2021, the patient experienced THROMBOSIS (Thrombosis on left arm and chest) (seriousness criterion medically significant) and VASODILATATION (""you can see all of the veins""). At the time of the report, THROMBOSIS (Thrombosis on left arm and chest) and VASODILATATION (""you can see all of the veins"") had not resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Concomitant product include an unspecified neuralgia medication. No treatment information was reported. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1220980-1	Blood clot; Left leg pain; Discoloration (blue-purple); Swelling; This spontaneous case was reported by an other health care professional and describes the occurrence of THROMBOSIS (Blood clot), PAIN IN EXTREMITY (Left leg pain), SKIN DISCOLOURATION (Discoloration (blue-purple)) and SWELLING (Swelling) in a 78-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 011J20A and 013L20A) for COVID-19 vaccination. The patient's past medical history included Clot blood in 2010. Concurrent medical conditions included Drug allergy (Dalaudid, Ambien and Demerol) and Hypertension. Concomitant products included DAPAGLIFLOZIN PROPANEDIOL MONOHYDRATE (FARXIGA), GLIMEPIRIDE (AMARYL), ACETYLCARNITINE HYDROCHLORIDE (NEUROTIN [ACETYLCARNITINE HYDROCHLORIDE]), COLECALCIFEROL (CALTRATE VITAMIN D DAILY), MINERALS NOS, VITAMINS NOS (CENTRUM A TO ZINC) and VITAMIN D NOS for an unknown indication. On 11-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 09-Mar-2021, the patient experienced PAIN IN EXTREMITY (Left leg pain) (seriousness criterion hospitalization), SKIN DISCOLOURATION (Discoloration (blue-purple)) (seriousness criterion hospitalization) and SWELLING (Swelling) (seriousness criterion hospitalization). On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion hospitalization). At the time of the report, THROMBOSIS (Blood clot), PAIN IN EXTREMITY (Left leg pain), SKIN DISCOLOURATION (Discoloration (blue-purple)) and SWELLING (Swelling) outcome was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Patient was sent to hospital to remove clots. The patient stated that the clot was 2/3 size of thigh. The patient scheduled to have a second procedure in May to remove blockages from iliac. Action taken with mRNA-1273 was not applicable. Based on the current available information and temporal association between the use of the product and the onset date of the reported events, a causal relationship cannot be excluded. Prior medical history of blood clot is considered a significant risk factor.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the onset date of the reported events, a causal relationship cannot be excluded. Prior medical history of blood clot is considered a significant risk factor.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1220994-1	tested positive for COVID- 19; Trouble breathing; blood clots in lungs; Chills; fever; body aches; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clots in lungs) and COVID-19 (tested positive for COVID- 19) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 038A21A) for COVID-19 vaccination. The patient's past medical history included No adverse event (No medical history reported). On 16-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-Mar-2021, the patient experienced THROMBOSIS (blood clots in lungs) (seriousness criteria hospitalization prolonged and medically significant) and COVID-19 (tested positive for COVID- 19) (seriousness criterion hospitalization prolonged). At the time of the report, THROMBOSIS (blood clots in lungs) and COVID-19 (tested positive for COVID- 19) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No treatment information was provided. Based on the current available information which includes a strong temporal association between the use of the product and onset of the reported events, and excluding other etiologies, a causal relationship cannot be excluded. Fever, chills and myalgua are consistent with the product known safety profile. Reporter did not allow further contact; Sender's Comments: Based on the current available information which includes a strong temporal association between the use of the product and onset of the reported events, and excluding other etiologies, a causal relationship cannot be excluded. Fever, chills and myalgua are consistent with the product known safety profile.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221012-1	small intestine got blocked and due to blockage blood supply stopped; started having blood clot thrombosis both in his right leg as well as left leg; Missed the 2nd dose; Blocked small intestine; This spontaneous case was reported by a nurse and describes the occurrence of SMALL INTESTINAL OBSTRUCTION (Blocked small intestine), INTESTINAL ISCHAEMIA (small intestine got blocked and due to blockage blood supply stopped) and THROMBOSIS (started having blood clot thrombosis both in his right leg as well as left leg) in a 73-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 030M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse effect (No relevant medical history reported) and Surgery. Concomitant products included thyroid medication for an unknown indication, cholesterol medication. On 13-Feb-2021 at 10:30 AM, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 13-Feb-2021, the patient experienced SMALL INTESTINAL OBSTRUCTION (Blocked small intestine) (seriousness criterion medically significant). On 13-Mar-2021, the patient experienced INTESTINAL ISCHAEMIA (small intestine got blocked and due to blockage blood supply stopped) (seriousness criterion medically significant), THROMBOSIS (started having blood clot thrombosis both in his right leg as well as left leg) (seriousness criterion medically significant) and PRODUCT DOSE OMISSION ISSUE (Missed the 2nd dose). On 13-Mar-2021, PRODUCT DOSE OMISSION ISSUE (Missed the 2nd dose) had resolved. At the time of the report, SMALL INTESTINAL OBSTRUCTION (Blocked small intestine), INTESTINAL ISCHAEMIA (small intestine got blocked and due to blockage blood supply stopped) and THROMBOSIS (started having blood clot thrombosis both in his right leg as well as left leg) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant medications included thyroid medication and cholesterol medication. Patient called in to report that he received his 1st dose of Moderna vaccine on 13Feb2021 in left arm at 10.30 am and his 2nd dose of Moderna vaccine was scheduled on 13Mar2021. On 13Mar2021he went to Emergency Room (ER) and he had major surgery on small intestine, patient mentioned that his small intestine got blocked and due to blockage the blood supply was stopped and the intestine started to die. On that day, the patient underwent a surgical procedure which the doctor had to take out 12 inches of small intestine and stapled back. He was hospitalized for five days. Five to seven days after his surgery, he developed blood clot thrombosis in his right and left legs. Treatment for the event included blood thinner, which he has been on for 45 days. It was reported that the second dose of Moderna vaccine was missed due to hospitalization for the events and the blood thinner that he was on. Most recent FOLLOW-UP information incorporated above includes: On 13-Apr-2021: reporter added, product indication added, event added(missed dose), event assessment done, additional info added(Source Document and mail document added); Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221044-1	Blood clot in the right leg; Blood clot in the lungs; Blood clot in the hip; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot in the hip), THROMBOSIS (Blood clot in the right leg) and PULMONARY EMBOLISM (Blood clot in the lungs) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. Unknown and 015L20A) for COVID-19 vaccination. The patient's past medical history included No adverse event (No reported medical history). On 06-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 06-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. In February 2021, the patient experienced THROMBOSIS (Blood clot in the hip) (seriousness criteria hospitalization and medically significant). In March 2021, the patient experienced THROMBOSIS (Blood clot in the right leg) (seriousness criteria hospitalization and medically significant) and PULMONARY EMBOLISM (Blood clot in the lungs) (seriousness criteria hospitalization and medically significant). The patient was hospitalized on 16-Mar-2021 due to PULMONARY EMBOLISM, THROMBOSIS, THROMBOSIS At the time of the report, THROMBOSIS (Blood clot in the hip), THROMBOSIS (Blood clot in the right leg) and PULMONARY EMBOLISM (Blood clot in the lungs) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Patient states that 2-3 weeks after receiving the second shot they had a blood clot in the hip and a week later they had a blood clot in the right leg and in the lungs. Patient reports they had to be hospitalized for two nights, in the Medical City Fort Worth Hospital. Patient states they do not have any history of blood clots. Patient mentioned reading online, in forums, that people had taken Pfizer and Moderna vaccines and two weeks after had blood clots and ended in the hospital. Based on the current available information and temporal association between the use of the product and the start dates of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start dates of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221061-1	DBD blood clot in leg; Arm got little sore; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (DBD blood clot in leg) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history was reported.). Concomitant products included RIVAROXABAN (XARELTO) for an unknown indication. On 14-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 14-Mar-2021, the patient experienced PAIN IN EXTREMITY (Arm got little sore). On 05-Apr-2021, the patient experienced THROMBOSIS (DBD blood clot in leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (DBD blood clot in leg) outcome was unknown and PAIN IN EXTREMITY (Arm got little sore) had resolved. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Reportedly, the patient's sore arm went away pretty fast. The patient received treatment with Xarelto (Rivaroxaban) for blood clot in his leg. Company Comment Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Reporter did not allow further contact
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221095-1	possible stroke; small clot that went through and resolved on its own; Loss use of left leg/couldn't use her leg; tingling in left arm; This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (possible stroke), THROMBOSIS (small clot that went through and resolved on its own), MONOPLEGIA (Loss use of left leg/couldn't use her leg) and PARAESTHESIA (tingling in left arm) in a 55-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 046A21A) for COVID-19 vaccination. The patient's past medical history included Familial hypercholesterolemia, Peripheral arterial disease and Stent placement (has stents in her iliac ery). Concomitant products included ATORVASTATIN CALCIUM (ATORVASTATIN [ATORVASTATIN CALCIUM]), ACETYLSALICYLIC ACID (BABY ASPIRIN) and SERTRALINE for an unknown indication. On 26-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 27-Mar-2021, the patient experienced CEREBROVASCULAR ACCIDENT (possible stroke) (seriousness criterion hospitalization), THROMBOSIS (small clot that went through and resolved on its own) (seriousness criterion hospitalization), MONOPLEGIA (Loss use of left leg/couldn't use her leg) (seriousness criterion hospitalization) and PARAESTHESIA (tingling in left arm) (seriousness criterion hospitalization). The patient was hospitalized from 27-Mar-2021 to 28-Mar-2021 due to CEREBROVASCULAR ACCIDENT, MONOPLEGIA, PARAESTHESIA and THROMBOSIS. At the time of the report, CEREBROVASCULAR ACCIDENT (possible stroke), THROMBOSIS (small clot that went through and resolved on its own), MONOPLEGIA (Loss use of left leg/couldn't use her leg) and PARAESTHESIA (tingling in left arm) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In March 2021, Computerised tomogram: Inconclusive. In March 2021, Magnetic resonance imaging: Inconclusive. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No treatment information was provided. Patient reports an MRI and CT scan was done, and she was informed that they suspected she had a small cot that went through and resolved on its own. Company Comment: Based on the temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However patient's hx of familial hypercholesterolemia, peripheral arterial disease and stent placement are confounding factors that may play a possible contributory role.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221096-1	Pt started having headaches on 3/20/21. He was playing golf with a cardiologist and he recommended that he see his doctor.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221214-1	I woke up in the middle of night with chest pain and trouble breathing. I had a pulmonary embolism. The pain and difficulty breathing lasted several days. Since then, I have experience a flare of the antiphospholipid syndrome - low platelets (blood work), splinter hemorrhages, clots, and the like. Whether from the embolism or other, the pulmonary hypertension has gotten drastically worse - with increased shortness of breath, irregular heart beats, dropping pulse ox.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1221428-1	Ended up in healthcare facility on 3-1 dehydrated and released Back to healthcare facility on 3-4 stayed until 3-7 Started hallucinating and given meds. very dehydrated Released and then family came to pick her up. Hallucinating on drive back to where she would be staying. 3-8 back to healthcare facility because she was slurring words ---gave fluids and released 3-9 Unit had to come to get her very weak and fragile. Couldn't move her ourselves. Ended up admitting her until 3-12 At house she was supervised the whole time, weak, needed assistance, slurring words off and on. 3-22 Eye doc appt found blood clot behind eye. Sent her to healthcare facility to be examined by Neugolist. Did CT Scan and MRV/MRI found. They found clots and ended up sending her to healthcare facility. Was in city where they run tons of tests. mini strokes lead to the Thrombosis which caused the blood clots. Released on 4-2 into the care of a home where she started to hallucinate even more. Back to healthcare facility on 4-9 still currently there where they are treating her hallucinations, memory loss and more.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1222394-1	Bilateral pulmonary embolism, clot in leg, pneumonia developed after the vaccine. Patient was hospitalized for 3 nights.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1222969-1	Blood Clot that resulted in a stroke 25 hours after the shot. My heart went into A-Fib (first time ever)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223056-1	My wife started to fall and pass out had no strength to get up this happened two to three times cinch she got the shot the last time she past out in the shower and i rushed her to hospital were she pasted away from blood clots to the right side of the neck and stated bleed on the brain
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223220-1	On 4/6/2021 woke up with severe left muscle cramp in left leg. Next 3 to 5 days tried massage, ice, compression and Motrin - no relief. Did no treatment for the next couple of days. Went to primary care doctor and then got an ultra-sound on 4/14/2021 at clinic - by Doctor. Results showed had three blood clots in left leg. Started on Eliquis that day.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223222-1	A few days after vaccination I had a high fever and began coughing so hard that I was throwing up. This lasted for about a week, I then became short of breath and tacchychardic. I went to the ER and was admitted for 5 days where they said I had acute bronchitis that triggered reactive asthma. I?m now on steroids and a bunch of medication. This all followed my first dose of Moderna as I?ve never had breathing issues before hand. I?ve never had bronchitis or asthma in my life so this is all new. I am now experiencing nerve pain on my left side which is where I received the vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223282-1	Noticed painful right calf 3/09/2021. Went to urgent care at hospital on 3/10/2021. Was given an ultrasound and it was determined that I had a blood clot in my calf vein. Was given heparin treatment for seven days followed by xarelto. I have had 2 dvts in the past associated with surgery. This dvt was listed as unprovoked, however I believe it was caused by the second shot of modern. Was told by doctors to report this to veers.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223364-1	"I received vaccine doses on 2/16/21 and second on 3/16/21. Trying to complete good deeds for Lent, I donated Plasma for the first and only time on 3/7/21. On 3/8/21 my right calf starting seizing up and I was experiencing calf pain. I was advised to do stretches and use a heating pad. It persisted for over a week and I went for a Doppler ultrasound scan on 3/17/21. Blood clots were found. The Plasma donation center screening led me to believe that they were concerned with other vaccines but not the Covid19 vaccine. Writing report because I think there is a possible connection to the blood clots, vaccine and plasma donation. I had to ""do nothing"" for three weeks and have been placed on Eliquis, a blood thinner for four months. Follow-up with Dr. for blood work and Doppler in July."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223833-1	Bilateral pulmonary embolism, Left Brachial artery clot, abdominal clots
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1223961-1	Probably unrelated but I am on day 3 of my normal menstrual cycle (typically 5-7 days). Approximately 2 hours after receiving the vaccination I began passing large clots and my flow went from regular to heavy. I do have heavy cycles from time to time but found the abrupt change in cycle interesting. Other than being sore in the arm and tired I feel fine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1225782-1	blood clotting in brain
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1226028-1	Acute Saddle Pulmonary embolism, Hospital for 8 days, put on blood thinners (Xarelto) I had severe chest pain and shortness of breath.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1226073-1	Swollen left leg, shortness of breath, painful upper left thigh diagnosed with blood clots
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1226244-1	"Roughly 7 to 10 days after injection I presented with raised, red and itchy injection site. Roughly 13 days after the injection I presented with a tight right calf muscle that felt similar to a constant ""charlie horse"". It continued to worsen over the next 5 days until my right foot went numb and cold on Sunday April 4. I called my doctor the next day (Mon April 5) and they got me in that day and I went to the hospital the same day for an ultrasound. The hospital confirmed I had a blood clot in my lower leg."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1226448-1	Blood clot leading to heart attack
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1226954-1	Blood clot in left arm on 2/28/2021 (2 weeks after 2nd dose of Moderna). Diagnosed at Emergency Room.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1227369-1	2 lower- left leg Superficial blood clots; ~ a DVT Deep Vein Thrombosis per Ultrasound on 4/12/21, was diagnosis ; ER Dr. on 4/12/21 prescribed Eliquis 60 day Treatment.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1228468-1	He got his vaccine, and on Easter he thought that he had a cramp in his leg, and asked his wife and told her that he had this. By that afternoon he still had the cramping feeling, and started drinking water as his wife is a nurse and felt that he may be dehydrated. Throughout the week he had a Charlie Horse pain in his left calf, and felt that it was getting worse and not going away. The whole week he went to work and on the 7th day it started swelling and went to Hospital after he got off of work, and he saw a resident who said that his leg was swollen and told him that she was going to schedule an ultrasound. He had that on Friday on 4/16/21, and was scheduled for 4:00 but due to the pain went to the ER and gave him IV and checked his blood work. The ultrasound showed that he did have a blood clot and prescribed Eliquis and gave him a dose while in the ER. He told him that he would be on 2 a dat for 7 days and then 1 a day, but did not tell him how long. He is supposed to see the resident this afternoon for recheck. He is still having pain and wanting to know where he is going to do from this point. He does work out every day at his home gym and has not smoked, and now wanting to know how this is going to affect his life from this point into the future. He also jumps rope 3 times a day and now not able to do this.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1228483-1	He got his vaccine, and on Sunday he thought that he had a cramp in his leg, and asked his wife and told her that he had this. By that afternoon he still had the cramping feeling, and started drinking water as his wife is a nurse and felt that he may be dehydrated. Throughout the week he had a Charlie Horse pain in his left calf, and felt that it was getting worse and not going away. The whole week he went to work and on the 7th day it started swelling and went to Hospital after he got off of work, and he saw a resident Dr. who said that his leg was swollen and told him that she was going to schedule an ultrasound. He had that on Friday on 4/16/21, and was scheduled for 4:00 but due to the pain went to the ER and gave him IV and checked his blood work. The ultrasound showed that he did have a blood clot and prescribed Eliquis and gave him a dose while in the ER. He told him that he would be on that for a while. He is supposed to see the resident this afternoon for recheck. He is still having pain and wanting to know where he is going to do from this point. He does work out every day at his home gym and has not smoked, and now wanting to know how this is going to affect his life from this point into the future. He also jumps rope 3 times a day and now not able to do this.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1229402-1	Patient was hospitalized due to multiple blood clots, including jugular, brachial, and others. Patient was hospitalized for treatment on 4/10/21 and is still there.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1229596-1	Blood clot in left arm caused a stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1229975-1	"Reported by patients wife. She Reports blood clot in left leg ""from thigh to calf"". Treated with blood thinners since beginning of March 2021. Has taken Enoxaparin Sodium 100mg bid since onset and recently started on Xarelto by hematologist 04/16/2021."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1230555-1	Vaccination was on Wednesday, March 17, 2021 at 3 PM. Symptoms began 8 days later, and Emergency Room admission 3days later on Sunday, March 28, 2021, for Blood Clots in legs and lungs,(critical in Pulmonary Artery). Blood thinner started immediately was Zarelto, and continues daily. Pain, nausea, shortness of breath, and fatigue were the symptoms. The patient was admitted to hospital for 5 days and sent home to recuperate. 3 days later, on Sunday she was back into the emergency department with intestinal bleeding that had gone on for 2 days. She was admitted again and had a colonoscopy. Bleeding stopped and she went home after 3 more days. She is currently getting home health care.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1230841-1	Blood clot, stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1230927-1	Extremely heavy period bleeding started on February 20th out of no where. Bleeding contained big clots and was very heavy. Went to my GYN on March 15th who prescribed a five day supply of Tranexamic acid. On March 20th, my bleeding finally stopped.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1230972-1	Blood Clot in the Left leg; Pain in the Left Leg; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood Clot in the Left leg) in a 74-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 007B21A and 007B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Blood pressure high. Concomitant products included LISINOPRIL and HCTZ for Blood pressure high. On 27-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 27-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 2 dosage form. On 13-Apr-2021, after starting mRNA-1273 (Moderna COVID-19 Vaccine), the patient experienced PAIN IN EXTREMITY (Pain in the Left Leg). On 14-Apr-2021, the patient experienced THROMBOSIS (Blood Clot in the Left leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood Clot in the Left leg) and PAIN IN EXTREMITY (Pain in the Left Leg) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 14-Apr-2021, Ultrasound joint: ultrasound (abnormal) Blood Clot in the Left leg. Patient did not have a history of Blood Clots. She was Not taking any hormones (Estrogen, Progesterone)Treatment for event included Eliquis for blood clot. The patient received both scheduled doses of mRNA-1273 prior to the events; therefore, action taken with the drug in response to the events is not applicable. Based on the current available information and temporal association between the use of the product and the onset date of the reported event a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the onset date of the reported event a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1231122-1	Achy left ear, paralyzed left side of face, left side of mouth paralyzed, left eye would not close or blink. This started the day after the vaccine injection. The facial paralysis lasted 5 weeks. The leg pain started 3 weeks after injection.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1231366-1	Migraine High blood pressure Bruises Blood clot Dizziness Nauseous Fatigue Itching Chills Muscle aches
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1232427-1	She got her vaccine, initially she had bruising on her forearm that was blue/yellow with a swollen vein connected running down to her hand. She had tingling and numbness in her fingers in a select few on the left hand. She also had drooping of her left eye for a short period and a sensation on her left leg, swelling and pain. She went to the urgent care on 4/15/21, they told her to go home and monitor it. She knew that somewhat was not right and called a nurse at Hospital and went to the ER shortly thereafter and confirmed that she had a blood clot in her jugular vein IJ-DVT. In the ER they gave her a shot of Lovenox in the ER into her stomach and run several tests. They also gave her an anti-inflammatory in her right arm. She was sent home with Eliquis for the next 3 months. She has extreme exhaustion, and in addition to that she has had a lot of swelling on the left side of her leg, and visited the ER again on 4/17/21. Her left foot became blue, very swollen, so they ran a number of tests which all came back negative. She thinks these are side effects in relation to the previous. She is still continuing with stiffness in her neck as well. She also has a severe headache, fever of about 100, and severe achiness. She has not been able to work due to these symptoms since 4/15/21. The ER told her to have a lot of rest, no heavy listing/straining due to the clot.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1232500-1	Blood clot causing swelling, pain, redness in left leg from groin to ankle--sent for ultrasound from Vascular surgeon's office at.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1232513-1	Spoke with patient via telephone on 4/20/21. Patient reports brief syncopal episode 4/15/21 at 6pm; reports he went to urgent care then to hospital. Reports he had CT scan and that multiple blood clots and pulmonary embolism were diagnosed. Reports he was transferred via ambulance early am on 4/16/21 to more advanced hospital. States thrombectomy was performed and he was discharged home on 4/17/21. States he is recovering at home and was advised he still has a blood clot in the leg. Reports he is on blood thinning medication at this time.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1232686-1	Spotting (light bleeding) for 3 days turned into shedding huge blood clots vaginally from 4/3 - 4/5. I bled though pads in just 5-10 minutes. Clots ranged from the size of a quarter to the size of the palm of my hand. They were very large, very frequent, and very painful. I never experienced this before in my life. The bleeding lasted 12 days total.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1232919-1	Blood clot in lower leg. After ultrasound, Doctor gave treatment protocol. Still undergoing treatment course.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1233715-1	Event: Blood clot in right leg (superficial thrombosis) Treatment: Rest, elevation, heat and ibuprofen Outcomes: TBD

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1234466-1	Patient came to pharmacy today w/ his wife to pick up Rx for XARELTO. He has developed multiple DVT's in his leg. He reported symptoms started the day after (Saturday) his second dose of Moderna COVID 19 vaccine on Friday 4.9.21. His leg was swollen by Sunday 4.11.21. Symptoms would get worse and then get better. He finally went to see Dr. today 04.20.21. They found several blood clots in the leg via doppler. Patient was therefore prescribed Xarelto. Pt wife advised that Dr said clots were due to Moderna vaccine and they should let pharmacy know also. Their next appointment will be in 2 weeks.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1234509-1	On 3/4/21 noticed my left calf was swollen. I did not think much of it at first. A week later the swelling had progressed to include my ankle. Saw Dr. and ordered an ultrasound on both of my legs. Right leg was normal & left lower leg had a clot(s) in it. Dr. ran blood work then prescribed Eliquis. Saw him in follow-up on April 14. Swelling down but not gone. Renewed the prescription for Eliquis and set a followup visit for Friday June 18. He ordered additional blood work to be drawn on Wednesday June 16 to have results at the next visit.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1234908-1	Severe headache, has continued intermittently until present. Strange menstrual flow, was very heavy, with clotting, unusual per patient
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235517-1	blood clot in his pulmonary artery and several others in his legs; blood clot pulmonary artery; shortness of breath; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot in his pulmonary artery and several others in his legs) and PULMONARY EMBOLISM (blood clot pulmonary artery) in an 82-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Family history included Factor V deficiency, Factor II deficiency and MTHFR deficiency. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (blood clot in his pulmonary artery and several others in his legs) (seriousness criteria hospitalization prolonged and medically significant), PULMONARY EMBOLISM (blood clot pulmonary artery) (seriousness criterion medically significant) and DYSPNOEA (shortness of breath). At the time of the report, THROMBOSIS (blood clot in his pulmonary artery and several others in his legs), PULMONARY EMBOLISM (blood clot pulmonary artery) and DYSPNOEA (shortness of breath) outcome was unknown. Concomitant product use was not provided by the reporter. Treatment information was not provided. Action taken with mRNA-1273 in response to the event was not applicable. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235625-1	Consistent headache - Tylenol or Excedrine relieved Heavy period - began 4 days early. Excessive blood clots and amount of blood. Left arm pain- unable to move arm without pain x 3 days. Tylenol relieved Fever- began 2 days after. Tylenol reduces fever.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235672-1	blood clots in his right leg and left leg; Emergency Room at 4am because he had trouble breathing and walking; found blood on both his lungs; chills especially at night that lasted 2 nights; He states he is still in misery; Patient was supposed to get second dose on 04-Mar-2021, but he is not going to get it; shoulder pain/hip pain; Pain in arms, leg; trouble walking due to his pain; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots in his right leg and left leg), DYSPNOEA (Emergency Room at 4am because he had trouble breathing and walking), HAEMOPTYSIS (found blood on both his lungs) and CHILLS (chills especially at night that lasted 2 nights) in an 88-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 004M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Asthma since an unknown date. Concomitant products included MONTELUKAST and FLUTICASONE FUROATE, VILANTEROL TRIFENATATE (BREO ELLIPTA) for Asthma, METOPROLOL and APIXABAN (ELIQUIS) for an unknown indication. On 04-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 05-Feb-2021, the patient experienced THROMBOSIS (blood clots in his right leg and left leg) (seriousness criteria hospitalization and life threatening), DYSPNOEA (Emergency Room at 4am because he had trouble breathing and walking) (seriousness criterion hospitalization), HAEMOPTYSIS (found blood on both his lungs) (seriousness criteria hospitalization and medically significant), PAIN IN EXTREMITY (Pain in arms, leg), GAIT DISTURBANCE (trouble walking due to his pain) and ARTHRALGIA (shoulder pain/hip pain). On 04-Mar-2021, the patient experienced INTENTIONAL DOSE OMISSION (Patient was supposed to get second dose on 04-Mar-2021, but he is not going to get it). On an unknown date, the patient experienced CHILLS (chills especially at night that lasted 2 nights) (seriousness criterion hospitalization) and FEELING ABNORMAL (He states he is still in misery). The patient was hospitalized on 20-Feb-2021 due to DYSPNOEA, HAEMOPTYSIS and THROMBOSIS, and then for 2 days due to CHILLS. At the time of the report, THROMBOSIS (blood clots in his right leg and left leg), DYSPNOEA (Emergency Room at 4am because he had trouble breathing and walking), HAEMOPTYSIS (found blood on both his lungs), PAIN IN EXTREMITY (Pain in arms, leg), GAIT DISTURBANCE (trouble walking due to his pain), FEELING ABNORMAL (He states he is still in misery) and ARTHRALGIA (shoulder pain/hip pain) had not resolved and CHILLS (chills especially at night that lasted 2 nights) and INTENTIONAL DOSE OMISSION (Patient was supposed to get second dose on 04-Mar-2021, but he is not going to get it) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Chest X-ray: (abnormal) Found blood on both of his lungs. On an unknown date, SARS-CoV-2 test: (Negative) Negative. On an unknown date, Ultrasound scan: (abnormal) Found blood clots in his right leg and left leg. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Patient was taken to the ER (Emergency Room) on 20-Feb-2021 at 4 AM. Several tests including Chest X-Ray, MRI, Blood work, Ultrasound (groin and legs) were performed. Only few tests results were reported. Received treatment including Oxygen, Eliquis and Antibiotics. Patient has been on oxygen full time until 7Apr2021 Company comment: Based on the current available information and temporal association between the use of the product and the start date of these event, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of these event, a causal relationship cannot be excluded. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235676-1	Blood Clotting; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood Clotting) in a 59-year-old female patient who received mRNA-1273 for COVID-19 vaccination. The patient's past medical history included No adverse event. On 26-Feb-2021, the patient received first dose of mRNA-1273) (Intramuscular) 1 dosage form. On 05-Mar-2021, the patient experienced THROMBOSIS (Blood Clotting) (seriousness criteria medically significant). At the time of the report, THROMBOSIS (Blood Clotting) had resolved. Not Provided. The action taken with mRNA-1273 (Intramuscular) was unknown. Treatment information was not provided. Concomitant medications were not reported. Company Comment: Very limited information regarding the events has been provided at this time and is insufficient for causality assessment. Further information has been requested.; Sender's Comments: Very limited information regarding the events has been provided at this time and is insufficient for causality assessment. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235687-1	Memory lapse, difficulty with memory recall; Confused; Blood clots; Cerebral Ischemic Attack; Brain Fog; Headache; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of MEMORY IMPAIRMENT (Memory lapse, difficulty with memory recall), CONFUSIONAL STATE (Confused), THROMBOSIS (Blood clots) and TRANSIENT ISCHAEMIC ATTACK (Cerebral Ischemic Attack) in a 49-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 046A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Heart valve operation. Concomitant products included WARFARIN SODIUM (COUMADIN) for an unknown indication. On 12-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 09-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 10-Apr-2021, the patient experienced MEMORY IMPAIRMENT (Memory lapse, difficulty with memory recall) (seriousness criterion hospitalization), CONFUSIONAL STATE (Confused) (seriousness criterion hospitalization), THROMBOSIS (Blood clots) (seriousness criterion hospitalization) and TRANSIENT ISCHAEMIC ATTACK (Cerebral Ischemic Attack) (seriousness criterion hospitalization). On 15-Apr-2021, the patient experienced FEELING ABNORMAL (Brain Fog) and HEADACHE (Headache). The patient was hospitalized from 10-Apr-2021 to 12-Apr-2021 due to CONFUSIONAL STATE, MEMORY IMPAIRMENT, THROMBOSIS and TRANSIENT ISCHAEMIC ATTACK. At the time of the report, MEMORY IMPAIRMENT (Memory lapse, difficulty with memory recall), FEELING ABNORMAL (Brain Fog) and HEADACHE (Headache) had not resolved and CONFUSIONAL STATE (Confused), THROMBOSIS (Blood clots) and TRANSIENT ISCHAEMIC ATTACK (Cerebral Ischemic Attack) had resolved. Not Provided DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 10-Apr-2021, Computerised tomogram: abnormal (abnormal) revealed blood clots. On 10-Apr-2021, International normalised ratio: low (Low) Low. On 10-Apr-2021, Magnetic resonance imaging: abnormal (abnormal) revealed blood clots. On 15-Apr-2021, International normalised ratio: low (Low) Low. It was reported that the patient still has difficulty with memory recall, brain fog, headache, and still having trouble with low INR. The patient did not have any side effects after the first dose. Treatment for the events included Levonox injections and Coumadin (increased dose). Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, excluding other etiologies causal relationship cannot be excluded. Headache is consistent with the product known safety profile.; Sender's Comments: Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, excluding other etiologies causal relationship cannot be excluded. Headache is consistent with the product known safety profile.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235690-1	Blood clots; Kind of bleeding; arm began being extremely itchy; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) and HAEMORRHAGE (Kind of bleeding) in a 40-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No reported medical history). On 02-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 02-Apr-2021, after starting mRNA-1273 (Moderna COVID-19 Vaccine), the patient experienced PRURITUS (arm began being extremely itchy). On 05-Apr-2021, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant) and HAEMORRHAGE (Kind of bleeding) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) and HAEMORRHAGE (Kind of bleeding) outcome was unknown and PRURITUS (arm began being extremely itchy) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In April 2021, Pregnancy test: Negative. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not provided by the reporter. Treatment information was not provided. Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, excluding other etiologies causal relationship cannot be excluded; Sender's Comments: Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, excluding other etiologies causal relationship cannot be excluded
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1236676-1	Developed a blood clot in his right leg, inner thigh, up toward groin area. ER diagnosis on 4/20/21 at healthcare facility.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1236718-1	patient reports he spent many days in hospital after 2nd Moderna vaccine wtih severe blood clots. Informed he did not tell hospital that he had the 2nd Moderna shot but informed the other hospital. It was the other hospital that instructed the patient to contact the Health Department to inform of the same
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1236733-1	PER PATIENT REPORT, PAIN A WEEK LATER MY LEG WAS ACHING. I USED ICY HOT, MASSAGER. STARTED SWELLING, WENT TO WALK IN. THEY DID AN ULTRASOUND AND SAID I HAD A BLOOD CLOT IN MY LEG AND IT WAS TRAVELING TO MY LUNGS. THERE WERE A COUPLE SPECKS ON LUNG PER XRAY.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1237101-1	Like the reaction to the first dose, pain and swelling at injection site, swollen lymph node under armpit, fever of 101 F, headache , severe joint pain, fatigue and unusual and heavy menstrual bleeding starting two days after the injection. Bleeding so heavily medical care was sought after first injection. Thought it might be my IUD, which was removed after the first episode. Full gynecological exam indicated everything was normal. Menstrual bleeding happened again 2 days after second vaccine, characterized by early arrival, heavy flow and clotting.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1237551-1	Blood clot in Right leg and bilateral lungs. Patient also reported headache for >1 week, fatigue, shortness of breath and blood pressure that elevated above 210 systolic.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1237904-1	APRIL 9 EVENING LUMP IN RIGHT LEG, PAIN TENDERNESS AND REDNESS, SITE IS HOT AS WELL. SWELLED OUT, RUNS 2 INCHES, RIGHT ON INNER PART OF LEG, BELOW KNEE LEVEL, HAS HAD 2 NEW KNEE IMPLANTS ON RIGHT LEG.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1238013-1	Blood clot in left arm. Heavy menstrual clots bigger than a quarter
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1238361-1	Blood clots
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1238542-1	Left calf pain. US completed. Blood clot in left great saphenous vein and branches. Patient is being treated with Lovenox 60 mg BID. Outcome is unknown at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1240236-1	Blood clot in mouth on day 2, 9, and 16 from first injection
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1242013-1	8 Days after my 2nd Moderna Shot, Fatty Blood Clots started pouring out of my mouth(about 50 over a 24 hour period) . There was No cause or affect to it- meaning No cough or Nose Bleeding,etc. I went to emerency room on 03/09/2021 and was admitted for testing. While there they ALSO noticed my heart was Pausing. A 3 day ZIO Test was done. It came back extreme with 39 Pauses and a pulse as low as 35 BPM. A Heart Pace Maker had to be placed in me at Medical CTR on 04/20/2021.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1242396-1	Day after the shot on Tuesday morning woke up a little fatigue she did a little cooking and her chest started really getting tight she was sweating really bad went to bed slept all day tried walking on Wednesday put a heating pad on her legs with muscle spasms called me at 5 AM to get her to the ER she couldn?t walk her leg swelled up really bad
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1242501-1	Blood Clot in left leg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1243509-1	Reports she was exhausted and weak for a few days after the shot. on 4/20 she went to the ER to receive treatment for a blood clot in her right leg. Reports she had been having trouble with her leg for 5 days before going to the ER. She has started taking Xarelto. She went for treatment. She was not admitted.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1244061-1	Hospitalized for acute blood clot in each leg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1244283-1	Body aches, fever, high blood sugars, diarrhea & extreme nausea-could not eat or drink for 2 days No clotting during menstrual cycle, EXTREMELY heavy and long flow

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245371-1	irritated her heart condition; Clot blood; weakness in legs, couldnt walk with legs; Coma; couldn't breathe; stroke/massive stroke in left side of her brain; Irritated her heart condition, rapid heart beat; Headache; Tiredness; This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (stroke/massive stroke in left side of her brain), CARDIAC DISORDER (irritated her heart condition), DYSPNOEA (couldn't breathe), THROMBOSIS (Clot blood), COMA (Coma) and MUSCULAR WEAKNESS (weakness in legs, couldnt walk with legs) in a 95-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for Covid-19 Vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Heart disease, unspecified, Hypertension and AFib. Concomitant products included APIXABAN (ELIQUIS) for Anticoagulant therapy, SACUBITRIL VALSARTAN SODIUM HYDRATE (ENTRESTO) for Hypertension, METOPROLOL for an unknown indication. On 10-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 11-Mar-2021, the patient experienced HEADACHE (Headache) and FATIGUE (Tiredness). On 12-Mar-2021, the patient experienced MUSCULAR WEAKNESS (weakness in legs, couldnt walk with legs) (seriousness criterion hospitalization). On 13-Mar-2021, the patient experienced CARDIAC DISORDER (irritated her heart condition) (seriousness criterion hospitalization), THROMBOSIS (Clot blood) (seriousness criterion hospitalization) and HEART RATE INCREASED (Irritated her heart condition, rapid heart beat). On 15-Mar-2021, the patient experienced CEREBROVASCULAR ACCIDENT (stroke/massive stroke in left side of her brain) (seriousness criteria death and medically significant). On 16-Mar-2021, the patient experienced DYSPNOEA (couldn't breathe) (seriousness criterion hospitalization prolonged) and COMA (Coma) (seriousness criterion hospitalization prolonged). The patient was hospitalized on 13-Mar-2021 due to CARDIAC DISORDER, COMA, DYSPNOEA, MUSCULAR WEAKNESS and THROMBOSIS. The patient died on 17-Mar-2021. The reported cause of death was massive stroke in left side of her brain. It is unknown if an autopsy was performed. At the time of death, CARDIAC DISORDER (irritated her heart condition), DYSPNOEA (couldn't breathe), THROMBOSIS (Clot blood), COMA (Coma), MUSCULAR WEAKNESS (weakness in legs, couldnt walk with legs), HEART RATE INCREASED (Irritated her heart condition, rapid heart beat), HEADACHE (Headache) and FATIGUE (Tiredness) outcome was unknown. Action taken with mRNA-1273 in response to the events was not Applicable. This case concerns an 95 year old female patient, with medical history of A Fib, Heart disease, Hypertension who experienced a serious unexpected event of Death 8 days after receiving 1st dose of mRNA- 1273 . Very limited information regarding these events has been provided at this time. However, the patient's advance age, multiple co-morbidities, may remain as risk factors. Further information is requested. This case was linked to MOD-2021-074814 (Patient Link).; Sender's Comments: This case concerns an 95 year old female patient, with medical history of A Fib, Heart disease, Hypertension who experienced a serious unexpected event of Death 8 days after receiving 1st dose of mRNA- 1273 . Very limited information regarding these events has been provided at this time. However, the patient's advance age, multiple co-morbidities, may remain as risk factors. Further information is requested.; Reported Cause(s) of Death: massive stroke in left side of her brain
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245407-1	a blood clot right below his lung on the left side; his chest was hurting; he couldn't breath; Second shot three weeks later; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (a blood clot right below his lung on the left side) in a 32-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history reported). In February 2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (a blood clot right below his lung on the left side) (seriousness criterion medically significant), CHEST PAIN (his chest was hurting), DYSPNOEA (he couldn't breath) and INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (Second shot three weeks later). At the time of the report, THROMBOSIS (a blood clot right below his lung on the left side), CHEST PAIN (his chest was hurting) and DYSPNOEA (he couldn't breath) outcome was unknown and INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (Second shot three weeks later) had resolved. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter considered THROMBOSIS (a blood clot right below his lung on the left side), CHEST PAIN (his chest was hurting) and DYSPNOEA (he couldn't breath) to be related. No further causality assessment was provided for INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (Second shot three weeks later). Patient was taken to the ER (Emergency Room). Treatment Xarelto was started. Patient has Cardiologist appointment on 28-APR-2021. Company comment:Limited information regarding the blood clot, chest pain and difficulty breathing has been provided at this time and a causal relationship cannot be excluded. Inappropriate schedule of vaccine administered is unrelated to the vaccine and is not reported as specifically resulting in adverse events. This case was linked to MOD21-076660 (E2B Linked Report).; Sender's Comments: Limited information regarding the blood clot, chest pain and difficulty breathing has been provided at this time and a causal relationship cannot be excluded. Inappropriate schedule of vaccine administered is unrelated to the vaccine and is not reported as specifically resulting in adverse events. MOD21-076660:

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1245413-1	Blood clot in left leg and moved down to bottom of her heal; Hard Time Walking due to blood clot; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clot in left leg and moved down to bottom of her heal) in a 59-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history was provided.). Concomitant products included ACETYLSALICYLIC ACID (ASPIRIN 81) for an unknown indication. On 26-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 26-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 02-Apr-2021, the patient experienced THROMBOSIS (Blood clot in left leg and moved down to bottom of her heal) (seriousness criterion medically significant) and GAIT DISTURBANCE (Hard Time Walking due to blood clot). At the time of the report, THROMBOSIS (Blood clot in left leg and moved down to bottom of her heal) and GAIT DISTURBANCE (Hard Time Walking due to blood clot) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Patient reported that doctor prescribed prednisone to treat the symptoms. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1245430-1	"Veins was outside and sore; really big and sore red area; really big and sore red area; all the veins in his R leg was protuberate; two blood clot; DGThrombosis; Felt terrible; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (two blood clot) and THROMBOSIS (DGThrombosis) in a 77-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 016M20A and Unknown) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Leg injury (severe damage to right leg, hit by ""protellier"" small airplane.) in 1977, Surgery (""many surgeries"") and Clot blood. On 11-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 12-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 12-Mar-2021, the patient experienced MALAISE (Felt terrible). On 13-Mar-2021, the patient experienced THROMBOSIS (two blood clot) (seriousness criterion medically significant), THROMBOSIS (DGThrombosis) (seriousness criterion medically significant), VASCULAR PAIN (Veins was outside and sore), ERYTHEMA (really big and sore red area), PAIN IN EXTREMITY (really big and sore red area) and VEIN DISORDER (all the veins in his R leg was protuberate). At the time of the report, THROMBOSIS (two blood clot), THROMBOSIS (DGThrombosis), MALAISE (Felt terrible), VASCULAR PAIN (Veins was outside and sore), ERYTHEMA (really big and sore red area), PAIN IN EXTREMITY (really big and sore red area) and VEIN DISORDER (all the veins in his R leg was protuberate) outcome was unknown. Not Provided Concomitant medication included unspecified statin for high cholesterol. The patient reported that he is retired. He reported that his leg is functioning well, he has only one of the three arteries in the legs and many damaged veins. Treatment medication included Seroto to treat the blood clots. Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, a causal relationship cannot be excluded. This case was linked to MODERNATX, INC.-MOD-2021-077978 (E2B Linked Report).; Sender's Comments: Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, a causal relationship cannot be excluded. MODERNATX, INC.-MOD-2021-077978:"

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1245440-1	"blood clots; his arm felt sore, almost like he had just received the vaccine; doctor thinks it has something to do with the heart; low blood pressure; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clots) in a 64-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No reported medical history). On 24-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 08-Apr-2021, the patient experienced THROMBOSIS (blood clots) (seriousness criterion hospitalization), PAIN IN EXTREMITY (his arm felt sore, almost like he had just received the vaccine), CARDIAC DISORDER (doctor thinks it has something to do with the heart) and HYPOTENSION (low blood pressure). The patient was hospitalized on 08-Apr-2021 due to THROMBOSIS. At the time of the report, THROMBOSIS (blood clots), PAIN IN EXTREMITY (his arm felt sore, almost like he had just received the vaccine) and CARDIAC DISORDER (doctor thinks it has something to do with the heart) had not resolved and HYPOTENSION (low blood pressure) had resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not provided. On 08 Apr 2021, the patient's arm felt sore, ""almost like he had just received the vaccine"". He went to urgent care, and they sent him to the hospital and he was admitted. The patient said his doctor thinks the blood clots had something to do with the heart. The patient thought it had to do with the vaccine because he has ""always been healthy"" and never had any heart problems. The patient was still in the hospital at the time of this report. Treatment information was not provided. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Doctor thinks ""thrombosis"" has something to do with the heart"
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1245454-1	Small amount of blood clots; Potassium levels dropped; Increased hypertension; dizziness; Arm pain from her elbow to shoulder; Shortness of breath; Blood pressure increased; fatigue; chills; Hear arm was hot to touch; Itchiness; Red circle near injection site the size of a quarter; not to take the second dose; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Small amount of blood clots) and HYPOKALAEMIA (Potassium levels dropped) in a 44-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 026A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse reaction. On 15-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 23-Mar-2021, the patient experienced VACCINATION SITE ERYTHEMA (Red circle near injection site the size of a quarter). On 24-Mar-2021, the patient experienced VACCINATION SITE WARMTH (Hear arm was hot to touch) and VACCINATION SITE PRURITUS (Itchiness). On 25-Mar-2021, the patient experienced PAIN IN EXTREMITY (Arm pain from her elbow to shoulder), DYSPNOEA (Shortness of breath), BLOOD PRESSURE INCREASED (Blood pressure increased), DIZZINESS (dizziness), FATIGUE (fatigue) and CHILLS (chills). On 01-Apr-2021, the patient experienced THROMBOSIS (Small amount of blood clots) (seriousness criterion medically significant), HYPOKALAEMIA (Potassium levels dropped) (seriousness criterion medically significant) and HYPERTENSION (Increased hypertension). On an unknown date, the patient experienced PRODUCT DOSE OMISSION ISSUE (not to take the second dose). At the time of the report, THROMBOSIS (Small amount of blood clots), HYPOKALAEMIA (Potassium levels dropped), PAIN IN EXTREMITY (Arm pain from her elbow to shoulder), DYSPNOEA (Shortness of breath), BLOOD PRESSURE INCREASED (Blood pressure increased), HYPERTENSION (Increased hypertension), VACCINATION SITE WARMTH (Hear arm was hot to touch), DIZZINESS (dizziness), VACCINATION SITE PRURITUS (Itchiness), FATIGUE (fatigue), VACCINATION SITE ERYTHEMA (Red circle near injection site the size of a quarter) and CHILLS (chills) outcome was unknown and PRODUCT DOSE OMISSION ISSUE (not to take the second dose) had resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. On 01-Apr-2021 patient went to the ER for 2 hours. The HCP at the ER stated she had increased hypertension, potassium levels dropped and there was a small amount of blood clots. Her primary care physician said not to take the second dose. Patient inquired about what she can do to manage her symptoms. Concomitant drugs were not reported. Treatment information was not provided. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. However, Further information is requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. However, Further information is requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245460-1	I had a stroke; Blood clot; Speak was bad, couldn't form words, speak was impaired; Missed her dose due to to the decision not to get it; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of CEREBROVASCULAR ACCIDENT (I had a stroke) and THROMBOSIS (Blood clot) in a 73-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 010M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No medical history was reported.). On 10-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 13-Feb-2021, the patient experienced CEREBROVASCULAR ACCIDENT (I had a stroke) (seriousness criterion hospitalization). On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion medically significant), SPEECH DISORDER (Speak was bad, couldn't form words, speak was impaired) and INTENTIONAL DOSE OMISSION (Missed her dose due to to the decision not to get it). On 16-Feb-2021, CEREBROVASCULAR ACCIDENT (I had a stroke) had resolved. At the time of the report, THROMBOSIS (Blood clot) and SPEECH DISORDER (Speak was bad, couldn't form words, speak was impaired) outcome was unknown and INTENTIONAL DOSE OMISSION (Missed her dose due to to the decision not to get it) had resolved. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not provided. The patient was receiving treatment with clopidogrel (Plavix) 75mg since the stroke (which is a stronger blood thinner TPA drug that resolve the blood clot). Reportedly, the patient was hospitalized for 3 days. Patient also stated that she was nervous about getting her 2nd dose so she missed her dose due to the decision not to get it with regards to the fear of having another stroke. Company comment Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked; Sender's Comments:
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245476-1	Blood clots; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clots) in an 82-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 013A21A and 031L209) for COVID-19 vaccination. Concurrent medical conditions included Cancer (The patient was in recovery from cancer). On 03-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 05-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. In March 2021, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In March 2021, Computerised tomogram: blood clots (abnormal) Blood clots. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant medications were not provided. Description: The patient developed blood clots (medically significant) two weeks after receiving his second dose of the vaccine . The patient was in recovery from cancer and was required to have computerized tomography (CT) scans. He had a computerized tomography (CT) scan within two weeks of taking his second dose and the blood clots were discovered. The patient had never had blood clots before. The patient had already been in contact with his oncologist and was prescribed apixaban. The patient received both scheduled doses of mRNA-1273 prior to the event, therefore action taken with the drug in response to the event is not applicable.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245770-1	Blood clot in right arm
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1246432-1	4 days (4/20/2021) post vaccination pain in right calf was experienced. As the pain did not resolve over the next week and felt similar to a DVT experienced post a spinal fusion in 2018 I consulted, on 4/16/2021, with the vascular surgeon, who treated my post surgery DVT in 2018 . He performed an ultrasound and identified a clot in my right calf. On 4/21/2021 he prescribed a course of Eliquis (initial dose 20 mg per day) to resolve the clot. As of today (4/23/2021) the pain has lessened but not resolved. In addition to the DVT I experienced a significant level of fatigue for 3 days post the vaccination. While the fatigue has lessened it has not completely resolved.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1246568-1	I had a pulmonary embolism on 3/17/2021. Prior to that a blood clot in my left calf.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1246721-1	Fever over 100, 3 blood clots discovered in hospital where I was sent by the Hematologist after fever started.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1247150-1	Pt presented to the ED from home for b/l lower extremity swelling x 2 weeks. She was reportedly in her recliner for approximately 12-18 hrs prior to being brought in via EMS. In the ED her b/l LE extremity was cold to touch and mottled. Her toes were white and nail beds blue. No pulses were found on Doppler on either extremity. She was also noted to be in a.fib with RVR. Doppler of LE showed extensive occlusive thrombus. Bedside echo showed the RV severely dilated with mobile thrombus and EF 20-25%. Given these findings there is a very high probability of massive PE, however given current situation of AKI and instability CT was unable to be performed. She was given TPA and admitted to ICU. Pt was started heparin gtt, as well as levophed and dobutamine for obstructive and cardiogenic shock as her troponin, kidney function, lactate and liver function all were elevated. Of note pt lives in a hoarding situation and lives a sedentary lifestyle and reportedly only gets up from recliner to use the bathroom. Pt also has a 96 pack year history but was noted in her medical chart to have quit smoking in 2017.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1248534-1	I because confused and unsteady. I then fell. My husband helped me up but he knew something was wrong. He called 911 an ambulance took me to the hospital but I was transferred to another hospital. In case I needed surgery. It was touch and go. I was suffering a stroke. A blood clot was also found. I was in the hospital ICU for 3 days then transferred to another room. For another day.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1249568-1	blood clot on the left leg; blood clot on the right leg; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot on the right leg) and THROMBOSIS (blood clot on the left leg) in a 30-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 038321A) for COVID-19 vaccination. The patient's past medical history included No adverse event (no medical history reported). On 11-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 08-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 05-Apr-2021, the patient experienced THROMBOSIS (blood clot on the right leg) (seriousness criterion medically significant). On 12-Apr-2021, the patient experienced THROMBOSIS (blood clot on the left leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (blood clot on the right leg) and THROMBOSIS (blood clot on the left leg) outcome was unknown. Not Provided The patient has not provided with any concomitant medication. No treatment medications are listed. Company Comment Based on the current available information which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, a causal relationship cannot be excluded. This case was linked to US-MODERNATX, INC.-MOD-2021-043122 (E2B Linked Report). Reporter did not allow further contact; Sender's Comments: US-MODERNATX, INC.-MOD-2021-043122:1st dose

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1249676-1	massive blood clot in the right lung; painful knot under the skin on his arm; threw up about 20 times in 6 hours, and kept vomiting all the next day; massive blood clot in the lung ended up in different parts of the lung causing extreme amount or pain\ Left lung is beginning to hurt again and his chest is killing him; next day when he got up and could barely walk; This spontaneous case was reported by a consumer and describes the occurrence of PULMONARY EMBOLISM (massive blood clot in the right lung), PULMONARY PAIN (massive blood clot in the lung ended up in different parts of the lung causing extreme amount or pain\ Left lung is beginning to hurt again and his chest is killing him) and THROMBOSIS (painful knot under the skin on his arm) in a 45-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 027A21A and 007B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Autoimmune disorder NOS in September 2020 and Broken bones. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 19-Mar-2021, the patient experienced GAIT DISTURBANCE (next day when he got up and could barely walk). On 26-Mar-2021, the patient experienced PULMONARY EMBOLISM (massive blood clot in the right lung) (seriousness criteria hospitalization, medically significant and life threatening). On 04-Apr-2021, the patient experienced PULMONARY PAIN (massive blood clot in the lung ended up in different parts of the lung causing extreme amount or pain\ Left lung is beginning to hurt again and his chest is killing him) (seriousness criterion medically significant). On an unknown date, the patient experienced THROMBOSIS (painful knot under the skin on his arm) (seriousness criterion medically significant) and VOMITING (threw up about 20 times in 6 hours, and kept vomiting all the next day). On 19-Mar-2021, GAIT DISTURBANCE (next day when he got up and could barely walk) outcome was unknown. On 28-Mar-2021, PULMONARY EMBOLISM (massive blood clot in the right lung) had not resolved. On 04-Apr-2021, PULMONARY PAIN (massive blood clot in the lung ended up in different parts of the lung causing extreme amount or pain\ Left lung is beginning to hurt again and his chest is killing him) had not resolved. At the time of the report, THROMBOSIS (painful knot under the skin on his arm) and VOMITING (threw up about 20 times in 6 hours, and kept vomiting all the next day) outcome was unknown. Not Provided Treatment information provided as blood thinners (IV heparin) and antibiotics were used to treat the events. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked to US-MODERNATX, INC.-MOD-2021-081632 (E2B Linked Report).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. US-MODERNATX, INC.-MOD-2021-081632:2nd dose
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1249678-1	Left lung is beginning to hurt again; chest is killing him; can barely walk; pain all over; the knot on the arm is blood clot; knot under the skin on his arm; painful knot under the skin; nauseated; kept vomiting all the next day, threw up about 20 times; This spontaneous case was reported by a patient and describes the occurrence of THROMBOSIS (the knot on the arm is blood clot) in a 45-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 007B21A and 027A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Broken bones. Concurrent medical conditions included Autoimmune disorder in September 2020. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 15-Apr-2021, the patient experienced NAUSEA (nauseated) and VOMITING (kept vomiting all the next day, threw up about 20 times). On 18-Apr-2021, the patient experienced SKIN MASS (knot under the skin on his arm) and PAIN OF SKIN (painful knot under the skin). On 19-Apr-2021, the patient experienced THROMBOSIS (the knot on the arm is blood clot) (seriousness criterion medically significant). On 20-Apr-2021, the patient experienced PULMONARY PAIN (Left lung is beginning to hurt again), CHEST DISCOMFORT (chest is killing him), GAIT DISTURBANCE (can barely walk) and PAIN (pain all over). On 17-Apr-2021, NAUSEA (nauseated) and VOMITING (kept vomiting all the next day, threw up about 20 times) had resolved. At the time of the report, THROMBOSIS (the knot on the arm is blood clot), PULMONARY PAIN (Left lung is beginning to hurt again), CHEST DISCOMFORT (chest is killing him), GAIT DISTURBANCE (can barely walk), SKIN MASS (knot under the skin on his arm), PAIN OF SKIN (painful knot under the skin) and PAIN (pain all over) outcome was unknown. No concomitant medications were reported. Treatment for the events included Elquis and I.V heparin. This case was linked to MOD-2021-081600 (Patient Link).; Sender's Comments: Based on the information provided which includes a strong temporal association between the use of mRNA-1273 vaccine and onset of the reported events, a causal relationship cannot be excluded. Nausea and vomiting are consistent with the product safety profile.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1250826-1	heart attack from blood clot

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1250891-1	Blood clot in right leg with rash mainly on arms and legs but included body.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1250998-1	Feb 6th pain in right calf. Feb 9th 2021 had ultrasound of leg and diagnosed with blood clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1254028-1	Patient passed away; patient had blood clots in his brain/ Legs/ Lungs/ Arms; Severe hypotension; This spontaneous case was reported by a physician (subsequently medically confirmed) and describes the occurrence of DEATH (Patient passed away), THROMBOSIS (patient had blood clots in his brain/ Legs/ Lungs/ Arms) and HYPOTENSION (Severe hypotension) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient's past medical history included No adverse event. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced DEATH (Patient passed away) (seriousness criterion death), THROMBOSIS (patient had blood clots in his brain/ Legs/ Lungs/ Arms) (seriousness criterion death) and HYPOTENSION (Severe hypotension) (seriousness criterion death). The cause of death was not reported. It is unknown if an autopsy was performed. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. Treatment information was not provided. Company comment: This case concerns the death of a patient of unknown age and gender after administration of mrna-1273 ()LOT UNKNOWN). Very limited information regarding this event/s has been provided at this time. Further information has been requested. Critical details such as the mRNA-1273 date of administration, onset of any signs an d symptoms, and date of death is lacking.; Sender's Comments: This case concerns the death of a patient of unknown age and gender after administration of mrna-1273 ()LOT UNKNOWN) . Very limited information regarding this event/s has been provided at this time. Further information has been requested. Critical details such as the mRNA-1273 date of administration, onset of any signs an d symptoms, and date of death is lacking.; Reported Cause(s) of Death: Unknown cause of death
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1256399-1	Second vaccine given 3-4-21 on 4-21-21 was treated at the ER for blood clot in right calf . My calf was tight and a bit swollen. My heart beat was rapid with shoulderblade pain and tightness in chest. Tightness and blade pain started on 4/20 pm with a hard calf. Morning pain worse and leg was more swollen and warm
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1256866-1	I had tender areas on the inside of my left leg starting about mid or late March in the upper thigh region and some aching in the inner aspect of the calf portion of the leg. This became worse over the Easter weekend and I went to clinic on April 6, 2021. I had a superficial blood clot just above the knee on the inner aspect of my left leg. After that appointment the redness and soreness and very hard feeling of the veins extended up my leg 8 to 10 inches. I did not have a blood clots in the deep veins. I have had some varicose vein?s and surgery for such about five years ago. The clinic did not say the clot was due to the vaccine but I had no injuries, no recent surgeries and nothing unusual that would have caused the clot other than the vaccine
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1258878-1	Received Moderna Covid Vaccine on 3/3/21 and 4/8/2021. Nted onset 4/24 of right calf discomfort, swelling and firmess and at same time began to have exertional dyspnea with chest heaviness. U/S RLE - extensive DVT. CTA - multiple segmental vessels with clot, dilated right ventricle
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1259563-1	Patient called and reported she had a blood clot in her leg confirmed by doctor on 4/14/21. Leg pain started 4/13/21. Patient reported her doctor informed her the clot was due to the COVID vaccine she received. Doctor put her on Elliquis and instructed her to elevate leg. She follows up with doctor once a week.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1260559-1	Left underarm has a painful lymph node. Blood clot on the same arm above the elbow.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1261538-1	Blood clots; Gangrene; Missed her second dose as the hospital where the patient was admitted did not have Moderna vaccine; patient was sent to rehabilitation center; surgery (a leg was amputated); two clots were removed; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clots), THROMBECTOMY (two clots were removed), GANGRENE (Gangrene) and LEG AMPUTATION (surgery (a leg was amputated)) in a 50-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 025J20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Clot blood (blood clots in heart and lungs for last 15 years) and Thrombectomy (40 clots removed from heart and lungs from past 15 years). Concurrent medical conditions included Systemic lupus erythematosus (bad lupus). On 15-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 10-Feb-2021, the patient experienced THROMBOSIS (Blood clots) (seriousness criteria hospitalization and medically significant). On 01-Mar-2021, the patient experienced THROMBECTOMY (two clots were removed) (seriousness criterion hospitalization prolonged) and LEG AMPUTATION (surgery (a leg was amputated)) (seriousness criterion hospitalization prolonged). On 12-Mar-2021, the patient experienced REHABILITATION THERAPY (patient was sent to rehabilitation center). On an unknown date, the patient experienced GANGRENE (Gangrene) (seriousness criterion hospitalization prolonged) and PRODUCT ADMINISTRATION INTERRUPTED (Missed her second dose as the hospital where the patient was admitted did not have Moderna vaccine). The patient was hospitalized on 10-Feb-2021 due to THROMBOSIS, then on 27-Feb-2021 due to LEG AMPUTATION and THROMBECTOMY. The patient was treated with Surgery for Gangrene. On 01-Mar-2021, THROMBECTOMY (two clots were removed) and LEG AMPUTATION (surgery (a leg was amputated)) had resolved. At the time of the report, THROMBOSIS (Blood clots), PRODUCT ADMINISTRATION INTERRUPTED (Missed her second dose as the hospital where the patient was admitted did not have Moderna vaccine) and REHABILITATION THERAPY (patient was sent to rehabilitation center) outcome was unknown and GANGRENE (Gangrene) had resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. As per the patient's doctor, it was too late and dangerous to remove clots from the patient's leg and eventually, the doctor had to amputate the patient's leg because of gangrene. No concomitant medications were reported. Company comment: Based on current available information and the temporal association between product use and het start date of the events a causal relationship cannot be excluded.; Sender's Comments: Based on current available information and the temporal association between product use and het start date of the events a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1261946-1	Blood clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) in a 52-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 044A21A and 031B21a) for COVID-19 vaccination. No Medical History information was reported. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) outcome was unknown. No treatment information was provided. No concomitant medication was reported. The patient received both scheduled doses of mRNA-1273 prior to the event(s); therefore, action taken with the drug in response to the event(s) is not applicable. Company Comment: Very limited information regarding this event/s has been provided at this time. Further information has been requested. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender's Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1261984-1	"VTE legs, large intestine, & lungs Narrative: Patient reports received his first COVID vaccine dose of Moderna on 2/17/21 and reports in the early morning of 2/23/21 he began feeling pain in his right side. Patient states he thought his appendix had burst and he went to get checked out. Patient states they ran test and could not determine what was wrong, so he was sent to hospital. Patient states they found clots in his legs, large intestine, and lungs. States he underwent surgery to remove the clot from his large intestine and he now on Eliquis. Patient states he was ""perfectly healthy"" before receiving the COVID vaccine. Active meds: Ibuprofen 800mg TID for pain, Lidocaine 5% patch; 1 patch daily for 12 hours, and Sodium fluoride 1.1% oral cream; apply to mouth 1-2 times daily when brushing teeth"
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1263633-1	On Friday 4/9 I experienced chills, fever and body aches. I called the on call doctor and he informed me my symptoms would be better in 24 hours and to call back if my symptoms worsened. On Saturday 4/10 I started having problems breathing and pain in my chest which lasted all day. Sunday 4/11 I went to the emergency room of which I was admitted and was told I had Pulmonary Embolism. I am currently being treated with Eliquis.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1263807-1	I completed my Moderna vaccine series on 2/5/21 to protect myself against COVID-19. I also tested positive for COVID-19 in mid-December. This past week I had a menstrual cycle, and I discharged a high volume of blood blots. I am taking an oral contraceptive and my periods are are normal frequency, length and flow, until this last period. The clots could fill an entire maxi pad that needed to be changed every 2 hours, for about 48 hours. After that single clots were discharged every 2 hours for another 2 days (and still occurring as I write this). I sent my PCP (an OBGYN) a direct message in patient portal also reporting symptoms.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1264696-1	Blood clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No medical history was provided by the reporter. On 03-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. In March 2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) was resolving. The patient took the second vaccine on the 29th or 30th of March Company Comment Very limited information regarding this event has been provided at this time. Further information has been requested. This case was linked to MOD-2021-076604 (Patient Link). Reporter did not allow further contact
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1265444-1	2 hrs after he received his vaccine he experienced a sore arm - shaking - chills - and a temp of 101. I gave him Tylenol. He received his vaccine on a Wednesday. Thursday - He experienced Chills - a temp of 103.4 - nose bleeds and spitting up blood. Clots of blood coming from his nose and exhausted. I continued with the Tylenol for the fever Friday - A temp of 100 - his arm still sore - exhausted - very little nose bleeds. By evening time - low grade temp - nose bleeds had stopped - exhausted. Continued with Tylenol. Saturday - just exhausted and sore arm
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1266459-1	The patient came in for her second dose of Moderna and had concerns because she stated that she developed a nose bleed the day after she received her first dose. She stated that she also passed clots and her nose bled for more than 1 day. She stated that she went to the ER but was only given a nasal spray to use. She denied a history of nosebleeds and does not take blood thinners.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1267593-1	Headache, Increasing fatigue and difficulty breathing over two weeks before 2nd dose -- exacerbated by 2nd dose. Worsening condition led to visit to Primary Care In office tests indicated presence of blood clots and need for emergency hospital treatment. Emergency surgery to remove blood clots throughout her body and attempt to put her on ECMO were unsuccessful and patient succumbed at 12:06 AM 3/3/21.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1267830-1	4/7/21 around 630pm began to feel ill and had sore arm. By about 1030pm, had significant body aches, headache, clammy skin, vomiting, diarrhea, near passing out, fast heart beat, with fever. These symptoms lasted about 48 hours. The next 48 hours just had ill feeling with extreme fatigue. Come 4/12/21, I had nausea, headache, fever of about 100.4 and body aches. By 4/14/21, I ended up going to the ER for shortness of breath, chest tightness, extreme fatigue and voice hoarseness. Those doctors performed several labs to include Chest CT which showed elevated D-Dimer levels. An EKG was performed which was baseline. Ultrasound of my veins was performed. They also did a COVID test which was negative. They sent me home and told me to rest and drink lots of fluid. On 4/16/21, I had a video call with my doctor who didn't like the sound of my voice and straining to catch my breath and I was sent to urgent care/ER. They admitted me as an ER visit due to my continued shortness of breath, cough, sore throat, night sweats, dizziness, chest tightness along with my voice being hoarse. Another COVID test was performed and was negative. During their exam, my oxygen levels were dropping into the low 80s and my heart rate was in the 120s and 130s. Their CT scan was down so they sent me to another ER where another CT scan was performed. Another COVID test was performed and was negative. My D-Dimer was elevated, liver enzymes was slightly elevated and again my oxygen levels were lower than would like to see and elevated heart rate. I ended up being admitted into the hospital for an overnight stay to monitor my symptoms. They treated me for suspected blood clot, shortness of breath, chest tightness. I was given 2 shots of heparin in the stomach, IV fluids, multiple vitamins in attempt to boost my immune system, along with ADVAIR and Albuteral. My discharge diagnose from the hosptial was Acute Bronchitis, Anxiety disorder, Recent Moderna 2nd dose COVID vaccine with significant side effects in the form of fever, generalized weakness and dizziness.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1268350-1	a week after receiving the vaccine I started feeling shortness of breath. It progressively got worse and on April 23, 21 was admitted into Hospital with blood clots to the left leg and to my lungs.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1268431-1	Pt awoke on 02/28/2021 with increased lower extremity edema went to Urgent care had CXR and was told to increase Lasix to 40 mg daily for 4 days. Swelling became worse over next 2 days and patient became short of breath and on 03/02/2021 she went to Hospital and had a CT scan of chest which showed a pulmonary embolus and a bilateral lower extremity ultrasound which showed a right femoral DVT and popliteal occlusive thrombus and a posterior tibial and peroneal vein occlusive thrombus. She was started on Lovenox and admitted to the hospital . she was discharged on 03/05/2021 and had a right femoral thrombectomy as an outpatient on 03/ 08 2021 with complete resolution . She was discharged home on Lovenox and later switched to Eliqus. She is home and doing well at time ofreport.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1268957-1	Shortness of breath, fatigue, blood clots in right lung diagnosed in ER, Medical Center
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1269286-1	Clots; This spontaneous case was reported by a health care professional (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Clots) in a 38-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 25-Apr-2021, the patient experienced THROMBOSIS (Clots) (seriousness criteria hospitalization and medically significant). The patient was hospitalized on 25-Apr-2021 due to THROMBOSIS. At the time of the report, THROMBOSIS (Clots) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No concomitant medications reported. No treatment information provided.; Sender's Comments: Limited information regarding the event has been provided at this time and a causal relationship cannot be excluded
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1270199-1	acute superficial thrombus
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1270689-1	Entered hospital on 3/22/21 At around 8am in the emergency room. Had ultrasound of left leg and was placed in ICU for a massive blood clot in the left leg. Had surgery at around 4pm on 3/22/21 and placed back in ICU until the next surgery on 3/23/21 to remove the blood clot. Was released on 3/25/21.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1271346-1	Blood clot behind left knee found on 3/11/21 and traveled to lungs. Admitted to hospital for Heart Failure from Chemo, PE and pneumonia.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1271452-1	Severe pain to lower left leg without injury hx. Went to doctor on 3/17/21 and was examined and doppler to left leg showing no blockage. Dr prescribed nsaid as needed for pain and monitor. Monitored and pain continued until approx 4/9 and noticed swelling starting and some redness to skin slightly warm. Returned to dr on 4/13 and was placed on antibiotics. On 4/14 noticed slight shortness of breath on exertion and on 4/15 shortness of breath worsened and some irregular heartrate. I went to hospital. Bloodwork, EKG, Chest xray and CT showed multiple pulmonary embolisms and several DVTs in left leg. Admitted to hospital and placed on Heparin drip and monitored for three days. Released from hospital on 4/19 on Eliquis and advised to follow up with personal physician and report severe breathing problems or CP.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1271834-1	Blood Clot and pulmonary embolism from first shot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1271920-1	3 weeks after Patient got this shot he got blood clots in his leg and amputated next week .The problem is he has ITP platelets were 12,000 He can't make blood clots!!!

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1272201-1	Pulmonary embolism; died - autopsy showed multiple blood clots all over his body - pelvic area, hearts,arteries, lungs; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (died - autopsy showed multiple blood clots all over his body - pelvic area, hearts,arteries, lungs) and PULMONARY EMBOLISM (Pulmonary embolism) in an 83-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 039A21A and 003A21A) for COVID-19 vaccination. Concurrent medical conditions included Diabetes, Prostate cancer and Memory loss. Concomitant products included METFORMIN for Diabetes. On 05-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 02-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 04-Apr-2021, the patient experienced THROMBOSIS (died - autopsy showed multiple blood clots all over his body - pelvic area, hearts,arteries, lungs) (seriousness criterion death). On an unknown date, the patient experienced PULMONARY EMBOLISM (Pulmonary embolism) (seriousness criterion death). The patient died on 04-Apr-2021. The reported cause of death was Pulmonary embolism. An autopsy was performed. The autopsy-determined cause of death was multiple blood clots. Action taken with mRNA-1273 in response to the event was not applicable Other concomitant medications were reported included unspecified medications for memory loss and diabetes. No treatment information was provided.; Sender's Comments: This is an 83-year-old, male patient who received mRNA-1273 Vaccine who experienced multiple thrombosis and died, 2 days after receiving second dose of vaccine. Medical history includes Diabetes, and Prostate cancer. Conmeds including some unspecified medications for memory loss and diabetes. The autopsy-determined cause of death was multiple blood clots. Very limited information has been reported at this time. Further information is expected,; Reported Cause(s) of Death: Pulmonary embolism; Autopsy-determined Cause(s) of Death: multiple blood clots
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1272205-1	Blood clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) in a 68-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 011J20A and 038K20A) for COVID-19 vaccination. No Medical History information was reported. On 01-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 05-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) outcome was unknown. Not Provided No concomitant medications were reported. The patient had no problems after her second dose. Treatment information was not provided. Action taken with mRNA-1273 in response to events was not applicable. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1272645-1	"03/31/2021: began feeling aches/pain/feverish about 10 hours after receiving the first Moderna vaccine and had a temperature of 100.3 F. 04/01/2021: continued feeling aches/pains/feverish and had a temperature of 99 F. 04/02/2021: aches/pains/fever subsided. 04/03/2021: left leg began to hurt and kept feeling worse throughout the day; took ibuprofen for pain. 04/04/2021: left leg continued to hurt; took ibuprofen for pain. 04/05/2021: visited Urgent Care clinic and an ultrasound confirmed the presence of a blood clot in the left leg. Urgent Care Doctor prescribed Eliquis for treatment. See Continuation Page for subsequent events/treatments. 04/07/2021: Meeting with Primary Care Physician (PCP). PCP changed treatment for the blood clot [acute deep vein thrombosis (DVT)] from Eliquis to Lovenox/Coumadin. Additional tests ordered: CBC with Differential; PTT; Comprehensive Metabolic Panel; Protime INR; Sedimentation Rate. 04/08/2021: Meeting with PCP to discuss DVT. PCP ordered Noninvasive Vascular Imaging - External Scan. 04/15/2021: Meeting with PCP to discuss DVT. PCP instructed to continue treatment with Lovenox and keep appointment with hematologist on 04/20/2021. PCP ordered the following tests: CBC no Differential; Comprehensive Metabolic Panel; Protime INR. 04/20/2021: PCP changed recommended treatment of DVT from Lovenox to Eliquis. 04/20/2021: Meeting with hematologist to discuss DVT. Dr. agreed with PCP to change treatment from Lovenox to Eliquis. Dr. ordered the following immediate tests: Antiphospholipid Ab Panel; CK Total; Comprehensive Metabolic Panel; Dilute Russel Viper Venom Confirm; Fibrinogen; Lupus Anticoagulant Profile; NT-PRO BNP; Phospholipid, Hexagonal Phase; VAS Porto Hepatic Duplex. Additional future tests were ordered as follows: Antiphospholipid Ab Panel, CK Total; Fibrinongen; Lupus Anticoagulant Profile; NT-PRO BNP. 04/23/2021: Meeting with hematologist to discuss DVT. Dr. suspects the DVT is related to pre-existing conditions (includes vitamin K deficiency) plus the effect of the first Moderna vaccine. Dr. concluded that vitamin deficiencies (due to malabsorption issues) are the root cause of several ills. Dr. ordered an injection of vitamin K and recommended continued treatment with Eliquis. Dr. also stated ""I think no to second vaccination #2. She might be vaccinated later if vitamin def are resolved."" Dr. plans to review the case in three months to determine if Eliquis treatment can be discontinued. Dr. also ordered an ""ECHO Complete"" to rule out additional clots in the heart/lungs."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1273828-1	"Received 2nd Moderna vaccine on 4/15/21. Woke up on Saturday 4/17/21 with significant ""cramp"" in left calf. Pain persisted, so patient went to doctor on 4/19/21. Blood work did not ""rule out"" clot. Doctor ordered venous doppler which confirmed clot. Prescribed 3 month course of Xarelto."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1274591-1	Extreme pain and cramping, blood clots and still having my period since the first of April.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1274672-1	Body aches, fever, nose blister, headaches swollen and arm diarrhea. 0n 3/2021 I statdetedbtonget pain on my left foot and calf. I went to the hospital and on 3/22/2021 I had a ultrasound and the results showed that I have a blood clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1274828-1	Had three blood clots within 36 hours. One in left leg, one right leg and one in groin. Also, rheumatological problems in tendons in arms and legs.Hearing in left ear diminished by 10%.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1275047-1	"About 6 ? 7 days w/ side effects ? Significant Malaise, tiredness. Unable to function or concentrate. Likely a ""self inflicted"" situation by unknowingly sitting in chair in a daze ultimately developing blood clots (pulmonary edema), blood clots in legs, leading to mild heart attack, hospitalization. I believe there may be a direct connection to a medical procedure (vedolizumab (Entyvio) 300 mg intravenously infusion) that preceded the vaccination on the previous Friday (2/19/21)."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1276686-1	Blood Clot; left knee had red marks above it; pain in thigh; swelling from the knee up to the thigh; left knee was in pain tremendously; difficulaty walking; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood Clot) in a 68-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 013620A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 14-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Jan-2021, the patient experienced GAIT DISTURBANCE (difficulaty walking) and ARTHRALGIA (left knee was in pain tremendously). On 16-Jan-2021, the patient experienced ERYTHEMA (left knee had red marks above it), PAIN IN EXTREMITY (pain in thigh) and JOINT SWELLING (swelling from the knee up to the thigh). On an unknown date, the patient experienced THROMBOSIS (Blood Clot) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood Clot), GAIT DISTURBANCE (difficulaty walking), ERYTHEMA (left knee had red marks above it), PAIN IN EXTREMITY (pain in thigh), JOINT SWELLING (swelling from the knee up to the thigh) and ARTHRALGIA (left knee was in pain tremendously) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant medications like high blood pressure medication, cholesterol medication and medication for irregular heartbeat were reported. On 15 Jan2021, the patient said that his left knee was in pain tremendously and had difficulty walking. On 16 JAN 2021, the left knee had Red marks above it. He also had swelling from the knee up to the thigh. He also experienced pain in the thigh and knee. He went to the doctor on 20 JAN 2021 and was sent for an ultrasound. The doctor thought it could be related blood clot. No laboratory details were provided. On 20-Jan-2021, the patient was sent for an ultra sound, results unknown. Treatment included ibuprofen and paracetamol. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1276717-1	blood clots in both legs; heart attack/blood clot in heart; Blood clots in lungs/shortness of breath; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots in both legs), MYOCARDIAL INFARCTION (heart attack/blood clot in heart) and PULMONARY EMBOLISM (Blood clots in lungs/shortness of breath) in a 72-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 8077727310 and 012LZ0A) for COVID-19 vaccination. No Medical History information was reported. On 25-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 09-Apr-2021, the patient experienced THROMBOSIS (blood clots in both legs) (seriousness criteria hospitalization prolonged and life threatening), MYOCARDIAL INFARCTION (heart attack/blood clot in heart) (seriousness criteria hospitalization prolonged and life threatening) and PULMONARY EMBOLISM (Blood clots in lungs/shortness of breath) (seriousness criteria hospitalization prolonged and life threatening). The patient was hospitalized on 09-Apr-2021 due to MYOCARDIAL INFARCTION, PULMONARY EMBOLISM and THROMBOSIS. At the time of the report, THROMBOSIS (blood clots in both legs), MYOCARDIAL INFARCTION (heart attack/blood clot in heart) and PULMONARY EMBOLISM (Blood clots in lungs/shortness of breath) outcome was unknown. No concomitant medications were reported. Treatment of the events included Xarelto and unspecified IV medications. Action taken with mRNA-1273 in response to the events was not applicable. Company Comment: Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1276790-1	having a blood clot down whole right leg / up right side into right chest / under right breast / several clots in her lower abdomen; she almost died; urinary tract infection; right leg is very swollen; Kept losing consciousness/Passed out; hard time walking; had to use a walker; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (having a blood clot down whole right leg / up right side into right chest / under right breast / several clots in her lower abdomen), LOSS OF CONSCIOUSNESS (Kept losing consciousness/Passed out) and FEELING ABNORMAL (she almost died) in a 56-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 002b21a and 004m20a) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included LEVOFLOXACIN, RIVAROXABAN (XARELTO), LISINOPRIL, HYDROCHLOROTHIAZIDE, OXYCODONE and LEVOTHYROXINE SODIUM (SYNTHROID) for an unknown indication. On 12-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 12-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 29-Mar-2021, the patient experienced GAIT DISTURBANCE (hard time walking) and WALKING AID USER (had to use a walker). On 12-Apr-2021, the patient experienced THROMBOSIS (having a blood clot down whole right leg / up right side into right chest / under right breast / several clots in her lower abdomen) (seriousness criteria hospitalization prolonged, medically significant, life threatening and intervention required) and LOSS OF CONSCIOUSNESS (Kept losing consciousness/Passed out) (seriousness criterion medically significant). 12-Apr-2021, the patient experienced FEELING ABNORMAL (she almost died) (seriousness criteria hospitalization and medically significant). On 26-Apr-2021, the patient experienced PERIPHERAL SWELLING (right leg is very swollen). On an unknown date, the patient experienced URINARY TRACT INFECTION (urinary tract infection). The patient was hospitalized from 12-Apr-2021 to 19-Apr-2021 due to FEELING ABNORMAL and THROMBOSIS. On 12-Apr-2021, LOSS OF CONSCIOUSNESS (Kept losing consciousness/Passed out) had resolved. At the time of the report, THROMBOSIS (having a blood clot down whole right leg / up right side into right chest / under right breast / several clots in her lower abdomen) was resolving and FEELING ABNORMAL (she almost died), URINARY TRACT INFECTION (urinary tract infection), GAIT DISTURBANCE (hard time walking), PERIPHERAL SWELLING (right leg is very swollen) and WALKING AID USER (had to use a walker) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 12-Apr-2021, Computerised tomogram: blood clots (abnormal) blood clot. On 12-Apr-2021, Ultrasound scan: blood clots (abnormal) blood clots. Action taken with mRNA-1273 in response to the events was not applicable. Patient stated she was in the ICU24-48 hours then transferred to another area of the hospital where she spent another 5-6days. She is still having to use a walker. Her right leg is very swollen. She has to walk and keep her right leg elevated. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. However, Further information has been requested. This case was linked to MOD-2021-090647 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. However, Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1276794-1	Blood clot in leg; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot in leg) in a 67-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 002B21A and 002A21A) for COVID-19 vaccination. No Medical History information was reported. On 26-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 26-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 2 dosage form. On 02-Apr-2021, the patient experienced THROMBOSIS (Blood clot in leg) (seriousness criteria hospitalization and medically significant). The patient was hospitalized from 02-Apr-2021 to 06-Apr-2021 due to THROMBOSIS. The patient was treated with Surgery (Angioplasty) for Thrombosis. On 06-Apr-2021, THROMBOSIS (Blood clot in leg) had resolved. Not Provided Concomitant product use was not provided by the reporter. The patient received both scheduled doses of mRNA-1273 prior to the events; therefore, action taken with the mRNA-1273 in response to the events was not applicable.; Sender's Comments: Based on current available information and the temporal association between product use and the stat date of the events a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1276796-1	blood clot; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (blood clot) (seriousness criterion death). The reported cause of death was Clot blood. It is unknown if an autopsy was performed. The reporter stated that his wife had recently passed away from a blood clot after receiving a second dose of the Moderna COVID-19 vaccine. Treatment information was not provided. Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Clot blood
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1277626-1	I got blood clots. I do not know if they were related to the vaccine or not. I was on Eliquis and took it as prescribed. I was put on another blood thinner.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1277897-1	Normal menstrual cycle (i.e. week 4/inactive pills) on birth control pill ran April 4-10, but menstruation continued for 1 further week while taking the first week of active BC pills, April 11-17, 2021. Everything seemed normal for the second week, but menstruation began again during the third week of active BC pills, April 25-today. Menstrual flow has been heavier than normal, and I've noticed many blood clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1279122-1	Blood clot on 2/9/2021, just thought it was a severe calf cramp and took ibuprofen for several days until the calf felt somewhat better. Went to ER on 3/20 for dual pulmonary embolisms.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1280726-1	Heavy bleeding and clotting
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1280735-1	Blood clotting in fingers.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1281264-1	Superficial blood clot in right calf.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1281547-1	We noticed my mother's foot was swollen, after alerting her nurse practicioner they ordered a scan to be done on my mother. It confirmed that she had a blood clot between her groin and below her knee. They prescribed eloquis.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1281899-1	Developed blood clot diagnosed 5/1
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1282105-1	CVA with thrombosis on 4/23/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1283056-1	He noticed a hard lump in his bicep this morning and it appeared to be in a vein. He went in to see his doctor and the doctor diagnosed him with a blood clot. The doctor said it was in a superficial vein and didn't give him any treatment. Told him to keep a watch on it and if it got swollen or turned red or he developed shortness of breath to come back in and be seen.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1284721-1	Superficial blood clot; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Superficial blood clot) in a 58-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 014C21A and 018B21A) for COVID-19 vaccination. No Medical History information was reported. On 25-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 23-Apr-2021, the patient experienced THROMBOSIS (Superficial blood clot) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Superficial blood clot) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 24-Apr-2021, Ultrasound scan: abnormal (abnormal) Ultrasound of left arm was done and determined a blood clot in that arm.. Patient went to the emergency room on 24 Apr 2021, approximately 7 pm. The patient was a healthy individual and used to do bike ride 11 miles per day with average speed 16mph. He did bike ride the evening of his second covid shot. No relevant concomitant medications were reported. Treatment information included anti-inflammatory medicine. Action taken with mRNA-1273 in response to the events was not applicable. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Most recent FOLLOW-UP information incorporated above includes: On 26-Apr-2021: Additional information was received on 26 Apr 2021. Added lab data and treatment; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1284727-1	heart attack; Bood clot; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of MYOCARDIAL INFARCTION (heart attack) and THROMBOSIS (Bood clot) in a 67-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 23-Mar-2021, the patient experienced MYOCARDIAL INFARCTION (heart attack) (seriousness criteria death and medically significant) and THROMBOSIS (Bood clot) (seriousness criterion death). The patient died on 23-Mar-2021. The reported cause of death was Heart attack and Clot blood. It is unknown if an autopsy was performed. Not Provided No concomitant medication were reported. No treatment information was provided.; Reported Cause(s) of Death: Heart attack; Clot blood
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1285213-1	On 2/28/21, had first COVID Shot. On 3/16/21 had Hysterectomy. On 3/30/21 had second COVID shot. On 4/19/21, was diagnosed with blood clot. The doctor reported that it was not the norm for someone my age and with my good health to get a blood clot after a hysterectomy.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1285525-1	Menstrual cycle came two weeks early, no cramping at all (which is abnormal) thicker/ clotty, lasted a normal amount of time. I have had hx of abnormal menstrual cycles in the past, but not that come this early.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1285611-1	My periods since receiving the vaccination have had significantly more and larger clots than normal.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1286241-1	The patient developed a blood clot in the left calf 2 weeks after the first dose of Moderna COVID vaccine, which required treatment with Eliquis 5 mg twice daily. The patient also reported hearing loss that needed treatment with a tapering steroid protocol. The sense of hearing did not recover after the most recent loss event.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1286548-1	The blood vessels to the pt's spleen burst due to a clot causing removal of her spleen. During surgery she was given 2 units of blood. Due to blood clots (albeit in other parts of the body) with other vaccines, the lack of splenic trauma preceeding this event, the pt's lack of history re: blood clots, and in the interest of furthering this emerging COVID-19 vaccine technology/progress, the pt's PCP and this POA thought it worth noting.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1286682-1	My Father received the first dose of CV19 on 03/02/21 and the second dose on 03/03/2021 in the morning, and at night he had a stroke, we called 911 they hospitalized him within a week he was out of danger and had physical therapies for a month at facility he got out on the 31st of March 2021 and on April 4th he once again went into the hospital for lack of oxygen, they detected blood clots in his legs and his lungs and also an aneurysm in his heart.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1287643-1	Severe rash covering scalp, face, neck, chest all the way down the body that got worse every day, despite being on Zyrtec daily, and then taking benedryl every 4-6 hours. Rash continues to spread over even more of the body. Is red, slightly raised. My entire body burned and itched, even on benedryl, creams to stop itching. Body temp of 99.6. Started getting blood clots when blowing nose starting Thursday morning. Rash was even inside my ears. Was worst 72 hours after 2nd dose, and has gotten a little bit better each day. Still have mild rash and itching even 1 week after shot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1289821-1	I got sick to my stomach almost instantaneously, I lost my taste for 3 days, I had fever and body aches for 3 days, my period became completely extreme I had blood clots which is not ordinary for me and it lasted an extra 4 days, my ears were pluged as well as not being able to move my arm for 3 days really at all, my throat was also sore and I had glands swell up

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1289890-1	I had a couple of blood clots earlier in the week which I found weird. On Saturday morning I went to the bathroom and wiped and there was very bright red blood. I called and was told as long as I did not bleed through a tampon or pad not to worry and if I was still having issues to contact my primary on Monday. I went through 2 super tampons on Saturday. The color was very dark, almost black in color and sticky to the touch. I did not need another tampon that night and it was gone the next day. I am and always have been VERY regular. My period is not due to start until the 15th of this month.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1290770-1	Blood clot; Couldn't breathe; Heart palpitations; Hives; vertigo; This spontaneous case was reported by a health care professional and describes the occurrence of THROMBOSIS (Blood clot) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Medical History not reported. On 15-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Mar-2021, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion medically significant), DYSPNOEA (Couldn't breathe), PALPITATIONS (Heart palpitations), URTICARIA (Hives) and VERTIGO (vertigo). At the time of the report, THROMBOSIS (Blood clot), DYSPNOEA (Couldn't breathe), PALPITATIONS (Heart palpitations), URTICARIA (Hives) and VERTIGO (vertigo) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. On 15 Mar 2021, within 1 hour of receiving first dose of the Moderna COVID-19 vaccine, the patient couldn't breathe, she had heart palpitations, hives, vertigo, and a blood clot. The patient stated that she was able to fell the blood clot. At the time of the report it was unclear if the patient had seen a physician for the events. Treatment for the event was cetirizine, and the patient stated that she has no hives when she takes cetirizine, but once she stops the cetirizine the hives return. Treatment provided with cetirizine for hives.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1291244-1	"1. Bulging veins appeared the day following first vaccination. From inside the left nostril onto the upper lip. on right temple above end of eye brow and between hair line, legs, behind the right knee. Behind the left knew were several large clots. 1.a. Had vein work done from November 2020 to January 2021. Clots had disappeared. 1.b. Feb. 27, 2021 clot on right temple was ""in"" or on top of area treated. There was pain that lasted for three days. 1.c. Told primary care doctor and saw Vein Doctor. Then there was not much concern about clots. 2. Eyes: 02/26/2021 through 02/27/2021 shooting pains around and through the eyes. Started right after vaccination. 3. Two bulging painful masses Center of forehead right and left of center of nose. 4. Lega and feet swelled 5. Toes: 03/03 toes separated and cramped on feet. Each foot two toes went left stuck together and two went right. 6. Feet 03/03/21 sore spots on top of feet. Pain made it hard to walk. Pain extended toward and over ankles and wrapped around heel.. I use a walker 7. Dizzy when looking down, shortness of breath. Contacted cardiology 8. Belching and air bubbles originating in the chest area 9. Sleep: Slept almost around the clock for first 5 days. 10. Very thirsty even after drinking water 11. Urinated more than usual. some burning. Reported to doctor. 12. Vaginal etching and dryness for 2 weeks 13. Joint Dryness; All joints but very painful and extra dry rough feeling in right hip. For 3 days 14. Stubbled when waling March 11, 12, 13, and 14 2021 15. Mind and body not working to"
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1293703-1	Blood clot, left leg
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1294101-1	Blood clot in calf deep vein thrombosis
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1295984-1	Systemic: BACK ACHE, CLOT IN LUNG-Severe, Systemic: Vomiting-Severe, Additional Details: PER PT: PT WENT TO URGENT CARE 2 DAYS AFTER ADMINISTRATION OF VACCINE. PT EXPERIENCED BACK ACHES AND STARTED COUGHING UP BLOOD/ AT URGENT CARE PT GOT A CT SCAN AND CLOT WAS SEEN IN LUNGS. PT WAS PRESCRIBED ELIQUIS AND IS WAITING FOLLOWUP BY HEMATOLOGIST
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1296659-1	patient says they got a STEMI (from a clot) and is now in the hospital. also left arm bruise where shot is.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1296700-1	Severe flu like symptoms occured approximately 12 hours after 2nd dose of Moderna vaccine. Blood Clot formed approximately 2 weeks later and created stroke on 4/23/2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1298761-1	Amnesia; Cardiac disorder; Thrombosis; Transient ischaemic attack; This case was received via another Manufacturer (Reference number: vsafe) on 27-Apr-2021 and was forwarded to Moderna on 27-Apr-2021. This regulatory authority case was reported by an other health care professional and describes the occurrence of AMNESIA (Amnesia), CARDIAC DISORDER (Cardiac disorder), THROMBOSIS (Thrombosis) and TRANSIENT ISCHAEMIC ATTACK (Transient ischaemic attack) in a 70-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 027L20A) for COVID-19 vaccination. The patient's past medical history included Glaucoma and Hypothyroidism. Concurrent medical conditions included Stroke. Concomitant products included LEVOTHYROXINE, BRIMONIDINE, LATANOPROST and TIMOLOL for an unknown indication. On 13-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 14-Jan-2021, the patient experienced AMNESIA (Amnesia) (seriousness criterion hospitalization), CARDIAC DISORDER (Cardiac disorder) (seriousness criterion hospitalization), THROMBOSIS (Thrombosis) (seriousness criteria hospitalization and medically significant) and TRANSIENT ISCHAEMIC ATTACK (Transient ischaemic attack) (seriousness criteria hospitalization and medically significant). At the time of the report, AMNESIA (Amnesia), CARDIAC DISORDER (Cardiac disorder), THROMBOSIS (Thrombosis) and TRANSIENT ISCHAEMIC ATTACK (Transient ischaemic attack) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 14-Jan-2021, Catheterisation cardiac: inconclusive. On 14-Jan-2021, Echocardiogram: inconclusive. On 14-Jan-2021, Magnetic resonance imaging: inconclusive. On 14-Jan-2021, Ultrasound scan: inconclusive. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. The next day, the patient had short term memory loss, patient went to the hospital, and they said patient had some type of heart episode. The patient was taken by ambulance to the hospital and was told had a blood clot in brain and had suffered a mini stroke. The patient was hospitalized for 4 nights. The patient was given blood thinner and got improved. Company comment: Based on the information provided which includes a temporal association between the use of mRNA-1273 vaccine, the onset of the reported events, a causal relationship cannot be excluded. However patient had an underlying risk in the past history of stroke.; Sender's Comments: Based on the information provided which includes a temporal association between the use of mRNA-1273 vaccine, the onset of the reported events, a causal relationship cannot be excluded. However patient had an underlying risk in the past history of stroke.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1299869-1	Blood clot. PT had blood clot removed at hospital.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1299873-1	Right leg burning, red, swollen. Doppler showed XL blood clot and numerous small clots in right leg. Put in Eliquis for2-3 wk. brought to hospital on 5.6.21 with worse swelling, red, and right foot very painful. CT showed clots in legs and in lungs. Still in hospital as of 5.8.21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1300080-1	I felt sluggish after my second Moderna shot on April 15th. On the evening of April 27th, I really started to feel horrible and was scared as it was hard to breathe and my chest pains were just horrible. I head to ER and am told I have blood clots in both lungs and 2 blood clots in lower leg. Taken by ambulance to the hospital.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1300351-1	symptoms lasting for over 25 days: Major Headache , extreme exhaustion, trouble breathing (SOB), loss of taste/smell, worsening arthritis, congestion in lungs and head, diarrhea, blood clot in finger (just was seen in the ER for confirmed blood clot in finger)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1300626-1	Day 15 post-vaccination, woke up with a vaginal bleeding similar to menstruation. It was very heavy, lasted 6 days, and I was passing clots. No medical help was needed or sought, but it was uncomfortable. I am in premature menopause and do not menstruate. I take HRT. Following my first Moderna shot, I had a very similar bleeding as well, starting the morning after. I have not had any vaginal bleeding following these two (after the first and second Moderna shot). Besides these effects, within hours of this second dose, I was feeling tired. I woke up the next morning with body aches similar to that of influenza, a temperature of 99.5, which is quite elevated for me, and overall tiredness. These symptoms resolved by next morning, when I woke up with a throbbing headache. It resolve following one dose of 200mg generic ibuprofen.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1300702-1	2-3 days after the first injection, I developed a blood clot in my lower right leg.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1301096-1	32 hours after the vaccine, he suffered a massive heart attack due to a blood clot and a massive clot in the right arm. He went to the hospital and under went surgery at 12 am on Saturday. The coronary surgeon mentioned that this was not from heart disease as all vessels were clear. Rather, this was due to an embolism due to an unknown origin. A further work up concluded that he also had blood clots throughout his body (both lungs, one kidney, heart, liver, legs, arms, and possibly brain). By 12 p.m. we were asked back to the hospital as he was worsening. Treatment options included a heparin drip, which caused a brain hemorrhage. He was just at the doctors two times this week, on Tuesday and Wednesday and was fine no issues found. They went through with the 2nd dose and came to our home on Thursday.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1301483-1	Blood clots in lungs. Had trouble breathing for 3 days before it was diagnosed in the er.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1302550-1	patient received 1st Moderna dose on 4/30/21. The next morning on 5/1 he felt pain on his left leg. He waited to see if would get better or worse. On the afternoon at an unknown time, his leg had gotten worse. He had severe pain on his left leg; it was painful to the touch; it was swollen and was hard as a rock. He called 911. He was taken to Hospital. Pt wanted me to report that he did not receive any treatment for his blood clot. He received a Lidocaine patch on his knee and was given a prescription for Voltaren Gel. (he was expecting to receive TPA to dissolve the clot like a few times before when he had blood clots but this particular time he did not)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1302589-1	After both shots of the vaccine my menstrual period was extremely heaving with golf ball size blood clots, extreme cramping and extremely heavy flows. The first period after my first shot lasted eight days compared to my usual five days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1303479-1	I am in menopause and had a cycle, lots of clotting and cramping like when I was a young woman. I have not had a cycle in 3 yeas.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1304176-1	Menstrual cycle abnormalities - no stop bleeding since first shot, became significantly worse with second shot. 2 months have passed and I still have excessive menstrual bleeding and very large menstrual blood clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1305011-1	"This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of ATRIAL FIBRILLATION (atrial fibrillation), PULMONARY EMBOLISM (blood clots in her lungs) and THROMBOSIS (blood clots everywhere) in an 89-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No medical history was provided by the reporter. Concomitant products included VITAMINS NOS for an unknown indication. On 16-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 17-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 17-Feb-2021, the patient experienced INFLUENZA LIKE ILLNESS (felt like she was getting the flu), DIARRHOEA (diarrhea), BLOOD PRESSURE ABNORMAL (no blood pressure), VACCINATION SITE PAIN (arm still hurts from time to time) and FATIGUE (felt tired). On 03-Mar-2021, the patient experienced ATRIAL FIBRILLATION (atrial fibrillation) (seriousness criterion hospitalization), PULMONARY EMBOLISM (blood clots in her lungs) (seriousness criterion hospitalization) and THROMBOSIS (blood clots everywhere) (seriousness criterion hospitalization). At the time of the report, ATRIAL FIBRILLATION (atrial fibrillation), PULMONARY EMBOLISM (blood clots in her lungs), THROMBOSIS (blood clots everywhere), INFLUENZA LIKE ILLNESS (felt like she was getting the flu), BLOOD PRESSURE ABNORMAL (no blood pressure) and FATIGUE (felt tired) outcome was unknown and DIARRHOEA (diarrhea) and VACCINATION SITE PAIN (arm still hurts from time to time) had not resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Computerised tomogram: normal (normal) haven't found anything. On an unknown date, Endoscopy: normal (normal) haven't found anything. Treatment medication included apixaban. Action taken with mRNA-1273 in response to the events was not Applicable. The patient had an endoscopy and CAT scan where they haven't found anything at when she was at the clinic. Consumer was vaccinated with first dose on 16JAN2021 (was not able to locate lot/expiration on vaccination card). She got her second dose on 17FEB2021 (was not able to locate lot/expiration on vaccination card) and she felt like she was getting the flu, felt tired and on 3MAR2021, she was rushed to the hospital because she had ""no blood pressure"" and presented atrial fibrillation and blood clots in her lungs and everywhere. Her arm still hurts from time to time and continues to have diarrhea. She states she is a fairly healthy person does not take any medications, only Centrum vitamins. Since then she has gone to the clinic, had an endoscopy and CAT scan where they haven't found anything. She is now on Eliquis. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked to MOD-2021-101609 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1306326-1	epiploic appendages; nausea; dizziness; blood clot; pain on left side of abdomen; This spontaneous case was reported by a nurse (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clot) in a 63-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. D17B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 19-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 30-Apr-2021, the patient experienced THROMBOSIS (blood clot) (seriousness criterion medically significant) and ABDOMINAL PAIN (pain on left side of abdomen). On 03-May-2021, the patient experienced DIZZINESS (dizziness) and NAUSEA (nausea). On an unknown date, the patient experienced EPIPLOIC APPENDAGITIS (epiploic appendages). At the time of the report, THROMBOSIS (blood clot), ABDOMINAL PAIN (pain on left side of abdomen), DIZZINESS (dizziness), EPIPLOIC APPENDAGITIS (epiploic appendages) and NAUSEA (nausea) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In April 2021, Computerised tomogram: epiploic appendagitis Epiploic Appendagitis. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Treatment given included antibiotic amoxicillin/clavulanic acid. Company Comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However, Very limited information regarding this events has been provided at this time. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However, Very limited information regarding this events has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1307025-1	Saturday 30 minutes after administration patient felt nausea lethargy and stiff neck and back. Next morning Sunday patient woke with a fever of 99.6 and large blood clot in mouth. Two days post vaccine Monday patient work with large blood clot on pillow. That evening patient had 2 more blood clots spontaneously coming from teeth. Tuesday morning upon arising from chair pt experienced an intense metallic taste. Patient then experienced several 8 mouth fulls of blood on 3 occasion in same evening. Patient states it was pushing out from gums.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1307671-1	pt states his right leg had pain and tightness from the inner thigh to the knee. It was painful to touch. Pt went to his PCP Dr. on 2/22/2021. He had blood work and ordered a sonogram for legs to check for blood clot. He was told to use a topical cream Voltaren which did not help. The sonogram showed that he had multiple blood clots on this inside of this leg down to his calf and was prescribed Xarelto. The blood clots disappeared after a couple of weeks.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1307687-1	on post vaccine day #9 pt developed light bruising r antecubital [vaccine was l deltoid] and knots. she has had ongoing mild nontraumatic bruising here and sq nodular lesions ? clot. pt also has had generalized malaise and some mild shortness of breath with this and has generally felt sick.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1308237-1	Presents to ED 4/21 with complaints of L side CP starting at approx 2100. Pt reports shooting pain to L side of chest and armpit with palpating area. Pt reports recently diagnosed with LLE blood clot on 4/13, put on eliquis and has been taking for the past 11 days. Pt reports resting makes pain better, palpating makes pain worse and standing makes pain worse in LLE. Pt denies SOB, DOB or cough. Pt denies history of PE, reports multiple bilat lower extrem DVT's. Patient CBC is normal, metabolic panel is normal, troponin is normal. CTA of the chest was obtained and shows no signs of pulmonary embolism. There is some prominence of his pulmonary arteries bilaterally. BNP is also negative. The patient's pain is mostly reproducible on that left chest, doubt ACS in this patient. Patient will be discharged home have him follow-up with his doctor as an outpatient. Patient is stable for discharge.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1309286-1	vertigo; Migraine type headaches; Flu like symptoms; Fever; Blood clot in right upper arm; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of VERTIGO (vertigo) and THROMBOSIS (Blood clot in right upper arm) in a 61-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 022M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 16-Feb-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 28-Feb-2021, the patient experienced VERTIGO (vertigo) (seriousness criterion hospitalization). On an unknown date, the patient experienced THROMBOSIS (Blood clot in right upper arm) (seriousness criterion medically significant), MIGRAINE (Migraine type headaches), INFLUENZA LIKE ILLNESS (Flu like symptoms) and PYREXIA (Fever). The patient was hospitalized on 28-Feb-2021 due to VERTIGO. At the time of the report, VERTIGO (vertigo), THROMBOSIS (Blood clot in right upper arm), MIGRAINE (Migraine type headaches), INFLUENZA LIKE ILLNESS (Flu like symptoms) and PYREXIA (Fever) outcome was unknown. No concomitant medications were reported. No corrective treatment was provided. 28-Feb-2021, he was hospitalized for vertigo for 3 days. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Very limited information regarding these events have been provided at this time. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Very limited information regarding these events have been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1310190-1	Blood clot in my eye causing a hemmorage and now I can't see out of lower part of my eye. Vision is expected to come back in 2-4 weeks. My lymph nodes in my armpits are swollen and I have exhaustion. I was sick and my body ached the day after but that was it.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1310621-1	Renal Infarct (blood clots to both kidneys)
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1311087-1	Blood Clot Right Calf
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1311396-1	Leg started swelling the day of the vaccine. Became painful and eventually became unable to move, according to patient. She was given a prescription for Xarelto,
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1312642-1	My mother had her first Moderna COVID-19 vaccination on March 1st. On March 10th, she suffered a blood clot in her eye that caused the eye to lose vision. Her eye doctor said she is now permanently legally blind in that eye.... I do not know if the blood clot was caused by the vaccine, but I am reporting it in case there is a correlation.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1312860-1	thought she was suffering from a stroke; anemia; double vertigo; blood clots, thick blood; dissociated; liver spazzed; slurring words; could not speak; possible underlying autoimmune disorder where genetic composition may have triggered this; malabsorption; looked very pale; heart disease; blood pressure was elevated, blood pressure was 190/130, uncontrolled and unspecified hypertension; severe vasculitis on the right side of brain; Thyroid calcification/thyroid had a lot of calcium build up; interstitial markings on lungs/doctor stated looked like COVID lungs; hands were so swollen that could not clench it; face was swollen; electrical shooting pain from the right brain stem of the neck; vision was blurry; stuttering; paranoid; confused; not able to recognize her daughter; hands were very stiff; stomach was bloated/looked 6 months pregnant; lethargic; bile dumps which then leads to inflammation of her intestines; clots the size of golf balls/soaked through a 24 count box of super tampons in 24 hours/period was very bright red; stomach was inflamed; period started 10 days early, which is completely abnormal for her menstrual cycle; severe ovarian pain that was up very high; could feel her hormones whooshing through her veins/foggy head/feeling funky/ felt stoned/ daydream dazes; irritable; migraine; stabbing pain; kidney pain; period pain; lightheaded; pimple in the back of her head; headache; chills; fatigue, tired; fever of 102.6F/fever which lasted another 12 hours; nausea; body aches, muscles started to ache; diarrhea; significant abdominal pain from the cramping due to heaving from the nausea; cramping/lot of cramping in the groin and thigh area; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of CEREBROVASCULAR ACCIDENT (thought she was suffering from a stroke), HYPERTENSION (blood pressure was elevated, blood pressure was 190/130, uncontrolled and unspecified hypertension), CENTRAL NERVOUS SYSTEM VASCULITIS (severe vasculitis on the right side of brain), AUTOIMMUNE DISORDER (possible underlying autoimmune disorder where genetic composition may have triggered this), THROMBOSIS (blood clots, thick blood) and INTERSTITIAL LUNG DISEASE (interstitial markings on lungs/doctor stated looked like COVID lungs) in a 33-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 030B21A and 010A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Gallbladder removal. On 03-Mar-2021, the

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				past medical history included Gallbladder removal. On 05-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 09-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 09-Apr-2021, the patient experienced MUSCLE SPASMS (cramping/lot of cramping in the groin and thigh area), ABDOMINAL PAIN (significant abdominal pain from the cramping due to heaving from the nausea), DIARRHOEA (diarrhea), PYREXIA (fever of 102.6F/fever which lasted another 12 hours), NAUSEA (nausea), MYALGIA (body aches, muscles started to ache), CHILLS (chills) and FATIGUE (fatigue, tired). On 10-Apr-2021, the patient experienced HEADACHE (headache). On 12-Apr-2021, the patient experienced ACNE (pimple in the back of her head). On 13-Apr-2021, the patient experienced MENSTRUAL DISORDER (period started 10 days early, which is completely abnormal for her menstrual cycle), ADNEXA UTERI PAIN (severe ovarian pain that was up very high), FEELING ABNORMAL (could feel her hormones whooshing through her veins/foggy head/feeling funky/ felt stoned/ daydream dazes), IRRITABILITY (irritable), MIGRAINE (migraine), PAIN (stabbing pain), RENAL PAIN (kidney pain), DYSMENORRHOEA (period pain), DIZZINESS (lightheaded), GASTRITIS (stomach was inflamed), GASTROINTESTINAL INFLAMMATION (bile dumps which then leads to inflammation of her intestines) and HEAVY MENSTRUAL BLEEDING (clots the size of golf balls/soaked through a 24 count box of super tampons in 24 hours/period was very bright red). On 14-Apr-2021, the patient experienced LETHARGY (lethargic). On 15-Apr-2021, the patient experienced MUSCULOSKELETAL STIFFNESS (hands were very stiff), ABDOMINAL DISTENSION (stomach was bloated/looked 6 months pregnant), PERIPHERAL SWELLING (hands were so swollen that could not clench it), SWELLING FACE (face was swollen), NECK PAIN (electrical shooting pain from the right brain stem of the neck), VISION BLURRED (vision was blurry), DYSPHEMIA (stuttering), PARANOIA (paranoid), CONFUSIONAL STATE (confused) and MEMORY IMPAIRMENT (not able to recognize her daughter). On 16-Apr-2021, the patient experienced CEREBROVASCULAR ACCIDENT (thought she was suffering from a stroke) (seriousness criterion hospitalization), HYPERTENSION (blood pressure was elevated, blood pressure was 190/130, uncontrolled and unspecified hypertension) (seriousness criterion hospitalization prolonged), CENTRAL NERVOUS SYSTEM VASCULITIS (severe vasculitis on the right side of brain) (seriousness criterion medically significant), INTERSTITIAL LUNG DISEASE (interstitial markings on lungs/doctor stated looked like COVID lungs) (seriousness criterion medically significant), PALLOR (looked very pale), CARDIAC DISORDER (heart disease) and THYROID CALCIFICATION (Thyroid calcification/thyroid had a lot of calcium build up). On 19-Apr-2021, the patient experienced AUTOIMMUNE DISORDER (possible underlying autoimmune disorder where genetic composition may have triggered this) (seriousness criterion medically significant) and MALABSORPTION (malabsorption). On an unknown date, the patient experienced THROMBOSIS (blood clots, thick blood) (seriousness criteria disability and medically significant), ANAEMIA (anemia), VERTIGO (double vertigo), DISSOCIATION (dissociated), LIVER DISORDER (liver spazzed), DYSARTHRIA (slurring words) and APHASIA (could not speak). The patient was hospitalized from 16-Apr-2021 to 17-Apr-2021 due to CEREBROVASCULAR ACCIDENT and HYPERTENSION. On 10-Apr-2021, PYREXIA (fever of 102.6F/fever which lasted another 12 hours) and CHILLS (chills) had resolved. On 12-Apr-2021, MUSCLE SPASMS (cramping/lot of cramping in the groin and thigh area), MYALGIA (body aches, muscles started to ache) and FATIGUE (fatigue, tired) had resolved. At the time of the report, CEREBROVASCULAR ACCIDENT (thought she was suffering from a stroke), CENTRAL NERVOUS SYSTEM VASCULITIS (severe vasculitis on the right side of brain), AUTOIMMUNE DISORDER (possible underlying autoimmune disorder where genetic composition may have triggered this), THROMBOSIS (blood clots, thick blood), INTERSTITIAL LUNG DISEASE (interstitial markings on lungs/doctor stated looked like COVID lungs), IRRITABILITY (irritable), PAIN (stabbing pain), DIZZINESS (lightheaded), MUSCULOSKELETAL STIFFNESS (hands were very stiff), ABDOMINAL DISTENSION (stomach was bloated/looked 6 months pregnant), GASTRITIS (stomach was inflamed), PERIPHERAL SWELLING (hands were so swollen that could not clench it), SWELLING FACE (face was swollen), NECK PAIN (electrical shooting pain from the right brain stem of the neck), VISION BLURRED (vision was blurry), DYSPHEMIA (stuttering), PARANOIA (paranoid), CONFUSIONAL STATE (confused), MEMORY IMPAIRMENT (not able to recognize her daughter), PALLOR (looked very pale), CARDIAC DISORDER (heart disease), MALABSORPTION (malabsorption), ANAEMIA (anemia), VERTIGO (double vertigo), THYROID CALCIFICATION (Thyroid calcification/thyroid had a lot of calcium build up), GASTROINTESTINAL INFLAMMATION (bile dumps which then leads to inflammation of her intestines), LETHARGY (lethargic), ACNE (pimple in the back of her head), DIARRHOEA (diarrhea), DISSOCIATION (dissociated), LIVER DISORDER (liver spazzed), DYSARTHRIA (slurring words), APHASIA (could not speak), NAUSEA (nausea) and HEADACHE (headache) outcome was unknown, HYPERTENSION (blood pressure was elevated, blood pressure was 190/130, uncontrolled and unspecified hypertension) and FEELING ABNORMAL (could feel her hormones whooshing through her veins/foggy head/feeling funky/ felt stoned/ daydream dazes) had not resolved, MENSTRUAL DISORDER (period started 10 days early, which is completely abnormal for her menstrual cycle), ADNEXA UTERI PAIN (severe ovarian pain that was up very high), RENAL PAIN (kidney pain), DYSMENORRHOEA (period pain), HEAVY MENSTRUAL BLEEDING (clots the size of golf balls/soaked through a 24 count box of super tampons in 24 hours/period was very bright red) and ABDOMINAL PAIN (significant abdominal pain from the cramping due to heaving from the nausea) had resolved and MIGRAINE (migraine) was resolving. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				if available): On 09-Apr-2021, Body temperature: 102.6 (High) fever of 102.6F. In April 2021, Neurological examination: abnormal (abnormal) neurological damage to the hippocampus of the central cranial. On 16-Apr-2021, Blood pressure measurement: 190/130 (abnormal) 190/130 and 175/99 (abnormal) 175/99 after treatment with Metoprolol. On 16-Apr-2021, Chest X-ray: interstitial markings of her lungs (abnormal) interstitial markings of her lungs. On 16-Apr-2021, Computerised tomogram: interstitial markings of her lungs (abnormal) interstitial markings of her lungs. On 16-Apr-2021, Magnetic resonance imaging: severe vasculitis on the right side of her brain (abnormal) severe vasculitis on the right side of her brain. On 16-Apr-2021, Neurological examination: 3/5 tests failed (abnormal) 3/5 tests failed. On 19-Apr-2021, Stool analysis: abnormal (abnormal) showed she was putting out too much potassium, magnesium. On 28-Apr-2021, Blood pressure measurement: 145/90 (abnormal) 145/90. On 28-Apr-2021, Heart rate: 90 (High) was not coming under 90 bpm. On 28-Apr-2021, Magnetic resonance imaging: blood clots, thick blood, anemia (abnormal) blood clots, thick blood, anemia. Action taken with mRNA-1273 in response to the events was not Applicable. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked to MOD-2021-095842 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1313667-1	Before vaccine went for sonogram, no signs of blood clot Vein doctor did more advanced sonogram and was negative again for clots. A few days later clots started appearing on varicose veins. First clot started inside of knee, thought it was a boil, it was a big red figure with a dark center, doctor diagnosed with superficial blood clots (about 10 days after vaccine) On the 10th when seeing doctor blood pressure shot up, had to double lisinopril. Patient reports history of smoking.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1314351-1	PATIENT CAME TO THE PHARMACY WITH SWOLLEN LIPS AND END UP HAVING ANAPHYLATIC REACTION AND BLOOD CLOT
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1314785-1	Less than 1 month after receiving vaccine, I experienced weakness and shortness of breath. Went to ER. Was hospitalized with multiple pulmonary embolisms in right lung. I am now on blood thinner Eliquis, and oxygen therapy which may be permanent
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1315834-1	had two very large blood cloths moved from her leg to her lung; blood clot was pressuring her heart; >42 days received the second vacicne; This spontaneous case was reported by a patient (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (had two very large blood cloths moved from her leg to her lung) and CARDIAC DISCOMFORT (blood clot was pressuring her heart) in a 77-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 16-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 08-Mar-2021, the patient experienced THROMBOSIS (had two very large blood cloths moved from her leg to her lung) (seriousness criteria hospitalization, medically significant and life threatening) and CARDIAC DISCOMFORT (blood clot was pressuring her heart) (seriousness criteria hospitalization and life threatening). On 06-Apr-2021, the patient experienced INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (>42 days received the second vacicne). The patient was hospitalized from 08-Mar-2021 to 12-Mar-2021 due to CARDIAC DISCOMFORT and THROMBOSIS. On 06-Apr-2021, INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (>42 days received the second vacicne) had resolved. At the time of the report, THROMBOSIS (had two very large blood cloths moved from her leg to her lung) and CARDIAC DISCOMFORT (blood clot was pressuring her heart) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No concomitant medications were reported. The patient was given Heparin in the hospital for about 2 days and was taking Apixaban at the time of this report. Very limited information regarding these events has been provided at this time. Further information required.; Sender's Comments: Very limited information regarding these events has been provided at this time. Further information required.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1315855-1	Went hiking and got a blood clot; beyond 36 days since first dose without receiving second dose; Interfered with work; Glands got super swollen; Trunk Rash; Excruciating headaches for three days; Extremely fatigued; High temperature; Nauseous; Ringing in ears; Taste buds changed; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Went hiking and got a blood clot) in a 48-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 017B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Grover's disease. Concurrent medical conditions included Allergy NOS (allergies for which she has an epi pen) and Premenopausal symptoms (Rashes with premenopausal condition). On 31-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 31-Mar-2021, the patient experienced TINNITUS (Ringing in ears), TASTE DISORDER (Taste buds changed), LOSS OF PERSONAL INDEPENDENCE IN DAILY ACTIVITIES (Interfered with work), LYMPHADENOPATHY (Glands got super swollen), RASH (Trunk Rash), HEADACHE (Excruciating headaches for three days), FATIGUE (Extremely fatigued), PYREXIA (High temperature) and NAUSEA (Nauseous). On 30-Apr-2021, the patient experienced INTENTIONAL PRODUCT USE ISSUE (beyond 36 days since first dose without receiving second dose). On 01-May-2021, the patient experienced THROMBOSIS (Went hiking and got a blood clot) (seriousness criterion medically significant). On 30-Apr-2021, INTENTIONAL PRODUCT USE ISSUE (beyond 36 days since first dose without receiving second dose) had resolved. At the time of the report, THROMBOSIS (Went hiking and got a blood clot), TINNITUS (Ringing in ears), TASTE DISORDER (Taste buds changed), LOSS OF PERSONAL INDEPENDENCE IN DAILY ACTIVITIES (Interfered with work), LYMPHADENOPATHY (Glands got super swollen), RASH (Trunk Rash), HEADACHE (Excruciating headaches for three days), FATIGUE (Extremely fatigued), PYREXIA (High temperature) and NAUSEA (Nauseous) was resolving. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, SARS-CoV-2 test: negative (Negative) Negative. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not reported. Treatment information was not provided. Company Comment Very limited information regarding these events has been provided at this time. The events are probably related to the patient's comorbidities. A causal relationship cannot be excluded for the events of rash, fatigue, pyrexia, lymphadenopathy and headache.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1318546-1	Blood clot formed.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1321140-1	My mom, got her first Moderna shot on Feb. 4th, 2021 and developed trombosis in the legs about a week later. She went to the doctor, who at first didn't want to see her and told her to elevate her leg. Pt. had a scheduled eye appointment on Feb. 12th with an other doctor and when she asked if she had any other issues, pt. showed her her leg. The eye doctor immediately told her to go to urgent care. When she did she received blood thinners and started injections in her stomach for 5 days. Feb. 18th pt. suffered a mini stroke and went to the hospital emergency room. She was transferred over to another facility in the afternoon and kept overnight. She had a phone appointment with the Dr. on Feb. 24th and on the 26th she received a heart monitor to wear for 2 weeks. She also had a check up about her leg, which was still swollen. On March 4, pt. received her second Moderna shot and she was feverish and very tired for days and weeks after. By March 18th her condition didn't improve at all and in fact she was felling really bad over night, so on March 19th she was admitted again to the hospital. She was monitored and kept over night, but during the early morning she suffered a stroke and a heart attack and passed way the next day.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1321295-1	patient said 4 to 5 days after the vaccine, she went to the hospital and had blood clots in the lungs and legs. she said she's ok now and dr told her to get her second vaccine today. she is now on Eliquis.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1321426-1	blood clots in both legs were detected on 5/11/2021, 18 days after receiving the 2nd dose of Moderna vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1321678-1	blood clot, stroke
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1321745-1	He suffered a blot clot on may 11th and a stroke.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1327562-1	Right after injection, I got really hot and flushed. I thought I am not a fan of needles - so I thought it was anxiety. My throat got really dry. They gave me a Dixie cup of water and I asked for more. After I had finished the one cup, I was clearing my throat and it felt like I couldn't swallow and my tongue had swollen up and I couldn't swallow any water from next Dixie cup. I couldn't swallow - it felt like it would get stuck at the back of my throat. The nurse said she was going to stab me with the epi-pen but had me lay down first and my face started to feel slightly numb and I told her that and she stabbed me in right hip with the epi-pen. They called the ambulance to take me to hospital and after the epi-pen was given they had me sit up and put a mask on my face and I was in a state of panic. I wore an oxygen full face mask and was breathing into it. The paramedics came in and put me on stretcher - monitored for six hours. Because of the adrenaline from the epi- pen - I had high blood pressure and heart rates - everything was really high. After a few hours, she told me that my tongue was looking better and there was better color in my tongue. I was sitting there every six hours - I was monitoring as how to I was doing. They gave me Benadryl and I think they gave me something else but I can't remember what it was. I came home and it was late at night - 11:00 or 11:30 and from there until February 1st, I was extremely sick. I went back to ER on 23rd; and did telehealth visits - with my doctor (right after ER visit from Injection was first one; and then I went to the hospital on the 23rd; Telehealth on 24th or 25th and I - I was running fevers of 104, 103 and went up to 105 almost- on 23rd of January - 3 days after injection - I'm weak, I can't move - everything hurt. From head to toe. I couldn't sleep or do anything. I was so weak, I couldn't even move. It was difficult just to use the bathroom. I had the fevers that was causing my chills and my body aches were bad. I had been trying to alternate Tylenol and Ibuprofen; I was finally prescribed Hydro Codone just to sleep. I was told to take this during the day and the Tylenol and Ibuprofen combo at night. Shoulder surgery was about a month after the vaccine - 18th of February - And on the 22nd of February I went to the hospital again for left side chest pains. - I couldn't breathe - I was discharged February 25th - I was on blood thinners for three months. - the surgeon said that I developed clots and I developed these shortly after surgery and ended up in the hospital for three days. The clots are unknown - blood clots started in the legs and then developed the fluid/clot issue in the lungs - I had a pulmonary embolism in my left lung. Surgeon said it could have been caused by vaccine as he did not work on the lower part of my body.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1327655-1	4/24-4/28/2021 pain in left leg hurts to walk or put weight on it 4/29/2021 swelling begins in left lower leg and foot 4/30/2021- 5/4/2021 swelling continues to worsen left leg, can't walk, can't move or feel toes, purple discoloration of foot 5/5/2021 go to Emergency Room Admitted into hospital due to extensive blood clot in left leg 5/6/2021 Surgery for removal of blood clot with Dr. 3 hour procedure 70 percent removed 5/7/2021 Released with blood thinner med 5/7/2021 revisit emergency room due to pain swelling and numbness moving into left upper leg and thigh 5/8-512/2021 swelling continues to progress 5/13/2021 follow up at vascular Dr. Readmitted into hospital 5/13 surgery for removal of clot Not all removed 5/16/2021 released from hospital on blood thinner
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1329170-1	Small Pulmonary Embolism; blood clots in upper arm, arm pit, neck, across right side of chest; Opposite arm blew up like a baloon; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PULMONARY EMBOLISM (Small Pulmonary Embolism) and THROMBOSIS (blood clots in upper arm, arm pit, neck, across right side of chest) in a 45-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 18-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-Apr-2021, the patient experienced PULMONARY EMBOLISM (Small Pulmonary Embolism) (seriousness criteria medically significant and life threatening), THROMBOSIS (blood clots in upper arm, arm pit, neck, across right side of chest) (seriousness criteria medically significant and life threatening) and PERIPHERAL SWELLING (Opposite arm blew up like a baloon). At the time of the report, PULMONARY EMBOLISM (Small Pulmonary Embolism), THROMBOSIS (blood clots in upper arm, arm pit, neck, across right side of chest) and PERIPHERAL SWELLING (Opposite arm blew up like a baloon) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant medication information was provided by the reporter. Treatment medications for the events included Xarelto (rivaroxaban). Reporter states patient went to Urgent Care on 22-Apr-2021 due to the events and was then told to go to the Emergency Room. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1329172-1	multiple subsegmental pulmonary emboli; blood clot; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PULMONARY EMBOLISM (multiple subsegmental pulmonary emboli) and THROMBOSIS (blood clot) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced PULMONARY EMBOLISM (multiple subsegmental pulmonary emboli) (seriousness criterion medically significant) and THROMBOSIS (blood clot) (seriousness criterion medically significant). At the time of the report, PULMONARY EMBOLISM (multiple subsegmental pulmonary emboli) and THROMBOSIS (blood clot) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant and treatment medications were provided. Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1331012-1	"Per client, ""I received the second dose of Moderna in my right arm on 03/19/2021. The next morning on 03/20/2021 around 9:00 AM, I noticed my left arm swelling. I got to feeling real bad. On 03/25/2021, I was feeling so bad that I went to Family Medicine to see my doctor, MD. He sent me straight to the Emergency room. They gave me the clot buster for my left arm. I was in there for 3 or 4 days. I was discharged. I have blood clots in my lungs. I had a follow up appointment. I also started seeing a new doctor, a pulmonologist, for the blood clots in my lungs. I lost a lot of weight, 32 pounds."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1332504-1	possibility of a blood clot; Legs were numb; Rash on arm; Whole arm was totally swollen; whole arm Itchy; Fell down; weakness post-vaccination; fever; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (possibility of a blood clot), HYPOAESTHESIA (Legs were numb), RASH (Rash on arm) and PERIPHERAL SWELLING (Whole arm was totally swollen) in a 33-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 3020B1A and 044BZ1A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 01-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 30-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 30-Apr-2021, the patient experienced THROMBOSIS (possibility of a blood clot) (seriousness criteria hospitalization and medically significant), HYPOAESTHESIA (Legs were numb) (seriousness criterion hospitalization), RASH (Rash on arm) (seriousness criterion hospitalization), PERIPHERAL SWELLING (Whole arm was totally swollen) (seriousness criterion hospitalization), PRURITUS (whole arm Itchy), ASTHENIA (weakness post-vaccination) and PYREXIA (fever). 30-Apr-2021, the patient experienced FALL (Fell down). The patient was hospitalized on 30-Apr-2021 due to HYPOAESTHESIA and PERIPHERAL SWELLING. At the time of the report, THROMBOSIS (possibility of a blood clot), HYPOAESTHESIA (Legs were numb), RASH (Rash on arm), PERIPHERAL SWELLING (Whole arm was totally swollen), PRURITUS (whole arm Itchy), FALL (Fell down), ASTHENIA (weakness post-vaccination) and PYREXIA (fever) outcome was unknown. No concomitant medications were reported. Treatment for the events included IV antibiotics, medication to stop itching, oral cephalexin, and Tylenol. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked to MOD-2021-121103 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1333233-1	large blood clot to lower extremity
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1333360-1	"5-19-21 Patient self-reported to that 8 days after receiving his first dose of Moderna (3/18/21), he began coughing up blood, was taken to the hospital, transferred to the ICU with thrombosis. He states he was in the ICU for 6 days. He then received second dose on 04/09/21 and was ill for 5 days again. States he currently remains fatigued, short of breath and ""just not himself."" Patient is concerned that the blood clots were related to his COVID Vaccines."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1333782-1	Extreme depression after vaccine Delayed period, spotty dark blood, very large blood clots and heavy cramping for over 3 days

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1334600-1	"Patient passed away on 4/30/21. Wife reports that on 4/29/21, she found him in the garage in a ""pool of blood, face down"" after a fall. Upon finding him on 4/29/21, she took him to the emergency room where he had a CT scan. Patient was diagnosed with a ""concussion"" and sent back home, and was instructed to follow-up with his primary care provider on 4/30/21. On 4/30/21, patient was ""sleepy all day."" On his way out of the house around 3:30, he suddenly put his hand on his wife and proceeded to collapse. The wife also fell, and by the time she gathered herself and went to check on her husband, he had passed away. An ambulance came to the house; CPR was performed for 40 minutes but ""they could not get him back."" He was not transported to a medical center. An autopsy was not done, but the coroner informed patient's wife that he likely died from a ""clot, maybe a PE."" Wife wonders how this could be, since he was on Plavix for 5 years. Wife does not feel that the COVID vaccine is to blame, but thought she should report the issue just in case. This writer is assisting wife of patient to fill out electronic report."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1334847-1	The morning of April 13,2021, I was taken to the emergency room with heart attack like symtems. After blood test a CT was done and multipul blood colts were found in both my lungs. I was then placed on blood thinners a low salt diet. I remained in the hospital until the April 16,2021.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1338628-1	Although I have heart disease, and had triple bypass surgery three years ago, I consistently take my blood pressure. My blood pressure has been in a normal range since the operation, and following receiving the first dose of the vaccine it became elevated and I needed to go back on an old medication. Even then, my blood pressure was erratic. Two weeks after my 2nd COVID shot I had a heart attack due to a blood clot. This clot was able to be removed by a heart cath-lab. This could be a coincidental flare up due to an old issue, but the timing aligns with both doses of the vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1340276-1	This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clot in each lung) in a 67-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 039K20A and 029L20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included CALCIUM FRUCTOBORATE, CHONDROITIN SULFATE SODIUM, GLUCOSAMINE HYDROCHLORIDE, HYALURONIC ACID (MOVE FREE JOINT HEALTH) for Joint disorder NOS, MINERALS NOS, VITAMINS NOS (ONE A DAY [MINERALS NOS;VITAMINS NOS]) for an unknown indication. On 21-Jan-2021 at 11:00 AM, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 18-Feb-2021 at 9:30 AM, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 28-Feb-2021, the patient experienced THROMBOSIS (blood clot in each lung) (seriousness criteria medically significant and life threatening). On 28-Feb-2021 at 6:00 PM, the patient experienced DYSPNOEA (shortness of breath). At the time of the report, THROMBOSIS (blood clot in each lung) outcome was unknown and DYSPNOEA (shortness of breath) had resolved with sequelae. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 28-Feb-2021, Computerised tomogram: blood clot (abnormal) Blood clot in both lung. On 21Jan2021, patient had received the first dose of Moderna Covid-19 vaccine and about 4-5 days later, he began to noticed that his breathing was labored. At the time of the report, the patient did not think much about the labored breathing since he was older in age and thought it was due to exercise. Patient received second dose of Moderna COVID-19 Vaccine on 18-Feb-2021 and he reported that his breathing became more laboring where he needed to stop walking to catch his breath. He had 2 scan which revealed that he have a blood clot at each lung and he was prescribed Xarelto 15 mg twice a day, then he will begin Xarelto 20mg once a day on 20-MAY-2021. Patient is on Kirkland C 500mg as a concomitant medication as well. Most recent FOLLOW-UP information incorporated above includes: On 13-May-2021: Significant FU- outcome of the event.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1340294-1	Hospitalization involving some serious health conditions; blood clot removed from legs; second dose has passed over 28 days; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of HOSPITALISATION (Hospitalization involving some serious health conditions) and THROMBOSIS (blood clot removed from legs) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Cardiac transplant in 2004. Concomitant products included TACROLIMUS and MYCOPHENOLATE MOFETIL (MYCOPHENOLATE) for Prophylaxis against heart transplant rejection, APIXABAN (ELIQUIS) for Thrombosis prophylaxis. On 18-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced HOSPITALISATION (Hospitalization involving some serious health conditions) (seriousness criterion hospitalization), THROMBOSIS (blood clot removed from legs) (seriousness criterion medically significant) and PRODUCT DOSE OMISSION ISSUE (second dose has passed over 28 days). The patient was treated with Surgery (blood clot removed) for Thrombosis. At the time of the report, HOSPITALISATION (Hospitalization involving some serious health conditions) and THROMBOSIS (blood clot removed from legs) outcome was unknown and PRODUCT DOSE OMISSION ISSUE (second dose has passed over 28 days) had resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Treatment information not provided. It was reported that, the patient missed an appointment for his second dose due to hospitalization involving some serious health conditions. The patient was in the hospital for extended period and underwent surgery to get blood clot removed from legs during the hospitalization. Very limited information has been provided for the event at this time. However, details of hospitalization and compliance with anti-coagulant therapy is required for further evaluation. This report also This report refers to a case of product dose omission issue for mRNA-1273, lot # unknown with no associated AEs.; Sender's Comments: Very limited information has been provided for the event at this time. However, details of hospitalization and compliance with anti-coagulant therapy is required for further evaluation. This report also This report refers to a case of product dose omission issue for mRNA-1273, lot # unknown with no associated AEs.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1343572-1	Clotting that lead to a stroke. lost right side vision in both eyes
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1343580-1	Blood Clot after moderna vaccine
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1345811-1	blood clots in both lungs; blood clots in left leg; shortness of breath; after the administration her arm hurt really bad; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PULMONARY THROMBOSIS (blood clots in both lungs) and THROMBOSIS (blood clots in left leg) in a 51-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 037A21B) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 31-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 31-Mar-2021, the patient experienced VACCINATION SITE PAIN (after the administration her arm hurt really bad). On 15-Apr-2021, the patient experienced PULMONARY THROMBOSIS (blood clots in both lungs) (seriousness criteria hospitalization and medically significant) and THROMBOSIS (blood clots in left leg) (seriousness criteria hospitalization and medically significant). On an unknown date, the patient experienced DYSPNOEA (shortness of breath). The patient was hospitalized from 15-Apr-2021 to 18-Apr-2021 due to PULMONARY THROMBOSIS and THROMBOSIS. On 01-Apr-2021, VACCINATION SITE PAIN (after the administration her arm hurt really bad) had resolved. At the time of the report, PULMONARY THROMBOSIS (blood clots in both lungs), THROMBOSIS (blood clots in left leg) and DYSPNOEA (shortness of breath) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant medications were reported. The patient received Apixaban as a treatment medication for blood clots. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. Further information has been requested. .

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1345817-1	suspected blood clot; When walking experiences traumatic pain in legs; her legs went numb while at store; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (suspected blood clot) in a 73-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 12-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 16-Apr-2021, the patient experienced PAIN IN EXTREMITY (When walking experiences traumatic pain in legs) and HYPOAESTHESIA (her legs went numb while at store). On an unknown date, the patient experienced THROMBOSIS (suspected blood clot) (seriousness criterion medically significant). The patient was treated with IBUPROFEN (MOTRIN [IBUPROFEN]) for Pain, at a dose of unknown. On 16-Apr-2021, HYPOAESTHESIA (her legs went numb while at store) had resolved. At the time of the report, THROMBOSIS (suspected blood clot) outcome was unknown and PAIN IN EXTREMITY (When walking experiences traumatic pain in legs) had not resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. No relevant concomitant medications were reported. Company Comment: Limited information regarding the events has been provided at this time and a causal relationship cannot be excluded.; Sender's Comments: Limited information regarding the events has been provided at this time and a causal relationship cannot be excluded
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1345821-1	Blood clots around the heart, 70% blockage; Pneumonia; Can barely walk, need walker or somebody need to hold; Hardly talk; Barely sees; Brain stem stroke; This spontaneous case was reported by a consumer and describes the occurrence of BRAIN STEM STROKE (Brain stem stroke), THROMBOSIS (Blood clots around the heart, 70% blockage) and PNEUMONIA (Pneumonia) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 046B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Stent placement on 29-Apr-2021. On 28-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 28-Apr-2021, the patient experienced BRAIN STEM STROKE (Brain stem stroke) (seriousness criterion medically significant). On 29-Apr-2021, the patient experienced THROMBOSIS (Blood clots around the heart, 70% blockage) (seriousness criterion medically significant), PNEUMONIA (Pneumonia) (seriousness criterion medically significant), GAIT DISTURBANCE (Can barely walk, need walker or somebody need to hold), SPEECH DISORDER (Hardly talk) and VISION BLURRED (Barely sees). At the time of the report, BRAIN STEM STROKE (Brain stem stroke), THROMBOSIS (Blood clots around the heart, 70% blockage), PNEUMONIA (Pneumonia), GAIT DISTURBANCE (Can barely walk, need walker or somebody need to hold), SPEECH DISORDER (Hardly talk) and VISION BLURRED (Barely sees) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No concomitant medication, No treatment information provided. The patient had brain stem stroke after 5 hours of the vaccination. The next day, on 20Apr2021, the doctor found that patient had 70% blockage, blood clot around the heart and they have to put stent. Company comment: The event of pneumonia is of an infective etiology, hence a causal association is unlikely. However, based on strong temporal association between the remaining events and the administration of mRNA-1273, a causal relationship cannot be excluded.; Sender's Comments: The event of pneumonia is of an infective etiology, hence a causal association is unlikely. However, based on strong temporal association between the remaining events and the administration of mRNA-1273, a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1345830-1	blood clot was discovered in the left leg; excruciating pain in the left leg below the knee; area is swollen; Area of the left leg below the knee is red; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot was discovered in the left leg) in an 82-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 026B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 06-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form(s). On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 06-Apr-2021, the patient experienced THROMBOSIS (blood clot was discovered in the left leg) (seriousness criterion medically significant), PAIN IN EXTREMITY (excruciating pain in the left leg below the knee), PERIPHERAL SWELLING (area is swollen) and ERYTHEMA (Area of the left leg below the knee is red). At the time of the report, THROMBOSIS (blood clot was discovered in the left leg), PAIN IN EXTREMITY (excruciating pain in the left leg below the knee), PERIPHERAL SWELLING (area is swollen) and ERYTHEMA (Area of the left leg below the knee is red) had not resolved. The patient's concomitant medication was not reported. The information regarding the patient's treatment medication was not provided. Company comment: Based on the current available information and temporal association between the use of the product and the start date of these events, a causal relationship cannot be excluded. This case was linked to MOD-2021-116642 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of these events, a causal relationship cannot be excluded

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1345839-1	Spontaneous blood clots; very swollwn leg; Fever; Chills; leg being lightly swollen; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Spontaneous blood clots) in a 75-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 031B21A and 030A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included No adverse event (No reported medical history). Concomitant products included DONEPEZIL HYDROCHLORIDE (ARICEPT), EMPAGLIFLOZIN (JARDIANCE), LEVOTHYROXINE, LISINOPRIL, METFORMIN, PRAVASTATIN, METOPROLOL, ARIPIPRAZOLE (ABILIFY), VENLAFAXINE HYDROCHLORIDE (EFFEXOR), FENTANYL, APIXABAN (ELIQUIS) and TESTOSTERONE for an unknown indication, CEFIXIME (FLEXERIL [CEFIXIME]). On 24-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 24-Apr-2021, the patient experienced PERIPHERAL SWELLING (leg being lightly swollen), PYREXIA (Fever) and CHILLS (Chills). On 25-Apr-2021, the patient experienced PERIPHERAL SWELLING (very swollwn leg). On an unknown date, the patient experienced THROMBOSIS (Spontaneous blood clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Spontaneous blood clots), PYREXIA (Fever) and CHILLS (Chills) outcome was unknown and PERIPHERAL SWELLING (leg being lightly swollen) and PERIPHERAL SWELLING (very swollwn leg) had not resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In 2021, Ultrasound scan: blood clots blood clots. On 25-APR-2021, patient experienced a very swollen leg and was scheduled an ultrasound which reveled blood clots on an unspecified date in 2021. No treatment information was provided. Action taken with mRNA-1273 in response to events was not applicable. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. For the event Thrombosis concomitant medication interactions are a confounder.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. For the event Thrombosis concomitant medication interactions are a confounder.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1346923-1	prolonged menstrual period, excessive blood clots on menses
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1347141-1	Heavy menstrual cycle bleeding; discharged was gushy and with clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1347435-1	Feb. 26-2 days after 1st shot, unconscious, blood sepsis, nausea, on a ventilator, severe pneumonia in right lung. In ICU for 3 days, normal ward 3 days, treated with antibiotics Mar. 26-2 days after 2nd shot, unconscious, blood sepsis, blood clot, heart failure, kidney & liver damage, treated with Eliquis and antibiotics. At hospital in ICU for 2 days, normal ward 3 days. Continuing with Eliquis, increased Metoprolol. Follow-up with kidney specialist and Cardiologist PA.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1347759-1	THROMBOSIS-THROMBOCYTOEPNIA SYNDROME--- LEFT LEG DVT
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1349111-1	"Dizziness (After 2nd Vaccine); Labored Breathing (After 2nd Vaccine); Low Blood Pressure (After 2nd Vaccine); Multiple Blood Clots Red area under the injection site, Symptoms were returning (After 2nd Vaccine); Spider Veins going from Arm to Chest (After 2nd Vaccine); Hot to the touch left Arm (After 2nd Vaccine); Protruding Red Veins on Left Arm (After 2nd Vaccine); Purple Swollen entire left arm from under the Injection Site to fingers (After 2nd Vaccine); Red band (3 inches wide and 4-5"" long) Below the Injection Site (After 2nd Vaccine); Swollen 1/2 inch below the Injection Site; Fever (After 2nd Vaccine); Chills (After 2nd Vaccine); Left Arm Hurt (After 2nd Vaccine); This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Multiple Blood Clots Red area under the injection site, Symptoms were returning (After 2nd Vaccine)), PERIPHERAL SWELLING (Purple Swollen entire left arm from under the Injection Site to fingers (After 2nd Vaccine)), SPIDER VEIN (Spider Veins going from Arm to Chest (After 2nd Vaccine)), DIZZINESS (Dizziness (After 2nd Vaccine)), DYSPNOEA (Labored Breathing (After 2nd Vaccine)), HYPOTENSION (Low Blood Pressure (After 2nd Vaccine)), SKIN WARM (Hot to the touch left Arm (After 2nd Vaccine)), PAIN IN EXTREMITY (Left Arm Hurt (After 2nd Vaccine)), SUPERFICIAL VEIN PROMINENCE (Protruding Red Veins on Left Arm (After 2nd Vaccine)), VACCINATION SITE ERYTHEMA (Red band (3 inches wide and 4-5"" long) Below the Injection Site (After 2nd Vaccine)), VACCINATION SITE SWELLING (Swollen 1/2 inch below the Injection Site), PYREXIA (Fever (After 2nd Vaccine)) and CHILLS (Chills (After 2nd Vaccine)) in a 49-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 045B21A and 017B21A) for COVID-19 vaccination. Concurrent medical conditions included Ulcerative colitis. Concomitant products included MESALAZINE (LIALDA), CETIRIZINE HYDROCHLORIDE (ZYRTEC [CETIRIZINE HYDROCHLORIDE]), MULTIVITAMIN [VITAMINS NOS] and VITAMIN C [ASCORBIC ACID] for an unknown indication. On 27-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 24-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Vaccine) (Intramuscular) dosage was changed to 2 dosage form. On 25-Apr- **Adverse Event Description**
				2021, the patient experienced PAIN IN EXTREMITY (Left Arm Hurt (After 2nd Vaccine), PYREXIA (Fever (After 2nd Vaccine)), and CHILLS (Chills (After 2nd Vaccine)) . On 26-Apr-2021, the patient experienced VACCINATION SITE ERYTHEMA (Red band (3 inches wide and 4-5"" long) Below the Injection Site (After 2nd Vaccine), and VACCINATION SITE SWELLING (Swollen 1/2 inch below the Injection Site) .On 27-Apr-2021, the patient experienced PERIPHERAL SWELLING (Purple Swollen entire left arm from under the Injection Site to fingers (After 2nd Vaccine)). On 28-Apr-2021, the patient experienced SPIDER VEIN (Spider Veins going from Arm to Chest (After 2nd Vaccine)), SKIN WARM (Hot to the touch left Arm (After 2nd Vaccine)) and SUPERFICIAL VEIN PROMINENCE (Protruding Red Veins on Left Arm (After 2nd Vaccine)). On 02-May-2021, the patient experienced THROMBOSIS (Multiple Blood Clots Red area under the injection site, Symptoms were returning (After 2nd Vaccine)) (seriousness criteria medically significant), DIZZINESS (Dizziness (After 2nd Vaccine)) DYSPNOEA (Labored Breathing (After 2nd Vaccine)) and HYPOTENSION (Low Blood Pressure (After 2nd Vaccine)). On 02-May-2021, DIZZINESS (Dizziness (After 2nd Vaccine)) and HYPOTENSION (Low Blood Pressure (After 2nd Vaccine)) outcome was unknown, DYSPNOEA (Labored Breathing (After 2nd Vaccine)) had resolved. On 14-May-2021, PYREXIA (Fever (After 2nd Vaccine)) and CHILLS (Chills (After 2nd Vaccine)) had resolved. At the time of the report, THROMBOSIS (Multiple Blood Clots Red area under the injection site, Symptoms were returning (After 2nd Vaccine)), PERIPHERAL SWELLING (Purple Swollen entire left arm from under the Injection Site to fingers (After 2nd Vaccine)), SPIDER VEIN (Spider Veins going from Arm to Chest (After 2nd Vaccine)), SKIN WARM (Hot to the touch left Arm (After 2nd Vaccine)), PAIN IN EXTREMITY (Left Arm Hurt (After 2nd Vaccine)), SUPERFICIAL VEIN PROMINENCE (Protruding Red Veins on Left Arm (After 2nd Vaccine)), VACCINATION SITE ERYTHEMA (Red band (3 inches wide and 4-5"" long) Below the Injection Site (After 2nd Vaccine)) and VACCINATION SITE SWELLING (Swollen 1/2 inch below the Injection Site) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 02-May-2021, Blood pressure measurement: low (Low) low. On 10-May-2021, Full blood count: wbc very high (High) WBC very high. On an unknown date, Full blood count: normal (normal) normal. On an unknown date, Ultrasound scan: multiple blood clots in the red area (abnormal) Multiple Blood Clots in the Red area under the injection site. The patient did not have a medical history of blood clots. On 25APR2021 the patient experienced fever, chills, and her arm hurt. On 26APR2021 her left arm hurt really bad and was swollen 1/2 inch below the injection site (3 inches wide and 4-5"" long) and also had a red band wrapped around the left arm below the injection site. On 27Apr2021 the patient's entire arm from under the Injection site to her fingers became swollen and purple. The patient visited the emergency room (ER) and was told to elevate it and place ice on the area and was prescribed Advil. On 28APR2021 the patients veins were protruding, were red and hot to the touch and she experienced spider veins started going to her chest. On 30Apr2021, she went back to the ER . The doctor gave her a steroid and suggested an ultrasound. 02MAY2021 she started experiencing labored breathing, dizziness, and low blood pressure. She visited the ER again. The doctor performed an ultrasound, an EKG, a chest x-ray, and a hot scan of the heart. The ultrasound of the left arm showed multiple blood clots in the red area under the injection site. The doctor started her on Eliquis. On 10May2021 the symptoms were returning. A complete blood count (CBC) showed white blood cells (WBC) were very high but a second blood level showed, on an unknown date showed normal WBC. On 14May2021, the swelling on the arm had come down and her arm was still purple but a lighter color purple with dark protruding veins and spider veins still visible. The patient continued taking Eliquis. Most recent FOLLOW-UP information incorporated above includes: On 14-May-2021: Significant F/U Case & Email ID Add. Conmeds., medical history, Lab data, Product dsetails, iNarrative updated; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1349112-1	Embolism arterial; Thrombosis; Herpes zoster; Limb discomfort; Pain in extremity; Feeling hot; This case was received via VAERS on 11-May-2021 and was forwarded to Moderna on 11-May-2021. This regulatory authority case was reported by an other health care professional (subsequently medically confirmed) and describes the occurrence of EMBOLISM ARTERIAL (Embolism arterial), THROMBOSIS (Thrombosis), HERPES ZOSTER (Herpes zoster), LIMB DISCOMFORT (Limb discomfort), PAIN IN EXTREMITY (Pain in extremity) and FEELING HOT (Feeling hot) in a 71-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 030A21A) for COVID-19 vaccination. Concurrent medical conditions included COPD and Asthma. Concomitant products included ALBUTEROL [SALBUTAMOL], BUDESONIDE, FORMOTEROL FUMARATE (SYMBICORT) and TIOTROPIUM BROMIDE MONOHYDRATE (SPIRIVA) for an unknown indication. On 24-Mar-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) at an unspecified dose. On 31-Mar-2021, the patient experienced EMBOLISM ARTERIAL (Embolism arterial) (seriousness criteria hospitalization and medically significant), THROMBOSIS (Thrombosis) (seriousness criteria hospitalization and medically significant), HERPES ZOSTER (Herpes zoster) (seriousness criterion hospitalization), LIMB DISCOMFORT (Limb discomfort) (seriousness criterion hospitalization), PAIN IN EXTREMITY (Pain in extremity) (seriousness criterion hospitalization) and FEELING HOT (Feeling hot) (seriousness criterion hospitalization). At the time of the report, EMBOLISM ARTERIAL (Embolism arterial), THROMBOSIS (Thrombosis), HERPES ZOSTER (Herpes zoster), LIMB DISCOMFORT (Limb discomfort), PAIN IN EXTREMITY (Pain in extremity) and FEELING HOT (Feeling hot) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 31-Mar-2021, Full blood count: unknown (Inconclusive) Unknown. On 31-Mar-2021, Ultrasound Doppler: unknown (Inconclusive) Unknown. On 31-Mar-2021, Whole body scan: unknown (Inconclusive) Unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. The patient went to hospital because of blood clots and he had an emergency surgery. It was a huge arterial clot and developed shingles. Action taken with mRNA-1273 in response to the events was not applicable. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1350390-1	"Patient received first dose Moderna vaccine on 3/31/2021. Approximately two weeks after receiving the first Moderna vaccine patient states he started to have pain in the lower leg which was not normal. Pain increased to a point that patient seeked care and was evaluated for a blood clot which he was told was ""negative"" as no clot was initially found. On 4/28/2021 Patient presented to pharmacy to receive second dose. Pharmacist asked how patient tolerated first dose to which he replied he had no issues, so the second dose was administered. AFTER the second dose, patient did state he was seen by a CNP for leg pain but a clot was ruled out. Approximately 3 weeks after the second dose of Moderna vaccine the patients leg pain increased to a point that he presented to the local ER and it was determined at that visit that he did indeed have a large blood clot in the leg and a second smaller clot in the upper thigh. He was sent home on Eliquis. He subsequently presented to another ER approximately one week later for uncontrolled leg pain due to massive size of blood clot in leg. At this point, patient is at home recovering."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1350684-1	Patient seen in ER with pulmonary embolism , (L) lower extremity blood clot
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1350784-1	High fever of 104.3 Heart Palpitations with chest pain, shaking/chills throughout body and splitting headache (not migraine). Heavy mentrual bleeding with clots for long periods of time
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1351106-1	Heavy menstrual bleeding (menorrhagia), massive clots followed by bleeding, ongoing 11 days. Abnormally high compared to any menstrual cycle in entire life.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1351921-1	"""Moderna COVID-19 Vaccine. Vaccine on 2/25. Took him to an urgent care on 3-7 because of swollen ankle, foot and lower leg. Had xray but nothing was broken. thought he had sprained ankle. Given prednisone. Ankle continued to be swollen for following days so we followed up with family dr. on 3-22 who sent him for ultrasound which showed an extensive blood clot - DVT. Sent to emergency room. Put on blood thinner shots for a week and sent to doctor on 3-24. He did many blood tests. He started eliquis on 3-30. Saw doctor again on 4-13. His blood tests had shown he has Factor 5--Leiden. On May 11 we saw doctor again. Remain on Eliquis. Another ultrasound and more bloodtests show clot is chronic, but blood thinner is doing its job. No active clotting."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1352139-1	Great difficulty breathing, diagnosed with pulmonary embolism and one clot in right leg on May 15. First went to hospital with extreme breathing difficulty on March 15 but no diagnosis was made. this was one month after first Moderna vaccination. Two months later, with same issues (also extremely swollen ankles) and same tests, the embolisms were seen on CT test with contrast.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1353068-1	Passed out multiple times; Blood clots; This spontaneous case was reported by a consumer and describes the occurrence of LOSS OF CONSCIOUSNESS (Passed out multiple times) and THROMBOSIS (Blood clots) in a 32-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 047a21a) for COVID-19 vaccination. No Medical History information was reported. On 22-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 31-Mar-2021, the patient experienced LOSS OF CONSCIOUSNESS (Passed out multiple times) (seriousness criterion medically significant) and THROMBOSIS (Blood clots) (seriousness criterion medically significant). At the time of the report, LOSS OF CONSCIOUSNESS (Passed out multiple times) and THROMBOSIS (Blood clots) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not provided by the reporter. Treatment information was not provided. Patient was taken to hospital where it was discovered that he had blood clots that needed to be removed. Very limited information regarding the events has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding the events has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1353119-1	lower leg was swollen and started to get bigger and bigger; blood clot in the right leg; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clot in the right leg) in a 70-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 02AL20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No medical history were reported. Concomitant products included PANTOPRAZOLE for GERD, TRAZODONE for Sleeplessness, SIMVASTATIN and CHLORMEZANONE (RESTORIL [CHLORMEZANONE]) for an unknown indication. On 28-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 18-Feb-2021, after starting mRNA-1273 (Moderna COVID-19 Vaccine), the patient experienced THROMBOSIS (blood clot in the right leg) (seriousness criterion medically significant). On an unknown date, the patient experienced PERIPHERAL SWELLING (lower leg was swollen and started to get bigger and bigger). The patient was treated with RIVAROXABAN (XARELTO) for Thrombosis, at an unspecified dose and frequency. At the time of the report, THROMBOSIS (blood clot in the right leg) and PERIPHERAL SWELLING (lower leg was swollen and started to get bigger and bigger) outcome was unknown. Not Provided The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant medication includes Benapraxine for to simmer down. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1353773-1	Clot; Stroke; Swelling began in the groin / swelling extending up to my upper thigh / swelling extending to the back; Problem with walking; Got mad and walked out of hospital byhimself; Nauseous; Pins and needles sensation; Felt chilly; Dizzy; Whole body(left side) went numb; This spontaneous case was reported by a consumer and describes the occurrence of SWELLING (Swelling began in the groin / swelling extending up to my upper thigh / swelling extending to the back), THROMBOSIS (Clot) and CEREBROVASCULAR ACCIDENT (Stroke) in a 68-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 030A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 06-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 06-May-2021, the patient experienced SWELLING (Swelling began in the groin / swelling extending up to my upper thigh / swelling extending to the back) (seriousness criterion hospitalization), FEELING COLD (Felt chilly), DIZZINESS (Dizzy), HYPOAESTHESIA (Whole body(left side) went numb), PARAESTHESIA (Pins and needles sensation) and NAUSEA (Nauseous). On 07-May-2021, the patient experienced THROMBOSIS (Clot) (seriousness criteria hospitalization and medically significant), CEREBROVASCULAR ACCIDENT (Stroke) (seriousness criteria hospitalization and medically significant), GAIT DISTURBANCE (Problem with walking) and TREATMENT NONCOMPLIANCE (Got mad and walked out of hospital byhimself). The patient was hospitalized on 07-May-2021 due to CEREBROVASCULAR ACCIDENT and THROMBOSIS. At the time of the report, SWELLING (Swelling began in the groin / swelling extending up to my upper thigh / swelling extending to the back), THROMBOSIS (Clot), CEREBROVASCULAR ACCIDENT (Stroke), FEELING COLD (Felt chilly), DIZZINESS (Dizzy), HYPOAESTHESIA (Whole body(left side) went numb), GAIT DISTURBANCE (Problem with walking), PARAESTHESIA (Pins and needles sensation), TREATMENT NONCOMPLIANCE (Got mad and walked out of hospital byhimself) and NAUSEA (Nauseous) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 07-May-2021, Blood test: unknown (Inconclusive) Unknown. On 07-May-2021, Computerised tomogram: unknown (Inconclusive) Unknown. On 07-May-2021, Magnetic resonance imaging: unknown (Inconclusive) Unknown. On 07-May-2021, X-ray: unknown (Inconclusive) Unknown. Company comment: Although a temporal association exists, provided information is not adequate to assess the causal association between the events of thrombosis, stroke, swelling, hypoesthesia, paresthesia, gait disturbance and mRNA-1273. The detailed medical history and diagnostic report has not been provided. Based on the current available information and temporal association between the use of the product and the start date of the events of dizziness, feeling cold, nausea, a causal relationship cannot be excluded. The causality assessment for the event of Treatment noncompliance remains Not applicable. This case was linked to MOD-2021-133656 (Patient Link).; Sender's Comments: Although a temporal association exists, provided information is not adequate to assess the causal association between the events of thrombosis, stroke, swelling, hypoesthesia, paresthesia, gait disturbance and mRNA-1273. The detailed medical history and diagnostic report has not been provided. Based on the current available information and temporal association between the use of the product and the start date of the events of dizziness, feeling cold, nausea, a causal relationship cannot be excluded. The causality assessment for the event of Treatment noncompliance remains Not applicable.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1354626-1	with this 1st of 2 shots I began getting a headache that began within 1 hour after the shot. The headache immediately became intense and stayed. Advil did not do anything for it. That same evening I began having intense abdominal pain on my right side, it became so bad I was scared my appendix was going to burst. I had my son on standby to take me to the hospital. It was so severe I could not stand. The only way that I can describe the pain would be: after natural birth and you begin nursing, the insanely intense contractions of the uterus to shed the afterbirth is how intense the pain was. I then started my menstrual cycle but was not supposed to for another week and 1/2. The pain stayed, but did dull some, it wasn't as sharp but still intense. My flow was extraordinarily heavy and I lost count of how many huge blood clots I passed. Some clots were the size of my palm. This cycle lasted a full 7 days. My normal cycle starts with a little spotting, then 1 or 2 days of medium flow, then a couple days of spotting. I also do not have my cycle every 30 days, more like 38-40. I figure its pre-menopause.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1354753-1	Day after second vaccine blood clots in legs and lungs, mesh inserted, started bleeding vaginally and now in intensive care almost on life support.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1355341-1	As reported by NP that called with event: Pt has had 3 hospital stays after receiving vaccine. Pt initially went in on 3/31/2021. Stays include: 3/31/21-4/5/21 4/12/21-4/16/21 4/29/21-5/5/21 Symptoms/diagnoses include: IJ thrombosis, PE, esophageal strictures d/t nausea/vomiting
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1355401-1	About 3 weeks after receiving the first Moderna shot I developed, what i believed to be a severe charlie horse in my left lower leg that lasted about. week and disopated. A few weeks later May 9,2021; I noticed discomfort and swelling in the same lower left leg and my, medical alert service dog, kept alerting to it. I had a regular appointment with my cardioligist on May 10,2021 and advised of my issue and he ordered an ultrasound. The ultra sound revealed blood clots on my llower left leg. I have never had clotting issues in my life and absolutely believe that this colt was caused by the Moderna vaccine, since i have been on anticoagulants and asprin theray since 2008.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1355563-1	4/29/21: HAD BLURRED VISION (ALL DAY UNTIL MORNING OF NEXT DAY) + PRESSURE IN HEAD (ALL DAY INTO EVENING OF NEXT DAY) ABOUT 40 MINUTES AFTER RECEIVING VACCINE. 4/29/21: EVER SINCE VACCINATION, EXPERIENCING EXCESSIVE THIRST. ONGOING 5/1/21: PATIENT DESCRIBED SYMPTOM AS SKULL BONE PAIN (TENDER SCALP WHEN TOUCHED) UNTIL 5/5/21 5/3/21: ONGLING BLADDER IRRITATION. NO BLOOD IN URINE, BUT SOME SPOTTING (~2TSP) ON PADDING 5/17/21 . NO BLOOD PRESENT SINCE 5/4/21 MORNING: LYMPH NODE SWOLLEN UNDER THE LEFT ARM (INJECTION WAS IN RIGHT ARM). SIZE OF A TENNIS BALL, VERY PAINFUL, AND PAIN RADIATED TO AROUND THE LEFT BREAST AREA. PAIN LASTED UNTIL THE AFTERNOON OF 5/6/21. 5/9/21: BLOOD CLOT IN LEFT LEG REACTIVATED IN THE MORNING, ACCOMPANIED BY EXTREME PAIN, SWELLING, WARM TO THE TOUCH. PAIN AND SWELLING HAS BEEN MINIMAL BEGINNING 5/24/21, AND HAS NOT BEEN AS WARM TO THE TOUCH AS IT HAS BEEN. TREATMENTS: TOOK IBUPROFEN AND TYLENOL AS NEEDED FOR PAIN. USED ARNICA GEL ON THE ARM AT THE INJECTION SITE + MASSAGE+ CONSTANT MOVEMENT FOR PAIN/STIFFNESS. USED ARNICA GEL ALSO ON LYMPH NODE UNDER LEFT ARM ELEVATED CLOTTED LEG AND STAYED ACTIVE TO OPTIMIZE BLOOD FLOW, USED LIGHTLY HEATED PAD TO EASE THE LEG PAIN
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1355774-1	Already having blood clots in my left leg and
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1358033-1	2 days after second shot blood clot in left arm. Hit while walking in my home. Could not lift my arm. 5 days later heart attack. Pilot with EKG yearly. Last EKG less than one month from my heart attack on April 29, 2021
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1359991-1	Blood clots under skin; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots under skin) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots under skin) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots under skin) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. No concomitant medications reported by reporter. No treatment medications provided by the reporter. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1361434-1	Uterine hemorrhage, severe heavy menstrual with large clots
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1361440-1	Blood clot in right temporal lobe
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1361642-1	Deep Vein Thrombosis (DVT) in right lower leg. Symptoms initially occurred about two days after feeling sick with a high fever. Initially, the pain in the right calf muscle was thought to be a muscle or tendon strain. On May 22nd, during a campout, there was pain along the entire right leg, but at the time I thought it was just related to lower back pain from sleeping on the ground. On May 27 and 28th there was sharp pain at the bottom of my right foot, then on May 29th I noticed sweeling and a hard warm spot just above the inside right ankle. I reported to a local urgent care clinic first then was referred to Medical ER.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1362330-1	Spinal fusion on January 12, 2021 one week after first vaccine. Second vaccine on Feb 2, 2021. Developed blood clot in right calf & pulmonary embolism in right lung the week of Feb 15th. May have been post op problem but I had no issue until after the 2nd vaccine. Surgeon felt that post op pts usually get a blood clot within 2 weeks after surgery- that's why I feel it was from the 2nd vaccine.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1362340-1	After my terrible chills and fever were gone I got my period 2 weeks early. It was a normal period however only 5 days later I got ANOTHER period. This was was much heavier than normal with done quarter sized blood clots. It lasted 7 days.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1363975-1	Developed blood clot; Sever pain in his stomach; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Developed blood clot) and ABDOMINAL PAIN UPPER (Sever pain in his stomach) in an 83-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 032M20A and 016L20A) for COVID-19 vaccination. Concomitant products included RIVAROXABAN (XARELTO), AMLODIPINE, NEBIVOLOL HYDROCHLORIDE (BYSTOLIC) and AZILSARTAN MEDOXOMIL, CHLORTALIDONE (EDARBYCLOR) for an unknown indication. On 06-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 06-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 27-Apr-2021, the patient experienced ABDOMINAL PAIN UPPER (Sever pain in his stomach) (seriousness criterion hospitalization). On an unknown date, the patient experienced THROMBOSIS (Developed blood clot) (seriousness criteria hospitalization and medically significant). The patient was hospitalized from 28-Apr-2021 to 04-May-2021 due to ABDOMINAL PAIN UPPER and THROMBOSIS. The patient was treated with Surgery (leading to two and half feet of his small intestine removed) for Thrombosis. At the time of the report, THROMBOSIS (Developed blood clot) and ABDOMINAL PAIN UPPER (Sever pain in his stomach) outcome was unknown. Patient started having severe pain in his stomach on 27-Apr-2021. He was taken to the hospital on 28-Apr-2021, patient's doctor said that he had a clot in his intestine and that caused the pain, was operated removed to two and half feet of his small intestine. Patient was discharged from the hospital on 04-May-2021. The patient received both scheduled doses of mRNA-1273 prior to the events, therefore action taken with the drug in response to the events was not applicable. Company Comment : Very limited information regarding this events has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this events has been provided at this time. Further information has been requested.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1363977-1	Clots in left arm after taking 2 shots; Swollen arm after 2 vaccine shots; pain in left arm; 50 days gap between 2 vaccine doses; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Clots in left arm after taking 2 shots) in a 50-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Idiopathic thrombocytopenic purpura, Spina bifida and Psoriasis. On 06-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 26-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Clots in left arm after taking 2 shots) (seriousness criterion medically significant), PERIPHERAL SWELLING (Swollen arm after 2 vaccine shots), VACCINATION SITE PAIN (pain in left arm) and INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (50 days gap between 2 vaccine doses). The patient was treated with APIXABAN (ELIQUIS) at a dose of 25 mg twice a day and APIXABAN (ELIQUIS) at a dose of 5 mg twice a day. At the time of the report, THROMBOSIS (Clots in left arm after taking 2 shots), PERIPHERAL SWELLING (Swollen arm after 2 vaccine shots) and VACCINATION SITE PAIN (pain in left arm) outcome was unknown and INAPPROPRIATE SCHEDULE OF PRODUCT ADMINISTRATION (50 days gap between 2 vaccine doses) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Ultrasound scan: abnormal (abnormal) clot in the left arm. Patient went to emergency after 3 weeks of taking Moderna COVID-19 second Vaccine. No relevant concomitant medications were reported Reportedly, the treatment with Apixaban (Eliquis) needed to continue for 6 months. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1363983-1	Heart Attack; Panic attack; Couldnt breathe; Felt like chest caved in; Chest pain; Blood clot; Arm Pain; This spontaneous case was reported by a consumer and describes the occurrence of MYOCARDIAL INFARCTION (Heart Attack), PAIN IN EXTREMITY (Arm Pain), PANIC ATTACK (Panic attack), DYSPNOEA (Couldnt breathe), CHEST DISCOMFORT (Felt like chest caved in), CHEST PAIN (Chest pain) and THROMBOSIS (Blood clot) in a 34-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 048B21A) for COVID-19 vaccination. The patient's past medical history included Flu vaccination (received about 5 weeks ago before he got the Covid Shot and had chest pain) and Sinusitis (the patient stated that he violently threw up and he was sick for 2 weeks; the doctors then said that it may have been a stroke or a heart attack but he definitely had sinusitis.). Concurrent medical conditions included Diabetes, Cholesterol high and Blood pressure high. On 13-May-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 13-May-2021, the patient experienced PAIN IN EXTREMITY (Arm Pain) (seriousness criterion hospitalization). On 15-May-2021, the patient experienced PANIC ATTACK (Panic attack) (seriousness criterion hospitalization), DYSPNOEA (Couldnt breathe) (seriousness criterion hospitalization), CHEST DISCOMFORT (Felt like chest caved in) (seriousness criterion hospitalization), CHEST PAIN (Chest pain) (seriousness criterion hospitalization) and THROMBOSIS (Blood clot) (seriousness criteria hospitalization and medically significant). On 15-May-2021 at 4:00 PM, the patient experienced MYOCARDIAL INFARCTION (Heart Attack) (seriousness criteria hospitalization and medically significant). The patient was hospitalized for 3 days due to CHEST DISCOMFORT, CHEST PAIN, DYSPNOEA, MYOCARDIAL INFARCTION, PAIN IN EXTREMITY, PANIC ATTACK and THROMBOSIS. The patient was treated with ACETYLSALICYLIC ACID (BABY ASPIRIN) ongoing since an unknown date at an unspecified dose and frequency; LISINOPRIL ongoing since an unknown date at a dose of 5 mg; METOPROLOL SUCCINATE ongoing since an unknown date at a dose of 50 mg; NITROGLYCERIN ongoing since an unknown date at a dose of 10 mg as required; PRASUGREL ongoing since an unknown date at a dose of 10 milligram; SPIRONOLACTONE ongoing since an unknown date at a dose of 25 mg and ROSUVASTATIN ongoing since an unknown date at a dose of 80 milligram. At the time of the report, MYOCARDIAL INFARCTION (Heart Attack), PAIN IN EXTREMITY (Arm Pain), PANIC ATTACK (Panic attack), DYSPNOEA (Couldnt breathe), CHEST DISCOMFORT (Felt like chest caved in), CHEST PAIN (Chest pain) and THROMBOSIS (Blood clot) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant medications were not reported. The patient reported that he had a heart attack on 15 May 2021, after football practice Patient was in the hospital for 3 days. Treatment also included catheter placement and then a stint in heart. Company Comment : Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1363987-1	blood clot from ankle the groin; flu like symptoms; difficulty walking; soreness in Left arm; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clot from ankle the groin) in a 56-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 005C21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 10-May-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 10-May-2021, the patient experienced THROMBOSIS (blood clot from ankle the groin) (seriousness criterion medically significant), INFLUENZA LIKE ILLNESS (flu like symptoms), GAIT DISTURBANCE (difficulty walking) and MYALGIA (soreness in Left arm). The patient was treated with RIVAROXABAN (XARELTO) ongoing since an unknown date for Clot blood, at an unspecified dose and frequency. At the time of the report, THROMBOSIS (blood clot from ankle the groin), INFLUENZA LIKE ILLNESS (flu like symptoms), GAIT DISTURBANCE (difficulty walking) and MYALGIA (soreness in Left arm) outcome was unknown. No concomitant medications were provided. The patient went to the ER after experiencing the blood clot (from her ankle to the groin area) and was provided treatment with Xarelto. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1364001-1	pain in calf; Deep vein thrombosis; Blood Clot; This spontaneous case was reported by a physician (subsequently medically confirmed) and describes the occurrence of DEEP VEIN THROMBOSIS (Deep vein thrombosis) and THROMBOSIS (Blood Clot) in a 39-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 immunisation. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Deep vein thrombosis. On 07-May-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 13-May-2021, the patient experienced DEEP VEIN THROMBOSIS (Deep vein thrombosis) (seriousness criterion medically significant) and THROMBOSIS (Blood Clot) (seriousness criterion medically significant). On an unknown date, the patient experienced PAIN IN EXTREMITY (pain in calf). The patient was treated with APIXABAN (ELIQUIS) for Clot blood, at a dose of 1 dosage form. At the time of the report, DEEP VEIN THROMBOSIS (Deep vein thrombosis), THROMBOSIS (Blood Clot) and PAIN IN EXTREMITY (pain in calf) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 14-May-2021, Scan: blood clot (abnormal) blood clot. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Concomitant medications included unspecified anti-coagulant and antibiotic. Company comment Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1364019-1	clotting in body; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (clotting in body) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (clotting in body) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (clotting in body) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant medications were reported. No treatment details were provided. The adverse event was submitted. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1364028-1	Stroke; Blood clots; Brain bleed; Wife found him on the floor after he collapsed; Still struggling to move left hand; Felt really weird; Acting differently; Kept rubbing his arm and saying it was burning; Wife found him on the floor after he collapsed and fell off the couch; Unable to move or talk; Can't speak / unable to talk; Can not stand up; Arm hurting like crazy / arm was killing him; This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (Stroke), THROMBOSIS (Blood clots), CEREBRAL HAEMORRHAGE (Brain bleed), LOSS OF CONSCIOUSNESS (Wife found him on the floor after he collapsed) and HEMIPARESIS (Still struggling to move left hand) in a 54-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. unknown) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced CEREBROVASCULAR ACCIDENT (Stroke) (seriousness criteria hospitalization and medically significant), THROMBOSIS (Blood clots) (seriousness criteria hospitalization and medically significant), CEREBRAL HAEMORRHAGE (Brain bleed) (seriousness criterion medically significant), LOSS OF CONSCIOUSNESS (Wife found him on the floor after he collapsed) (seriousness criterion medically significant), HEMIPARESIS (Still struggling to move left hand) (seriousness criterion medically significant), FEELING ABNORMAL (Felt really weird), ABNORMAL BEHAVIOUR (Acting differently), BURNING SENSATION (Kept rubbing his arm and saying it was burning), FALL (Wife found him on the floor after he collapsed and fell off the couch), HYPOKINESIA (Unable to move or talk), APHONIA (Can't speak / unable to talk), DYSSTASIA (Can not stand up) and MYALGIA (Arm hurting like crazy / arm was killing him). At the time of the report, CEREBROVASCULAR ACCIDENT (Stroke), THROMBOSIS (Blood clots), CEREBRAL HAEMORRHAGE (Brain bleed), LOSS OF CONSCIOUSNESS (Wife found him on the floor after he collapsed), HEMIPARESIS (Still struggling to move left hand), FEELING ABNORMAL (Felt really weird), ABNORMAL BEHAVIOUR (Acting differently), BURNING SENSATION (Kept rubbing his arm and saying it was burning), FALL (Wife found him on the floor after he collapsed and fell off the couch), HYPOKINESIA (Unable to move or talk), APHONIA (Can't speak / unable to talk), DYSSTASIA (Can not stand up) and MYALGIA (Arm hurting like crazy / arm was killing him) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Concomitant product use was not provided by the reporter. It was reported that, the patient felt really weird and was acting differently. Four days after the patient received the vaccine, his wife found him on the floor after he collapsed and fell off the couch and was unable to move or talk. He was taken to hospital and was in ICU for 5-6 days. Treatment information was not provided. Very limited information regarding this event/s has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event/s has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1364809-1	Had shortness of breath and admitted to hospital for blood clot/pulmonary embolism on 4/28/21 * Hematologist suggested I should report for information purposes. Not stating this is believed to be the cause of the event - just reporting as requested
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1365171-1	Superficial Venous Thrombus
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1366867-1	blood clot in the left lower leg; left ankle swelling; left foot pain; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clot in the left lower leg) in a 70-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included METOPROLOL for Hypertension. On 26-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 23-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 26-Mar-2021, the patient experienced JOINT SWELLING (left ankle swelling) and PAIN IN EXTREMITY (left foot pain). On 12-May-2021, the patient experienced THROMBOSIS (blood clot in the left lower leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (blood clot in the left lower leg), JOINT SWELLING (left ankle swelling) and PAIN IN EXTREMITY (left foot pain) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 12-May-2021, Ultrasound scan: abnormal (abnormal) Diagnosed with blood clot in the left lower leg. Other concomitant medication include statin for cholesterol. Treatment medication for events included Xarelto. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (unknown) was not applicable. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1366871-1	blood clots; leg pain due to clots; chills; diarrhea worse than when I had covid; chest pains; nerve sensations up and down back and arms; blood pressure went up; higher fever than when got covid; headaches; This spontaneous case was reported by a patient and describes the occurrence of THROMBOSIS (blood clots) in a 57-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 011J20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included COVID-19 (Patient had Covid-19 last year in April and experienced blood clot.) in April 2020 and Thrombosis (Patient had Covid-19 last year in April and experienced blood clot.) in April 2020. Concurrent medical conditions included Hypothyroidism, Blood pressure and Allergy. Concomitant products included CETIRIZINE HYDROCHLORIDE (CETRIZINE) and FEXOFENADINE HYDROCHLORIDE (ALLEGRA) for Allergy, HYDROCHLOROTHIAZIDE and METOPROLOL for Blood pressure, LEVOTHYROXINE for Hypothyroidism, POTASSIUM for Potassium, VITAMIN B NOS, VITAMIN D NOS and ASCORBIC ACID, CALCIUM, MINERALS NOS, RETINOL, TOCOPHERYL ACETATE, VITAMIN B NOS, VITAMINS NOS, ZINC (CENTRUM SILVER [ASCORBIC ACID;CALCIUM;MINERALS NOS;RETINOL;TOCOPHERYL ACETATE;VITAMIN B NOS;VITAMINS NOS;ZINC]) for Vitamin supplementation, SAMBUCUS NIGRA (ELDERBERRY EXTRACT) for an unknown indication. On 19-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 25-Feb-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 25-Feb-2021, the patient experienced BLOOD PRESSURE INCREASED (blood pressure went up), PYREXIA (higher fever than when got covid) and HEADACHE (headaches). On 15-May-2021, the patient experienced DIARRHOEA (diarrhea worse than when I had covid). In May 2021, the patient experienced CHEST PAIN (chest pains) and PARAESTHESIA (nerve sensations up and down back and arms). On an unknown date, the patient experienced THROMBOSIS (blood clots) (seriousness criterion medically significant), PAIN IN EXTREMITY (leg pain due to clots) and CHILLS (chills). On 16-May-2021, DIARRHOEA (diarrhea worse than when I had covid) had resolved. At the time of the report, THROMBOSIS (blood clots), CHEST PAIN (chest pains), PARAESTHESIA (nerve sensations up and down back and arms), BLOOD PRESSURE INCREASED (blood pressure went up), PAIN IN EXTREMITY (leg pain due to clots), PYREXIA (higher fever than when got covid), HEADACHE (headaches) and CHILLS (chills) outcome was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Treatment information was not reported. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1366915-1	No blood clot but testing shows signs of blood clot; Unable to eat; Chest pains; Red spots; Muscles feel like needles; She has not taken second dose; Fainted; hand turned red; Palpitations; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (No blood clot but testing shows signs of blood clot) and SYNCOPE (Fainted) in a 26-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 02-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 02-Apr-2021, the patient experienced SYNCOPE (Fainted) (seriousness criterion medically significant), ERYTHEMA (hand turned red) and PALPITATIONS (Palpitations). On an unknown date, the patient experienced THROMBOSIS (No blood clot but testing shows signs of blood clot) (seriousness criterion medically significant), FEEDING DISORDER (Unable to eat), CHEST PAIN (Chest pains), RASH MACULAR (Red spots), MYALGIA (Muscles feel like needles) and INCOMPLETE COURSE OF VACCINATION (She has not taken second dose). At the time of the report, THROMBOSIS (No blood clot but testing shows signs of blood clot), SYNCOPE (Fainted), ERYTHEMA (hand turned red), PALPITATIONS (Palpitations), FEEDING DISORDER (Unable to eat), CHEST PAIN (Chest pains), RASH MACULAR (Red spots), MYALGIA (Muscles feel like needles) and INCOMPLETE COURSE OF VACCINATION (She has not taken second dose) had not resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On an unknown date, Coagulation test: signs of blood clot (abnormal) signs of blood clot. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. It was reported that within 3 minutes of receiving dose, patient's hand turned red and she fainted. Patient was brought to the hospital and was released a couple hours later when palpitations stopped. No medication given to treat symptoms. The patient was receiving unspecified acid reflux medication. Company comment Very limited information regarding these events has been provided at this time. Further information has been requested. Company causality for events Thrombosis, Syncope, Erythema, Palpitations, Feeding disorder, Chest pain, Rash macular and Myalgia cannot be excluded, while company causality for Incomplete course of vaccination is assessed as not applicable.; Sender's Comments: Very limited information regarding these events has been provided at this time. Further information has been requested. Company causality for events Thrombosis, Syncope, Erythema, Palpitations, Feeding disorder, Chest pain, Rash macular and Myalgia cannot be excluded, while company causality for Incomplete course of vaccination is assessed as not applicable.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1366919-1	blood clots in her legs; blood lot traveled to lungs, had a very severe case of blood clots; severe pain in her legs; could not walk; weak; feeling fatigued; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clots in her legs), PULMONARY EMBOLISM (blood lot traveled to lungs, had a very severe case of blood clots) and PAIN IN EXTREMITY (severe pain in her legs) in a 73-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 15-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 22-Mar-2021, the patient experienced PAIN IN EXTREMITY (severe pain in her legs) (seriousness criterion hospitalization), GAIT INABILITY (could not walk), ASTHENIA (weak) and FATIGUE (feeling fatigued). On 31-Mar-2021, the patient experienced THROMBOSIS (blood clots in her legs) (seriousness criteria hospitalization and medically significant) and PULMONARY EMBOLISM (blood lot traveled to lungs, had a very severe case of blood clots) (seriousness criteria hospitalization and medically significant). The patient was hospitalized on 31-Mar-2021 due to PULMONARY EMBOLISM and THROMBOSIS. At the time of the report, THROMBOSIS (blood clots in her legs), PULMONARY EMBOLISM (blood lot traveled to lungs, had a very severe case of blood clots), PAIN IN EXTREMITY (severe pain in her legs), GAIT INABILITY (could not walk), ASTHENIA (weak) and FATIGUE (feeling fatigued) outcome was unknown. No concomitant and treatment medications were reported. The patient had no side effects from the first dose of Moderna vaccine. The patient remained hospitalized at the time of report. Based on the current available information which shows a temporal association between the use of mRNA-1273 and the onset of the reported events, a causal relationship cannot be excluded. Fatigue is consistent with the Known safety profile of the product.; Sender's Comments: Based on the current available information which shows a temporal association between the use of mRNA-1273 and the onset of the reported events, a causal relationship cannot be excluded. Fatigue is consistent with the Known safety profile of the product.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1368322-1	I woke up and 1/2 my mouth and my tongue felt numb and tingling as if I had Novocain and it was wearing off. Woke up the next day and it was still there so I called the NP. She said to go and get a CAT scan in the morning and it didn't show anything. Later in the afternoon I noticed I was dropping things and dragging my feet and got concerned it was a stroke so I drove down to a larger medical center and they did a contrast CAT scan and that showed nothing. So then they did an MRI and they found a clot. So they put me on Plavix, blood thinner and all that stuff. After a couple of days my hand and foot were fine but the numbness and tingling in my mouth is still there till now. All of this wasn't real pronounced so I didn't think it was a stroke but it was. They sent me to a Neurologist and Cardiologist.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1368387-1	Two blood clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1368486-1	Moderna Dose 1 3/5/21 (029A21A) Moderna Dose 2 4/16/21 (027B21A) COVID Positive 5/20/21 5/20/21: Presented to ED. The patient presents with fever. 61 year old male presents to the emergency department for a fever with an onset of 1 day. Patient was diagnosed with pancreatic cancer in April 2021. Patient states he has had 2 rounds of chemotherapy but states he could not have his 3 round last week because his white count was too low. Patient is also complaining of dull chest pain that is exacerbated with deep breathing, cough, and a headache but denies vomiting, diarrhea, neck pain, dysuria, hematuria, and nausea. Patient was diagnosed with blood clots on 05/09/2021 but states that he is not having similar symptoms to when he was diagnosed with blood clots. Patient is taking Xarelto. He has never had COVID but is vaccinated. Patient sees Dr. (Name) at the UI Oncology clinic. 5/25/21: 61-year-old male on chemotherapy for pancreatic cancer, also recently diagnosed pulmonary embolism on rivaroxaban, presents with 1 day of fever. Initially, did not have any cough or shortness of breath, was admitted to the hospital, where a chest x-ray, urinalysis and examination failed to reveal an exact source of the fever. His absolute neutrophil count was less than 500, so he was admitted for neutropenic fever. Cultures were obtained. IV antibiotics started. COVID-19 was tested and, surprisingly, came back positive. The patient has actually had both of his COVID-19 vaccines and basically, does not have pneumonia, shortness of breath, or hypoxemia. We did not use any medications for COVID while in the hospital, watching for any signs of respiratory compromise, but none developed. Blood cultures were negative. He was switched from ceftazidime to doxycycline at time of discharge. Will follow up with Dr. (Name) for his next cycle of chemotherapy. Because of his COVID-19 positivity, we will not schedule any immediate followup appointments, although the patient will call if he has persistent fever, dyspnea, or signs of hypoxemia. For the past 2 days, the T- max has been 99.9, and most the time 98.6. He was seen the day of discharge, doing quite well. Blood pressure 153/80, pulse 90, regular, oximetry 95% on room air. Glucose was 210. Discharged on Rivaroxaban 15 mg b.i.d., doxycycline 100 mg b.i.d., Robitussin AC as needed for cough, Lantus 28 units at bedtime, atorvastatin 80 mg daily, aspirin 81 mg daily. The patient did get 1 dose of Neupogen 480 mcg subcu while in the hospital.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1368663-1	Patient is hospitalized for blood clot a month after getting second moderna shot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1369075-1	patient reported a couple weeks after 1st moderna vaccine (she was unsure of date, I guessed on the first page of the report because I had to put an exact date). she began to feel short of breath. eventually went to hospital where she was diagnosed with blood clots in both the legs and lungs. she said dr is saying it was due to the vaccine and told her not to get the second dose. she stayed 5 days in the hospital according to her. had surgery to place a filter for the clots and is now home on oxygen around the clock.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1370335-1	Blood clots in lungs, legs and brain; She cant speak or stand on her own/in diaper; This spontaneous case was reported by a non-health professional (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clots in lungs, legs and brain) and PHYSICAL DISABILITY (She cant speak or stand on her own/in diaper) in a 51-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 048A21A and 022M20A) for COVID-19 vaccination. No Medical History information was reported. On 25-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 25-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 26-Mar-2021, the patient experienced THROMBOSIS (Blood clots in lungs, legs and brain) (seriousness criteria hospitalization prolonged and medically significant) and PHYSICAL DISABILITY (She cant speak or stand on her own/in diaper) (seriousness criterion hospitalization prolonged). The patient was treated with HEPARIN (intravenous) at a dose of UNK dosage form. At the time of the report, THROMBOSIS (Blood clots in lungs, legs and brain) and PHYSICAL DISABILITY (She cant speak or stand on her own/in diaper) outcome was unknown. Concomitant medications were not provided. Reporter stated that hospital is trying to send the patient to a nursing home. No further treatment information was not provided. Company Comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1370351-1	Heart attack; Stroke; Blood clots; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of MYOCARDIAL INFARCTION (Heart attack), CEREBROVASCULAR ACCIDENT (Stroke) and THROMBOSIS (Blood clots) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient experienced MYOCARDIAL INFARCTION (Heart attack) (seriousness criteria hospitalization and medically significant) and CEREBROVASCULAR ACCIDENT (Stroke) (seriousness criteria hospitalization and medically significant). an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criteria hospitalization and medically significant). The patient was treated with Surgery for Thrombosis. At the time of the report, MYOCARDIAL INFARCTION (Heart attack), CEREBROVASCULAR ACCIDENT (Stroke) and THROMBOSIS (Blood clots) outcome was unknown. Concomitant product use was not provided. It was reported that the patient experienced heart attack, stroke, and uncontrollable blood clotting after 5 days of vaccine. Patient was also admitted at hospital for surgery. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1371668-1	Pain, swelling, stiffness and blood clot in left arm
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1371851-1	"06/02/21 - Moderna vaccine administered at 3:30pm. Waited 15 mins at the HD without any reported side effects. When driving home, client states she felt dizzy. Developed fatigue around 5pm. At 8pm went to have a bowel movement and large amounts of red blood with small clots passed from rectum. Client states she had to flush 3 times d/ t the amount of blood. No stool passed. No other episodes reported. Client does state that around 2-3 am she was up to use the restroom and noted small amount of dried blood spots in her undergarment. Client states that she does have a h/o hemorrhoids since the birth of her baby but states this ""definitely"" was not related. Client reported feeling hot during the night with mild abdominal cramping but states she is to start her menstrual cycle soon. PHN questioned if she felt this was her menstrual cycle and client clearly states the blood was coming from her rectum. No further episodes of passing blood reported and states that she is feeling better today. Client states that she does not have a h/o of blood in stool except on occasion scant amounts from hemorrhoids. PHN encouraged client to report this finding to her PCP as well and that the PHN was going to report this to VAERS. No other s/s of bleeding reported to the PHN."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1374014-1	Blood clot in leg; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clot in leg) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot in leg) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clot in leg) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. No relevant concomitant medications were reported. No treatment information was provided. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1374020-1	blood clot; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (blood clot) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (blood clot) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (blood clot) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. No concomitant medications were reported. No treatment medications were reported. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1374282-1	The vaccine is affecting young people; My son felt like he had an aneurysm in his neck/He had pain in the large artery on the right side of his neck like a myocardial inflammation or something; The right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes; On Tuesday, on the right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes; shoulder pain; On Saturday night he was feeling tired and cold; feeling cold; On Saturday his legs were very heavy and had pain, like he had blood clots or thrombosis in his veins; On Saturday his legs were very heavy and had pain; This spontaneous case was reported by a patient and describes the occurrence of THROMBOSIS (On Saturday his legs were very heavy and had pain, like he had blood clots or thrombosis in his veins) and ANEURYSM (My son felt like he had an aneurysm in his neck/He had pain in the large artery on the right side of his neck like a myocardial inflammation or something) in a 23-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 027L21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included BUPROPION HYDROCHLORIDE (ZYBAN) for an unknown indication. On 21-May-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-May-2021, the patient experienced THROMBOSIS (On Saturday his legs were very heavy and had pain, like he had blood clots or thrombosis in his veins) (seriousness criterion medically significant), PAIN IN EXTREMITY (On Saturday his legs were very heavy and had pain), FEELING COLD (feeling cold) and FATIGUE (On Saturday night he was feeling tired and cold). On 25-May-2021, the patient experienced ANEURYSM (My son felt like he had an aneurysm in his neck/He had pain in the large artery on the right side of his neck like a myocardial inflammation or something) (seriousness criterion medically significant), NECK PAIN (The right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes), SWELLING (On Tuesday, on the right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes) and ARTHRALGIA (shoulder pain). On an unknown date, the patient experienced VACCINATION COMPLICATION (The vaccine is affecting young people). At the time of the report, THROMBOSIS (On Saturday his legs were very heavy and had pain, like he had blood clots or thrombosis in his veins), ANEURYSM (My son felt like he had an aneurysm in his neck/He had pain in the large artery on the right side of his neck like a myocardial inflammation or something), VACCINATION COMPLICATION (The vaccine is affecting young people), PAIN IN EXTREMITY (On Saturday his legs were very heavy and had pain), NECK PAIN (The right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes), SWELLING (On Tuesday, on the right side of his neck, between the shoulder and neck, on the inside, he felt swelling, pressure pain, dull pain that comes and goes), FEELING COLD (feeling cold), ARTHRALGIA (shoulder pain) and FATIGUE (On Saturday night he was feeling tired and cold) outcome was unknown. No treatment information was provided. Company comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. This case was linked to MOD-2021-182543 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1374298-1	Presented with clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Presented with clots) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) .5 milligram. On an unknown date, the patient experienced THROMBOSIS (Presented with clots) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Presented with clots) outcome was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. No concomitant medications was provided by the reporter. No treatment information was provided. Action taken with mRNA-1273 in response to the events was not applicable. Company comment: Very limited information regarding this event has been provided at this time. Further information cannot be requested.; Sender's Comments: Very limited information regarding this event has been provided at this time. Further information cannot be requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1375081-1	Trouble Breathing, blood clot, other issues. Next day. Spent 4 days in hospital
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1375342-1	Only nine days after my second Moderna shot, I broke my ankle. I then had to wear an immobilizing boot for 5-6 weeks. Despite taking one baby aspirin a day during the immobilization period, I developed a blood clot in my paroneal calf vein 5 weeks into the immobilization. I am wondering if the proximity of my immobilization to my Moderna vaccination put me at an increased risk of developing a blood clot.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1375724-1	Blood clots; Severe allergic reactions; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots) and HYPERSENSITIVITY (Severe allergic reactions) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots) (seriousness criterion medically significant) and HYPERSENSITIVITY (Severe allergic reactions) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clots) and HYPERSENSITIVITY (Severe allergic reactions) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Concomitant medication and treatment information were not reported. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Reporter did not allow further contact; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1375821-1	Blood clot in right eye.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1376365-1	Patient presented with premature rupture of membranes and passed a clot at 20wks and 2 days approximately 18 days post vaccination for COVID19 (second dose). She is G2P1, no complications with current pregnancy and no history of preterm labor. 10/22/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1376594-1	clot behind right knee, had ultra sound done, sent me o emergency room was given a pill of xarelto and prescription ans sent home, have to take pills for the next 31 days.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1377126-1	She got her vaccine, she wasn't feeling good right there and didn't know what it was going to progress to. She then at 2:00 AM the next morning she rolled out of the bed and fell to the floor and got stuck between the bed and was not able to stand up, and crawled down 14 stairs and then couldn't move. She could not tell him why she couldn't and she was not able to stand up (large women). Her husband did not see any signs of stroke. They called his son and noticed that her face was sagging and he said that it looked like a stroke. They called the paramedics and they sent her to hospital and she was flown to Clinic and diagnosed with a stroke. She had surgery and they tried to remove the blood clot and her blood pressure went high and they cut a huge part of her skull to remove the pressure, but not able to remove the clot that he is aware of. She is in rehab right now, and it's not really showing any signs of promise. She has a lot of stroke symptoms of being combative and she has a blood clot that has moved to her arm as well. She had 6 blood clots on her brain when they went into it, and now has blood clots in each arm. She is aware of her surroundings to some degree and thinks that she is somewhere else and feels like she has wet herself. Her husband said that the rehab does not look promising. She is still in the hospital and they are having a meeting tomorrow regarding how long they feel she may be there. She has very little headway in rehab. She cannot move her left side, hand or leg. She is able to move her right side though. Husband feels it has destroyed her life and it will be a miracle if she comes out of it.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1377366-1	heavy bleeding, large blood clots, spotting and two periods in the month following first moderna vaccine ear hissing - tinnitus

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1378167-1	The night after receiving the vaccine, the patient began having continuous fevers and coughs. It was later discovered that there were blood clots in the lung and legs. Patient health rapidly declined during hospitalization period. Mobility, apetite and overall strength and energy decreased. Patient was fully bedridden and soon lost eye sight.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1378203-1	Patient was first diagnosed with bursitis at his pcp and sent home. Patients elbow was swollen and referred to urgent care by the pharmacist pharmacy. Patient was seen by Dr, and found to have a blood clot when doctor tried to aspirate the elbow. Patient has fully recovered at returned to work
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1378401-1	a blood clot from the left groin to the foot. desolved using blood thinner process. Still receiving treatment as of today.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1378992-1	On the 19th of March, one day after the two week period after the second shot, my heart began to race and my breathing became difficult. My husband took me to the ER and at that time the doctors were unable to find the problem. After several ER visits, visit with neuro due to a nerve problem that came along with the heart and breathing problems , my primary care doctor finally suggested an ultra sound be conducted due to the extreme leg/thigh pain that had made it impossible for me to walk. The results of the ultra sound indicated severe extreme multiple blood clots. At this point, I had become wheel chair bound. The nerve injury was pinching on the optcitical nerve which has now caused my vision to be impaired; my vision is no longer sharp (blurred vision). At this point, I can only walk a few feet without the assistance of a wheel chair or some other support. My thighs remain huge/swolllen. I continue experience leg/thigh pain. The nerve injury in the left side of my brain seems to be causing some crippling of my right hand which is my dominant side.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1380721-1	lots of people who have had blood clots.; have experienced numbness in their hand 4 hours; Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (lots of people who have had blood clots.) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (lots of people who have had blood clots.) (seriousness criterion medically significant) and HYPOAESTHESIA (have experienced numbness in their hand 4 hours). At the time of the report, THROMBOSIS (lots of people who have had blood clots.) and HYPOAESTHESIA (have experienced numbness in their hand 4 hours) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. The concomitant medications on use were not provided. No laboratory data was provided. No treatment information was provided. Caller reported she is on a forum and lots of people who have had blood clots have experienced numbness in their hand 4 hours after receiving the vaccine. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Very limited information regarding this events has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this events has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1380728-1	Blood clot in left leg; Not feeling like herself since second dose; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clot in left leg) in a 52-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 046B21A and 018B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included LEVOTHYROXINE, HYDROCHLOROTHIAZIDE, FLUTICASONE PROPIONATE (FLONASE [FLUTICASONE PROPIONATE]), FEXOFENADINE, LOSARTAN and SERTRALINE for an unknown indication. On 25-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 22-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 22-Apr-2021, the patient experienced FEELING ABNORMAL (Not feeling like herself since second dose). On 16-May-2021, the patient experienced THROMBOSIS (Blood clot in left leg) (seriousness criterion medically significant). The patient was treated with RIVAROXABAN (XARELTO) ongoing since an unknown date for Clot blood, at an unspecified dose and frequency. At the time of the report, THROMBOSIS (Blood clot in left leg) outcome was unknown and FEELING ABNORMAL (Not feeling like herself since second dose) had not resolved. It was reported that the patient was not feeling like herself since the second dose but was unsure if it was a result of the vaccination. Company Comment : Very limited information regarding this event has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information regarding this event has been provided at this time. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				blood clots in her legs and lungs; blood clots in her legs and lungs; back and legs were hurting; couldn't walk.; back and legs were hurting; alertness declining; activity declining; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots in her legs and lungs), PULMONARY EMBOLISM (blood clots in her legs and lungs), BACK PAIN (back and legs were hurting), GAIT DISTURBANCE (couldn't walk.), PAIN IN EXTREMITY (back and legs were hurting), DEPRESSED LEVEL OF CONSCIOUSNESS (alertness declining) and DECREASED ACTIVITY (activity declining) in a 20-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No medical history was reported. On 22-Feb-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 18-Mar-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to .5 milliliter. On 27-May-2021, the patient experienced DEPRESSED LEVEL OF CONSCIOUSNESS (alertness declining) (seriousness criteria hospitalization, disability and medically significant) and DECREASED ACTIVITY (activity declining) (seriousness criteria hospitalization and disability). On an unknown date, the patient experienced THROMBOSIS (blood clots in her legs and lungs) (seriousness criteria hospitalization, disability and medically significant), PULMONARY EMBOLISM (blood clots in her legs and lungs) (seriousness criteria hospitalization, disability and medically significant), BACK PAIN (back and legs were hurting) (seriousness criteria hospitalization and disability), GAIT DISTURBANCE (couldn't walk.) (seriousness criteria hospitalization and disability) and PAIN IN EXTREMITY (back and legs were hurting) (seriousness criteria hospitalization and disability). The patient was hospitalized on 16-May-2021 due to BACK PAIN, GAIT DISTURBANCE, PAIN IN EXTREMITY, PULMONARY EMBOLISM and THROMBOSIS. The patient was treated with HEPARIN for Coagulopathy, at a dose of 1 dosage form; Surgery (thrombectomy and Intubated) for Thrombosis and Surgery (thrombectomy and Intubated) for Pulmonary embolism. At the time of the report, THROMBOSIS (blood clots in her legs and lungs), PULMONARY EMBOLISM (blood clots in her legs and lungs), BACK PAIN (back and legs were hurting), GAIT DISTURBANCE (couldn't walk.), PAIN IN EXTREMITY (back and legs were hurting), DEPRESSED LEVEL OF CONSCIOUSNESS (alertness declining) and DECREASED ACTIVITY (activity declining) had not resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): In May 2021, Blood creatinine: 5.37 (High) high. In May 2021, Blood potassium: high (High) high. In May 2021, Glomerular filtration rate: 9.5 (Low) low. In May 2021, Imaging procedure: abnormal (abnormal) had new imaging and she was still developing clots. In May 2021, Ultrasound abdomen: inconclusive (Inconclusive) inconclusive. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. She was in the cardiac intensive care unit (ICU). She had a thrombectomy and was on heparin. They intubated her and put her on the ECMO machine on 19 May 2021. The patient got off ECMO on 24 May 2021 and the intubation tube was removed 25 May 2021. They were having trouble obtaining a non-hemolyzed sample from her since 27 May 2021. The patients concomitant medication information included birth control pills. Very limited information regarding these events have been provided at this time. However subject's use of birth control pills can be a confounding factor to the events thrombosis and Pulmonary embolism. No further information is expected.; Sender's Comments: Very limited information regarding these events have been provided at this time. However subject's use of birth control pills can be a confounding factor to the events thrombosis and Pulmonary embolism. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1380740-1	
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1380756-1	blood clot on the shaft of their penis; Bruise; This spontaneous case was reported by a patient family member or friend and describes the occurrence of THROMBOSIS (blood clot on the shaft of their penis) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On an unknown date, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (blood clot on the shaft of their penis) (seriousness criterion medically significant) and CONTUSION (Bruise). At the time of the report, THROMBOSIS (blood clot on the shaft of their penis) and CONTUSION (Bruise) outcome was unknown. No concomitant or treatment medications were reported. Reporter did not allow further contact; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However, very limited information regarding this event/s has been provided at this time. Further information cannot be requested because Consent to a follow up from safety was not granted. The reporter did not provide further contact information.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1381749-1	Swollen ankle left leg Saturday afternoon After midnight shortness of breathe and heart started racing and I thought I was going to die Sunday leg really swollen having trouble walking Sunday night heart was pounding. Called GYN and was instructed to go to the Emergency Room.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1382937-1	Patient notified me 6/3/21.... Patient states that he developed a blood clot behind his right eye. and the result of this clot caused him to lose site in his right eye.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1384351-1	Superficial blood clot; it was bigger after the first dose; Very painful to touch/arm was in pain; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Superficial blood clot) in an 83-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 013A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 02-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Superficial blood clot) (seriousness criterion medically significant), VACCINATION SITE SWELLING (it was bigger after the first dose) and VACCINATION SITE PAIN (Very painful to touch/arm was in pain). At the time of the report, THROMBOSIS (Superficial blood clot), VACCINATION SITE SWELLING (it was bigger after the first dose) and VACCINATION SITE PAIN (Very painful to touch/arm was in pain) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant medications was reported. Treatment information was not provided. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1384353-1	blood clots; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots) in a male patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 18-May-2021, the patient experienced THROMBOSIS (blood clots) (seriousness criteria hospitalization and medically significant). At the time of the report, THROMBOSIS (blood clots) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Concomitant product use was not reported. Treatment information included blood thinners. This a case of a male patient of unknown age who had blood clots and was hospitalized after receiving a dose of Moderna COVID Vaccine. Very limited information has been provided at this time.; Sender's Comments: This a case of a male patient of unknown age who had blood clots and was hospitalized after receiving a dose of Moderna COVID Vaccine. Very limited information has been provided at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1385129-1	Pulmonary embolism in Right lung. Subsequent finding was multiple blood clots throughout both lungs. Gave blood 3 weeks after second Moderna injection, blood clotting would not allow a full donation (right arm). Upon checking in to the ER for chest pain on April 17, blood clotting would not allow IV hook up in right arm.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1386054-1	DEATH FROM BLOOD CLOT
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1386411-1	severe vaginal bleeding with large clots starting at 2 am 6/9/21
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1386844-1	Patient has no clotting history. He developed body aches, left arm pain, and fevers immediately after immunization, but improved. Two days later on 6/2, he developed left flank pain and fevers. On 6/4, he was evaluated in the ED, with CT scan showing partially occlusive thrombosis of left renal artery with areas of renal infarction. Cr remained stably at baseline. Initial workup was otherwise notable for WBC 17 and negative infectious workup. He was initiated on therapeutic Lovenox. On 6/5, CTA chest/abdomen/pelvis was performed to evaluate for other clots. Left renal artery was complete thromboses with evolving foci of renal infarction. No other clots were detected. He continued to have fevers. By 6/8, his leukocytosis, fever, and flank pain resolved. Renal ultrasound with doppler showed restoration of some flow in the left renal artery. He is to be discharged 6/9 on Lovenox 1mg/kg BID without patient Hematology follow-up. His coagulability workup was notable for positive LAC, FVII deficiency, and PET/CT without evidence of occult malignancy.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1387515-1	Blood clots in lungs, heart and right leg; felt like someone punched me in the chest; severe shortness of breath; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood clots in lungs, heart and right leg), CHEST DISCOMFORT (felt like someone punched me in the chest) and DYSPNOEA (severe shortness of breath) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On 19-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clots in lungs, heart and right leg) (seriousness criteria hospitalization, medically significant and life threatening), CHEST DISCOMFORT (felt like someone punched me in the chest) (seriousness criterion hospitalization) and DYSPNOEA (severe shortness of breath) (seriousness criterion hospitalization). The patient was hospitalized for 7 days due to CHEST DISCOMFORT, DYSPNOEA and THROMBOSIS. At the time of the report, THROMBOSIS (Blood clots in lungs, heart and right leg), CHEST DISCOMFORT (felt like someone punched me in the chest) and DYSPNOEA (severe shortness of breath) outcome was unknown. No concomitant medications were reported. Treatment information was unknown. Action taken with mRNA-1273 in response to the event was not applicable. After spending 7 days in ICU patient was found to have blood clots in lungs, heart and right leg. This was caused by the moderna vaccination. As per doctors if patient had waited 1 more day patient would have died. Company Comment Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Most recent FOLLOW-UP information incorporated above includes: On 28-May-2021: Event thrombosis added.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1387904-1	Blood clots in her leg and lungs; She is black and blue; She felt pain above her knee and on the inside of her thigh toward the groin that felt like she had been wearing a tight sock. Also felt pain midway between her knee and ankle; She also felt pain midway between her knee and ankle; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (Blood clots in her leg and lungs) in a 71-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 017B21A and 022M20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Dyslipidaemia. Concurrent medical conditions included Blood pressure high, Depression, Acid reflux (oesophageal) and Allergy. On 05-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 02-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 12-May-2021, the patient experienced GROIN PAIN (She felt pain above her knee and on the inside of her thigh toward the groin that felt like she had been wearing a tight sock. Also felt pain midway between her knee and ankle) and ARTHRALGIA (She also felt pain midway between her knee and ankle). On 18-May-2021, the patient experienced THROMBOSIS (Blood clots in her leg and lungs) (seriousness criteria hospitalization and medically significant). On an unknown date, the patient experienced CONTUSION (She is black and blue). The patient was hospitalized from 21-May-2021 to 22-May-2021 due to THROMBOSIS. The patient was treated with APIXABAN (ELIQUIS) on 22-May-2021 at an unspecified dose and frequency. At the time of the report, THROMBOSIS (Blood clots in her leg and lungs), CONTUSION (She is black and blue), GROIN PAIN (She felt pain above her knee and on the inside of her thigh toward the groin that felt like she had been wearing a tight sock. Also felt pain midway between her knee and ankle) and ARTHRALGIA (She also felt pain midway between her knee and ankle) outcome was unknown. The patient thought there were bug bites on her leg on 12-May-2021. Treatment also included IV blood thinner while she was hospitalized. The action taken with mRNA-1273 in response to the event was not applicable. Company Comment: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However, Further information has been requested.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. However, Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1388050-1	First symptoms were severe back pain bent over when walking. Later pain in left shoulder and shortness of breath climbing stairs with heartbeat racing. Later developed pain in right thigh and calf. On May 30 emergency room determine blood clots in right leg and both lungs. First Symptoms appeared about one week after second dose on April 19
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1388147-1	patient received the Moderna covid-19 vaccine and two days later states she was admitted to the ER with blood clots. patient did not report the even to the pharmacy until the week of June 1st
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1389188-1	After 3 weeks of getting the vaccine, I developed a blood clot in my arm and severe swelling.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1390367-1	"Right calf pain with leg swelling and heaviness over a couple of days then right medial calf pain with what appeared to be a bruise the size of my fist and a small central painful lump. Urgent Care appt. US done and found a medial distal thigh venous clot. started Aspirin 325mg once daily. On day 4 of taking this medication, my right leg completely resolved of the swelling and heaviness and the ""bruise"" completely resolved without any discoloration. This makes me think the ""bruise"" was engorged capillary vessels that were not flowing due to the clot."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1390722-1	Blood clot; This case was received via an unknown source (no reference has been entered for a health authority or license partner) on 01-Jun-2021 and was forwarded to Moderna on 02-Jun-2021. Based on the current case data, this case has been classified as invalid. This spontaneous case was reported by a patient and describes the occurrence of THROMBOSIS (Blood clot) in a patient of an unknown age and gender who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. Unknown) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On an unknown date, the patient experienced THROMBOSIS (Blood clot) (seriousness criterion medically significant). At the time of the report, THROMBOSIS (Blood clot) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. Treatment medication was not provided. Concomitant product use was not provided by the reporter. Company comment Very limited information has been provided at this time. Further information has been requested.; Sender's Comments: Very limited information has been provided at this time. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1391461-1	Multiple blood clots in lungs, and kidney fuction has declined . Was in hospital for 3 days for treatment. I am currently taking xerelto for clots.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1391500-1	Patient had his 2nd Moderna vaccine at pharmacy at 2:00 and was struggling to breathe a few hours later. We were at the emergency room around 6:00pm and he was placed on a ventilator between midnight and 1:00am. He died 10 days later after experiencing clots that led to seizures and congestive heart failure. I have a death certificate if needed.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1394058-1	This spontaneous case was reported by a patient family member or friend (subsequently medically confirmed) and describes the occurrence of DEATH (She passed away) and THROMBOSIS (Clots in her leg) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 037A21B) for COVID-19 vaccination. No Medical History information was reported. On 17-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced DEATH (She passed away) (seriousness criteria death, hospitalization and medically significant) and THROMBOSIS (Clots in her leg) (seriousness criteria hospitalization and medically significant). The patient died on an unknown date. The cause of death was not reported. It is unknown if an autopsy was performed. At the time of death, THROMBOSIS (Clots in her leg) outcome was unknown. No concomitant medications reported. Description: The patient had clots in her leg and was hospitalized. There was talk of amputation, but she passed away before that occurred. No treatment information provided. Company Comment: Very limited information regarding these events have been provided at this time. Further information cannot be requested.; Sender's Comments: Very limited information regarding these events have been provided at this time. Further information cannot be requested.; Reported Cause(s) of Death: Unknown cause of death
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1394064-1	Heart Failure; Bilateral DVT; Blood Clot; Exacerbation of pre existing medical conditions; This spontaneous case was reported by a nurse (subsequently medically confirmed) and describes the occurrence of CARDIAC FAILURE (Heart Failure), DEEP VEIN THROMBOSIS (Bilateral DVT) and THROMBOSIS (Blood Clot) in a 77-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Deep vein thrombosis ('had an episode of DVT 30 years ago'). On 02-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 07-Apr-2021, the patient experienced CARDIAC FAILURE (Heart Failure) (seriousness criterion medically significant), DEEP VEIN THROMBOSIS (Bilateral DVT) (seriousness criterion medically significant), THROMBOSIS (Blood Clot) (seriousness criterion medically significant) and CONDITION AGGRAVATED (Exacerbation of pre existing medical conditions). At the time of the report, CARDIAC FAILURE (Heart Failure), DEEP VEIN THROMBOSIS (Bilateral DVT), THROMBOSIS (Blood Clot) and CONDITION AGGRAVATED (Exacerbation of pre existing medical conditions) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular), the reporter did not provide any causality assessments. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested. Of note, the patient's age and pre-existing medical conditions are confounding factors that may play possible contributory roles.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested. Of note, the patient's age and pre-existing medical conditions are confounding factors that may play possible contributory roles.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1394069-1	9 blood clots; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (9 blood clots) in a 50-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. In February 2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. In March 2021, the patient experienced THROMBOSIS (9 blood clots) (seriousness criteria hospitalization and medically significant). At the time of the report, THROMBOSIS (9 blood clots) was resolving. No concomitant medications were provided. The patient was in the ICU (intensive care unit) for 3 days. No additional treatment information was provided. The reporter stated that the patient thought that her Moderna COVID-19 vaccinations caused the blood clots.; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. No further information is expected at this time.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1394070-1	blood clots from the knee to top of her leg.; feeling uncomfortable; This spontaneous case was reported by a patient (subsequently medically confirmed) and describes the occurrence of THROMBOSIS (blood clots from the knee to top of her leg.) in a 69-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 004M20A and 030A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Hypercholesterolemia. Concurrent medical conditions included Diabetes and Hypertension. On 30-Mar-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 30-Mar-2021, the patient experienced THROMBOSIS (blood clots from the knee to top of her leg.) (seriousness criterion medically significant) and MALAISE (feeling uncomfortable). At the time of the report, THROMBOSIS (blood clots from the knee to top of her leg.) and MALAISE (feeling uncomfortable) outcome was unknown. Very limited information regarding this event/s has been provided at this time. Further information is not expected. Reporter did not allow further contact; Sender's Comments: Very limited information regarding this event/s has been provided at this time. Further information is not expected.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1394618-1	Blood clotting during periods. Never experienced anything like this before.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1396298-1	Developed blood clot in leg.. having lysis surgery today 6/14/2021
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1396407-1	On 6/9/2021 patient was transferred to local Hospital from another Hosptial for chest pain, worsening EF and concerns of ischemic cardiomyopathy. She was visiting from out of town (attending her brother's funeral who also reportedly died after receiving the Covid vaccine). On 5/31 she had been hospitalized with ischemic colitis, was discharged on 6/6/2021 then returned to the the hospital on 6/9/2021 with increasing shortness of breath, confusion. A repeat TTE showed anterior wall mothion abnormalities and decreased EF, resulting in the decision to transfer to another hospital. She had a cardiac cath, was improving, but had a sudden change in condition and expired. A throbotic event was suspected. She is currently awaiting an autopsy.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1398524-1	"large blood clot in the right lung/smaller blood clot in the left lung; blood clot in the calf of the left leg; Baker's cyst behind the left knee; ""I still didn't feel right or good""; ""problems with my kidneys""; dull pain; ""really bad chest pains""; ""knee got swollen up""; This spontaneous case was reported by a patient and describes the occurrence of PULMONARY EMBOLISM (large blood clot in the right lung/smaller blood clot in the left lung) and THROMBOSIS (blood clot in the calf of the left leg) in a 69-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 042L20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No medical history was reported. Concomitant products included LOSARTAN, NEBIVOLOL HYDROCHLORIDE (BYSTOLIC), OMEPRAZOLE, EZETIMIBE, CETIRIZINE HYDROCHLORIDE (ZYRTEC ALLERGY), VITAMIN D [VITAMIN D NOS], CALCIUM, MULTIVITAMINS [VITAMINS NOS] and APIXABAN (ELIQUIS) for an unknown indication. On 28-Jan-2021, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 25-Mar-2021, the patient experienced JOINT SWELLING (""knee got swollen up""). On 26-Apr-2021, the patient experienced PULMONARY EMBOLISM (large blood clot in the right lung/smaller blood clot in the left lung) (seriousness criteria hospitalization and medically significant) and CHEST PAIN (""really bad chest pains""). On an unknown date, the patient experienced THROMBOSIS (blood clot in the calf of the left leg) (seriousness criterion medically significant), SYNOVIAL CYST (Baker's cyst behind the left knee), MALAISE (""I still didn't feel right or good""), RENAL PAIN (""problems with my kidneys"") and PAIN (dull pain). At the time of the report, PULMONARY EMBOLISM (large blood clot in the right lung/smaller blood clot in the left lung), THROMBOSIS (blood clot in the calf of the left leg), JOINT SWELLING (""knee got swollen up""), SYNOVIAL CYST (Baker's cyst behind the left knee), MALAISE (""I still didn't feel right or good""), CHEST PAIN (""really bad chest pains""), RENAL PAIN (""problems with my kidneys"") and PAIN (dull pain) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. Company comment: Very limited information regarding this events has been provided at this time. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested.; Sender's Comments: Very limited information regarding this events has been provided at this time. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Further information has been requested."
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1399422-1	02/25/2021: Patient received 1st dose of Moderna 03/17 ? 3/19: 20 days after 1st dose, c/o progressively worsening bilateral lower extremity weakness of several weeks? duration. Patient had also suffered from recurrent falls over the preceding several days prior to admission. subsequent CT scan of the abdomen and pelvis revealed severe hepatic steatosis and a CT angiogram of the chest revealed bilateral pulmonary emboli. She subsequently had a duplex ultrasound of the lower extremities that revealed a right lower extremity DVT. MRI of the lumbar spine was performed for further evaluation of her bilateral lower extremity weakness that revealed some moderate bilateral neuroforaminal narrowing with mild spinal stenosis at L3-L4 and moderate spinal stenosis at L4-L5 with some degenerative endplate changes at similar levels. Dx: Pulmonary embolism - discharged home on Xarelto for complete treatment of her DVT/PE. 3/25: 2nd dose of Moderna given at Primary care - referred to ER for complaints. 3/25: ER encounter - She has been experiencing heavy vaginal bleeding, passing large clots at times, for the past six days. She reports that she usually has regular menstrual cycles, regulated on oral contraceptive pills. Dx: Menorrhagia with regular cycle, Anticoagulated ? treated and discharged on TXA 3/28 ? 4/4: 3 days after 2nd dose. presents with ongoing vaginal bleeding complicated by lightheadedness with standing, generalized weakness and fatigue, and dyspnea on exertion. Patient states she has been bleeding for almost 2 weeks vaginally. Dx: severe symptomatic anemia due to dysfunctional uterine bleeding / Menorrhagia and need for anticoagulation, Folate deficiency anemia, Venous stasis. Rx: Transfused 3 units, Provera, Due to recent VTE in stable hemoglobin cautiously started heparin, Oral contraceptives stopped, resumed Xarelto, folic acid supplement.
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1399516-1	Sweating, soreness, vomiting
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1400278-1	After having the vaccine I have had nose bleeds and severe headaches. I have never had a nose bleed in my life. I have had 12-13 in about a 3 week period. I have only had one in the last week. I can stand on the toilet and it will just pour out of my nose. The first time that it happened there was a big clot but after that I have not seen another clot. I work for a doctor and I asked him about it after I have had about 6 nose bleeds. He kind of checked me out a little bit and told me to keep a watch on it and see what happens. He did not see anything when he checked me. I continued to have them and he suggested that I go with an EMG.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1401716-1	Blood Clots; Altered Speech; Body Is Stiff; Word Finding Difficulties; Left Eye Muscle Feels Gummy; Forgets What They Are Talking About; Nausea; periodic Stumbling/Left Leg Is Dragging; Unable To Draw A Clock For A Cognitive Exam; Afraid Of Being Alone; Lack Of Appetite; Depth Perception Issues; Headaches Everyday; This spontaneous case was reported by a consumer and describes the occurrence of THROMBOSIS (Blood Clots) in a 48-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 039B21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concomitant products included WARFARIN for Clot blood, BUTALBITAL, CAFFEINE, PARACETAMOL (FIORICET), SUMATRIPTAN, CELECOXIB (CELEXA [CELECOXIB]), ETHINYLESTRADIOL, ETONOGESTREL (NUVARING) and BACLOFEN for an unknown indication. On 20-Apr-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 20-Apr-2021, the patient experienced HEADACHE (Headaches Everyday). On 23-Apr-2021, the patient experienced ILLUSION (Depth Perception Issues), FEAR (Afraid Of Being Alone) and DECREASED APPETITE (Lack Of Appetite). On 30-Apr-2021, the patient experienced THROMBOSIS (Blood Clots) (seriousness criteria hospitalization and medically significant), GAIT DISTURBANCE (periodic Stumbling/Left Leg Is Dragging) and COGNITIVE DISORDER (Unable To Draw A Clock For A Cognitive Exam). On an unknown date, the patient experienced DYSARTHRIA (Altered Speech), MUSCULOSKELETAL STIFFNESS (Body Is Stiff), APHASIA (Word Finding Difficulties), ABNORMAL SENSATION IN EYE (Left Eye Muscle Feels Gummy), MEMORY IMPAIRMENT (Forgets What They Are Talking About) and NAUSEA (Nausea). The patient was hospitalized on 30-Apr-2021 due to THROMBOSIS. The patient was treated with ONDANSETRON ongoing since an unknown date for Nausea, at a dose of 1 UNK and HEPARIN for Clot blood, at an unspecified dose and frequency. At the time of the report, THROMBOSIS (Blood Clots), DYSARTHRIA (Altered Speech), MUSCULOSKELETAL STIFFNESS (Body Is Stiff), ILLUSION (Depth Perception Issues), GAIT DISTURBANCE (periodic Stumbling/Left Leg Is Dragging), APHASIA (Word Finding Difficulties), ABNORMAL SENSATION IN EYE (Left Eye Muscle Feels Gummy), FEAR (Afraid Of Being Alone), DECREASED APPETITE (Lack Of Appetite), MEMORY IMPAIRMENT (Forgets What They Are Talking About), COGNITIVE DISORDER (Unable To Draw A Clock For A Cognitive Exam), HEADACHE (Headaches Everyday) and NAUSEA (Nausea) outcome was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. The patient reports going to the emergency room on 25Apr2021 and was discharged with medication for the treatment of headaches. During the hospital stay, a MRV (Magnetic Resonance Venography) was conducted, however the results were not provided. Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded. This case was linked to MOD-2021-211608 (Patient Link).; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the event, a causal relationship cannot be excluded.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Seizures; Bood clots; Stuggling to breathe; Struggling to walk; ARDS; Acute hypoxic respiratory failure; Acute exacerbation of heart failure and sepsis; Acute exacerbation of heart failure and sepsis; Cardiac arrest; Blood infection; Fever; This spontaneous case was reported by a patient family member or friend (subsequently medically confirmed) and describes the occurrence of ACUTE RESPIRATORY DISTRESS SYNDROME (ARDS), ACUTE RESPIRATORY FAILURE (Acute hypoxic respiratory failure), SEPSIS (Acute exacerbation of heart failure and sepsis), CARDIAC FAILURE ACUTE (Acute exacerbation of heart failure and sepsis), CARDIAC ARREST (Cardiac arrest), SEIZURE (Seizures), THROMBOSIS (Bood clots), DYSPNOEA (Stuggling to breathe) and GAIT DISTURBANCE (Struggling to walk) in a 53-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Obesity. On 16-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-May-2021 at 2:00 PM, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) dosage was changed to 1 dosage form. On 15-May-2021, the patient experienced ACUTE RESPIRATORY DISTRESS SYNDROME (ARDS) (seriousness criteria death and medically significant), ACUTE RESPIRATORY FAILURE (Acute hypoxic respiratory failure) (seriousness criteria death and medically significant), DYSPNOEA (Stuggling to breathe) (seriousness criterion hospitalization prolonged) and GAIT DISTURBANCE (Struggling to walk) (seriousness criterion hospitalization prolonged). In May 2021, the patient experienced SEPSIS (Acute exacerbation of heart failure and sepsis) (seriousness criteria death and medically significant), CARDIAC FAILURE ACUTE (Acute exacerbation of heart failure and sepsis) (seriousness criteria death and medically significant), CARDIAC ARREST (Cardiac arrest) (seriousness criteria death and medically significant), INFECTION (Blood infection) and PYREXIA (Fever). On 17-May-2021, the patient experienced SEIZURE (Seizures) (seriousness criteria hospitalization prolonged and medically significant) and THROMBOSIS (Bood clots) (seriousness criteria hospitalization prolonged and medically significant). The patient was hospitalized from 15-May-2021 to 25-May-2021 due to DYSPNOEA, GAIT DISTURBANCE, SEIZURE and THROMBOSIS. The patient died on 25-May-2021. The reported cause of death was Cardiac arrest, Acute hypoxic respiratory failure, ards and acute exacerbation of heart failure and sepsis. It is unknown if an autopsy was performed. At the time of death, SEIZURE (Seizures), THROMBOSIS (Bood clots), DYSPNOEA (Stuggling to breathe), GAIT DISTURBANCE (Struggling to walk), INFECTION (Blood infection) and PYREXIA (Fever) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 15-May-2021, Blood pressure measurement: elevated (High) Twice normal. On 15-May-2021, Glycosylated haemoglobin: elevated (High) High. On 15-May-2021, Oxygen saturation: low (Low) Required intubation and was placed on ventilator.. On 15-May-2021, SARS-CoV-2 antibody test: negative (Negative) 1 negative COVID antibody test. On 15-May-2021, SARS-CoV-2 test: negatie (Negative) 3 negative COVID antigen tests. No concomitant medications were provided. It was reported that within few hours after second dose of vaccine patient was struggling to walk and to breathe. The reporter and her husband went to see the patient and took him to an emergency room (ER). In the ER his blood pressure was markedly elevated (twice normal) and his oxygen level was low and by midnight he required intubation and was placed on a ventilator. His HbgA1c was noted to be elevated. Two days after admission (17-May-2021), he developed seizures which the medical team believed to be due to blood clots and he was anti-coagulated. It was determined that he did not have blood clots on his heart valves as the source of the clots. The source of the blood clots was never identified. He later developed fever and was diagnosed with a blood infection and treated with antibiotics. At one point his oxygen requirement on the ventilator went down to 65% but later returned and remained at 100%. He ultimately died on 25-May-2021 and as per copy of his death certificate he had 4 causes of death as cardiac arrest, acute hypoxic respiratory failure, acute respiratory distress syndrome (ARDS) and acute exacerbation of heart failure and sepsis. Treatment included, anticoagulants, antibiotics, intubation and ventilator. Very limited information regarding these events has been provided at this time. Further information has been requested. This case was linked to MOD-2021-219410 (Patient Link).; Sender's Comments: Very limited information regarding these events has been provided at this time. Further information has been requested.; Reported Cause(s) of Death: Cardiac arrest; Acute hypoxic respiratory failure; ARDS; Acute exacerbation of heart failure and sepsis
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1401732-1	
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1402629-1	Patient developed blood clots in her legs, heart, and lungs within 48 hours after receiving her second dose of the Moderna vaccine. She was subsequently treated with Eliquis (apixaban).
THROMBOSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1403614-1	I was diagnosed with a blood clot (Deep Vein Thrombosis) in my left leg after the event of my vaccine. No family history and clean bill of health. Am now on Eliquis blood thinners to ease the pain and reduce my blood clot. Seeing a hematologist in a few days

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1405282-1	Stroke; Blood clot; This spontaneous case was reported by a consumer and describes the occurrence of CEREBROVASCULAR ACCIDENT (Stroke) and THROMBOSIS (Blood clot) in a female patient of an unknown age who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. No Medical History information was reported. On an unknown date, the patient received dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 01-Jun-2021, the patient experienced CEREBROVASCULAR ACCIDENT (Stroke) (seriousness criterion medically significant) and THROMBOSIS (Blood clot) (seriousness criterion medically significant). At the time of the report, CEREBROVASCULAR ACCIDENT (Stroke) and THROMBOSIS (Blood clot) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. The concomitant medications on use were not provided. No laboratory data was provided. No treatment information was provided. The patient did not have any family history of stroke. The patient received the vaccine 3 months prior to AEs. Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded. Reporter did not allow further contact; Sender's Comments: Based on the current available information and temporal association between the use of the product and the start date of the events, a causal relationship cannot be excluded.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1406795-1	Top of left hand formed a blood clot
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1406923-1	Vaccine was given on second day of menstruation cycle on 4th day of mensuration when normally my cycle is finished or tapering off. I had an increase amount of bleeding with large blood clots. At the same time my birth control brand was switched to another brand so didn't know if the change in bleeding/clots was related to change in birth control or moderna vaccine. I reached out to my gyne and made her aware she stated that it was not related to the birth control as it was the same med/dosage. That more than likely was related to moderna vaccine and if the bleeding didn't stop or if it increased to seek medical attention. My cycle lasted for a total of 9 days vs 4-5 days. Clots were from day 4-9.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1406953-1	1st - severe body aches, severe headaches, lethargic, congestion, coughing and couldn?t sleep much. - medicine given was a z-pak, cough medicine and an inhaler. 2nd - severe body aches, vomiting, diarrhea, severe migraines, severe headaches, blood clots coming out of my nose without nose bleeds. Stomach pains behind my chest. So incredibly tired, - medicine given anti nausea Med, cough Med and pain Med. Went to Urgent Care then ER.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1407029-1	"""Moderna COVID-19 Vaccine EUA"" Initial mild pain in lower left calf lasting 3 days. Described as ""pulled muscle feeling."" Dissipated after 3 days. Then, mild, left sided middle back pain that progressively worsened over 3-4 days. Pain eventually felt across entire front and back of chest. Shortness of breath worsened. Difficulty pulling air into lungs and pushing it out. After a total of 5 days from onset of back pain, emergency room visit required."
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1407044-1	Covid Arm, Swollen Lymph nodes in neck, under arms and behind knees. After first vaccine, I went to emergency room to have lump behind knee checked - it was lymphacitis. I then went to an endocrinologist and had adrenals checked and. he said to wait a bit before getting next vaccine. After 2nd dose, I was sick for over 4 days with 104 fever. Very sick and adrenals swelled up bigger than last time. I went to emergency room after 6 weeks of lump behind knee and they found a blood clot behind knee. I am getting an ultrasound from vascular surgeon on June 30.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1407669-1	PT RECEIVED VACCINE 6/8 AND EXPERIENCED STANDARD INJECTION SITE SORENESS FOR 2 DAYS. SHE REPORTS BEING FINE UNTIL ONE WEEK LATER, ON 6/15, LATE IN THE DAY SHE DEVELOPED A SMALL BUMP AT THE INJECTION SITE. ON 6/16 (WED) HER ENTIRE ARM WAS SWOLLEN/INFLAMMED AND WARM TO THE TOUCH. SHE WENT TO ER AND WAS ADMITTED TO HOSPITAL AND TREATED FOR MULTIPLE BLOOD CLOTS. PATIENT REPORTED REACTION TO PHARMACY ON 6/17 AND WAS STILL IN HOSPITAL ON IV ANTICOAGULANTS. SHE IS POTENTIALLY GOING TO BE DISCHARGED ON 6/18 PER PATIENT.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1407804-1	As per wife's account, patient died approximately 9 days after receiving vaccine on 4/21 of massive, unknown (suspected cardiac) event. Wife explained that patient was diagnosed with Wolff-Parkinson-White (WPW) syndrome prior to being vaccinated. PCP unaware of diagnosis, and patient did not disclose that on vaccine consent form. Patient reported to his wife that he was feeling unwell after receiving vaccine including unable to move legs well. Patient later diagnosed with thrush and supposedly treated by physician. Patient did not discuss concerns regarding vaccine or WPW with physician. Patient called his wife the day of death and asked her to bring him to the doctor the following day. Wife reported that patient experienced seizures but not concerned as it was similar to childhood seizure. Wife was done at work and drove home after phone call, arriving at house approximately 10-15 minutes after call. Patient supposedly dead when wife arrived. Wife performed CPR for ~15 minutes until EMS arrived, and EMS attempted for another 40 minutes for any electrical activity. Wife declined full autopsy from ME, suspecting that it was massive MI or clot. Wife and patient unaware that WCW could place patient at risk of clots and wondering if the vaccine was partially responsible.
THROMBOSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1410329-1	Development of DVT in R axila

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0916795-1	Soreness at injection site 3 hours post injection. 30 hours post injection, pt experiences a huge blood clot from vagina, about the circumference of a clementine. was not old blood, looked like frank blood. over the next 30 hours, pt experiences more clots, much smaller, about the size of a pea. pt has a headache and chills throughout. 48+ hours, experiencing what feels like menstrual cramps. no blood clots 72 hours post injection
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0923031-1	Worsening arm pain after injection that eventually resulted in my being diagnosed with a blood clot in the arm that I recevied the injection on
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0934745-1	Resident had seizure like activity followed by a vagel response with large bowel movement. Resident then began to show signs of blood clot to left lower extremity. No pedal pulse, area on leg warm to touch. Left lower leg now cold to touch, stiff, purple and white in color. No other signs of modeling, body warm to touch, no fever noted. Respirations and pulse increased with low oxygen levels. Resident not responding to stimuli.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0938186-1	Heavy period with more bleeding. And cramping and multiple clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0946743-1	She said she received her First Covid vaccine on 12/22/2020. She said on 1/4/2021 she worked that evening and started having severe pain in calf of her left leg no redness ,no heat, just hurt to work 1/8/2021 She had surgery on her left leg. 2nd Covid Vaccine received on 1/12/2021 She said she had PT on her right leg with severe pain in her left calf She said she did not have any therapy on her left leg. Two ? three hours later she developed chills, temp 100.1 took temp again later is was 99. She had ultrasound that showed a large blood clot She said the PA told her the blood clot could be from receiving the Covid vaccine
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0949555-1	Received Pfizer vaccine, first dose on Wed. 01/13/21 between 12 and 1 P.M. Thurs. 01/14/21 in the afternoon he began to note that he had difficultly walking. Went to bed when he woke up at 5:48 A.M. he reported he had ataxia. Patient reported having to walk in tiny steps to stay upright. He went to the emergency room. Had CT scan of head and found blood clots. MRI performed. Stroke found in right PCA territory, but no loss in strength in left lower extremity. Sensation and vision intact. Strength in all four extremities is 5 out of 5.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0950996-1	Constant headache began approximately 1 hr after receiving vaccine. Awoke this morning with blood clots on my tongue
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0957555-1	I am a registered nurse at hospital. On 12/25, seven days after receiving the shot I started to get right lower leg pain and I kept complaining about it till New Years Day. I had no symptoms of a DVT. I triaged on 1/1/21 and the doctors ordered labs/imaging and the results were as followed: D-Dimer biomarker (+) , Ultrasound of the Rt lower leg (-) , CTA showed a PE (segmental right upper lobe pulmonary artery consistent with pulmonary embolus). I was discharged on Xarelto and advised to follow up with a hematologist. On 1/5/2021, I went to hematology and they did a whole bunch of labs. I was sent to get a ultrasound of the leg because the pain persist and they found a clot hidden by my soleus. The plan is to continue on the Xarelto for 6 months. Come back in 3 weeks to scan my leg again and get my lab results. On 1/12/2021, I received the 2nd shot of the Pfizer vaccination.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0957860-1	"""I received the Pfizer Covid vaccine Wed afternoon around 4pm. Thursday morning around 9:30 I started with severe pain in my left leg. The pain worsened through the day and my leg began swelling. No other symptoms at all. This morning my leg was twice the size of my right leg so I went to the ER. I live in so I'm at ED. I have a massive blood clot running the the length of my leg - from my thigh to my ankle. I'm very lucky I got here so fast! I?m a very healthy 49 year old with no history of DVT or blood clots so they dug further to find out why. A cat scan showed I have a congenital condition called May Thurner Syndrome. I?m so relieved to have an answer and it?s fixable! The vascular doctors are not 100% convinced that?s not all that was going on as I was born with the syndrome and I?ve gone this long without a clot. So they are doing lots of labs to see if anything else shows up. This is where we are at. I?m being admitted to take care of the clot."""
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0959469-1	Employee received her 2nd Pfizer vaccine on 1/12/2021. Twelve hours later she developed a fever with a T-max of 102 degrees. The fever lasted 18 hours. She also complained of severe malaise and fatigue lasting 36 hours. On day 3 she developed left lower extremity pain that persisted over the weekend. She was seen in Urgent Care today where she was diagnosed with a thrombosis. She also came to the ETC and was ruled out for a PE The employee does have a family history of antiphospholipid in her mother. She has a follow up with her PCP and will also receive a work up in hematology as she had never been tested for any genetic coagulation conditions after her mom's diagnosis.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0961282-1	Felt like a restless leg or blood clot but not as severe.; Felt like a restless leg or blood clot but not as severe.; Left below knee pain and discomfort. Some to the right leg as well.; Left below knee pain and discomfort. Some to the right leg as well.; Checked back of leg for warmth or bruising lasted day 2 and day 3.; Checked back of leg for warmth or bruising lasted day 2 and day 3.; Did feel some discomfort to left arm as a result of the shot in left arm, tender and could feel when holding arm up some discomfort; This is a spontaneous report from a contactable nurse reporting for herself. A 49-years-old female patient started to receive bnt162b2 (BNT162B2; Lot: ECO142) vaccine , intramuscular in the left arm on 29Dec2020 15:45 at single dose for Covid-19 immunisation . Medical history included hypertension, anaemia, blood cholesterol increased , depression, food allergy (whey casium) (taking doxepin for it with relief). Concomitant medication included doxepin (DOXEPIN), metoprolol (METOPROLOL), calcium ascorbate (VITAMIN C [CALCIUM ASCORBATE]), tocopherol (VITAMIN E [TOCOPHEROL]), cyanocobalamin (VITAMIN B-12) , atorvastatin (ATORVASTATIN), sertraline (SERTRALINE). The patient experienced felt like a restless leg or blood clot but not as severe. on 30Dec2020 07:00 with outcome of recovered , left below knee pain and discomfort. some to the right leg as well. on 30Dec2020 07:00 with outcome of recovered , checked back of leg for warmth or bruising lasted day 2 and day 3. on 30Dec2020 07:00 with outcome of recovered , did feel some discomfort to left arm as a result of the shot in left arm, tender and could feel when holding arm up some discomfort on 30Dec2020 07:00 with outcome of recovered. The event blood clot was considered serious (Important Medical Event). Course of the events The patient reported left below knee pain and discomfort. Some to the right leg as well. She checked back of leg for warmth or bruising lasted day 2 and day 3. She felt like a restless leg or blood clot but not as severe. She felt better to stand instead of sitting. Did feel some discomfort to left arm as a result of the shot in left arm, tender and could feel when holding arm up some discomfort. No shortness of breath, no nausea, no dizziness, no increased fatigue (baseline - not enough sleep - working 3 jobs).; Sender's Comments: A possible contributory effect of suspect BNT162B2 on reported leg thrombosis cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0975052-1	Blood clot, DVT of a ill art vein acute left
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0978544-1	First vaccine small bruise at her left knee and extreme pain in her neck, shoulder and left arm and missed work. Second vaccine she has 4 bruises and clots on her inner leg from ankle to thigh. So painful she can't sleep and she is having a hard time walking. Her left shoulder is limited due to extreme pain. She sought care from her physician and he is sending her for scans that the facility is requiring a deposit.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0979379-1	19th vaccinated; my wife was exposed to COVID on 26th and she developed symptoms on 29th; I developed on 31st and tested Positive on January 2 for COVID; January3, I was admitted to ER for shortness of breath, my oxygen saturation - between 88 and 94; I was in the hospital and discharged on January 8. Remdesivir and Decadron treated with and also Eliquis. I was on oxygen until Wednesday (three days) and then I went home on 8th and continued on Prednisone for last week (Friday). Haven't gone back to work - shortness of breath, fatigue and headaches continue.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0979775-1	Employee states that she received Dose #2 the day after the end of her menstrual cycle and that she began bleeding again after receiving her second dose and that it was a heavy and clotted cycle. She was advised to contact her OB/GYN
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0983350-1	Patient experienced a blood clot on January 11th. She is unsure if it is related to the vaccine or not.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0986749-1	5:30 am I found husband by bed, babbling, Called 911, Had Lt Hemi Stroke. Aphasic, Rt side limp, given TPA. Sent to ICU. Recovered within 2hrs, speech, movement of extremities. lt hemi clot found on ct angiogram & mri. 2nd mri found clot busted with residual. transfered to telemetry next nite. echo unconclusive. 02 sats low, venogram done 3days later show lt dvt, lung ct wnl. ld asa & b/p meds were given. blood work to be drawn for baseline prior to anticoagulent therapy. possible d/c 9/30.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				On day 04 After receiving the 2nd Covid 19 Pfizer vaccine, I experienced a deep pain in my upper inner right thigh around 0600 am of the 16th of January. At the same time my calf muscle of that same right leg was in pain. I thought it to be muscular pain as I was told by the reading material regarding symptoms of the covid 19 vaccine there could be aches an pains attributed to the 2nd shot. I used a deep heating rub on both areas of my right leg and put heat on them to treat the soreness. This occured over the next few days feeling this pain in my right leg and treated the same way applying heat. Within about 24 hours of the pain in my legs I started experiencing a shortness of breath while walking up and down st. On day 04 After receiving the 2nd Covid 19 Pfizer vaccine, I experienced a deep pain in my upper inner right thigh around 0600 am of the 16th of January. At the same time my calf muscle of that same right leg was in pain. I thought it to be muscular pain as I was told by the reading material regarding symptoms of the covid 19 vaccine there could be aches an pains attributed to the 2nd shot. I used a deep heating rub on both areas of my right leg and put heat on them to treat the soreness. This occured over the next few days feeling this pain in my right leg and treated the same way applying heat. Within about 24 hours of the pain in my legs I started experiencing a shortness of breath while walking up and down stairs as part of my daily activities. The symptoms of shortness of breath only seemed to appear when I was on any form of extended walking activity or physical movements or exercise this would of have been starting around the 17th of January. The right leg pain was masked by the heating rub while the shortness of breath continued for the next few days. On the 19th of January, I went to an Urgent Care Facility at 0800 am to see a Medical professional to discuss my symptoms I was previously experiencing and to figure out why I was having a shortness of breath and the pain in my right leg. The on staff Physician's Assistant had a Nurse conduct a Covid 19 Rapids test (negative) and a second swab was administered and sent to the Lab. Which produced a (negative for Covid 19) on the 20th of January. An Xray was not taken to determine my shortness of breath. The Dr listened to my lungs and heart, though I did let the PA know I had received both Pfizer shots and when they were administered. I was carrying My Shot record for the vaccine with the dates and lot number. He didnt appear to be interested in further diagnosis and made sure I had the paperwork to track the results of my Covid swab sent to the lab. The visit was completed and I was released to go back to work/home. I carried on the symptoms of the shortness of breath from the 19th of January to the 22nd of January monitoring my O2 (oxygen levels) with a pulse oximeter. They ranged from 90-93. On the evening of the 22nd of January I was becoming very uncomfortable with my breathing climbing the stairs in my home and monitored my O2 readings with the oximeter on my finger when walking upstairs and they dropped down to 60-65. My wife drove me to the emergency room at Hospital. I walked into the ER and checked myself in for shortness of breath and leg pain in my right leg. I was admitted into the emergency room and put on 15 litres of oxygen. The emergency room Dr ordered a chest Xray, Cat Scan of my chest and heart and a sonogram of my right leg. The testing results came back with a noted large pulmonary emboli on my lungs/heart area and blood clots throughout my right leg (right lower extremity DVT). Surgery was performed to remove the pulmonary emboli and I was put on a Heprin Drip to thin out my blood for the remaining clots in my chest and my right leg. The attending physician ordered a T.E.E. (transesophageal Echocardiogram) to observe my heart functions. I was told the TEE did not demonstrate any further recommend surgeries. I was administered a test for my cardio and heart functions prior to my discharge on 27 January 2021. I was prescribed Eloquis for maintenance medication to be monitored by followup visits to my cardiologist and Hemotologist within the next few weeks. I have not family or personal history of Blood Clots. I do perform administrative duties as part of my job, which requires sitting and working on a computer. Note however I move around very often during a work day and operate a farm tractor and have an active life style. I have not recently had any extended trips more than 2 hours nor have I flown within the past year or so.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	0988076-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	0990853-1	Pfizer Covid 19 vaccine treatment under Emergency Use Authorization(EUA): My menstural cycle has changed and I have started with old brown bleeding with clots 10 days prior to my normal cycle. I have never had this happen and have never had any issues with my periods. Will be seeking medical treatment on 1/27/21 with OBGYN.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	0992022-1	Superficial blood clot in right medial knee area. Aspirin, elevate right leg, compression and heat.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1000489-1	I may have a blood clot in my leg.; This is a spontaneous report from a contactable consumer (patient). A male patient of an unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not provided), via an unspecified route of administration on 20Jan2021 at a single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient got the first dose of the vaccine on 20Jan2021 and is going to visit his doctor because they think that he may have a blood clot on his leg on an unspecified date. He wanted to know if it was okay if they give him a shot of something like contrast to see the ultrasound. The outcome of the event was unknown. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1000666-1	confirmed about a blood clot/check me for a blood clot, I didn't have one but it was kind of like a thrombosis kind of thing; Both of my legs, lower part of my legs were swollen that night when I went to bed and they got really red, almost purply red; Both of my legs, lower part of my legs were swollen that night when I went to bed and they got really red, almost purply red; This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; lot number and expiry date: not known), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The consumer stated that after the first vaccine and it might be coincidental, the patient was not sure because the patient didn't know what to expect. Both of the patient's legs, lower part of the patient's legs was swollen that night (unspecified date) when the patient went to bed and they got really red, almost purply red and the patient stayed that way for a couple of days. The patient went to see the doctor two days later and doctor confirmed about a blood clot, so they did check the patient for a blood clot; the patient didn't have one, but it was kind of like a thrombosis kind of thing. So, the doctor put the patient on antibiotics for that. The patient asked if it is related to the shot, the patient didn't know it happened the same day the patient got the shot; so, the patient didn't know about that. The outcome of the events was unknown. Information on the Lot/Batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1000670-1	she was hurting at her chest/ Chest pain; on her left arm hurt real bad that's what the clot on her left arm; on her left arm hurt real bad that's what the clot on her left arm; She passed away; heart attack; This is a spontaneous report from a contactable consumer. An 87-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 19Jan2021 at single dose for COVID-19 immunisation. Medical history included diabetes mellitus, for which she was taking a pill like an hour before she would take her meal. On Monday (Jan2021) the patient experienced was hurting at her chest/ chest pain, her left arm hurt real bad as she had a blockage in her left arm/clot on her left arm, and they wanted to put in a stent and after the surgery it went well and she all go home in two days. The patient was hospitalized in Jan2021 due to the events. She had a heart attack and that the chamber between the dividers had a hole in it and her heart tissue was too thin so much thin she couldn't repair it. The patient passed away on 26Jan2021. The patient was tested negative for COVID-19 on unknown date. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: She passed away
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1004777-1	Headache the next morning after receiving vaccine that hasn?t gone away, fluid in left lung, blood clots in right lung and leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1012703-1	1/14/2021-0545, blood noted left and right ear. 0715, vomited x 1. Covid Antigen positive. Acute MD visit-basilar crackles right and coughing. Increased confusion.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1017411-1	patient developed blood clot in her left groin one week after getting first COVID19 vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1020333-1	Had swelling in leg for a couple of days prior to the early morning of 1/31 where the knee and entire lower leg was swollen and painful and made it very painful to walk. After sending photo to primary care physician had mom evaluated for a blood clot. It was determined based on blood test, vascular ultrasound that right non occlusive (mid) Femoral vein DVT.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1023638-1	Dose #2 - Nose bleeds following the 2nd Pfizer COVID-19 vaccine beginning a few days later and lasting at least (1) one week; on 2/2/2021 this patient coughed up a pea-sized bright red blood clot; no blood clots noted since that one time and no petechiae noted; severe body and joint aches and pain beginning the next day and lasting 24 hours. No fever noted on any day. Dose #1 - Nausea / diarrhea beginning a few days later and lasting two and one-half (2 1/2) weeks; sore arm same day and lasting at least 3 days. No fever noted on any day.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1024728-1	"On 2/11/2021, after about an hour the vaccine was administered to the patient and with about minutes left of his dialysis treatment. , the patient reported to staff that he was feeling nauseous and then spit out a 3/4 quarter sized blood clot. Pt was almost done with his dialysis treatment and his time was cut short by 3 minutes. Pt then verbalized that he felt ""fine."" The RN working with the patient notified the on-call nephrologist and 911 was called. Pt refused to go to the hospital. On following up with patient and RCM on 2/10/2021, patient reported he notified his nephrologist and CXR was scheduled. As of 2/11/2021, pt stated he is not feeling well and did not arrive for scheduled hemodialysis. RCM aware."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1026980-1	Patient reported to Emergency room on 01/23/2021 with complaint of nausea. According to ER record patient reported he received a COVID 19 vaccine Pfizer the day before. Work up in the ER (CT ABD PELVIS) reveal a clotted of SMA. CT CHEST REVEALED BILATERAL PULMONARY EMBOLUS. THE PATIENT WAS TRANSFERRED TO THE STATE HOSPITAL. HE WAS SCHEDULED FOR EMERGENT VASCULAR SURGERY WHICH WAS CANCELLED AS THE PATIENT DIED SHORTLY AFTER HIS ARRIVAL.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1029515-1	Superficial blood clot in right lower leg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1030521-1	I am 77 yrs old, male , , 150# and 5ft 6inches...Within 24 hours of receiving the shot, my body thru-up clots in my left leg. and I spent 4 days in the hospital (3 days in surgical ICU), breaking up the clots to save the leg. This is too much of a coincidence to ignore and I will not receive the 2nd shot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1031120-1	Fever, Diarrhea, 1/20/21 SYMPTOMS BEGAN. ADMITTED TO HOSPITAL & CONTACTED EOH 1/27/21 WITH FEVER, CHILLS,SOB, COUGH., DIARRHEA, BLOOD CLOTS, FLUID AROUND HEART. Narrative: Other Relevant History:
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1032269-1	Blood clot right thigh
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1033682-1	L hand edema, hematoma which burst and caused bleeding sending pt to the ER for pressure dressing and 2 stitches. L hand and arm progressively got more edematous and bruised looking (severely black/blue/purple) and the hand continued to bleed and swell on 2/6/21. Severe arterial and venous issues and apparent blood clots. On 2/7/21 there were also lumps noted on left inner thigh. Pt. stopped eating or drinking on 2/8/21 and expired on 2/12/21.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1035850-1	Patient woke up on the morning of 2/6 with symptoms of a stroke. Rushed to hospital where clot found in brain. Recovered from initial stroke but then had another major stroke on 2/8 and never recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1035867-1	Noticed swelling in my left hand around Feb 6th, but by Feb 13th entire left arm was swollen and warm and observed blue/purple skin coloring on upper arm and lower arm was red. Went to ER on Feb 15th, and ultrasound revealed a blood clot near the junction of the subclavian vein and cephalic veins. Diagnosis was idiopathic subclavian deep vein thrombosis (DVT). Started on enoxaparin and warfarin on Feb 15th. Referred to hematology and coumadin clinic to monitor INR.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1036464-1	Headache, nausea, vomiting, sore arm, temp 102.8, chills began 1/08/2021 @ 9pm and lasted for 24 hours The next menstrual cycle was very heavy period, soaking heavy flow tampons every hour, large clots, and lasted 6 days from a typical mild flow 3 day period
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1037837-1	ultrasound revealed blood clots at right leg; Swollen ankle and foot after 5-6 days of first dose; Swollen ankle and foot after 5-6 days of first dose; This is a spontaneous report from a contactable consumer reported for herself. A 73-year-old female patient received first dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration in left arm on 20Jan2021 11:00 at single dose for COVID-19 immunisation. The patient was vaccinated in nursing home/senior living facility. Medical history included high blood pressure, monitored for lymph node growth. The patient had no known allergies. Prior to vaccination, the patient was not diagnosed with COVID-19. Concomitant medication included high blood pressure medication and clonazepam. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced swollen ankle and foot on 25Jan2021 12:00 AM after 5-6 days of first dose, ultrasound revealed blood clots at right leg. Doctor ordered to take apixaban (ELIQUIS) immediately during second week. The events resulted in Doctor or other healthcare professional office/clinic visit. Since the vaccination, the patient had not been tested for COVID-19. The outcome of events was not recovered. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1038473-1	1 week after receiving first dose of Pfizer COVID vaccine (received 1/26/2021), develop worsening shortness of breath. Presented to ER on 2/4/2021. Found to have submassive pulmonary embolism with evidence of right heart strain, US showed left lower extremity DVT. Also found to have descending aortic thrombus with extensive clot burden. Was hypoxic 89% on room air.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1039940-1	Blood clot in her leg; This is a spontaneous report from a non-contactable consumer (patient). This female patient of unspecified age (reported only as 72) received the first single dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unknown date, for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient believed she may have a blood clot in her leg since an unspecified date. She had the second vaccine dose scheduled on an unknown date. Event outcome was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1040552-1	experienced crippling lower abdominal pain followed by very very heavy bleeding and clotting; experienced crippling lower abdominal pain followed by very very heavy bleeding and clotting; experienced crippling lower abdominal pain followed by very very heavy bleeding and clotting; This is a spontaneous report from a contactable healthcare professional (patient). A 32-years-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; batch/lot number and expiration date unknown), via an unspecified route of administration in the left arm on 05Jan2021 10:15 at a single dose for COVID-19 immunization at a hospital. Medical history included idiopathic hypersomnia and allergies to eggplant. The patient has no covid prior to vaccination. Concomitant medications included methylphenidate hydrochloride (CONCERTA), methenamine, levonorgestrel (MIRENA), propranolol hydrochloride (PROPRANOL) and vitamin c [ascorbic acid] (VITAMIN C [ASCORBIC ACID]). The patient had no other vaccine in four weeks. The patient previously took cefazolin and levaquin and experienced allergies to both. The patient has a Mirena IUD and never get her period at all. Within 6 hours of her first Pfizer injection (02Feb2021 16:00), the patient experienced crippling lower abdominal pain followed by very very heavy bleeding and clotting. Resolved within 48 hours as quickly as it came and didn't occur with second dose 1 week out. The patient was not tested for Covid post vaccination. The events were not treated. The outcome of the events was recovered on 04Feb2021. Information on the lot/batch number has been requested.; Sender's Comments: Based on available information and known drug profile it is unlikely that the reported events were causally related to BNT162B2. These are intercurrent conditions. A contributory role of the patient's Mirena IUD should be evaluated. Case will be reassessed if additional information is received. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1044020-1	Flu like symptons about 4 hrs after vaccine(Thurs)...left arm, muscles, joints hurt, chills, tired, not feeling well. most symptoms disappeared after 24 hr however right ankle and leg continue to hurt (FRI and SAT) and by Sun. calf on right leg hot red and swollen. Mon called my physician and went to hospital for sonogram on right leg. Results were a blood clot on the back of my leg from ankle to knee. Xarelto was prescribed as blood thinner and taken. Tuesday appt. with Doctor for consultation and blood work was prescribed with follow up appointment March 2 . My concern is taking second dose of Pfizer vaccine on Mar 4. My doctor and I will discuss. On internet others have reported this same side affect of blood clots from 2 days to 2 wks after. This needs to be addressed.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1044071-1	I had a major stroke on 12/25/2020, 9 days after receiving the vaccine. I had a clot in my MCA. My left arm was completely flaccid, with left facial droop and garbled speech. Thankfully I was able to receive TPA that day, which resolved all of my symotoms.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1044102-1	Shortness of Breath causes by multiple large blood clots, put on blood thinners, seems to be fine, but has follow up appointment in March
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1044799-1	Blood clots in left leg diagnosed in Hospital, , emergency room 02/08/2021, about 5 hours after first vaccine. Treated with eliquis. History of prior embolus and blood clots. Referred to primary physician who referred me to a vascular surgeon and a hematologist. For about 3 days extremely tired and slept quite a bit. I already have rotator cuff injury of arm where I got shot so pain was not able to be separated from rotator cuff pain. No temperature, rash, redness or swelling of injection site.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1045120-1	extreme muscle pain for 24 hrs . started immediately as minimal and increase to extreme within 3 hrs. face felt numb for about 20 minutes after receiving shot that lasted 15-20minutes. there was a tingling sensation present during the numbness in the center of my forehead. possibly related: experienced light nosebleed day after shot. three days after shot experienced a heavy nosebleed with very large clotting.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1048205-1	he has chronic myeloid leukemia - white blood cell count is higher; His oxygen is low; Blood clot in his legs; first dose on 19Jan2021 and second dose on 02Feb2021; first dose on 19Jan2021 and second dose on 02Feb2021; This is a spontaneous report from a contactable nurse reported for her father that an 82-year-old male patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 02Feb2021 at single dose for covid-19 immunisation. Medical history included chronic myeloid leukemia (CML). Concomitant medications were not reported. Patient previously received first dose of BNT162B2 on 19Jan2021 at single dose for covid-19 immunisation. Patient received Pfizer-BioNTech COVID-19 Vaccine first dose on 19Jan2021 and second dose on 02Feb2021. Two days after second dose (04Feb2021) he had to go to the hospital - he had chronic myeloid leukemia - white blood cell count was higher (Feb2021), he had been in the hospital for the last 3 days as of 08Feb2021, his oxygen was low, had a blood clot in his legs, had a bone marrow biopsy, all on an unspecified date in Feb2021. The nurse is taking its related to this vaccine and assessed this case as not serious. Outcome of the events was unknown. Information on the lot/batch number has been requested.; Sender's Comments: A possible contributory effect of suspect BNT162B2 on reported events cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1048691-1	White blood cells went up greatly; Blood clots in his legs; oxygen went down/oxygen levels went down; This is a spontaneous report from a contactable consumer (patient's daughter). A male patient of an unspecified age received second dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot/batch number and expiry date were not provided), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. Medical history included he had Leukemia and was on treatment. The patient's concomitant medications were not reported. The patient took first dose bnt162b2 for COVID-19 immunization. Caller says that her father got his second COVID vaccine from Pfizer. He wound up in the hospital 3 days later. His white blood cells went up greatly and he has blood clots in his legs. They do not know what is happening. They are very worried about him. He is getting a bone marrow test today (08Feb2021). His oxygen went down yesterday (07Feb2021) and is now on oxygen. She is freaking out about it. She really thinks it has to do with vaccine. It was asked if there was any information on other people with Leukemia getting the vaccine. If what research there was on this. They need all the help and information they can get. Every minute counts as everything getting worse. He is now on oxygen as his oxygen levels went down. He is getting worse every hour. The patient underwent lab tests and procedures which included oxygen saturation: his oxygen went down yesterday/oxygen levels went down on 07Feb2021, white blood cell count: his white blood cells went up greatly on an unknown date, biopsy bone marrow: unknown results on 08Feb2021. Reporter seriousness for White blood cells went and Blood clots in his legs up greatly was hospitalization. Therapeutic measures were taken as a result of oxygen went down. The outcome of the events was not recovered. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1049035-1	On Sunday, February 21 at 07:42 AM I received a call from patient advising me she had called for an ambulance, she was awoken from her sleep with a rapid heart beat and was also suffering from shortness of breath. Patient was taken to Hospital where she was admitted due to a Pulmonary Embolism, a small clot was found in her lung.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1049150-1	Feb 9, patient was light-headed, as if he was going to faint. He did not have appetite. Evening of Feb 9, started vomiting large amounts of blood. Ambulance took him to hospital. CT scans showed abnormally enlarged pancreas. Patient aspirated blood and was put on a ventilator for 48 hrs. Endoscopic ultrasound showed ulcers in stomach that appear to have been bleeding, which were clipped and shot with epinephrine. After being treated for ulcers, patient developed blood clots in leg and lungs. It is almost two weeks since initial emergency, and patient is still showing sings of internal bleeding (low blood pressure, low hemoglobin, blood in stool). Still no firm explanation for continued bleeding. Before adverse event on Feb 9, patient did not report other symptoms from shot, however, he did show unusual signs of large bruising on his arm. Patient is currently at Hospital. Blood thinners are being discontinued, but patient still has blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1049931-1	2 days after vaccination right thighs and right leg started to have pain similar to pain I got with polymyalgia rheumatica that I have had in past and I though it was a flare up. A couple of days after that right leg started to swell and was warm. Went to ER on 1/29/2021 and had doppler which showed blood clots from groin area to below the knee. I was placed on Xarelto at that time. Got 2nd dose and about 24 hours latter back of thighs started to hurt again. Called doctor and applied heat and elevation and had no new swelling but I remain on Xeralto. Not sure if it was related but I have had no history of blood clots. 2 years previously broke my right patella and was with brace and no weight bearing for about 8 weeks and brace for 12 weeks without clot issues. Traveled by car a couple thousand miles last year and no issues.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1051004-1	blood clot; This is a spontaneous report from a contactable consumer. A 6-decade-old female patient (in her 50's) received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date at a single dose as vaccine. Medical history and concomitant medications were not reported. It was noted that one of the reporter's co-worker developed a blood clot after receiving the first dose of the Pfizer vaccine on an unspecified date. Any information or data on someone developing blood clots? The outcome of the event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1052242-1	Patient reported as being altered, GCS 6 with noted aphasia around 1415.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1052844-1	Pfizer-BioNTech COVID-19 Vaccine EUA: five days after vaccination patient presented to emergency department with left-sided facial droop, right gaze preference, and left hemi-paresis. Patient diagnosed with right middle cerebral artery occlusion, likely thrombotic etiology, admitted to hospital, and underwent thrombectomy and recanalization with symptom improvement. Discharged to home improved, stable, with vital signs within normal ranges two days after arrival to emergency department.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1053606-1	Within a week after receiving the first round of vaccine I have developed two blood clots in my left leg.; This is a spontaneous report from a contactable consumer (patient) A 53-year-old male patient received first dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number: EL3248) on 21Jan2021 at single dose via an unspecified route of administration on right arm for COVID-19 immunization. The patient didn't have medical history or concomitant medications. On an unspecified date in Jan2021 within a week after receiving vaccine patient developed two blood clots in his left leg. As treatment patient was currently on blood thinners. At the time of the reporting event outcome was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1054591-1	Stroke on January 20th, 2021. Unknown cause for blood clot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1054874-1	Blood clot large in leg DVT and PE both lungs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1059421-1	After the second vaccine dose she reported not feeling well with unspecified symptoms for a few days. On February 18th, 2021 she visited her doctor with numbness in her hand. They thought it may be carpal tunnel and sent her home. The morning or March 18th , 2021 she had a severe stroke and was transferred to Hospital and then to other hospital. She was in the hospital until Tuesday March 23rd when she was transferred back to her home for hospice care. She died on March 26th, 2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1060991-1	Blood clot in left leg, inside next to the knee
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1061209-1	Sharp pain behind right knee. Subsided by evening. Went to The ER Sunday morning, Feb. 21. They x-rayed leg and found nothing unusual for a woman my age. Then they conducted a scan and discovered a blood clot behind my knee. Since then, I have had intermittent pain. The Doctor referred me to a hematologist which I will see this week. Prior to the vaccination have never had a blood clot. It occurred three days after the vaccination. Coincidence? I think not.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1065118-1	Blood clot in lower left leg; This is a spontaneous report from a contactable consumer (patient herself). A 58-year-old female patient received BNT162B2 (Pfizer-BioNTech COVID-19 mRNA vaccine), via an unspecified route of administration on the left arm, at age 58 years, first dose on 23Jan2021 14:00 at single dose for COVID-19 vaccination. Medical history was reported as none. Patient had no known allergies. The patient is not pregnant. There were no concomitant medications. There were no other vaccines administered in four weeks and no other medications taken in two weeks. On 30Jan2021, patient experienced blood clot in lower left leg. ER doc indicated that patient had no reason to have formed a clot given the medical history, health, weight, age, diet, no surgeries, no prior injury, etc. Facility type vaccine was administered at a Public Health Clinic/Veterans Administration facility. Event resulted in emergency room/department or urgent care. Therapeutic measures were taken as a result of the event includes rivaroxaban (XARELTO) as blood thinner. The outcome of the event was not recovered. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1065435-1	blood clot; death cause: Heart Problems; tired; nauseous; This is a spontaneous report from a contactable consumer. An 81-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (Lot number EL3248), via an unspecified route of administration at single dose in the left arm on 19Jan2021 14:00 for covid-19 immunisation. Medical history included heart problems, pacemaker. Concomitant medication included heparin. The patient experienced death cause: heart problems on 20Jan2021, blood clot on an unspecified date with outcome of unknown that required hospitalization, tired on 19Jan2021 with outcome of unknown, nauseous on 19Jan2021 with outcome of unknown. The patient was hospitalized for blood clot from 16Jan2021 to 18Jan2021. The patient died on 20Jan2021. An autopsy was not performed. The events were described as follows: The patient was tired and nauseous about 3 hours after her vaccine. She had been in the hospital 16Jan2021 to 18Jan2021 for a blood clot. The patient died at her home on 20Jan2021 between 4 and 7 pm. No treatment required. The vaccine was administered at Hospital Facility. Prior to vaccination, the patient was not diagnosed with COVID-19 and since the vaccination, the patient had not been tested for COVID-19.; Reported Cause(s) of Death: death cause: Heart Problems
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1066342-1	I ended my normally scheduled menstrual cycle on 02/06/2021, and exactly 6 days later (02/15/2021) I had severe menstrual bleeding with blood clots that has continued to this day (03/02/2021). I went to urgent care and was prescribed Provera to stop the bleeding to no avail. I then proceeded to go to urgent care again because the bleeding had not stopped. At my second Doctors visit, I had an abdominal and a transvaginal ultrasound done, and both of them came back completely normal. I have now scheduled an upcoming obgyn appointment, but cannot be seen until 03/09/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1068264-1	"bilateral blood clots in legs that then went to lungs then went into BLAST crisis; bilateral blood clots in legs that then went to lungs then went into BLAST crisis; bilateral blood clots in legs that then went to lungs then went into BLAST crisis; Patient with myeloid leukaemia (CML) and WBCs abnormal received BNT162B2; Patient with myeloid leukaemia (CML) and WBCs abnormal received BNT162B2; This is a spontaneous report from a contactable consumer (patient). An 82-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) Solution for injection (lot number and expiration date unknown) via an unspecified route of administration in the left arm on 02Feb2021, 13:00PM (at 82 years old) at a single dose for COVID-19 immunization. The patient was vaccinated in the Nursing Home/Senior Living Facility. The patient's medical history included myeloid leukaemia (CML) diagnosed a few months earlier, WBCs abnormal, blood pressure, and heart surgery valve repair. The patient has no known allergies. Concomitant medications included atorvastatin, carvedilol, vitamin d nos, dasatinib monohydrate (SPRYCEL) from an unknown date to help get WBCs back in normal range; cyanocobalamin (VITAMIN B12), and cetirizine hydrochloride (ZYRTEC), all were received within 2 weeks of vaccination. At 82 years old, the patient received the first dose of BNT162B2 (lot number and expiration date unknown) via an unspecified route of administration in the left arm on 12Jan2021 at a single dose for COVID-19 immunization. The patient was not diagnosed with COVID prior to vaccination and did not receive any other vaccines within 4 weeks prior to BNT162B2. It was unknown if patient was tested for COVID post vaccination. On Friday (unknown date in Feb2021), patient's WBC was 150,000 and on Saturday (unknown date in Feb2021), patient had bilateral blood clots in legs that then went to lungs then went into BLAST crisis. The adverse events resulted in emergency room/department or urgent care as well as hospitalization due to life threatening illness (immediate risk of death from the events). The patient was hospitalized for 12 days. Therapeutic measures which include steroids, blood thinners, and lots of other meds were administered. Outcome of the events ""patient's WBC was 150,000"" and ""bilateral blood clots in legs that then went to lungs then went into BLAST crisis"" was recovering. Information on the lot/ batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1068762-1	"DEATH Narrative: patient's wife reported he had gone in an outside hospital, had held his brilinta as advised anticipating shoulder surgery ""and he threw a big clot and died.""""

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1070714-1	"she was hospitalized 2.5 days after having symptoms of a ""massive heart attack"" 2 days after the vaccine; blood clot; pain on the left side/pain so bad/ pain was so severe couldn't bent over and couldn't get up; pain was in the heart and underneath the rib; pain was in the heart and underneath the rib; had a little trouble breathing; broke out in a sweat; This is a spontaneous report from a Pfizer-sponsored program Pfizer First Connect from a contactable consumer (patient). A 77-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number/expiration date unknown), via an unspecified route of administration on an unspecified date in Feb2021 at a single dose for COVID-19 immunization. Medical history included asthmatic, had double pneumonia years ago, has ongoing atrial fibrillation and a pacemaker to control it as the only problem she has in her heart. The patient's concomitant medications were not reported. On an unspecified date in Feb2021, the patient was hospitalized for 2.5 days after having symptoms of a massive heart attack two days after the vaccine. It was reported that two days post vaccination, the patient was rushed to the hospital by ambulance as they thought she was having a massive heart attack. She added her symptoms lasted for hours, she had pain so bad that she was bent over and had a little trouble breathing. She said the pain was in the heart and underneath the rib, on the left side. She later mentioned she did not feel like it was AFib. She said they did every test possible and listed the following ones: ultrasound, stress test, x-ray (unknown results), and blood work in Feb2021. She specified that her blood work indicated a blood clot. She mentioned she was told by her personal friend, who was a doctor, that by having the shot sometimes it indicated you have blood clots when you really don't. The patient also specified that she broke out in a sweat, the pain was so severe that she could not bend over and could not get up; therefore, they treated her as if she was having a massive heart attack for 12 hours at the hospital. It was reported that they kept giving her stuff to stop the effect with her heart. She mentioned her second dose was scheduled on 25Feb2021. She asked if her experience was reported as a side effect to the vaccine and should she get the second dose of the vaccine. She explained the doctors did not know what she had, and she needed to determine if she can receive the second dose. She also asked if we could notify her if a similar reaction is reported. The outcome of the events was unknown. Information on lot/batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1070750-1	very high Troponin levels (4.79 & 12.49); repeat episodes of chest pain; heart attack; thrombus formation; coagulopathy; myocardial infarction; This is a spontaneous report from a contactable nurse (reporting for herself). A 44-years-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot# ELI283), via an unspecified route of administration single dose on 12Jan2021 for covid-19 immunisation. First dose was received on 22Dec2020 10:00 left arm (lot# EJI695). Medical history included chronic skin condition. No other vaccine was received in four weeks. Concomitant medications included oxycodone for pain, colecalciferol (VITAMIN D). The patient experienced myocardial infarction on 10Feb2021 20:00, very high troponin levels (4.79 & 12.49) on 11Feb2021, repeat episodes of chest pain on 10Feb2021 20:00, heart attack on 10Feb2021 20:00, thrombus formation on 10Feb2021 20:00, coagulopathy on 10Feb2021 20:00. The patient was hospitalized due to the events from 11Feb2021. Patient reported to be a healthy 44 year old woman with no history of hypertension or high cholesterol. She was now on 7 different medications to protect her heart, including blood thinners and antihypertensives that result in daily headaches and fatigue. She had had several repeat episodes of chest pain which has required taking sublingual nitroglycerin. She was off work for an undetermined amount of time. Her primary care physician was baffled by her case and reached out to a number of experts who have agreed that her heart attack was caused by a thrombus formation/coagulopathy most definitely related to the Covid vaccines she received. The events outcome was recovered with sequelae.; Sender's Comments: Based on available information, a possible contributory role of the subject product, BNT162B2 vaccine, cannot be excluded for the reported event of myocardial infarction and other events. However, the reported events may possibly represent intercurrent medical conditions in this 44-years-old patient. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics including EKG at baseline and during subject drug therapy, echocardiogram, electrolytes, chemistry panel and coronary angiogram, and concomitant medications. This case will be reassessed once additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1070763-1	large knot right above the injection site/size of a half dollar/size of a silver dollar and where the needle went in was right at the base of that knot/like egg under the skin or clot; felt like she got a flu shot and could tell her arm was very sore; swelling started almost immediately after the shot/swelling was exactly underneath the injection site, just in the pronounced area/about 4 inches wide and about 2 inches high; thought maybe someone has hit a vein because it did bleed and ran down her arm; This is a spontaneous report from a contactable consumer. A 66-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for injection, lot number: EN6201, unknown expiration), via an unspecified route of administration on 18Feb2021 at 04:30 at a single dose for COVID-19 immunization. Medical history reported as none. Concomitant medications included tolterodine and adalimumab (HUMIRA). The patient reported that she received the COVID vaccine on 18Feb2021 at around 4:30. Firstly it was fine, she felt like she got a flu shot and could tell her arm was very sore. What concerned her was the swelling started almost immediately after the shot on 18Feb2021. She knew that was one of the symptoms. This morning and yesterday though (18Feb2021), she noticed some of the symptoms. She got a large knot that was right above the injection site. Yesterday, it was about the size of a half dollar, today it is about the size of a silver dollar and where the needle went in was right at the base of that knot. It feels like an egg under the skin or a clot. Like a knot (a hard spot underneath the skin). The patient was wondering if it was something she should be concerned about because the swelling was exactly underneath the injection site, just in the pronounced area. It was a rectangle and is about 4 inches wide and about 2 inches high and then right above that was where the needle went in. The large circle or knot was right above that and she was thinking possibly when she was given the injection she has never really bleed before. So, she thought maybe someone has hit a vein because it did bleed and ran down her arm. She got no problem. That happens at times, but she started thinking if this was a clot or it was something that is right above there because it is so pronounced, and it really hurts. The patient stated that she took some Aspirin last night and is going to take an ibuprofen in a little while. She mentioned that she takes this one pill and it has nothing to do with anything. The patient had lab work done (unknown results) that was about 3 weeks ago (2021). She also stated that she also take Humira which is a shot every 2 weeks and it said in the fact sheet that she should be concerned or be sure that she did not take any medicine that affects the immune system and Humira does weaken the immune system but she was not ask that prior to. The patient wanted to know if is that something she should be concerned about. Outcome of the events was unknown. No follow-up attempts are possible. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1072556-1	right leg showed blood clot in lower back of leg from ankle to knee; right leg calf was red swollen and throbbed; Right leg ankle still hurt; flu symptoms appeared; Left arm, muscles, joints hurt; Left arm, muscles, joints hurt; Left arm, muscles, joints hurt; tired; chills; not feeling well; This is a spontaneous report from a contactable consumer (patient). A 70-year-old female non-pregnant patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EM9810), via an unspecified route of administration on 11Feb2021 10:00 at single dose in arm left for COVID-19 immunisation. There was no medical history and no known allergies. Concomitant medication included colecalciferol (D3) and multi vitamin. The patient did not have COVID prior vaccination. There was no other vaccine in four weeks. After about 4 hours after first Pfizer shot on a Thurs (11Feb2021), the patient experienced flu symptoms appeared, Left arm, muscles, joints hurt, chills, tired, not feeling well, on 11Feb2021 16:00. Right leg ankle still hurt after 24 hours Friday and Saturday by Sunday. Right leg calf was red swollen and throbbed on 14Feb2021 16:00. Monday (on 15Feb2021) the patient called physician and had sonogram on right leg which showed blood clot in lower back of leg from ankle to knee. Doctor put immediately on rivaroxaban (XARELTO) blood thinner for blood clot. Blood work has been done on 15Feb2021. The patient would follow up on 02Mar2021. She was concerned about taking second COVID vaccine dose on 04Mar2021. The adverse events resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. Covid was not tested post vaccination. The event outcome was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1073786-1	"The following day, she complained of headache, fatigue, dizziness, and lightheadedness for 3 days. On 2/19/2021 she experienced hives on her abdomen and back that were itchy. No medicine was taken. On the 2/20/2021, she complained of further headache, fatigue, dizziness. She also had shortness of breath. The next day, she presented to the urgent care. She was given an intravenous fluid bolus for dehydration. She was given a ""migraine cocktail."" She was told that her D-dimer was positive. She had a CT scan that showed sinusitis and her chest was clear. The following day, she complained of progressively worsened ascending pain in her left arm. She had red patches of a rash in her left arm as well. One week later, she went back to the urgent care. An ultrasound was performed and a blood clot was found at. She is now taking Eliquis daily."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1075308-1	still have sore muscles in my shoulders; still feeling bad with all the above symtoms; 23Jan2021 went to get covid test but due to blood in my nose received antibody test which was neg; bad headache; winded; vomited; diaherra; coughing/cough; large clots gushed from my rt nostril went to er stopped bleeding after 2 hrs; large clots gushed from my rt nostril went to er stopped bleeding after 2 hrs; This is a spontaneous report from a contactable other hcp (patient). A 61-year-old female patient received their first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; lot number: EL1283, expiry date not reported), via an unspecified route of administration on the left arm on 20Jan2021 18:45 at single dose for COVID-19 immunization. Medical history included diabetes mellitus and high BP from an unknown date and unknown if ongoing. Concomitant medications included levothyroxine, losartan potassium and sodium fluoride (CREST CAVITY PROTECTION). The patient previously took septra and experienced allergies. The patient reported that on 21Jan2021, they woke up and brushed their teeth then at 6:40am blood with large clots gushed from their right (rt) nostril. The patient went to ER and stopped bleeding after 2 hrs. On 22Jan2021, the patient woke up with bad headache, winded, vomited, diaherra (as reported), coughing. On 23Jan2021, they went to get covid test but due to blood in their nose, they received antibody test which was negative. On 24Jan2021, the patient was still feeling bad with all the above symptoms. Then on 25Jan2021, they went to ENT doctor who cauterized their nose and stated that since antibody was negative to take the maderna vaccine since they have no antibody in their system, the Pfizer vaccine did not work. From 26Jan2021 to 13Feb2021, the patient felt same started feeling better 14Feb2021 but still have sore muscles in their shoulders and cough. The patient underwent lab tests and procedures which included sars-cov-2 antibody test: negative on 23Jan2021. The action taken in response to the event(s) for bnt162b2 was not applicable. Therapeutic measures were taken as a result of epistaxis and clot blood which include the doctor cauterized their nose. The patient recovered with sequel from the events.; Sender's Comments: Based on available information, a possible contributory role of the subject product, BNT162B2 vaccine, cannot be excluded for the epistaxis and other reported events due to temporal relationship. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics including CBC and coagulation panel, counteractive treatment measures and concomitant medications. This case will be reassessed once additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1075857-1	Blood clot on thigh
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1081108-1	Cellulitis in left leg may be acting up/Might have an infection in area of cellulitis in left leg; Cellulitis in left leg may be acting up/Might have an infection in area of cellulitis in left leg; Questioned if she might have a blood clot in left leg; Might have an infection in area of cellulitis in left leg; pain in the left leg; This is a spontaneous report from a contactable consumer. A 72-year-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/lot number: EL9262, expiration date 31May2021), via an unspecified route of administration on 01Feb2021 at left upper arm around 19:00 or a little later at single dose (at the age of 72-years-old) for covid-19 immunization. Medical history included bad knees and got injections for this and recurrent cellulitis of legs (cellulitis was not active at time of vaccine; but anytime her leg swells with bad knees she had a little episode with the cellulitis down around her ankles on both legs but left leg was more predominant. She had been keeping it at bay), mineral supplementation, bad knee pain, blood pressure medication (abnormal), urine output control (abnormal), dehydrated and dizzy. Concomitant medication included lisinopril as blood pressure medication, potassium for Mineral supplement, solifenacin succinate for urine output control, paracetamol (TYLENOL 4) for bad knee pain. She had taken potassium before and it calmed it down: she was little dehydrated, little dizzy, so was drinking some Pedialyte, water, and taking potassium. The patient had it once before but it kind of snuck up on her again. The patient called to ask if anyone had reported any type of blood clot on the same side of the body that the Pfizer COVID-19 Vaccine was administered. She questioned if she might have a blood clot in her left leg after having been administered the Pfizer COVID-19 Vaccine. She had recurrent cellulitis in both legs prior to Pfizer COVID-19 Vaccine. The doctor thought the cellulitis in left leg may be acting up and that she might have an infection in that area of cellulitis. She reported pain in the left leg when she rested her leg on a pillow; but did not feel any pain when she was standing and walking. The pain became a little too hard for her to bear so the doctor prescribed her Cephalexin 500m capsule every 12 hours-she was on the 3rd capsule now. Onset date for the events was approximately 03Feb2021 or 04Feb2021. The events were better since started Cephalexin. Second dose date scheduled for 22Feb2021 but did not give her time. The outcome of the events was recovering. Follow-up (23Feb2021): New information received from the product quality complaint group includes confirmation of lot number (EL9262) and new expiry date.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1085254-1	Severe abdominal pain unable to eat or sleep for 36 hours. He went by ambulance to the Hospital emergency room. They tried to pump his stomach but he aspirated and and went into cardiac arrest. He was revived but never regained consciousness. (The ICU Dr said that he had blood clots in his abdomen from a recent stroke. We were unaware of him having a stroke other than in 2026. The same Dr. said that he had necrosis in his lungs from aspirating. The necrosis was from his bowel dying) He was put on a ventilator and given drugs to increase his heart rate. On 3-5-21 the heart drugs were reduced and he died. I was with him when he recieved the vaccination and he was healthy, just old. I think that the shot killed him.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1087883-1	"I had my first shot of Pfizer vaccine last Friday morning (March 5). In the evening, I felt fatigued. So I went to sleep early. When I got up Saturday morning, I felt dizzy. The dizziness went away in a few minutes. In the next 36 hours, I had the angina feelings - tightness in the left chest and I felt heart be squeezed occasionally. When I stand up and walk, I felt better. Finally Sunday evening (March 7), I took a walk outside and felt symptom free. Maybe the walk helped, maybe the ""thrombosis"" had run its course before my walk. I have not had the severe angina symptoms since my PCI procedure in Feb 2018. I am sure the recent occurrence is related to the vaccine. Besides reporting the adverse event, I want to get advices whether I should receive the second shot. Thank you!"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1090240-1	Cardiac arrest; Pulmonary embolus; Renal failure; Fever; Dehydration; Not eating or drinking; COVID-19 confirmed by positive COVID-19 test / COVID pneumonia; blood clot; blood pressure was low; Respiratory arrest; Respiratory failure; Hypoxemia; ventricular tachycardia; This is a spontaneous report from a contactable nurse reporting on behalf of the husband. A 71-year-old male patient received the first single dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot EL9264) on 10Feb2021 at about 19:00 (at the age of 71 years), in left deltoid, for COVID-19 immunisation. No other vaccines were given on the same day or within 4 weeks. The patient declined flu vaccine and pneumococcal vaccine (PNEUMOVAX), he had never had another vaccine except maybe his childhood vaccines. Medical history included rotator cuff surgery and cataract removed in 2020. The patient exercised regularly, he was healthy, he walked for miles and didn't eat any non-sense, he did not eat out, he did not smoke. The patient's mother was 100 years old and fully competent. The patient had two sisters older than him, the oldest one had hypertension the second sister did not have anything that they were aware of. The patient's father lived until he was 98 years old. The patient concomitant medications were none. The patient was told to take vitamin D 50,000 units but didn't even take them (he still had 9 of them in the bottle and they gave him 13). The patient experienced fever on 11Feb2021, renal failure on 14Feb2021, pulmonary embolus on 28Feb2021, cardiac arrest on 04Mar2021, dehydration and not eating or drinking on an unspecified date in Feb2021. These events required ER visit and were reported as serious as involved hospitalization from 14Feb2021 to 04Mar2021 and as fatal events. The patient died on 04Mar2021. Clinical course of the events included the following information. The patient received the first vaccine on 10Feb2021, the next day he developed a fever. The reporter spoke with the patient's doctor who told to give the patient paracetamol (TYLENOL) thinking the fever was from the vaccine. On 12Feb2021 and 13Feb2021, the patient's temperature was 102. Then the doctor advised to take the patient to the hospital. The patient's temperature was still 102, he was in renal failure, and they had to dialyze him. The patient was otherwise healthy, the patient's last physical was in Dec2020 and the only thing it showed was that his A1C was 5.7. The patient had no cholesterol or hypertension. The doctor advised the patient to decrease sugar and carbs because the holidays were coming up. The patient's follow up was scheduled on Mar2021. The reporter felt that the vaccine has something to do with the patient renal failure. The reporter spoke with the doctors at the hospital who didn't want to commit to anything. The reporter believed this was an adverse event. The caller mentioned that she had her vaccine before and she was fine. The patient was admitted on 14Feb2021 and by Wednesday he was not eating or drinking, he was dehydrated. The patient's admitting diagnoses was elevated temperature and ruling out COVID. The patient tested positive for Covid on 14Feb2021 (COVID-19 PCR test). The patient's temperature was 99.8 and then kept creeping up, on Saturday it was 102. The caller gave the patient Tylenol cold and flu (lot T0CL001021, expiry date Oct2021) took the edge off but in three hours the temperature was back up again. The patient never complained of pain and didn't want to take Tylenol. On 15Feb2021 the patient's numbers were getting better after the fluid challenge and then his numbers kept creeping up after that. The patient had the fever a week until they had it under control. The fever went away, it was gone for like 5 days, then it spiked again. The patient was started on piperacillin/tazobactam (ZOSYN) for like 3 or 5 days and the fever went away but then it kept getting worse. On 28Feb2021, the medical personnel thought the patient had a pulmonary emboli but because of the renal failure, they couldn't do a computerized tomography on the patient. The doctors mentioned that the patient was in renal failure and they thought they heparinized the patient and he had a blood clot who led to pulmonary embolus, cardiac arrest, and death. The patient was diagnosed with a pulmonary emboli on 28Feb2021. The patient started de-saturating and the doctors intubated and sedated him that whole time until this. Dialysis was started on 01Mar2021 and the patient received it every day except 04Mar2021. The patient's blood pressure was normal, it hardly ever went above 120. The patient was on the medical floor from 22Feb2021 to 04Mar2021. When the patient was on the medical surgical floor, he was on high flow 5 liters. After the patient started desaturating, he went to the intensive care unit and was put on a non-rebreather on 45%. The patient's highest heart rate was after intubation was

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				rebreather on 45%. The patient's highest heart rate was after intubation was 135, but the patient's blood pressure was low so they started him on some vasopressors. They did the fluid challenge on the patient and his labs were a little better than the labs kept creeping up until the doctor inserted a shiley catheter for dialysis. Respiratory: Respiratory arrest and then cardiac arrest. Respiratory failure, they intubated the patient. The reporter assumed dyspnea because the patient was intubated. Tachypnea was when the patient was in the intensive care unit already intubated. Hypoxemia, they intubated the patient so the caller guessed it was for the oxygen saturation drop. Covid pneumonia: yes. Chest x-ray showed mild pneumonia. The caller requested a follow up x-ray and the doctors said they were going to do another one but the caller is unsure if they did or when. The patient received additional therapies for COVID-19: remdesivir. Other radiological investigations: unable because of the patient's kidney function. They were looking at the D dimer and BMP to come up with the embolus since the patient couldn't have the scan. ARDS: no. Cardiovascular: The patient had a heart attack on 04Mar2021. The reporter thought it was from the pulmonary embolus which led to cardiac arrest. Arrhythmia: the caller guessed so, the patient was being worked on for 10 minutes before the caller got there. The caller saw a rhythm strip which showed a flat line and then she noticed ventricular tachycardia, then a flat line. The patient did not have SARS-CoV2 antibodies at diagnosis. Gastrointestinal/Hepatic, neurological, hematological, dermatological: none. Vascular: pulmonary embolus: yes, deep vein thrombosis, limb ischemia, vasculitis: no. Renal: renal failure: yes, acute kidney injury: no. The patient was scheduled for his second vaccine dose on 03Mar2021 at 04:15 but did not receive it. Time of death was 4:15 in the afternoon on 04Mar2021. The reporter considered renal failure, fever, dehydration, not eating or drinking, cardiac arrest and pulmonary embolus as fatal and related to the suspect vaccine. The outcome of the other events was unknown. Cause of death was unknown. No autopsy was performed.; Sender's Comments: Based on current information available, the company considered there is a possibility that all reported events are consequence of COVID-19 pneumonia on the basis of advanced age. The positive COVID-19 test occurred 4 days after the first injection of suspect vaccine BNT162B2. No complete effect can be achieved for short time interval. The COVID-19 is more likely pre-existing colonization or intercurrent condition, unrelated to suspect vaccine BNT162b2. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.; Reported Cause(s) of Death: Renal failure; Fever; Dehydration; Not eating or drinking; Cardiac arrest; Pulmonary embolus
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1092549-1	blood clots dx'd via ultrasound on 2-20-21
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1093361-1	Acute right index finger digital ischemia after initial complete numbness from PIP joint distally absent any previous symptoms ever. Recurrence Jan 20 of same symptoms. Suspect antibody complex clot blocking terminal branches of digital arteries.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1093491-1	Blood clot, swelling noticed a few hours aftee vaccine. Probably coincidence but felt it should be reported. Im not sure if the doctor she saw for blood clot knew she got the vaccine. It was a different clinic.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1096852-1	About 27-28 hours after injection felt strong pain in left lung area. Got progressively worse, especially at night when lying in horizontal position. The pain became extreme, could hardly breathe (inhalation causing extreme pain) and couldn't really walk on morning of March 6 (about 66 hours after vaccination. Called General Practicioner and he said to go to Emergency Rm. A catscan showed a small blood clot in each lung. Given blood thinner injection. Subsequent Ultrasound of legs showed no clots there. Released 48 hours later, put on Eliquis -- 5mg -- two each time, twice a day for 6 days, then one each time twice a day. for 3 months.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1096940-1	lost total vision in left eye, took about 2 seconds from onset. Recovered within 30 minutes. Suspect small clot. No other symptoms. No detectable lasting vision impact.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1098030-1	Woke up with sore upper right thigh Tuesday March 9. Very tender to the touch. Visited Dr. Wednesday March 10, ordered ultrasound which was completed on Thursday March 11 and diagnosed a superficial blood clot. Prescribed Xarelto 10mg.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1102512-1	I received my first dose Pfizer vaccine on 03-09-2021. I have a history of blood clot, 10 years ago. I am currently 9 weeks pregnant. I called my doctor's office who encouraged me o go to the ER since I was having symptoms of a blood clot. I went to the emergency room at Hospital where they did some blood testing. I then went to Hospital where they did more lab work and blood testing, they did an D-dimer test ad my level was at 3.23 and explained to me it could be due to my pregnancy. They ran some more tests showing I have a clot in the superficial femoral/ popliteal vein. They placed me on Lovanox medication for two days. I will be seeing a maternal fetal doctor tomorrow for further evaluation. They also monitored my baby and baby is doing fine.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1102699-1	"First Vaccine Dose on 12/26/2020 felt extremely fatigued. After 2nd vaccine dose on 01/16/2021 developed fever, chills, weakness, fatigue, body aches approximately 10 hours after received the 2nd dose. This all lasted about 24 hours. Then on 01/19/2021 which was the 2nd day of mine menstrual cycle, I had a very abnormal menstruation. I was passing ""fist"" size blood clots which made me very weak. this lasted again about 24 hours then it stopped. I had some labs drawn on 02/27/2021 which came back with very low Iron, Hematocrit and Hemoglobin levels. I usually do show signs of anemia in previous blood work but this time the levels dropped lower so i was referred to see a Hematologist."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1102925-1	I started to have tingling sensation to my left leg. I have history of DVT, hemochromatosis, hyperthyroidism but is all managed. The issue is that the tingling sensation I was experiencing was moving in my left leg, was painful but then it went away. I went to the emergency room 02-25-2021 that started with a blood clot and they placed me on Xarelto medication. My PCP encouraged me to go to the ER. they did an U/S of my left lower extremity which showed blood clot. I am already on blood thinner medication and no one can explain or give me an answer as to why I have a blood clot after my vaccine when i am on blood thinner medication.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1103171-1	Pain in left leg - outside 2/7/21 went to clinic 2/8/21 Given an ultrasound Diagnosis: Blood Clot (Dvt) Given immediate shot in belly And prescription for Eliquis Followed up with Dr. (internist) 2/11/21
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1103260-1	On Sunday Feb. 21st, in the evening before bed, I noticed that my left lower leg and ankle were swollen, (as complared to my right leg.) I was also experiencing shortness of breath.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1103410-1	Developed a DVT blood clot in right leg calf area -after the shot-
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1104339-1	I am retired handicap-MS. I was doing light housework. I started getting pain in right lung, over course of 2 hours pain became very severe. Only in right lung. Felt like blood clot or pulmonary embolism (which I had 15 yrs ago). Went to hospital, check X-ray no clot/embolisms. Blood work showed no clotting factor. Was admitted 4 days. We treated possibility of MS reaction (solu-methyl) as well as heparin in event of unknown clotting issue. Here?s why this is strange: If this was MS flare/relapse, I would have muscle lockup, drop foot, sight issues, sever balance issues ...(i always have sight issues).. with this even it was only severe lung pain. Did not appear to be MS driven issue. Being educated ?high analytic person, in reflection, My ms is very territorial, it fights anesthesia, colds/flus/covid when my husband had it. My MS gets cranky when my immune system is attacked. I suspect the second covid injection was doing its thing, dna modification, and I had some lung inflammation and my MS compounded the reaction with crippling issue. I was in hospital 4 days, IV fluids, heparin, and steriod IV. Within first 36 hours the extreme lung pain disappeared and I got better. I do have cats scan, mri, X-rays from stay. Cat scan clear, X-ray, normal, mri-classic ms history. Was this Ms event? I?m not sold as I didn?t have other normal MS flare symptoms. Was this injection aggreviated? Perhaps... I?m sharing this because I think tracking auto immune disease people is important data gathering. Feel free to call if more dialogue is necessary
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1104565-1	BLOOD CLOT. Began feeling ill 2 days after 1st vaccine...headache, cold and tired. Developed pain in my abdomen area quite noticeable 4 days later. 6 days later had major pain, vomiting, bloody diarrhea. Went to ER morning of the 7th day....spent 4 days in hospital. Had a blood clot--Portal Vein Thrombosis. Put on Heparin drip in hospital...and Eliquis blood thinners after released. I have never been sick like that...never been on any prescription...always been healthy, walking 3-5 miles per day. My family feels that there must be a connection with the vaccine. One of my doctors said no...another said didn't think so, but unknown at this time.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1104607-1	Stroke on 2/11/2021 caused by a blot clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1105456-1	Lymph gland under left arm size of golf ball, smaller glands swollen underarm, (soft), neck swollen, left foot swollen, weakness, small blood clot in myopic of middle finger, small modules in upper thigh, sore on lower lip right side
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1105500-1	"Pfizer-BioNTech Covid vaccine, first shot developed a blood clot in a superficial vein the day after the vaccine, was diagnosed by a doctor as ""superficial thrombophlebitis, pain swelling in leg. Treatment of ibuprofren started. Wondering if I should take my second shot which is scheduled for MArch 26"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1105661-1	Blood Clot's small in distal region of lungs bilateral 3 days after injection of vaccine

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1105772-1	My mother died on February 19, 2021. She had her 2nd dose vaccine on 2/11, on 2/12 it was noted that she was not able to walk, on 2/13 she was walking at 30%, on 2/14 she was walking with difficulty, on Monday 2/15 she was throwing up violently and her blood pressure dropped, so she was sent to Clinic. My sister was told she was just constipated and she had A Fib (never reported before to us). My sister was then told on 2/16 early a.m. that she had a blood clot that destroyed her colon. Due to age surgery would likely not be successful. She then died on the Friday. We are reporting in the event that the Pfizer vaccine was somehow a contributing factor to the A fib or to the Clot. She has no history of A fib or clotting prior to this incident. She was 93, and did have dementia, but was able to eat normal foods prior to this. What was unusual was the challenge in walking the day after the shot. Other than that no difference was observed until the day she was admitted to the hospital emergency room. She was a resident at Assisted Living, Memory Care, and that is where she received the vaccine. The mailing address I provided is her mailing address prior to death.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1105932-1	On Saturday patient woke up saying he was in extreme pain with his left arm. At 5 o'clock that night he lost his equilibran and I had to call 911. They came and after an initial examination took him to Hospital's emergency room. They did c-scans, MRI. and they said that a blood clog was showing in his brain and that he had atrium filibration of his heart. His vision has been affected, though the hospital thinks it will improve. The left side of his body is not function right and he needs a walker to walk. He has been in ICU since Sunday. This is late Tuesday.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1106331-1	developed two blood clots in his right calf; pain in his leg; This is a spontaneous report from a contactable consumer reporting for himself. A 70-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE) (lot number EN9581/expiration date: not provided), via an unspecified route of administration, on 04Feb2021 (at the age of 70 years old) as a single dose for COVID-19 IMMUNIZATION. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE) (lot number EL3249/expiration date: not provided), via an unspecified route of administration, on 15Jan2021 (at the age of 70 years old) as a single dose for COVID-19 IMMUNIZATION. Relevant medical history included was not provided. Concomitant medication included acetylsalicylic acid (BABY ASPIRIN), meloxicam, an allergy medication and anti-depressants. On 19Feb2021, the patient reported that he developed two blood clots in his right calf, one towards his ankle and one towards his knee and pain in his leg which required hospitalization on 22Feb2021. The patient was discharged on the same day, 22Feb2021. Relevant lab data included: blood test on 22Feb2021 were done to determine what medication to put him on. The results of the blood test was unknown. Treatment received for the event thrombosis included abixaban (ELIQUIS) tablets as a blood thinner. The patient reported he never had blood clots in his life. The outcome of the events thrombosis and pain in leg was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1106369-1	Pain for 4 or 5 days in left leg (that leg also has varicose veins), some swelling and tender to the touch . Fifth day became raised and warm to the touch, so went to Urgent Care Diagnosis: Occlusive and nonocclusive thrombus within a superficial vein underlying the region of concern
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1106925-1	I received my first vaccination on February 11, 2021. On February 17th I was rushed to my local ER with a massive heart attack. I was rushed by helicopter to second hospital where I underwent an emergency Heart Catherization. It was discovered that I will need a triple bypass surgery. However, there is a HUGE blood clot that formed in my heart, and further treatment is on hold for now, until the massive clot has safely dissolved.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1107421-1	I had sudden malaise, lost consciousness and my wife says was not breathing briefly. I revived and was taken to the hospital, and treated for 2 1/2 days with heart catheterization and TPA injected into my clot in the pulmonary arteries with good resolution. I did have cor pulmonale with acute severe right heart failure as part of that, Dr. thought the clot came from my right calf by his examination. Diagnosis was confirmed by CAT scan cardiac Echo and right heart catheterization.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1108193-1	Pain in calf of left leg. Two blood clots lower left leg, being treated with blood thinner Xeralta
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1110877-1	"EE admitted to hospital 3/18/2021 with ""blood clots"""
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1111010-1	Development of DVT of right leg and PE of right lung. Leg pain started on Friday 01/15/21. Care sought at Urgent Care on 01/18/21. Started on Eliquis 01/18/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1111548-1	Pfizer-BioNTech COVID-19 Vaccine EUA Approximately three weeks after this injection, I began to feel a pain in my left leg. I assumed it was a cramp. I then received a second dose (Lot # EN6206) on 3/13/21. Over that weekend the pain in my leg grew worse and the leg began to swell. on 3/15, I went to my primary care physician who said it looked like a blood clot. He sent me to a radiologist for an ultrasound (?) of the leg. After confirming the clot, I went to a local ER. They started me on Eliquis to treat the clot

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1111554-1	Multiple strokes resulting from blood clots starting 10 days past 2nd shot. Fuzzy headed with headache prior to strokes.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1111680-1	Developed severe back pain. which she went to the ER to have evaluated. Upon evaluation, she was diagnosed with a PE and blood clots in her spleen
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1111720-1	Pulmonary embolism: multiple blood clots in all five lobes of lungs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1112775-1	blood clot leading to brain stem stroke, intubation, shortness of breath and chest pain
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1114378-1	multiple blood clots in lungs; multiple blood clots in legs; feeling chest pain; short of breath; This is a spontaneous report from a contactable consumer (patient). A 68-year-old female patient (pregnant: No) received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL9269), via an unspecified route of administration in left arm on 06Feb2021 at 18:00 at single dose for covid-19 immunisation. The relevant medical history included DCIS Breast cancer from 2016, Known allergies: pencillin from an unspecified date. Concomitant medications included levothyroxine sodium (SYNTHROID). The patient previously received first dose of BNT162B2 on 16Jan2021 at the age of 68 years old (lot number: EL3249, at 06:15 PM, in Left arm) for covid-19 immunisation. The patient received 2nd Pfizer vaccine dose on Sat, 06Feb2021. On Wednesday 10Feb2021 she became short of breath. On 11Feb2021, she began feeling chest pain. On 12Feb2021, she was advised by primary care Dr to go to the emergency room. She was admitted on 12Feb2021 to ICU with multiple blood clots in her lungs and legs. She was told by the emergency Dr this was a sign of Covid, they did a Covid test, came back negative. She remained in the hospital till 18Feb2021. She was at the time of the report on Eliquis (10 mg/d). A hereditary blood clot test was done that came back negative. She had never had an issue before the vaccination with blood clots nor anyone in her family and she was in good health before the vaccination. Dr didn't think the vaccine caused the blood clots but her primary care Dr thought it did-which was why she was reporting it. The patient also stated that event multiple blood clots in her lungs and legs resulted in Emergency room/department or urgent care, Hospitalization and Life threatening illness. The patient had hospitalization for 7 days. Treatment Heparin 25,000unit in .45% NaCl Premix, Narcan received. No covid prior vaccination. The patient underwent lab test included Nasal Swab which showed negative on 12Feb2021; Hereditary blood clot test which showed negative on an unspecified date in 2021. Therapeutic measures were taken as a result of all the events. The outcome of the events was recovered with sequel.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1114514-1	Explained in previous section
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1114702-1	amaurosis fugax. Temporarily lost vision in top half of right eye - lasted about 15-20 seconds - blood clot in right eye. No past history of blood clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1118591-1	Hospotalive March 6th 2021. Numbness, loss of use left foot, leg, L-ARM, HAND. DAZED, COULDN'T CONSINTRATE, FUNCTION. TAKE TO EMERGENCY ROOM at Piedmont Athens Reg Med Ctr and admitted. Had cat scans, MRI's, ultrisounds, EKG's. Blood clots ediology unknown, caused by COVID 19 VACINES. PFIZER
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1118702-1	Had a stroke 36 hours after getting second vaccine. Lost ability to speak and see clearly, had word salad. Was identified quickly by my wife and was taken by ambulance to hospital where they gave me TPA clot buster infusion after identifying a clot in my left back side of brain and luckily I responded well and have all speech function back we believe so far.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1119208-1	10 days after my second shot of the Covid vaccine, I was short of breath and my heart rate was elevated. I went to the hospital and they found blood clots in both lungs and I had to be hospitalized.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1120490-1	I had experience of muscle cramps of left calf muscle in Feb / march and observed 3-4 times over a period of 2 months . I was thinking that could be due to dehydration . On 03/ 14 /21 morning , I got severe left leg pain and could not able to walk. I took pain killers and did not subside and that end up going to hospital emergency . I was diagnosed with 2 blood clots in left leg . I do not have family history of clots nor any h/o major medical issues . All my previous and recent blood reports normal . I am currently in use of blood thinners

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1120515-1	Patient had vaccine Wednesday and felt a little dizzy after the shot. Sunday morning she woke up very dizzy and ended up passing out in her kitchen while getting her morning coffee. She hit her head and was taken to the ER. Her blood pressure was found to be very high (systolic 218). CT scans revealed 2 blood clots in her lungs per pt. She was hospitalized for 2 days. She reports trouble walking now.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1121617-1	Plantar thrombosis - several SVT &DVT in multiple veins & artery of left foot.; several SVT &DVT in multiple veins & artery of left foot; several SVT &DVT in multiple veins & artery of left foot; Pain & swelling of foot; Pain & swelling of foot; This is a spontaneous report from a contactable consumer. This consumer (patient) reported that a 74-years-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in Arm Left on 12Feb2021 12:00 AM (Batch/Lot Number: EM9810) as SINGLE DOSE for covid-19 immunisation. Medical history included known allergies: Some antibiotics cause autoimmune issues, all sulphate drugs, narcotics. Historical vaccine included first dose of Pfizer COVID 19 vaccine (lot_number=EL3249) on 22Jan2021 12:00 AM administered in Right arm. Concomitant medications included thyroid (NATURE THROID) and multiple vitamin, taken for an unspecified indication, start and stop date were not reported. The patient experienced Plantar thrombosis - several SVT &DVT in multiple veins & artery of left foot. Pain & swelling of foot, all on 06Mar2021 01:00 AM with outcome of not recovered. Events resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Life threatening illness (immediate risk of death from the event). Treatment received for events included given Eliquis to avoid further clotting. No other vaccine in four weeks. No covid prior vaccination. No covid tested post vaccination.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1121620-1	chest pains; chest pains; blood clot; light headedness; low fever; blacked out for about 5 minutes; onset of major body ache and fatigue; onset of major body ache and fatigue; Body aches at injection site and into upper back; Body aches at injection site and into upper back; This is a spontaneous report from a contactable consumer (patient). A 67-year-old male patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: EN6200), via an unspecified route of administration, administered in left arm on 23Feb2021 10:30 AM (at age of 67 years old) as single dose for COVID-19 immunisation. Medical history included high cholesterol. Concomitant medications included atorvastatin; levobunolol (eye drops); latanoprost (LATANO). The patient previously received his first dose of BNT162B2 on 05Feb2021 for COVID-19 immunisation (brand=Pfizer; lot number: EL9269; administration time 03:30 PM; vaccine location=Left arm; dose number=1). 6 hours after injection on 23Feb2021, the patient experienced body aches at injection site and into upper back. After 24 hours after injection on 24Feb2021, started light headedness, low fever, blacked out for about 5 minutes, onset of major body ache and fatigue. Spent the next 36hrs in bed. 72hrs after injection recovered back to normal slowly. On 04Mar2021 at 8pm, started with chest pains called, admitted to Hospital with heart attack, immediately taken to Cardiac Cath Lab for coronary catherization in both legs, to partial remove blood clot and insert continuous balloon pump, IV Heparin infusion Troponin 6542.3 ng/l (critical), NO previous health issues for blood clots. Discharged 07Mar2021, Apixiban 5mg twice a day, Clopidogrel 75 mg /day. AE resulted in: Emergency room/department or urgent care, Hospitalization, Prolongation of existing hospitalization (vaccine received during existing hospitalization), Life threatening illness (immediate risk of death from the event. Number days hospitalization: 3 days. The outcome of events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1122738-1	blood clot formed in left arm; This is a spontaneous report from a contactable consumer (patient). A 62-year-old female patient, not pregnant at time of vaccination, received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EN6198), via an unspecified route of administration, on right Arm on 27Feb2021 0100, SINGLE DOSE for covid-19 immunisation. Medical history included Diabetic, High blood pressure, both from an unknown date. Concomitant medications included glipizide, metformin, fenofibrate. The patient previously took sumatriptan succinate (IMITREX) and experienced drug allergy. After vaccination 6 days later (05Mar2021) a blood clot formed in left arm. Shot was given in right arm. Hospitalization followed and now on blood thinners. AE resulted in: Emergency room/department or urgent care, Hospitalization, Life threatening illness (immediate risk of death from the event). The patient was hospitalized for 2 days. The outcome of the event was not recovered. Eliquis was given as treatment for the event. No covid prior vaccination. Not covid tested post vaccination.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1122743-1	severe thrombocytopenia; Bleeding at Impella insertion site; peripheral swelling in hands/feet; cardiogenic shock; myocarditis; hypoxic respiratory failure; mural thrombus; hypotensive despite pressors; fever; cough; myalgias; This is a spontaneous report from a contactable physician. A 46-year-old non-pregnant female patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Formulation: Solution for injection, Lot number and Expiration date was not provided), intramuscularly on 05Feb2021 as a single dose for COVID-19 immunisation. The patient's medical history included hyperlipidemia and COVID-19 pneumonia from an unspecified date in Jan2021 to an unspecified date in Jan2021 (the patient was diagnosed with COVID-19 pneumonia prior to the vaccination. Recovered. Returned to work on 25Jan2021). Concomitant medications included atorvastatin orally at 10 mg, once a day, acetylsalicylic acid (ASPIRIN) orally at 81 mg, once a day, colecalciferol (VITAMIN D); all the drugs were received within two weeks. The patient previously took clindamycin and experienced known allergies: Clindamycin. The patient did not receive other vaccine in four weeks. The patient developed fever, cough, myalgias on 19Feb2021 at 12:00 AM. She developed peripheral swelling in hands/feet on 24Feb2021, she was evaluated in the ER; admitted to (hospital name withheld) on 24Feb2021 with cardiogenic shock, myocarditis, hypoxic respiratory failure. The patient was started on IV vancomycin and Unasyn. TTE (transthoracic echocardiogram) demonstrated LVEF (left ventricular ejection fraction) 35%; reduced biventricular function; mural thrombus on 24Feb2021. Remained hypotensive despite pressors on 24Feb2021. Patient had elevated PCW with preserved cardiac index. Patient underwent VA ECMO (veno-arterial extracorporeal membrane oxygenation) and Impella placement on 25Feb2021. COVID-19 PCR was negative. Blood cultures were no growth. She developed severe thrombocytopenia and developed bleeding at Impella insertion site on 25Feb2021; required multiple, PRBC transfusions. Evaluated for HLH; Soluble IL2 receptor on 26Feb2021 elevated at 7232 pg/mL; ferritin 3054; CRP > 300. ECMO stopped 03Mar2021. The patient was treated with IV antibiotics, mechanical ventilation, pressor support, underwent VA ECMO and Impella placement. The patient was hospitalized from 24Feb2021 to 16Mar2021. Number of days of hospitalization was 20 days. The patient tested COVID post vaccination. The patient underwent lab tests and procedures which included blood pressure: hypotensive despite pressors, LVEF: 35 %, nasal swab: Negative on 24Feb2021, blood cultures: No growth, nasal swab: Negative on 25Feb2021, ferritin: 3054, HLH: Evaluated, Soluble IL2 receptor: 7232 pg/mL (elevated at 7232 pg/mL), CRP: > 300 on 26Feb2021, nasal swab: Negative on 11Mar2021, nasal swab: Negative on 14Mar2021. The events were considered as serious (hospitalization and life threatening) by the physician. The outcome of the events was recovering. Information about lot/batch number has been requested.; Sender's Comments: the events being serious, life threatening and hospitalisation ,medical intervention required are assessed as possibly related to the suspect drug __BNT162B2____ based on strong temporal association, but consider also possible contributory effects from patient's medical history and/or concomitant medications.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1124798-1	Blood Clot in the lower right
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1125931-1	Left leg started hurting, and swoll up from my knee to my foot, and the next day was told to go to hospital. They found a blood clot in my leg .
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1126808-1	Well he got the shot at 9 in the morning and by 4 in the a.m he had a stoke and it took him 5 hrs to get to his phone for help and called my mother
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1128059-1	"blood clots found on his arm and lung; blood clots found on his arm and lung; valve blockage; This is a spontaneous report from a contactable consumer (patient's friend) via Medical Information team. A male patient of an unspecified age received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; batch/lot number and expiration date were not reported), via an unspecified route of administration on 02Feb2021 as a single dose for COVID-19 immunization. Relevant medical history included COVID-19 infection from Sep2020 to an unknown date. The patient's concomitant medications were not reported. The reporter stated that the patient was due to receive his second dose the day prior reporting on 02Mar2021, and wanted to know if the patient should receive the second dose of the vaccine if he stayed longer than 3 weeks in the hospital. The patient was currently hospitalized; he received medical attention 2 weeks prior reporting on an unspecified date in Feb2021 due to some blood clots found on his arm and lung. The patient went into the hospital because he had a valve blockage on an unspecified date in Feb2021 and that the patient still had to go to rehab and might miss that 3 week window. The reporter wanted to know if a patient should restart the vaccination series if he could not receive the second dose 42 days after the first one. The patient was hospitalized for the events ""blood clots found on his arm and lung and valve blockage"" from an unspecified date in 2021. The outcome of the events was unknown. Information on the batch/lot number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1128546-1	Patient was admitted to the ED with left deep peroneal vein thrombosis on Feb. 9 2021 after patient noticed prolonged edema in both ankles. Patient started talking 10 mg of Eliquis twice a day on Feb. 11, 2021. And on Feb. 17 2021, 2 more blood clots were found in the patient's peripheral veins within the limbs. Patient descreased dose of Eliquis to 5mg twice a day on Feb. 18 2021. Patient continues to take 5mg of Eliquis twice a day.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1132430-1	The daughter of the patient stated that her mother is a cardiac patient and has a stent. The patient always has edemas on her legs but the day after the vaccination she observed edema and report pain on the left thigh. The following Monday she visited the MD and he order a venous Doppler. The result was a thrombosis on the right leg . The patient was put on medication (Xarelto)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1133506-1	Had a major nosebleed with blood clots; Had a major nosebleed with blood clots; This is a spontaneous report from a contactable consumer (the patient). A 54-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 1, intramuscular on 08Mar2021 (Batch/Lot Number: EN6199; Expiration Date: 30Jun2021) as single dose for COVID-19 immunisation. Medical history was not reported. There were no concomitant medications. The patient had a major nosebleed with blood clots on an unknown day in Mar2021. The patient reported that he never got nosebleeds before. There was no treatment taken for the event, it stopped on itself. The outcome of the events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1135488-1	blood clots right nostril; pain/discomfort right side at waistline going toward back; pain/discomfort right side at waistline going toward back; This is a spontaneous report from a contactable consumer (patient). An 82-year-old female patient received the 2nd dose of bnt162b2 (BNT162B2, Lot Number: EL9581), as single dose in left arm on 06Feb2021 10:30 for COVID-19 immunisation, administered at hospital. Medical history included artificial mitral valve. No known allergies. The patient was not pregnant. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient received concomitant medications in two weeks but were not reported. No other vaccine in four weeks. The patient received the 1st dose of bnt162b2 (BNT162B2), on ana unknown date for COVID-19 immunisation and experienced pulsatile tinnitus right ear. The patient experienced blood clots right nostril, pain/discomfort right side at waistline going toward back on 16Feb2021. The patient was hospitalized for blood clots right nostril. She went to Dr. for tinnitus, had MRI and CAT scan in 2021. All tests were negative. She received no treatment. COVID was not tested post vaccination. Right side got better after several days in 2021. Blood clots in nose lasted one morning and recovered on 06Feb2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1136780-1	Blood clots in lungs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1137006-1	Extreme fatigue, dizziness, lack of appetite increase in mucus, some blood clotting in mucus
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1137624-1	I noticed increasing episodes of SOB, dizziness, and PVCs. My blood pressure continued to increase. After 4 weeks of increasing symptoms I reported to the ED. I was admitted for Right Lower Lobe blood clot, hypertension, and PVCs.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1140696-1	Multifocal Intracerebral Hemorrhage; Disseminated Intravascular Coagulopathy; strokes, Ischemic and Hemorrhagic; strokes, Ischemic and Hemorrhagic; AML; Leukemia; Blood clot diagnosis; Sore lower leg; RDW Stand. Dev. H/RDW Coeff Var H; Platelet Count L, Platelet Vol L; Neutrophils L; Band Neutrophils H; Monocytes H; Metamyelocytes H; Myelocytes H; Absolute Neutrophils L; Other Cell Type Blast Like Cells H; This is a spontaneous report from a contactable consumer. A 70-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) lot number: EL9261, via an unspecified route of administration, administered in Arm Right on 02Feb2021 08:30 (Batch/Lot Number: EL9261) as SINGLE DOSE for covid-19 immunisation. Medical history included breast cancer (8 years ago no chemo just radiation). Historical vaccine included first dose of BNT162B2 (lot number: EL0140) on 11Jan2021 for Covid-19 immunization. Concomitant medication included vitamin c [ascorbic acid] (VITAMIN C [ASCORBIC ACID]), calcium citrate, colecalciferol (CALCIUM CITRATE + D3), glucosamine, magnesium citrate, docosahexaenoic acid, eicosapentaenoic acid, tocopheryl acetate (OMEGA 3 [DOCOSAHEXAENOIC ACID;EICOSAPENTAENOIC ACID;TOCOPHERYL ACETATE]) and curcuma longa (TURMERIC [CURCUMA LONGA]). On 04Feb2021, the patient's blood work result showed red cell distribution width (RDW) stand. dev. high; RDW coeff var high, platelet count low, platelet vol low; neutrophils low; band neutrophils high; monocytes high; metamyelocytes high; myelocytes high; absolute neutrophils low; other cell type blast like cells high. On 15Feb2021, the patient experienced sore lower leg. On 16Feb2021, the patient was diagnosed with blood clot. On 19Feb2021, the patient was diagnosed with leukemia. On 20Feb2021, the patient was diagnosed with acute myeloid leukemia (AML). On 21Feb2021, the patient had tow types of stroke, ischemic and hemorrhagic, the patient was intubated. On 23Feb2021, the patient was extubated and died due to multifocal intracerebral hemorrhage, disseminated intravascular coagulopathy, acute myeloid leukemia with blast crisis. The patient received chemotherapy and leukapheresis as treatment. The patient died on 23Feb2021. An autopsy was not performed.; Reported Cause(s) of Death: Disseminated Intravascular Coagulopathy; Acute Myeloid Leukemia With Blast Crisis; Multifocal Intracerebral Hemorrhage
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1141687-1	Blood clots a few weeks after 2nd dose. Currently being treated with Xarelto. No other changes in patient medical or prescription history. Covid vaccine was the only thing that was new. Patient had no prior history of blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1141723-1	My leg and foot swelled on Thursday. We elevated and treated with heating pad & Advil. The swelling went down, so we started for home, drove 4hrs. My leg & foot swelled back up. We went to the ER they gave me an ultrasound, and said I had 2 blood clots in my leg. They prescribed, Eliquis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1142935-1	1 week after second dose I developed a blood clot behind my left knee, in hospital for 9 days; This is a spontaneous report from a contactable consumer (patient). An 87-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 17Feb2021 09:00 (Lot Number: EN6201) as SINGLE DOSE for COVID-19 immunization. Medical history included high blood pressure, controlled from an unknown date and unknown if ongoing and penicillin allergy from an unknown date and unknown if ongoing. The patient previously had first dose of BNT162B2 in the left arm (lot number: EL3244) on 20Jan2021 for COVID-19 immunization. Concomitant medications included apixaban (ELIQUIS), metformin, caredevil and fish oil; all taken for an unspecified indication, start and stop date were not reported. On 24Feb2021 12:00 PM, it was reported that the patient 1 week after second dose, developed a blood clot behind left knee, in hospital for 9 days that also resulted in emergency room/department or urgent care. The event was treated with blood thinners. The patient had no COVID prior to vaccination and has not tested for COVID post vaccination. The outcome of the event was recovering.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1143469-1	"phlebitis in my Right leg; small blood clot; This is a spontaneous report from a contactable nurse (patient). A 66-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in right arm on 09Feb2021 (Batch/Lot Number: EL9269) as single dose for covid-19 immunisation. Medical history included many allergies. The patient has no COVID prior to vaccination. The patient had other medications in two weeks but no other vaccine in four weeks. The patient previously had the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot: EL9263) on 20Jan2021 at 10:00 administered in the right arm for Covid-19 immunization. On 20Feb2021 at 12:00 (also reported as ""in February""), the patient had a phlebitis in her right leg, a week later (Feb2021) a vascular surgeon office did an ultrasound and found a small blood clot. The events resulted in doctor or other healthcare professional office/clinic visit. The events were treated with Eliquis BID (twice a day) for 45 days. The patient was not tested for COVID post vaccination. The outcome of the events was not recovered.; Sender's Comments: A causal relationship between BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) and the event thrombosis cannot be excluded based on temporal association. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate ."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1143474-1	blood clots in couple of toes near nails; observed swollen; This is a spontaneous report from a non-contactable consumer. A 55-year-old male patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: EL3247), via an unspecified route of administration, administered in arm left on 24Feb2021 15:15 as a single dose for covid-19 immunisation . The patent has no medical history. The patient has no known allergies. The patient's concomitant medications were not reported. In Mar2021, the patient was observed with swollen and blood clots in couple of toes near nails. The events was not treated. The outcome of the events was not recovered. The patient has not had covid prior to vaccination and was not testes for Covid post vaccination. No follow-up attempts are possible. Information about Lot/Batch cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1143483-1	Blood clot; Unable to use arm at all; This is a spontaneous report from a contactable consumer (patient, self-reported). A 49-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot number and expiry dates were not provided), via an unspecified route of administration, in arm left, on 09Mar2021 02:15 AM, as single dose for COVID-19 immunization. Patient's medical history included Fibromyalgia. Concomitant medications were not reported. Patient was not diagnosed with COVID, prior vaccination. Patient has not been tested for COVID, post vaccination. Patient did not receive other vaccine in four weeks of vaccination. On an unknown date in Mar2021, patient had Blood clot and patient was unable to use arm at all. Patient resulted in Emergency room/department or urgent care, Disability or permanent damage in Mar2021. Patient did not receive treatment for adverse events. Seriousness of the events reported as hospitalization and disability. Outcome of the events was not recovered. Information about lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1143485-1	woke up with a blood clot lying in her mouth; This is a spontaneous report from a contactable consumer (patient's parent). A 23-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 13Mar2021 (Batch/Lot number was not reported) as single dose for Covid-19 immunization. The patient's medical history and concomitant medications were not reported. It was reported that the patient received her first dose of the Pfizer COVID-19 vaccine yesterday (13Mar2021), and this morning (14Mar2021) she woke up with a blood clot lying in her mouth. The reporter wanted to know if this side effect had been reported or of this could be associated with the Pfizer COVID-19 vaccine. The outcome of event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1143491-1	heavy bleeding with lots of blood clots; extended and heavy menstrual period/ heavy bleeding with lots of blood clots; This is a spontaneous report from a contactable consumer (patient). A 51-year-old non-pregnant female patient received second dose of BNT162B2 (Pfizer product), lot no. EN5318, via an unspecified route of administration (left arm) on 19Feb2021 at a single dose for COVID-19 immunisation at a workplace clinic. Medical history included high cholesterol and anxiety. No known allergies. No COVID prior to vaccination. Concomitant medications were not reported. Historical vaccine included first dose of BNT162B2 (brand: Pfizer, lot no. EL9261, left arm) for COVID-19 immunisation on 30Jan2021 at age of 50 years old. No other vaccine in four weeks. The patient was experiencing extended and heavy menstrual period. Her cycle began on 05Mar2021 and was still happening as of today 16Mar2021. She had heavy bleeding with lots of blood clots. This was not normal for her at all. She is calling her doctor tomorrow. No treatment received for the events. No COVID test post vaccination. The outcome of the events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1145050-1	2/2/2021 red scaly skin rash dorsum right hand. Next day pinpoint hemorrhagic 2-4 mm sized lesions on palmar surface of fingertips and sides of fingers, 21 in all. Diagnosed as clots by cardiologist and dermatologist on 2/25.Gradual resolution and healing by 3/14/2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1146761-1	"""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; ""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; ""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; Cardiogenic shock; Anterior myocardial infarction; This is a spontaneous report from a contactable consumer. An 81-years-old female patient received BNT162B2, dose 2 via an unspecified route of administration, administered in left arm on 06Feb2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunisation. Medical history included very sensitive to medication effects (usually takes only 1/2 dose with strong efficacy to avoid side effects). Breast cancer survivor (2014 onset, 2019 declared permanent remission) and mild blood pressure treated successfully with medication for about 10 years. Concomitant medications included spironolactone and valsartan, both taken for blood pressure. The patient previously received first dose of BNT162B2 on 16Jan2021 in left arm for COVID-19 immunization. The reporter's mother died 3 weeks and 6 days after having received the second dose of the Pfizer covid vaccine. The cause of death was a ""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; the event began about 11:45pm on 04Mar2021. The blockage was confirmed via cardiac cath procedure performed within 2 hours of the onset by Doctor, he removed the clot and placed a stent. However her heart was too damaged and could not recover. Doctor confirmed to us that she did not have excessive or evidence of any prior blockage and not excessive plaque. The blood clot likely came on and caused the cardiac event within roughly an hour, he explained. The patient had no prior symptoms and no comorbidities for blood clotting and was full of life and energy on 05Mar up to when she went to bed that night. She died 06Mar at 4:04 am at hospital. The strat date of the events was reported as 04Mar2021 at 11:45 PM. AE resulted in emergency room/department or urgent care, life threatening illness (immediate risk of death from the event). The patient died on 06Mar2021. An autopsy was not performed. The death cause: Triggered by the sudden 100% blockage of the LAD by a blood clot, the cause of death is listed as (A) Cardiogenic shock (B) Anterior myocardial infarction. Treatment was received for the events which included multiple resuscitations and angioplasty surgery. No covid prior vaccination, no covid tested post vaccination. The outcome of the events was fatal. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: ""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; ""widow maker"" type heart attack where the LAD artery suddenly became 100% blocked by a blood clot; ""widow maker"" type heart attack where the LAD artery"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1147267-1	Had vaccine about 9:35 am on 3/20/21. In the middle of the night about 2:30 am 3/21/21 He should signs of paralysis on right side of his body and was not able to walk. Went to the emergency room on 3/21/21. Admitted to the hospital. MRI was done about 8:00 pm on 3/21/21 and results were blood clot on left side of the brain caused a stroke. Paralysis of right side of his body and changes in speech. Transferred Rehabilitation Hospital 3/24/21. Possibility of second stroke, and another scan done 3/29/21. Waiting for results
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1147922-1	Off-cycle menstrual/vaginal bleeding. Today is 7th day of heavy bleeding, about twice as heavy as a heavy flow day for a normal menses. Many large blood clots through the day and night.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1147964-1	Within an hour of receiving the vaccine I felt a sharp pain shoot down the back of my knee to my heal. I continued to work with the pain. The next day my knee and calf was sizeably larger. I went to the urgent care on March 19, 2021. Then to the Emergency room on March 20, 2021. They performed an ultrasound of my leg and found a blood clot. It is in a superficial vein. Followed up with a vein doctor and will go again in three months for another follow up.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1148105-1	BLOOD CLOTS : Two days after receiving Pfizer dose 2 my mother had difficulty breathing and had chest pain. She was hospitalized for six days due to blood-clotting in the lungs.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1148782-1	Pain in left calf leading to development of blood clot above left knee cap near inside of quad; Pain in left calf leading to development of blood clot above left knee cap near inside of quad; This is a spontaneous report from a contactable consumer. A 38-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 2 via an unspecified route of administration, administered in Arm Left on 01Mar2021 12:00 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. Medical history included U/L fasciatomy in same leg with nerve damage, Blood transfusion, previous clot in femoral during recovery, Shellfish (allergy). The patient has not had COVID prior vaccination. The patient received first dose of BNT162B2 for COVID-19 immunization on an unspecified date (lot number: unknown) at 38-years-old. Concomitant medication included sulfamethoxazole, trimethoprim (BACTRIM) taken for an unspecified indication, start and stop date were not reported. On 04Mar2021 16:00, the patient experienced Pain in left calf leading to development of blood clot above left kneecap near inside of quad. AE resulted in: Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. The patient received treatment for the events: Originally treated as cellulitis with doxy. The outcome of the events was recovering. The patient was not tested for COVID post vaccination. Information about Lot/Batch number is requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1148783-1	Clot; Swelling and pain in left foot, ankle, and calf; Swelling and pain in left foot, ankle, and calf; This is a spontaneous report received from a contactable consumer (Patient's son). A 81-years-old female received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, batch/Lot number not reported), via an unspecified route of administration, administered in Arm Left on 02Feb2021 (believes in afternoon) at single dose for covid-19 immunisation. The patient medical history and concomitant medication were not reported. Patient previously took history vaccine included shingles shot received on Nov2020 and Flu shot typically gets every year. On 02Jan2021, patient received first dose of COVID vaccine (BNT162B2) for covid-19 immunisation. The patient experienced severe swelling and pain in foot, calf, and ankle on the left side shortly after the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) was administered. She was hospitalized and placed on Xarelto for treatment by her cardiologist and was scheduled to follow-up to have other procedures performed as deemed needed. Doctor advised patient had a clot and was put on Xarelto. An angio and cath was suggested to see if she had something closing up in her. Patient was going back in a few weeks to see about the clot. she never had this clotting concern before, and this was all new right after the second COVID vaccine. Swelling and pain in left foot, ankle, and calf was either 04Feb2021 or 05Feb2021. she first noticed left foot was swollen. she stated her foot was sore. It was unknown if it went to her ankle and then from there. It was stated that the pain and swelling was better. She was informed by the nurse administering the COVID vaccine that second dose would be a little bit stronger. The patient underwent lab tests and procedures which included MRI on 15Feb2021 and diagnosed the clot. The outcome of the event Clot was unknown. For other events was recovered on unspecified date in 2021. Information about Batch/Lot number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1150166-1	Stroke (blood clot in right brain), resulting in slurred speech and left-side weakness
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1150377-1	Superficial blood clot on left calf at the lower end of some varicose veins two days after injection; Superficial blood clot on left calf at the lower end of some varicose veins two days after injection; This is a spontaneous report from a contactable consumer (patient). A 62-year-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot: EN6207), via an unspecified route of administration in right arm on 10Mar2021 14:30 (at the age of 62-years-old) as single dose for covid-19 immunisation. The vaccine facility type was a pharmacy or drug store. The patient had no other vaccine in four weeks. The patient did not have Covid prior vaccination and was not Covid tested post vaccination. Medical history included macular. Concomitant medication included amitriptyline. The patient previously took erythromycin and statins and experienced drug allergies from these. The reported adverse events were superficial blood clot on left calf at the lower end of some varicose veins two days after injection (on 12Mar2021 at 10:00). It may be unrelated, but the patient wanted to let know. The events resulted in emergency room/department or urgent care. AE treatment included hot compress, support stockings and ibuprofen (MOTRIN). The outcome of the events was recovering.; Sender's Comments: Varicose veins most probably was a preexisting condition, unrelated to BNT162B2 vaccine. The reported superficial blood clots are considered a complication of varicose vein and unlikely related to BNT162B2.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1150383-1	blood clots in arm/Blood Clots in Left Arm; swollen lymph nodes /Swollen Lymph Nodes in Left Arm; nodules were found on her lungs/Nodules in Lungs; arm swelling /Left Arm Swelling; Left Arm Redness; Left Arm Discomfort; Soreness at the Injection Site; This is a spontaneous report from a contactable consumer (Patient's Mother) calling on behalf of daughter. A 50-year-old female patient received 1st dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, batch/lot number and expiration date unknown), via an unspecified route of administration, administered in Arm Left on 05Mar2021 13:30 at age of 50 years old at single dose for covid-19 immunisation. There was no medical history. There were no concomitant medications. The patient did not have prior vaccinations within 4 weeks. No adverse events following prior vaccinations. There were no additional vaccines administered on same date of the Pfizer suspect. The patient experienced the following after first dose. The patient experienced arm swelling /left arm swelling (hospitalization) on 09Mar2021 with outcome of recovering , blood clots in arm/blood clots in left arm (hospitalization) on 11Mar2021 with outcome of not recovered , swollen lymph nodes /swollen lymph nodes in left arm (hospitalization) on 11Mar2021 with outcome of not recovered , nodules were found on her lungs/nodules in lungs (hospitalization) on 11Mar2021 with outcome of unknown , soreness at the injection site on 05Mar2021 with outcome of not recovered , left arm discomfort on 08Mar2021 with outcome of not recovered , left arm redness on 09Mar2021 with outcome of not recovered. The patient was hospitalized from 11Mar2021 to 12Mar2021. On 05Mar2021 she began to have arm pain. 48 hours later, patient's arm started really hurting her, something about her arm being sore on the day of injection and reporter said she was going to be sore and when she got back from going away for the weekend, she thought uh oh, there was a problem here, this was not normal, not right. On 10Mar2021, she experienced swells arm swelling and went to urgent care which was diagnosed with swollen lymph nodes and given Cheplex prescription at urgent care. 11Mar2021, daughter was admitted to hospital by her provider. Daughter was found to have blood clots in arm and continued swollen lymph nodes. Daughter was given IV cheplex and Eliquis while in hospital. Daughter will have to be on Eliquis for next 3 months per reporter. The patient had her first vaccine, Pfizer, and became ill with blood clots, swollen lymph nodes, and wound up in the hospital. Reporter stated that she and the patient were worried about her getting the second shot. The patient was on blood thinners and was seeing a pulmonary doctor and hematologist. She didn't know about her getting the second shot and thought this should be reported. The patient was due for the second vaccine next week on 26Mar2021 and she was still sick, should it be postponed or what? The patient went the night before the hospitalization to the Urgent Care and the doctor prescribed Keflex, an antibiotic and saw her primary care doctor the next morning and she put her in the hospital. Reporter stated they were getting nervous and wanted to report for side effects. She stated the patient was home from the hospital but still sick really bad. The patient went to Urgent Care 10Mar2021, saw her primary care 11Mar2021 and was sent to the hospital 11Mar2021. Reporter later clarified that the patient was sent to the Emergency Room and after a few hours, she was sent upstairs to be admitted for a 24 hour hospital stay. The patient received Eliquis; Keflex as treatment. Fluids at the hospital, unknown type. Reporter stated they decided the patient wasn't doing really well and went to the Urgent Care that night and were given Keflex because they felt there was an infection. When they discovered it further that there were blood clots, the patient was told to stay on the antibiotic and given fluids and put on blood thinner, Eliquis, states patient had never been on a blood thinner before. Patient was admitted on 11Mar2021 for 24 hours. Reporter stated that they kept the medicine going and then felt she was stable enough to be discharged the next night at around 18:00 to continue with Keflex antibiotic and Eliquis for 3 months. Reporter stated they were told to contact a hematologist and pulmonary doctor for the nodules in her lungs and to contact her primary doctor if not doing better in one week, but they will probably contact her tomorrow. Reporter clarified that the patient had a chest x-ray a couple of months ago and everything appeared fine, so she thinks that the nodules developed after the vaccine. The patient had a chest x-ray on 11Mar2021 and nodules were found on her lungs, scheduled to follow up with lung doctor. Reporter stated whether they were there before or not was unknown. No follow up X ray yet. Reporter would like to know if daughter should get second dose? Second dose was scheduled 26Mar2021. Adverse events required visit to emergency room: Urgent Care the night before, follow up the next morning with primary doctor, the doctor got her in right away and said she needs to go to the hospital. Yes, patient went through the ER, doctor said that she would be coming and then after there a while, she was admitted to the hospital. Adverse events required visit to physician office. Patient saw the doctor on 11Mar2021 and was sent to the hospital. Information about lot/batch has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1150385-1	I had a return of a superficial thrombophlebitis on my left arm above where a catheter had been when I had surgery 7 weeks before. The blood clot returned again to the same level of pain and the same; I had a return of a superficial thrombophlebitis on my left arm above where a catheter had been when I had surgery 7 weeks before. The blood clot returned again to the same level of pain and the same; This is a spontaneous report from a contactable consumer (patient). A 67-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in right arm on 01Mar2021 13:30 (lot/batch number was not reported) as single dose for COVID-19 immunisation. The patient's medical history included high blood pressure. The patient had no known allergies. The patient's concomitant medications were not reported. On 04Mar2021 07:30, the patient had a return of a superficial thrombophlebitis on her left arm above where a catheter had been when she had surgery 7 weeks before. The blood clot returned again to the same level of pain and the same. No treatment was received for the events. Patient was not pregnant at the time of vaccination. Patient had no COVID prior to vaccination and not tested for COVID post vaccination. Vaccine facility type was other. There was no other vaccine in four weeks and there was other medications in two weeks. The outcome of the events were not recovered. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1150394-1	Two blood clots in left leg; This is a spontaneous report from a contactable consumer. A 77-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 2 via an unspecified route of administration, administered in left arm on 26Feb2021 (Batch/Lot Number: EN6203) as single dose for COVID-19 immunisation. Medical history included back disorder, ATTR amyloidosis, and allergies: caffeine from an unknown date. Concomitant medication included allopurinol (ELAVIL); meloxicam; tizanidine; tafamidis (VYNDAMAX) all taken for an unspecified indication, start and stop date were not reported. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL9269), on 05Feb2021, administration time: 06:00 PM, vaccine location: Left arm for COVID-19 immunisation. The patient experienced two blood clots in left leg on 03Mar2021 23:00. Treatment for the event includes Aspirin. The event resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. The patient had no COVID-19 prior vaccination. The patient was not tested for COVID-19 post vaccination. The outcome of the event was resolved on an unspecified date.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1151417-1	blood clots in his left legs and also some in his lungs.; blood clots in his left legs and also some in his lungs.; This is a spontaneous report from a contactable consumer (patient) received via Medical Information Team. A 64-year-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 02Mar2021 (Lot number was not reported) as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient reported that he had to go to the emergency room (ER) where they discovered blood clots in his left legs and also some in his lungs in Mar2021. The events were serious, the patient was hospitalized for 2 days. Therapeutic measures were taken as a result of the events and the patient was placed on blood thinners. The patient outcome of the events was unknown. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"if it was a lung issue or blood issue, such as blood clots; nervous system was affected; weakness on her leg; had very tight calf muscles; eye pain; have difficulty breathing; she was dragging her feet, could not take the next step.; had ""such a tight feeling"" of the calf, and a little bit of pain to it; nauseous; if it was a lung issue or blood issue, such as blood clots; whether the injection was done in the wrong location; her arm was very itchy and a red papule-like round rash developed, it was pigmented,; her arm was very itchy and a red papule-like round rash developed, it was pigmented,; her arm was very itchy and a red papule-like round rash developed, it was pigmented,; it was ""highlighted"" and ""dot-like""; had a sharp pain, and tingling or pain-like sensation at the fingertips/she was going ""through a world of pain"", and from the arm down to the side of body.; had a sharp pain, and tingling or pain-like sensation at the fingertips; a fever too, 99.4¦F at one time and over 100¦F for a day or so; This is a spontaneous report from a contactable consumer. This consumer (patient) reported for self that the female patient of an unspecified age received second dose of bnt162b2 (BNT162B2,PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: EN6200, Expiration Date: 30Jun2021) via an unspecified route of administration on 05Mar2021 at single dose for covid-19 immunisation. The patient medical history was not reported. The patient's concomitant medications were not reported. The patient previously took first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot#: EL9262, expiration date: 31May2021) for covid-19 immunization for covid-19 immunization, rabies vaccine and experienced pain. The patient experienced if it was a lung issue or blood issue, such as blood clots on Mar2021 with outcome of unknown, nervous system was affected on Mar2021 with outcome of unknown, weakness on her leg on Mar2021 with outcome of unknown, had very tight calf muscles on Mar2021 with outcome of unknown, eye pain on Mar2021 with outcome of unknown, have difficulty breathing on Mar2021 with outcome of unknown , she was dragging her feet, could not take the next step on Mar2021 with outcome of unknown , had ""such a tight feeling"" of the calf, and a little bit of pain to it on Mar2021 with outcome of unknown, nauseous on Mar2021 with outcome of unknown, if it was a lung issue or blood issue, such as blood clots on Mar2021 with outcome of unknown, whether the injection was done in the wrong location on Mar2021 with outcome of unknown, her arm was very itchy and a red papule-like round rash developed, it was pigmented on Mar2021 with outcome of unknown, her arm was very itchy and a red papule-like round rash developed, it was pigmented on Mar2021 with outcome of unknown, her arm was very itchy and a red papule-like round rash developed, it was pigmented on Mar2021 with outcome of unknown, it was ""highlighted"" and ""dot-like"" on Mar2021 with outcome of unknown , had a sharp pain, and tingling or pain-like sensation at the fingertips/she was going ""through a world of pain"", and from the arm down to the side of body on Mar2021 with outcome of unknown , had a sharp pain, and tingling or pain-like sensation at the fingertips on Mar2021 with outcome of unknown. The patient underwent lab tests and procedures which included body temperature: 99.4 fahrenheit on Mar2021, body temperature: 100 fahrenheit on Mar2021. The consumer reported that ""Who makes the vial label? What syringe did they use to give her the vaccine? How big was it? Where is the vaccine administered? Did we make a second batch with ""much less"" side effects? She thought there was an earlier version which should have more side effects. Did she have to share a vial with other people? Is it multiple doses? How much is the dose of a vaccine?"" Consumer stated the lot number was a ""so vague looking thing"" to come with the Pfizer vial. Upon clarification, said she did not see the lot number on the vial, but on the vaccination card, where the lot number was very small. She would include 5 labels with lot numbers to split to each person who gets the dose from the same vial. The clinical course was reported as follows: She got the second dose of Pfizer's COVID-19 vaccine on 05Mar2021 and has developed side effects. Her nervous system was affected, she has weakness on her leg, had very tight calf muscles, eye pain. When she was working, she started to have difficulty breathing, she was dragging her feet, could not take the next step. This was 2 weeks ago and went away for a while. Today she had ""such a tight feeling"" of the calf, and a little bit of pain to it, she was nauseous. She was wondering if it was a lung issue or blood issue, such as blood clots. She had a fever too, 99.4¦F at one time and over 100¦F for a day or so. The first dose didn't give her any side effects. She was also concerned about whether the injection was done in the wrong location, she thought it was in the deltoid, but 10 days later her arm was very itchy and a red papule-like round rash developed, it was pigmented, she wondered if it was a hair, it was ""highlighted"" and ""dot-like"". It was located in the 2/3 lower to the elbow from the shoulder, she thought it was ""really low"" and they missed the deltoid muscle, she had no idea if the nurse didn't know the exact location and putting it ""in the skin"" was the cause of her symptoms. Stated they put a bandage in the wrong location, there was nothing there. She additionally had a sharp pain, and tingling or pain-like sensation at the fingertips, yesterday she was going ""through a world of pain"", and from the arm down to the side of body. With the first dose she didn't have any pain, but it was ""very weird"", afterwards she had some ""sharp pain"" under the scapula and didn't know why. She thought it was ""echo"" through the nervous system. Stated when she got the vaccine, she saw 3 dark marks on the syringe and two thirds were filled with the liquid. She wanted to know if this dosage was correct. Mentioned a lot of people had side effects, a lot of times they are given in the wrong location, in an entirely different place. She thought there was an earlier version of the vaccine which should have more side effects. She had a ""very bad bad"" experience 4 years ago when she got a rabies vaccine, it was given at 45¦, the nurse gave the injection ""cutaneously"", after 5 min she was ""so in pain""."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1151451-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1151636-1	My father received his first Pfizer vaccine on 02/03/21. On 2/12/21 I rushed him to the ER. He was vomiting uncontrollably and had shortness of breath. Once arrived at ER, they immediately put him on oxygen. Vomiting lasted several hours. They tested him for Covid and did a chest x-ray. Tested positive for Covid. Chest x-ray showed Covid pneumonia. Was admitted. Stayed in hospital for 5 days and was then released to nursing home for physical and occupational therapy. He was very weak and on days experienced what they called Covid fog. After 2 weeks of therapy, he was released on 03/06/21 to go back home to his apartment, with extended visiting nurse therapy. On 3/10/21, was the first visiting nurse appointment. At 12:00 an RN came to his apartment from Home Health Care. She checked his vitals. She said his blood pressure was good, lungs sounded good and oxygen level was 98. She said he was doing good and that she would not need to continue to come out and check on him weekly. She left. At 2:30 the same day, a Physical Therapist from Home Health Care came. She asked him lots of questions and adjusted my fathers' walker for him. He showed her how he was doing using the walker. Walked approximately 15-20 feet in his apartment. She checked his vitals before she left. His oxygen level was now at 91. She had him take a few deep breathes until his oxygen level was up to 93. She left and said she would be back on Friday the 12th to begin the actual physical therapy then. Within 10 minutes after she left my father started shaking uncontrollably and was having difficulties breathing. I called 911. Paramedics arrived. My fathers' oxygen level was all the way down to 74. They took him to the ER. When getting him out of ambulance he began vomiting. Vomiting lasted for hours just like when he went to the hospital back in February. They tried 3 different drugs to control the nausea. They did EKG, chest and abdomen scans. Was found that he had multiple blood clots and inflammation in his lungs and a bacterial infection in his blood. After testing, bacteria was found to be E Coli. Treated him with heparin for clots and antibiotics for infection and had him on oxygen in nose. Every day thereafter, he felt worse. They switched him to a high flow oxygen mask to keep his oxygen levels up. By Saturday night (early morning Sunday) on 03/14, they had taken the high flow oxygen mask off and hooked him up to a BiPap oxygen machine because his oxygen levels were dropping too low. We were then told by the lung doctor, that the damage to his lungs was extreme and that the next step would be to put him on a ventilator and feeding tube. My father did not want this per his will and his discussion with Dr earlier in the week. Dr indicated that he would not get better just being on the BiPap machine and we then chose to have them take him off of the machine because he did not want to go on life support. My father passed away on Sunday, March 14th around 6:30pm.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1152001-1	Patient received second dose of Pfizer BioNTech COVID Vaccine on 2/23/21. On 2/25 patient called PCP office expressing swollen & painful lymph node to left clavicle/shoulder area . Same arm as injection. Pain 2-3/10. Pt denied any other symptoms. Managed with warm compress, however on 3/26/21 patient went to the emergency room with complaint of continued swelling and redness on left arm. Ultra sound of the upper extremity showed short segment Nonocclusive thrombus in the left proximal subclavian
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1153367-1	left leg was cold/left foot was 79-82 degrees Celsius; blood clots in his left leg/he had three/first one on 02Mar, second one 13Mar and the third one yesterday; he had the Pfizer vaccine on 15Feb and 26Feb; he had the Pfizer vaccine on 15Feb and 26Feb; This is a spontaneous report from a contactable consumer (patient). A 77-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), second dose via an unspecified route of administration on 26Feb2021 (Batch/Lot Number: EN6203) as single dose for covid-19 immunisation; tafamidis (VYNDAMAX), route of administration, start and stop date, batch/lot number and dose were not reported for an unspecified indication. The patient took the first dose of PFIZER-BIONTECH COVID-19 VACCINE on 15Feb2021 at age of 77 years old for covid-19 immunisation. The patient medical history and concomitant medications were not reported. A patient taking Vyndamax medication who stated that he had experienced blood clot in his left leg. Clarified that he had the Pfizer vaccine on 15Feb2021 and 26Feb2021. Clarified that on 02Mar2021 that he had blood clots in his left leg. Stated that he had three. Stated that he had the first one on 02Mar2021, second one 13Mar2021 and the third one yesterday (15Mar2021). Stated that he was told left leg was cold. Stated that he took his temperature with a thermometer and his left foot was 79-82 degrees Celsius. Stated that the right foot is 95-96 degrees Celsius. Stated that he filled it out for the Covid vaccination. Stated that he reported Vyndalink when it asked what other medications that he took. The action taken in response to the events for tafamidis was unknown. The outcome of events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1153487-1	Blood clots; This is a spontaneous report from a contactable consumer (reported for herself). A 65-years-old (non-pregnant) female patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Solution for injection, Lot Number: EN6199, expiry date not reported), via an unspecified route of administration, administered in left arm on 03Mar2021 10:45 at a single dose for covid-19 immunisation. Medical history included known allergies to sulpha. The patient had no Covid prior vaccination. Concomitant medications included atorvastatin (LIPITOR) and lisinopril, both taken for an unspecified indication, start and stop date were not reported. The patient had no other vaccine in four weeks. The patient had the vaccination on a public health clinic/veterans administration facility. On 12Mar2021, patient experienced blood clots that resulted in doctor other healthcare professional office/clinic visit, and emergency room/department or urgent care. The patient was not tested for Covid post vaccination. Therapeutic measures were taken as result of the event which included that the patient was put on blood thinners. The event blood clots

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1154074-1	Blood clot/very large blood clot from his thigh to his calf; Joint pain; chills; he can't even walk now; leg pain; This is a spontaneous report received from a contactable consumer (patient's wife). A 57-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot number: EN6202), via an unspecified route of administration, on 25Feb2021 (at the age of 57-years-old), at a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The consumer stated she's asking on behalf of her husband. He received his first dose of Pfizer-BioNTech COVID-19 Vaccine on 25Feb2021 and had Joint pain and chills after that (on unspecified date in 2021) he was diagnosed with Blood clot (unspecified date in 2021); she stated he had no history of similar health issues and wanted to know if it could be directly linked to this vaccine. The wife further reported that it's dramatically changing his life, he can't even walk now. He had the Covid vaccine on 25Feb2021 and he has no history of blood clots. He developed a leg pain (on unspecified date in 2021) which was odd and they thought that maybe it was some arthritis or something like that. They went to the orthopedist and he was being treated for what the orthopedist thought was a bakers cyst. But then they discovered it was a very large blood clot from his thigh to his calf. They would like to know about this. They have been looking into what he had experienced, and it seems very unusual. It just seems too coincidental that this happened after the Covid vaccine. The emergency room doctor of course shut them down immediately and said that the Covid vaccine never would have done this. But the patient's doctor yesterday said that there are too many unknowns to say for sure that the Covid vaccine didn't cause this. She is not saying that they are against the Covid vaccine, but they are concerned. The patient is on blood thinners now and they just need to know if there is a possible correlation. She had heard on the world news that there is a connection after receiving the (company name) Covid vaccine and now they are shutting this vaccine down in (region name). She saw on the news last night they were asking if there is a correlation with the Pfizer and (company name) Covid vaccines and this. It is very concerning. They have been trying to do research and there is nothing that you can find. The only thing that they did see online was about the doctor that died of a blood disorder in (state name), he had thrombocytopenia. The patient is going to be sent to see a vein doctor. She stated that he has a lot of things ahead of him and he is too young to be dealing with all of this. They are going to be following up with the vein doctor and she would like to have this information when the vein doctor calls later today. The reported adverse events resulted to an emergency room and physician office visit (went to the orthopedist). The patient received corrective treatment as response to the reported events. Outcome of the events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1154104-1	Clot blood; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as a single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient experienced clot blood on an unspecified date with outcome of unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1154356-1	Developed a blood clot behind the left knee. Symptoms started on 3/22/2021. Saw physician on 03/23/2021 for tests and diagnosis
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1154610-1	DVT Blood clot, followed by pulmonary embolism
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1158711-1	Blood clots; This is a spontaneous report from a non-contactable consumer through a Pfizer sales representative. An 80-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number and expiration date were not reported) as a single dose, with route of administration and therapy date unspecified, for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. On an unspecified date, the patient had blood clots. The patient was hospitalized on an unspecified date due to the reported event. The outcome of the event, blood clot, was unknown. No follow-up attempts are possible. Information about lot/batch number cannot be obtained. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1158868-1	Nausea, vomitting - Blood clot on kidney. No hx of clotting issues prior.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1159981-1	hemorrhagic bleeding (period), blood clot via bleeding. 1 week of bleeding, required transfusion cbc/esr/crp. now have +factors for antiphosophlid syndrome along CCP. along w/ joint pain in her right hands, fingers and right knee. have swelling to the fingers.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1160713-1	DETAILS OF HOSPITAL STAY: PRESENTING PROBLEM: Cardiac arrest (HCC) [I46.9] HOSPITAL COURSE: Patient is a 74 year old female who receives care through healthcare clinic and second healthcare clinic with past medical history of HTN, CKD, cardiomyopathy/congestive heart failure, atrial fibrillation on Pradaxa who presented to the ED 3/16 after suffering an out of hospital cardiac arrest at her dentist's office. Per report, patient had SBP in the 80s on arrival but was asymptomatic. Prior to start of any procedure (no reports of being given sedative medications), she became unresponsive. CPR was initiated and was found to be in asystole. She received 3 rounds of CPR with ROSC. CT head without acute abnormality. Chest XR showing mild vascular congestion and interstitial edema. Initial labs showing AKI, elevated liver enzymes, BNP >29,000, troponin 39, lactic acid of 11, INR of 6.6, PTT 62, APTT 87. UA with protein, nitrite, moderate blood. Urine culture ordered. Blood cultures ordered. In ED, patient was hypotensive requiring addition of vasopressors. Targeted temperature management was started. Ceftriaxone and flagyl started for possible urinary tract infection and aspiration. Patient with profound coagulopathy, INR increasing to 12.0 on arrival to the ICU. Two units FFP and vitamin K were given. Patient with escalating pressor requirements at this time so CT t/a/p was ordered showing multiple bilateral rib fractures, nondisplaced sternal fracture with small anterior mediastinal retrosternal hematoma, small right sided hemothorax, right chest wall hematoma, patchy bilateral airspace disease consistent with atelectasis/infiltrate/aspiration, diffuse GGO consistent with interstitial edema, enlarged pulmonary arteries consistent with pulmonary hypertension, cholelithiasis. FDP elevated and 2 units of cryoprecipitate given 3/16. Hemoglobin decreased to 5.9 3/17 with INR of 5.4. Two additional units of FFP and additional dose of vitamin K ordered. Two units RBCs ordered. CTA thorax and abdomen 3/17 re-confirmed hemothorax and chest wall hematoma but no active bleeding noted. CT bilateral LE showed no evidence of hematoma. Trauma consulted who recommended chest tube placement. Overnight 3/16-3/17, patient also noted to have seizure activity on EEG and patient loaded with Keppra. Head CT 3/17 negative for hemorrhage or other acute processes. Patient remained in status epilepticus 3/17am and additional Keppra load was given and neurology consulted. Received Praxbind for continued bleeding/coagulopathy. 3/17pm went into PEA arrest with 10 minutes of CPR with ROSC. Bronchoscopy following ROSC noted evidence of bleeding from multiple areas, clots removed. MRI brain showing diffuse anoxic brain injury. Propofol stopped 3/19am. After goals of care discussion this morning, all first degree relatives (daughter and son) all in agreement to transition to comfort care measures. I received call from bedside RN that patient had passed away. On exam, no heart or breath sounds appreciated upon auscultation for 2 minutes. No spontaneous movement or chest rise noted. No pulse palpated for two minutes. Pupils fixed and dilated. No response to noxious stimuli. Time of death 1400 3/20/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1160745-1	Patient complains of shortness of breath about 5-10 min after vaccine. BP was taken because patient stated she has high BP and a history of stroke. BP was158/114. 911 was called and ambulance arrived in 10 -15min. EMT assessed patient and asked her to smile. Patient's left face was not responding normally, Patient asked to lift both arms. Patient could not fully lift both arms. Patient transported to hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1162068-1	Swelling of vocal chords followed by need to blow nose which caused bleeding and clotting in nostrils.; Swelling of vocal chords followed by need to blow nose; Swelling of vocal chords followed by need to blow nose which caused bleeding and clotting in nostrils.; Causes constant congestion.; crusted scabs from nose for over two week timeframe.; This is a spontaneous report from a contactable consumer (patient). An 81-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number; EL9264), via an unspecified route of administration administered in left arm on 09Feb2021 as SINGLE DOSE for COVID-19 immunisation. The patient was not pregnant at the time of vaccination. The patient had no COVID prior to vaccination. Medical history included osteoarthritis, osteoporosis, neuropathy, hypothyroidism, thyroidectomy, hysterectomy, and two recent falls. The patient had no known allergies. Concomitant medication included influenza vaccine (FLU) taken for immunisation. Historical vaccine includes first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: EL3249) on 19Jan2021 at 04:30 PM administered in the right arm for COVID-19 immunization. The patient experienced swelling of vocal chords followed by need to blow nose which caused bleeding and clotting in nostrils. The patient was constantly blowing up bloody clots/crusted scabs from nose for over two week timeframe. At first, was a lot of clots, red blood and scabs. It got better over 14 days, but, still persists. Now clots and scabs are smaller but still frequent. Causes constant congestion. All events started on 05Mar2021 at 12:00 AM. The events resulted in doctor or healthcare professional visit/clinic visit. There was no treatment received for all events. The patient was COVID tested (nasal swab) post vaccination on 09Mar2021 which was negative. The patient had not yet recovered from all events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1162082-1	developed blood clots in my left leg that began swelling some on 12Mar2021 and by Monday 15Mar2021 it was huge and 16Mar2021 big and hurting; developed blood clots in my left leg that began swelling some on 12Mar2021 and by Monday 15Mar2021 it was huge and 16Mar2021 big and hurting; developed blood clots in my left leg that began swelling some on 12Mar2021 and by Monday 15Mar2021 it was huge and 16Mar2021 big and hurting; This is a spontaneous report from a contactable consumer (patient). A 71-year-old male patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Batch/Lot Number: EN6202), via an unspecified route of administration, administered in Arm Left on 02Mar2021 16:00 (at 71-year-old) as single dose for COVID-19 immunization. Medical history included Arthritis in knee. High blood pressure. Patient had no covid prior vaccination and no covid tested post vaccination. Patient has no known allergies. Concomitant medication included hydrochlorothiazide. The patient previously took first dose of BNT162B2 for COVID-19 immunization. The patient developed blood clots in my left leg that began swelling some on 12Mar2021 17:00 and by Monday 15Mar2021 it was huge and 16Mar2021 big and hurting. So he went to urgent care and they sent him to hospital ER. The patient was hospitalized for 3 days. Events resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Hospitalization. Events treatment included Ultra sound, CT scan, heparin blood thinner. The outcome of the events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1162139-1	"He was diagnosed with multiple blood clots in his lungs/ clots may have appeared in his leg and went to his lung; bed ridden; pain in left side; This is a spontaneous report from a contactable consumer (the patient). A 70 year old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH, COVID-19 MRNA VACCINE)), via an unspecified route of administration on 06Feb2021 (Lot number: UNKNOWN) as a single dose for COVID-19 vaccination. The patient medical history and concomitant medications were not reported. Historical vaccination included BNT162B2 (PFIZER-BIONTECH, COVID-19 MRNA VACCINE), dose 1 via an unspecified route of administration on 16Jan2021 (Lot number: UNKNOWN) as a single dose for COVID-19 vaccination. The patient reported that he received the first dose of the Pfizer-BioNtech Covid-19 vaccine on 16Jan2021. On 06Feb2021 he received the second dose of the vaccine. Twelve days later he was diagnosed with multiple blood clots in his lungs, and from ""15Feb2021 to 16Feb2021"" he was hospitalized. He was prescribed to take Eliquis (Apixaban) a blood thinner, at 5 mg tablets for 6 months. He also reported that during that period (first part of Jan and part of Feb) he was bedridden because he had a lot of pain in his left side and it would not let him get out of bed, so he was on bed for a while. His primary care doctor told him that the clots may have appeared in his leg and went to his lung. The patient would like to know if any participant developed blood clots during the clinical trials of the vaccine. The clinical outcome of the events was unknown. Information on the lot number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1162481-1	Two blood clots in my right calf that travelled up to my lungs; Shortness of breath; Pain in my right calf; Had difficulty just walking; This is a spontaneous report from a contactable consumer (patient). A 73-years-old non pregnant female patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), via an unspecified route of administration in left arm on 12Feb2021 19:30 (Batch/Lot number was not reported) as single dose for covid-19 immunisation. Medical history included heart (as reported) and known allergies to penicillin. There concomitant medications were not reported. On 15Feb2021 18:30 the patient experienced two blood clots in my right calf that travelled up to my lungs, shortness of breath, pain in my right calf, had difficulty just walking. It was reported that two nights after receiving the vaccine, she suffered shortness of breath and pain in her right calf. She tried to relax the next day, but had difficulty just walking. She went to a walk in medical facility. They sent her to the ER. After testing they found that she had two blood clots in her right calf that travelled up to my lungs. The event two blood clots in my right calf that travelled up to my lungs was considered as serious (hospitalization) and other event were considered as serious (hospitalization). The patient was hospitalized for all events for 4 days. The patient underwent lab tests and procedures which included blood test on 15Feb2021 with showed two blood clots in my right calf that travelled up to my lungs and sars-cov-2 test with negative results on Feb2021.Therapeutic measures were taken as a result of two blood clots in my right calf that travelled up to my lungs, shortness of breath, pain in my right calf and had difficulty just walking. AE treatment included blood thinner. The outcome of the events was recovering. No other vaccine was given to patient in four weeks and two weeks. No Covid prior vaccination. If Covid tested post vaccination: Yes. Patient was covid tested post vaccination in Feb2021 with negative results. AE resulted in: [Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Hospitalization]. Facility type vaccine was Public Health Clinic facility. No follow-up attempts are possible; information about lot/batch number cannot be obtained. No further information is expected
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1162715-1	I was 2 weeks post prolapse surgery and healing quite well. I thought that would be enough time to wait to heal and get my vaccine. I got the vaccine in the afternoon and that night at about 10 pm I started bleeding profusely with blood clots coming out of my vagina. Before this I had no bleeding post surgery or any complications. This lasted all night. I went to the surgeon next day and the bleeding and blood clots had stopped. 2 days later it started again all night long. I refrained, stupidly, to not go to ER because of the virus. It stopped next day and then 2 days later I woke up in blood and clots and it would not stop. I had to go to the ER my surgeon said to stop it. He added more stitches to my previous surgery.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1163369-1	Blood clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1163438-1	After the first vaccine on 2/6/2021, I did not have any reaction except a sore arm until 2/10/2021 when developed a bad headache causing me to leave work. When I awoke on 2/11/21 I felt worse as the day went on until I had complete flu like symptoms. On Friday, 2/12, I went to my primary care doctor and they did a COVID-19 test which was negative. Hence, I just waited for the reaction to go away, I was not better and able to resume normal activity until 2/15/21. I did decide to get the second shot as I was not entirely sure it was a result of the vaccine. On 2/27/2021, I received the second vaccination. Again, the immediate side effect was only a sore arm. On the following Monday (3/1/2021) see cont.I began to have what I thought was muscle aches and pains around the mid section on my body but they were manageable. On Tuesday and Wednesday (3/3/2021) the pain and area of the pain spread and intensified. I was unable to sleep or find a comfortable position and was losing strength and knew I needed medical attention as I was afraid it was a heart issue. My husband took me to the ER (3/4/2021). The ER doctor ran a CT scan with contrast and the result was a blood clot in my right lung and pneumonia in my left lung along with pleurisy. I was then admitted to Clinic on 3/4/2021. I remained in the hospital until 3/7/2021. There I was treated with IV antibiotics, warfarin, Lovenox (80mg), and pain medication for the pleurisy. I was also on a full time heart monitor and my heart was checked several times. Since returning home, I finished a dose of two antibiotics and am using an inhaler to help with breathing. I have also just finished a long dose of steriods. My INR is still not regulated to the therapeutic level. Prior to this reaction, I was in the therapeutic range and had been checked. As of 4/3/2021 I am at 1.8 not in the 2-3 range and have increased my dose from before the reaction at 7.5 mg to 9.0 mg daily.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1163548-1	10 days after my first shot my feet were swollen and I didn't feel well by 18 days after it was not getting better. I got my 2nd shot on day 21 and the day after I got an alert on my Watch regarding resting heart rate exceeding 125bpm and have been inactive for 10mins, so my husband call the doctor's office to make an appointment. They said we could go to the appointment or hospital, so we went to the hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1163914-1	I have blood and blood clot in my urine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1164912-1	Bloody nose, headache (same as first vaccine). My menstrual cycle started a week early (I?ve been obsessively charting for fertility reasons? I am like clockwork) and was full of large blood clots and painful.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1166696-1	right thigh pain Treatment : lovenox 80 sery
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1167081-1	Arm pain on first day Tendons stiffness on one arm and both wrists on second night 4th day - Early menstrual period (2 weeks early): sudden bright red bleeding without usual signs of menstrual periods (no cramps, very unusual), and large clots in the bleeding Nausea, 2nd day
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1168135-1	Pt developed intermittent shortness of breath around 3/7/21. Pt fell and had left arm and leg weakness. Pt had transient ischemic attack, multiple pulmonary emboli and deep venous thrombosis diagnosis on 3/27/21
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1168386-1	injection received 3/31/2021 at 12:55 pm. no event until next morning, noticed spotting, menstrual bleeding. heavy bleeding started 4/2/2021, 12-13 days before any menstruation expected. VERY SEVERE mental health issues, sadness, hopelessness, irritibility that I do not normally experience with PMS normally. I also am extremely regular with my menstruation and have tracked my period for years, and it is always consistent. as of 4/5 still having very heavy bleeding and clotting, some cramping. bleeding is showing no signs of letting up and my period expected start date is around 4/13. I almost didn't want to say anything because it could be chalked up to hormones or stress but this has not happened to me before, and the emotional state I was in was very bad, not even taking into account the bleeding. I am a very pro science and pro data person so I was hesitant to mention anything at all, but feel like IF this is related this could be a huge risk for folks who have a history of depression and anxiety or post partum conditions. The best way I can describe it was deep hopelessness, something I had not felt since having post partum depression when my daughter was born, 7 years ago.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1168453-1	Died 4 days after shot from blood clots in my lungs. No history of clotting before this. Did have Covid asyptomatic back in January. So not sure whether having had Covid caused the clotting. Having the vaccine or being out of work with torn meniscus. They think clots started in my legs and ended up in my lungs. I was brought back after coding for 6 minutes. They seem to think Covid caused it. Not sure if vaccine contributed or not. But it did happen 4 days after the shot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1169265-1	Acute DVT in left LE. Patient presented to ED on 3/22/21. She had 2 day h/o leg discomfort and leg was taught and painful on the day she went to the ED. Study was limited due to pt lack of cooperation. Lovenox started.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1169566-1	formation of a blood clot in right calf
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1169809-1	DVT / Blood Clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1171097-1	Blood clots (still experiencing... undergoing treatment)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1171196-1	After the shot, which I received on a Saturday, I developed pain and redness in my right leg. Two days later, on Monday morning, I contacted my primary care physician's office, who told me to go to the ER to be evaluated for a blood clot. At the ER, they performed a doppler ultrasound on my right leg and found a superficial clot in the greater saphenous vein of my right leg. There was no evidence of a DVT in the right leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1174114-1	blood clot; This is a spontaneous report from a contactable consumer(patient). A female patient of an unspecified age received BNT162B,(PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. prednisone (PREDNISONE), route of administration, start and stop date, batch/lot number and dose were not reported for allergies Medical history included weakened immune systems and ongoing allergies. It was reported that patient wants to know if there is any reported interactions between the use of prednisone and receiving the COVID-19 vaccine manufactured by Pfizer. She was informed that the vaccine might not work as well for patients taking prednisone. The patient has been on Prednisone for 10 years, caller heard that when the patient has been on Prednisone for that long the vaccine does not work. The patient is taking Prednisone to treat some allergies. Patient also has blood clots and wants to know if patients with a history of allergies should take the COVID-19 vaccine manufactured by Pfizer, and how could she know if she would have an allergic reaction towards the vaccine. The action taken in response to the event for Prednisone was unknown. The outcome of the event was unknown. Information on the batch/lot number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1174174-1	"I broke a pimple and came liquid after taking the pimple out, it came a little thing like clear and then it came like blood; it was like a clot; broke a pimple; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 1 via an unspecified route of administration on 06Mar2021 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. Consumer stated, ""I am kind of concerned because I was showering myself like 30 minutes ago, well I was cleaning my face and well you like night routine and I broke a pimple that I didn't have and came liquid after taking the pimple out, it came a little thing like clear and then it came like blood not too much for sure, then I realized that the blood that came out wasn't liquid, it was like a clot. I cleaned it with a t-shirt and I was able like to move it. So I am kind of concerned because I got my first dosage of the Pfizer vaccine on March the 6th, I am about to get my second dosage this Saturday and as you know and everyone should know right now what's going on with this (name) and (name) vaccine and then what everyone is talking about, it's impossible to not be concerned about the side effects."" The event clot blood was assessed as serious (medically significant). The outcome of the events was unknown. Information on lot/batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1174347-1	She had a stroke on 3/28/2021 due to a blood clot in her brain; She had a stroke on 3/28/2021 due to a blood clot in her brain; Administration_date=08/03/2021 number=1/administration_date=22/03/2021 number=2; Administration_date=08/03/2021 number=1/administration_date=22/03/2021 number=2; This is a spontaneous report from a contactable consumer reporting for herself. A 66-years-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: Unknown), via an unspecified route of administration, administered in Arm Right on 22Mar2021 17:45 (vaccinated at the age of 66 years old) as single dose for covid-19 immunisation. Patient received the first dose on 08Mar2021. Medical history included diabetes; obesity; high blood pressure, iodine allergy. The patient's concomitant medications were not reported. The patient experienced she had a stroke due to a blood clot in her brain on 28Mar2021 18:30, The patient was hospitalized for she had a stroke due to a blood clot in her brain (cerebrovascular accident) for 2 days. Patient visited Emergency room/department or urgent care for events a stroke due to a blood clot in her brain and received treatment. Outcome of event a stroke due to a blood clot in her brain was not recovered. Information about lot/batch number requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1175195-1	experiencing passing a ton of clots; I washed my face, it got extremely red; This is a spontaneous report from a contactable consumer (patient). A 20-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not reported) at the age of 20-year-old, via an unspecified route of administration, administered in the left arm on 19Jan2021 13:00 as single dose for COVID-19 immunisation. The patient was not pregnant at the time of vaccination. Medical history included lupus and idiopathic thrombocytopenic purpura (ITP). Concomitant medications included mycophenolate mofetil (CELLCEPT), esomeprazole magnesium (NEXIUM), medroxyprogesterone acetate (DEPOPROVERA), and propranolol; all taken for an unspecified indication, start and stop date were not reported. The patient has known allergies to rituximab and amoxicillin. As a woman, the patient was having a lot of blood clots being passed. The patient doesn't get a period because of depo, but for some reason, she was experiencing passing a ton of clots on an unspecified date. Also, after the first dose, when she washed her face, it got extremely red on an unspecified date in 2021 (She did the same in the days before and that didn't happen). The patient underwent lab tests and procedures which included Covid test: negative on an unspecified date. The outcome of the events was unknown. No treatment was received for the events. The patient received the second dose of BNT162B2 on 11Feb2021 for COVID-19 immunisation administered at the left arm. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1175716-1	1st symptom 3/19/21 at 5:30 am - Weakness in fingers on right hand. Called nurse line, they recommended calling EMT. EMT said no stroke but could go to urgent care for further evaluation. Urgent care recomended ER. ER recommended appointment with Neurology 2nd symptom 3/21/21 slight droop on right side of mouth. Went to ER in Hospital. MRI revealed clot. Additional ultra sound of heart and carotid arteries clear. Final diagnosis: Subacute CVA-Lt Centrum Semiovale w/ right facial droop and right hand deficits. Recomendations: Nuerology evaluation, Outpatient PT/OT for right hamd deficits. Hospitilization: 1 day Hospital Name: Unnamed City: Unnamed State: Unnamed
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1176884-1	Blood clot left arm.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1177170-1	6 days of heavy periods with large clots up to this morning. First time after several years of only light spotting (IUD/premenopausal). Lighter menstruation today.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1177443-1	Currently, developed 3 blood clots in right arm 4 days after receiving the vaccination. First was noticed on the morning of April 6, the second in the evening of April 6, and the third was noticed on the morning of April 7. First one has appearance of cellulitis located on the forearm. Doctor issued treatment for cellulitis on April 6. Reddish/purple skin colored with grey edges in a circle. Second clot is below the skin with slight bulge closer to the elbow than the first. The third indication is in the upper arm above the elbow on inside of arm. Appearance marked by 2 curved lines similar to varicose veins just below where the clot is bulging the skin. No discoloration of the 2nd and 3rd indications of clot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1178287-1	blood became a clot; pimple; This is a spontaneous report received from a Pfizer sponsored program received from a contactable consumer (patient). A male patient of an unspecified age received bnt162b2 (BNT162B2 reported as Pfizer Covid-19 vaccine), dose 1 via an unspecified route of administration on 06Mar2021 (Batch/Lot number was not reported) as a single dose for covid-19 immunization. The patient's medical history and concomitant medications were not reported. The patient reported that he received Pfizer Covid-19 vaccine first dose on 06Mar2021, and second dose will be on Saturday 27Mar2021. While showering, he broke his pimple on Mar2021 then tried to remove it using the tissue and the blood became a clot. The outcome of the events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1178288-1	he has thrombosis in leg; This is a spontaneous report received from a Pfizer sponsored program. A contactable consumer (patient himself) reported that a male patient of an unspecified age received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as a single dose for COVID-19 immunisation. Medical history included metastatic cancer in the lungs (he's taking chemotherapy) and heart problems from an unknown date. The patient's concomitant medications were not reported. The patient was due the next day for the 2nd dose of the Pfizer vaccine. He was asking if he can take his pre-scheduled medication, especially he's taking chemotherapy because he has metastatic cancer in the lungs. Due also to the nature of him having heart problems, he had thrombosis in leg and he's taking blood thinner since an unspecified date. He was asking if he should stop the medication or if he can take the blood thinner before the 2nd dose. Outcome of the event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1179672-1	DEVELOPED BLOOD CLOT IN RIGHT BREAST - CURRENTLY BEING TREATED

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1182724-1	On Apr 7th at 11:30 Pm my wife went to hospital. not being able to breathe sufficiently. This started the day before on the 6th while out walking. She is in the hospital being treated for embolisms of the lungs on both sides with multiple blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1182734-1	break in rash/rash in mouth and throat; sore throat; dizzy; weak; felt like sleeping all the time; pain in her arm; blood clot in right leg; couldn't walk; veins are swollen; right knee got swollen; ear ache; felt like burning/ sore in her mouth like she was burned; sore in her mouth like she was burned; nausea; headache; slight fever; This is a spontaneous report from a contactable consumer (patient). An 81-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration in right arm on 09Mar2021 (Lot Number: EL9266) at single dose for covid-19 immunization. Medical history included arthritis, respiratory issues, allergies, Eczema of the nipples and breast, arthroscopic surgeries, she lost her voice. Caller stated she exercised everyday, she did yoga. Caller reported the last 2 years she hadn't been able to climb the mountain on the stairs. Caller states she didn't catch a cold, she was fine, she was very active. She was good except arthritic conditions, she has had arthroscopic surgeries, that was a side effect of that. Caller reporting she was a first responder and had some respiratory issues. Caller stated she had some allergies, if with dust or smoke, her throat constricts, it was just small, like her throat. She cannot be around that. She got once in a while, when she got in a rush. Caller stated before the vaccine, she had eczema in her nipples, on her breast. Caller stated she never got sick, besides the rash she got once in a while. Caller states but she was very healthy otherwise, no medication, no smoking, no drinking. Caller stated she was from 9/11, this was what happened with her voice once in a while, she lost her voice. Caller stated she had had some respiratory issues and allergies and so on. Caller stated she was here for 56 years and her mother died of old age so she was not sure. There were no concomitant medications. The patient previously took advil. Caller stated she used to take Advil when she went to the gym and had problems but she hasn't taken Advil for a while. She didn't take meds. Caller stated she didn't drink or smoke because what it did was it gave you a release. Caller stated she was not taking none, she was pushing through and hoping it will heal itself. The patient experienced blood clot in right leg on 26Mar2021, break in rash on an unspecified date, headache on 09Mar2021, slight fever on 09Mar2021, veins are swollen on 12Mar2021, right knee got swollen on 12Mar2021, sore throat on an unspecified date, ear ache on 10Mar2021, nausea on 09Mar2021, rash in mouth and throat on an unspecified date, felt like burning/ sore in her mouth like she was burned on 10Mar2021, sore in her mouth like she was burned on 10Mar2021, dizzy on an unspecified date, weak on an unspecified date, felt like sleeping all the time on an unspecified date, couldn't walk on 12Mar2021, pain in her arm on an unspecified date. Was a first responder, sometimes break in rash once in a while. Got Pfizer 1st vaccine 09Mar, was scared to have difficulty breathing however had headache, slight fever, right knee and veins were swollen, sore throat, ear ache, nausea, rash in mouth and throat also felt like burning also was dizzy and weak and felt like sleeping all the time. Caller stated she was told by her doctor to take the shot on 09Mar2021. Caller stated she had headache, slight fever, then her head was ok the fever was gone but the headache persisted as of today. Caller stated she still had headache and ear ache and her knee got swollen, her right knee. Caller stated for a few days she couldn't walk. Caller reported she has arthritic conditions in knees, she exercises everyday and no problem, but she couldn't walk for a few days. Caller states now she is able to walk on something, but the veins in the knee were still there and she didn't know. Caller stated she was due for second next on Tuesday and she was wondering what to do. Caller stated she was honestly worried about this reaction after the first shot. Caller stated she was told the 2nd had a worse reaction, so she was really weary and worried. Caller stated she only slept 5 hours at night, but now she wanted to sleep all the time. Caller stated she had such a headache and earache. Caller stated it made her nauseous, like in a sea or something. She stated she didn't know how to explain. She didn't vomit like when you were nauseous, it wasn't that feeling. Caller stated it was just lingering like when you were in the ocean. Caller reported headache has gotten better. It has gotten worse and then gotten better. Caller stated the earache was so painful and was still there. Caller reported she got it during the day and when she lied down, but the severity had subsided. Caller stated she didn't measure fever. She wanted to say after that, her mouth was sore after fever from the heat or from the vaccination. Caller stated her mouth was still today, still sensitive. Caller reported it was like her throat and her mouth was so sore like a burn. Caller reported her headache was mostly on the left side, but went to the right side too. Caller reported she could feel it in her eye, it hurt and her ear hurt. Caller reported her sore throat, the upper part, was still sensitive. Caller stated she couldn't swallow, but it was clear, her mouth was clear, only the upper top she would say. Caller stated it's 99% gone, its healing. Nauseous, like in a sea, its not like she has it all day long- nauseous- maybe once a day, nothing severely like what she had for a few things. The patient had a pain in her arm and she had that symptom one night and the following day she had nothing. Her arm was cooperative. Since she had been diagnosed with a blood clot. She saw her doctor on Friday and was recommended that she took some medications. She continued to have headache which she reported, though not as bad. She also reported that she had a sore mouth and earache. She was still fighting symptoms and didn't know what to do with the blood clot. She was scared to take the blood thinner Eliquis and that she will have side effects that it will do her harm she felt Eliquis will do her harm if she received the second dose of the vaccine or takes Eliquis. She

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	was supposed to 20mg daily which she felt will kill her given her body was Adverse Event Description
				not healed from receiving the first dose of the Covid-19 vaccine. There was no prior vaccinations within 4 weeks. She was prescribed Eliquis Film-coated tablet (Lot Number: ABN4449S, Expiry date: Feb2023) 20mg daily for blood clot but had not started because she felt it will kill her was scared to take the second dose of the vaccine. The patient underwent lab tests and procedures which included Sonogram: blood clot in right on 26Mar2021. Therapeutic measures were taken as a result of blood clot in right leg. There was Physician Office Visit for all events. The outcome of the event blood clot in right leg, veins are swollen, right knee got swollen, couldn't walk was not recovered, of event slight fever was recovered on 10Mar2021, pain in her arm was recovered, of event ear ache, nausea, felt like burning/ sore in her mouth like she was burned, sore in her mouth like she was burned was recovering, of other events was unknown.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1182773-1	Massive blood clot in left calf leading to pulmonary embolisms in both lungs (multiple clots in each lung); Massive blood clot in left calf leading to pulmonary embolisms in both lungs (multiple clots in each lung); This is a spontaneous report from a contactable consumer (patient). A 58-year-old male patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration administered in left arm on 21Feb2021 14:00 (at the age of 58-year-old) as SINGLE DOSE for COVID-19 immunisation. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient was not diagnosed with COVID-19 prior to vaccination and it was unknown if the patient has been tested for COVID-19 since the vaccination. The patient's medical history was not reported. The patient had no allergies to medications, food, or other products. The patient had no concomitant medications received within 2 weeks of vaccination. On 10Mar2021 at 11:30 AM, the patient experienced massive blood clot in left calf leading to pulmonary embolisms in both lungs (multiple clots in each lung). The patient was hospitalized for the events for 4 days. The events were also considered as life-threatening. Therapeutic measures were taken as a result of massive blood clot in left calf leading to pulmonary embolisms in both lungs (multiple clots in each lung) which included intravenous anticoagulants and injected anticoagulant. The patient was recovering from the events. Information about the lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1183339-1	Heavy Nosebleed with blood clots - Nosebleed lasted 25 minutes -started in right side nose.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1183938-1	I began to bleed and pass blood clots from my vagina; I began to bleed and pass blood clots from my vagina; This is a spontaneous report from a contactable consumer (patient). A 32-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in the left arm on 11Mar2021 01:15 (at the age of 32 years) (Batch/Lot Number: EN6208) as a single dose for COVID-19 immunisation. Medical history included birth control. Concomitant medications were not reported. The patient is not pregnant. On 13Mar2021 04:00, the patient began to bleed and pass blood clots from her vagina. The event blood clots was assessed as serious (medically significant). The patient visited the doctor for the events and the events were given treatment. The outcome of the events was not recovered.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1184334-1	Two days after vaccine experienced shortness of breath and tachycardia. Went to ER. After testing, diagnosed with pulmonary emboli in each lung (one each). 1 blood clot also found in ankle.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1185441-1	13 days after receiving the second dose I was walking my trash can to the street and I could not breath very well. I went to the Urgent Care and was told I had blood clots in my lungs. I was hospitalized and given treatment for the blood clots.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1185467-1	Developed blood clots in my lungs. Had a pulmonary embolism performed to dissolve the clots.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1186057-1	swelling, redness, and warmth in shoulder of right arm starting about 7-8pm on 04/08, patient went to emergency room on 04/09 where they did an ultrasound and diagnosed her with a superficial blood clot. they told her to take aspirin every day for one month and then return to doctor for a re-evaluation.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1189666-1	Pulmonary Embolism. Emergency open heart surgery. Coded for 4 minutes. In ICU and Hospital. Kidney failure.....Total blockage of left lung. 9 inch clot in right ventricle.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1190415-1	Blood clot in brain causing a small stroke affecting right arm, wrist and hand

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1190936-1	Hypertensive crisis at 180/95; Vascular thrombosis; Head is spinning; This is a spontaneous report from a contactable consumer (patient). A 67-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot Number: EN6198), via an unspecified route of administration in left arm, on 16Mar2021, at a single dose, for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient had no known allergies to medications, food, or other products. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The most recent COVID-19 vaccine was administered in a hospital facility. In Mar2021, the patient's head was spinning, there was a hypertensive crisis at 180/95 for 2 times, and there may be a vascular thrombosis. It was unknown if treatment was received for the adverse events. The events were considered non-serious by the patient. Since the vaccination, the patient has not been tested for COVID-19. The outcome of the events was unknown.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1190955-1	blood clot in the lung; my left leg from the knee down started to swell; my left leg from the knee down started to swell, get really hard and became very painful/I could hardly walk and after and pain was so intense; my left leg from the knee down started to swell, get really hard and became very painful/I could hardly walk and after and pain was so intense; blood clot in the leg; This is a spontaneous report from a contactable consumer (patient). A 69-year-old female patient (patient was not pregnant at the time of vaccination) received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Lot number: EN3248, Expiry date: unknown) via an unspecified route of administration, administered in left arm on 16Feb2021, 15:30 PM at a single dose for covid-19 immunisation (age at vaccination was 69 years). Medical history included blood pressure and thyroid. Concomitant medication patient received within 2 weeks of vaccination included levothyroxine, valsartan, furosemide, acetaminophen. No other vaccine was received within 4 weeks prior to the COVID vaccine. On 16Feb2021, 3 hours after the patient received the vaccination at 18:30 the patient's left leg from the knee down started to swell, got really hard and became very painful. She could hardly walk and afterwards the pain was so intense that she went to the doctor. Doctor took X-rays of her left leg on 16Feb2021, then he diagnosed her with blood clots. He gave the patient 2 heparin shots and immediately sent her to the hospital for a deep vein ultrasound on 16Feb2021. The results showed blood clot in the leg. The patient was prescribed medication and sent home. Her leg continued to be rock hard and painful. On 02Mar2021, the patient was sent for a blood panel work up. There was no improvement. On 08Mar2021, the doctor sent her for a lung CT scan. On the same evening, the patient was called to the doctor's office and was sent to ER immediately as the results showed blood clot in the lung. The patient's dose of Eliquis was doubled. This has now been going for 5 weeks, the patient had not been without pain during this time. Her left leg hardness had gone down a little but was still twice the size of her right leg, knee was huge. The swelling never goes down, not even at night. No covid prior vaccination and no covid tested post vaccination was reported. This case was reported as serious. The outcome of all the events was not recovered. No follow-up attempts are possible. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1190964-1	the problem was a bloodclot; slight swelling of right foot; This is a spontaneous report from a contactable consumer, the patient. A 72-years-old non-pregnant female patient received the first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration in the left arm on 09Mar2021 13:30 (Batch/Lot number was not reported) as single dose for COVID-19 immunisation. Medical history included diabetes, high blood pressure and allergy to tape. Concomitant medications included levothyroxine; olmesartan; metformin; all were taken for an unspecified indication, start and stop date were not reported. The patient did not receive any other vaccine within four weeks prior to BNT162b2. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. On 13Mar2021 15:00, the patient noticed slight swelling of right foot and on 14Mar2021, the swelling increased so the patient texted the nurse who arranged appointment. The nurse said to contact the doctor and the patient spoke with registered nurse (RN) on duty who told the patient to go to the emergency room (ER) because the symptoms indicated the problem was a blood clot. The ER doctor examined and ordered an ultrasound and the results confirmed blood clot. The patient was given a prescription for blood thinner to take for 21 days (times/day, 15mg) and was told to make an appointment with the primary doctor. The patient's doctor was on spring break and she was walked in to see another doctor who said swelling looked improved and showed her where the clots were located. The patient saw her primary doctor today and she ordered a visit with a hematologist as soon as possible (ASAP) because of concerns about possible underlying issues. The outcome of events, slight swelling of right foot and the problem was a blood clot was recovering. The lot number for the vaccine, BNT162b2, was not provided and will be requested during follow up.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1190993-1	blood clots in his lungs; Lot of pain in his left side; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), dose 2 via an unspecified route of administration on 06Feb2021 as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient previously took bnt162b2 for COVID-19 immunisation and had lot of pain in his left side (1 st dose-During period (first part of Jan and part of Feb) he was bedridden). It was reported that caller received the first dose of the Pfizer-BioNtech Covid-19 vaccine on 16Jan2021, in the hospital. On 06Feb2021 he received the second dose of the vaccine. Twelve days later he was diagnosed with multiple blood clots in his lungs, from Feb2021 he was hospitalized, he was prescribed to take Eliquis (Apixaban; blood thinner) 5mg tablets for 6 months. During that period (first part of Jan and part of Feb) he was bedridden because he had a lot of pain in his left side and it would not let him get out of bed, so he was on bed for a while. His primary care doctor told him that the clots might have appeared in his leg and went to his lung. Caller would like to know if any participant developed blood clots during the clinical trials of the vaccine. Therapeutic measures were taken as a result of blood clots in his lungs. The outcome of events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1191609-1	Approximately one week after my second dose my right leg began swelling. For three weeks after the swelling began it would get better than get worse. On Tuesday, March 30, 2021 my entire leg swelled to twice its normal size at which time I went to the emergency room and a blood clot was diagnosed in my right leg. I completed a multitude of tests and bloodwork with a hematologist and there are no factors or reasons they can identify as to why I developed a blood clot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1191810-1	they had a ultrasound done and found out I had a number of clots in that position; a sore Leg, Upper part of the left calf 2 inches below the knee on the backside; This is a spontaneous report from a contactable consumer. A 55-years-old male patient received bnt162b2 (PFIZER-BIONTECH mRNA COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 18Mar2021 15:00 (Batch/Lot Number: En6207) as SINGLE DOSE for covid-19 immunisation. Medical history included Treatable Acid reflex, anxiety and mild depression. Concomitant medications included omeprazole, buspirone and alprazolam for unspecified indication. Patient had the Covid shot on 18Mar2021 at about 3 PM the next morning (on 19Mar2021 06:00) patient woke up with a sore Leg, Upper part of the left calf 2 inches below the knee on the backside. After it did not go away, he went to the doctor then they had a ultrasound done and found out he had a number of clots in that position. Treatment received for the adverse event includes they put patient on a blood thinner. Prior to vaccination, was the patient did not diagnose with COVID-19. Since the vaccination, the patient has been tested for COVID-19. patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient underwent lab tests and procedures which included sars-cov-2 test: negative on 24Mar2021. Covid test post vaccination done via nasal Swab, They had the results in 15 minutes, covid test result Negative, ultrasound scan: clots in that position on 19Mar2021 found out, had a number of clots in that position. Outcome of the events was recovering.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1191916-1	she had developed what she believes is a blood clot in her leg; This is a spontaneous report from a contactable other healthcare professional (nurse) via medical information team. The contactable consumer reported for her mother that a 77-year-old female patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection) intramuscularly on an unspecified date as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient previously took her first dose of bnt162b2, on an unspecified date for covid-19 immunisation. The patient's mother received the second dose of the covid vaccine yesterday morning. When the caller saw her mother last night, she had developed what she believes is a blood clot in her leg on an unknown date. The reporter would like to know whether a blood clot is a possible side effect from the covid vaccine. The patient did not receive any type of treatment for the adverse event. The outcome of the event was unknown. Information on the lot/batch number has been requested.; Sender's Comments: Based on available information, a possible contributory role of the subject product, BNT162B2 vaccine, cannot be excluded for the reported event of Thrombosis leg due to temporal relationship. However, the reported event may possibly represent intercurrent medical condition in this elderly patient. There is limited information provided in this report. Additional information is needed to better assess the case, including complete medical history, diagnostics including venous doppler of lower extremities and coagulation panel, counteractive treatment measures and concomitant medications. This case will be reassessed once additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1191960-1	Blood clot in LAD which caused a STEMI; Blood clot in LAD which caused a STEMI; This is a spontaneous report received from a contactable consumer (female). A 49-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EN6208), first dose via an unspecified route of administration, administered in Arm Left on 20Mar2021 08:15 (at the age of 49 years old), single dose for covid-19 immunisation. Medical history included hypothyroidism, Penicillin allergy, both from an unknown date. The patient was taking unspecified concomitant medications within 2 weeks of vaccination. The patient experienced blood clot in lad which caused a STEMI on an unspecified date (reported as 03Mar2021 15:15, for clarification since before vaccination date). The outcome of the events was unknown. The events was assessed as serious, causing hospitalization, disability and was life-threatening. Treatment for the events were Balloon in LAD / clot buster. The patient underwent lab tests and procedures which included COVID-19: negative on 03Mar2021. Patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Information on Lot/Batch number was available. Additional information has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1193707-1	Post-menopausal (15 years) heavy bleeding & clotting. Vaginal ultrasound showed mass in uterus
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1193782-1	Shortness of breath and chest pain began on Wednesday 02/10/2021 (2 days after vaccine given), Breathing and chest pain severe enough to go to Emergency Room on Saturday 02/13/2021 (5 days after vaccine given). CT scan in Hospital showed blood clotting in both lungs. Was kept overnight in hospital and released on 02/14/2021 with prescription for Eliquis (blood thinner).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1195044-1	PT STATES 4/7 AT 3PM SHE NOTICED SWELLING IN LEG AND WENT TO ER. DIAGNOSED AS SUPERFICIAL BLOOD CLOT IN LEFT LEG. SHE WAS TOLD TO COLD COMPRESS, WALK AND ELEVATE TO REDUCE SWELLING. TODAY SHE FELT DIZZY AND THOUGHT SHE MIGHT HAVE MORE BLOOD CLOTS, SHE WAS REFERRED TO ER AGAIN WHERE THEY GAVE HER ASPIRIN 81MG AND THERE WERE NO ADDITIONAL CLOTS, HER FIRST VISIT CLOT WAS HEALING.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1195233-1	Pain underneath right rib cage and difficulty breathing. It got worse over two days and then I was hospitalized. Coughed up blood. Tested with Heparin drip and now I?m on Xarelto. No previous history of blood clots. This happened three days after second vaccine
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1195691-1	One month after having my second covid vaccine, I had the worse menstrual cycle of my life. I had heavy bleeding for 4 days and horrible menstrual cramps. So bad that motrin/tylenol did not take away the pain. Also passed a lot of clots. Also after my period, I've had continual bleeding/spotting. I've even gone to the OBGYN to get checked. I'm still bleeding...
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1196101-1	Blood Clot On Friday, April 9th I started with a pain in my ankle around 4PM. It was getting worst and worst. It got red, swollen and I was unable to walk normal, I started limping. Around 2PM on Saturday, April 10th, I went to the ER and I had my ultrasound done. I have a blood clot. They prescribe me Eliquis; 10 mg twice a day for one week and 5 mg twice a day for two to six months...
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1196269-1	Patient received Pfizer COVID vaccines 1/26/21 and 2/16/21. Developed dyspnea and fatigue 2/2021. Admitted to hospital on 3/2/21 with extensive bilateral acute PE (5 lobes). Right acute occlusive DVT in femoral/popliteal veins and partial thrombosis in the left (age indeterminate) of the femoral and popliteal veins. TTE ok. Started on heparin and transitioned to eliquis (10 BID x7 days then 5 BID). Normal PSA (0.34)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1196279-1	Received shot on day 4 of period cycle when it was winding down. About 24 hours after receiving, flow began to increase dramatically and continue for 48 hours, with many clots and cramping. Never experienced something like this before.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1198698-1	Patient received his first Pfizer shot on March 20th. He began to have pain in the back of his right knee which sent him to the ED. Upon further investigation, he was found to have a blood clot and was given Eliquis as a blood thinner. His physician requested that he make a report as he received his first COVId shot 8 days prior to the blood clot. He denies any diabetes, chemotherapy that may have irritated his vessels. Gabapentin,
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1198759-1	Sever headache and nausea 24 hours after shot. Then 5 days after shot sever abdominal pains on the right side. Was hospitalized and they found a blood clot in a vein leading from my ovary on the ride side. Treatment was Eliquis and further testing.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1199422-1	Patient passed away from blood clot, did not feel well after 2nd shot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1199631-1	Blood clot. Right leg. Treatment Lovenox injections and Pradaxa blood thinner medication. Time on Pradaxa to be determined on resolution of blood clot.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1200335-1	Blood clot in right hand. Could not feel right thumb pointer and middle fingers and they turned purple. Went to hospital and treated with Heprin. Outcome - still have the blood clot. Put on Eliquis. Also developed high blood pressure from event and put on valsartan
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1200719-1	She had a stroke caused by blood clot in brain. Suffered a second stroke 4 weeks later. Has weakness left leg. Is now in Rehab Center
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201126-1	Blood clot, stroke, severe headaches, dizziness, vomiting
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201376-1	blood clot formed in left leg in surface vein, at intersection with deep vein.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201409-1	Delivered preterm infant (stillbirth) at 24 weeks on March 9, 2021. Infant had IUGR noted at 19 weeks. Placenta showed 2 blood clots restricting blood flow at delivery. Mother with no known clotting disorders. 2 prior healthy infants.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201500-1	Blood clot in leg. Began as pain and skin discoloration in leg, eventually went to emergency room on 07Apr2021 and diagnosed with ultrasound. Treated with Xarelto, ongoing. Acute right lower extremity superficial thrombophlebitis noted in the greater saphenous (above knee). This superficial thrombophlebitis extends more than 10 cm in length. It does not extend into the deep system but is about 1.3 cm from the origin of the common femoral vein.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201939-1	I was hospitalized due to severe blood clotting for four days following surgery to remove scar tissue from the urinary/prostate area and am now on blood thinner medication.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1202529-1	Blood clot in left Iliac vein
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1202540-1	Blood clot in right leg, with pain and swelling. Treatment with warfarin therapy
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1202729-1	On Sunday afternoon, March 14, I received the first dose of the Pfizer vaccination. On the morning of March 16 I had a sudden hit of fatigued and my left leg started going numb while doing some menial work in the yard, The numbness cleared when I went inside and laid down but some form of the fatigue persisted for several days. Then, on March 28, two weeks after the vaccine, I woke up to a leg severely swollen from toe to hip. I waited two days to see if it went down but it only got worse, swelling to double the size by volume. I went to the doctor and they sent me to the hospital for ultrasound, suspecting a blood clot. No blood clot was evident but there were multi swollen limp nodes in the upper leg area. The problem worsened and two days later I was sent in for a CTscan which now showed multi blood clots near the swollen limp nodes. I was barely able to walk as major pain persisted with no signs of decline. I am now on blood thinner. It has been 2 months since the vacination and my leg is still swollen the same amount (2X by volume).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1203153-1	pt says he had chills, fever, headache, body aches and pain. Lymph nodes were swollen under his left arm. A couple days after taking the vax he was passing blood clots when using the bathroom. Pt states this lasted about 4 days. By 4/3/2021 all symptoms had subsided.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1203267-1	58 year old white male with history of blood clots developed symptoms of leg pain during 4/11 early am . pt received COVID vaccine on 4/8 developed leg pain on 4/11 early am and went to er and was advised he was clotting in leg at point of previous procedure. he was given eliquis and was told to follow up with specialist
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1203725-1	10 days after receiving my first Pfizer shot I suffered a stroke that was caused by a blood clot. I was unable to speak for a short time the morning of 4/1/2021 and went to emergency room at. My speech had already returned and I was no longer having symptoms. There were numbers tests and scans run and it was confirmed that I had a stroke that was caused by a blood clot. There were no obvious issue present to why I would have had this happen.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1203817-1	Blood Clot detected liver vein. Dr. placed me on Xarelto

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1204579-1	Jan 13 first dose received...Jan 30th collapsed at night by herself. Was able to get up and go back to bed. Next day was taken to urgent care and sent by emergency to hospital. Suffered large blood clot in the leg that broke off and traveled to the lungs. Acute saddle pulmonary embolism with acute cor polmonale. Was put on blood thinners and released from hospital on 2/2. As soon as we got home from the hospital she suffered from a massive gastrointestinal intestinal bleed and had to be rushed back to the hospital. Two days later while still in the hospital she suffered a mild heart attack and had a stent inserted.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1204630-1	rt superficial lower extremity thrombosis- redness, tenderness, swelling rt lower extremity.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1204715-1	Blood clot in left leg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1204788-1	Diverticulitis arteries; I had a blood clot after ultra sound; sore arm/Feel sore arms after the first dose; spinal nerve damage/Spinal nerve pain; Legs are burning hot now; Pain felt in calf and thigh; Bone density test/Ultra sound bad; This is a spontaneous report from a contactable consumer(patient). An 84-years-old female patient received BNT162B2(PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), dose 1 intramuscular, administered in Arm Left on 01Feb2021 11:30 (Batch/Lot Number: EN5318) as SINGLE DOSE, dose 2 via an unspecified route of administration, administered in Arm Left on 22Feb2021 11:30 (Batch/Lot Number: EN6200) as SINGLE DOSE for covid-19 immunization. Medical history included angina from 2002 and ongoing, She says she has angina, and if she would have beating problems can she take a Nitro, cholesterol, depression from 2008 to an unknown date, blood pressure, anxiety from 2018 to an unknown date, arthritis from 1998 and ongoing Illness/AE: Arthritis Onset Date: 1998-2002 Stop Date: Ongoing , hypertension from 2002 and ongoing, blood cholesterol increased from 2002 and ongoing, ongoing urinary tract infection On and off, coronary artery disease from 2005 to an unknown date On and off, Family medical history included Mother coronary artery disease Died on 18Oct1967 and Father 1936 other. Concomitant medications included amlodipine taken for blood pressure abnormal from 2017 to an unspecified stop date; atorvastatin taken for blood cholesterol abnormal from 2002 to an unspecified stop date; bupropion taken for depression from 2008 to an unspecified stop date; buspirone taken for anxiety from 2008 to an unspecified stop date; carvedilol taken for blood pressure abnormal from 2002 to an unspecified stop date, 2x a day carvedilol 12.5. On an unspecified date, the patient experienced diverticulitis arteries, had a blood clot after ultrasound, sore arm/feel sore arms after the first dose, spinal nerve damage/spinal nerve pain, legs are burning hot now, pain felt in calf and thigh, bone density test/ultrasound bad. The patient underwent lab tests and procedures which included blood test: unknown results, computerized tomogram: diverticulitis arteries- On and off, investigation: unknown results. Comments: Going to pain clinic. On always, ultrasound scan: bad on Comments: Ultrasound bad, urine analysis: unknown results on comments: on and off. Treatment received for the events sore arm/feel sore arms after the first dose and spinal nerve damage/spinal nerve pain. Outcome of all events was unknown Follow-up (29Mar2021): This is a follow-up report combining information from duplicate reports 2021138557 and 2021162095. The current and all subsequent information will be reported under manufacturer report number 2021162095.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1205190-1	her veins were swollen and she had blood clot.; her veins were swollen and she had blood clot.; severe earache; severe headache; her mouth and throat were on fire.; her mouth and throat were on fire.; This is a spontaneous report from a non-contactable consumer (other HCP) via medical information team. A patient of unknown age and gender received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot number and expiration date were not reported) via unknown route of administration on 09Mar2021 as single dose for covid-19 immunisation. Reportedly, patient got the first dose of the Pfizer COVID-19 vaccine on 09Mar2021. After that, she had developed severe earache, severe headache, her mouth and throat were on fire. Few days later, her veins were swollen and she had blood clot. She was also afraid to take her medication, Eliquis, since there may be more adverse events from that medication. Patient was scheduled for her second shot of the Pfizer COVID-19 vaccine today, 20Mar2021. She was told that the adverse events after the second dose of the vaccine are worse than the first dose. She has also read that people die from the vaccine. She wants to cancel her appointment today. She wanted to know if the vaccine would be 80-85% effective after a single dose. She wanted to know if she would be ok with just one dose of the vaccine. No PQC was present. The outcome for the events was unknown. No follow-up attempts are possible. No further information is expected.; Sender's Comments: Based on the information currently available, a possible contributory role of the suspect vaccine BNT162B2 in triggering the onset of thrombosis and other events cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1205201-1	Three days after receiving the vaccine, I developed a blood clot in the lining of my stomach.; This is a spontaneous report from a contactable consumer. A 42-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for injection; Batch/Lot Number: EN6201), dose 1 via an unspecified route of administration, administered in Arm Left on 15Mar2021 11:30 as SINGLE DOSE for covid-19 immunization. The patient medical history was not reported. Concomitant medication(s) included ethinylestradiol, norgestimate (SPRINTEC) taken for an unspecified indication, on an unspecified date. On 18Mar2021 05:30 the patient experienced three days after receiving the vaccine, developed a blood clot in the lining of her stomach (thrombosis). The patient was hospitalized for three days after receiving the vaccine, developed a blood clot in the lining of her stomach for 3 days. Therapeutic measures with blood thinners were taken as a result of three days after receiving the vaccine, developed a blood clot in the lining of her stomach. The outcome of the event was recovered. Information on Lot/Batch number was available. Additional information has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1205261-1	"Diarrhea; Vomiting; Weak; Found Slumped Against the Wall; Low Potassium; MRI Found Multiple Small Emboli in Brain; thrown more clots; Aphasia; Difficulty Walking; potassium dropped and she went into A-Fib and threw clots; potassium dropped and she went into A-Fib and threw clots; has problems with recent memories; confusion; food sensitivities; This is a spontaneous report from a contactable physician (patient's daughter). An 86-year-old female patient received her second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date were not provided) at the age of 86-years-old, via an unspecified route of administration on 24Mar2021 at 14:00 at single dose for COVID-19 immunization. Medical history included high blood pressure diagnosed in 1971 (reported as 50 years ago) and ongoing; type 2 diabetes in 2011 (reported as 10 years ago) which resolved in 2019 (about 2 years ago) after the patient lost some weight; she had an episode of A-Fib in 2019 when her potassium was low (lower than the current hospitalization), once the potassium was corrected, her A-Fib resolved. The physician stated that patient has shrunk and was now about 5 foot 2 inches. Clarified that this has not recently occurred since the vaccine; stated it was happening just with age. The patient received her first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date were not provided), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The patient's concomitant medications were not reported. The patient did not have any issues with the first dose of the vaccine. She received her second vaccine dose on 24Mar2021. The physician spoke with the patient on Friday, 26Mar2021, and the patient was fine. On Saturday, 27Mar2021, the patient had an episode of diarrhea, vomiting and was weak. The lady cleaning the patient's house found her slumped against the wall. The physician met them at the hospital and said the patient seemed okay. They thought it was just an issue from her potassium being low on Mar2021, so they gave her that. The hospital did a CT scan that showed nothing on Mar2021. The hospital did an MRI of the head that showed multiple small emboli in the brain on Mar2021. The physician believed that the patient must have thrown more clots because she became aphasic on Mar2021 and could not name things. The patient also had difficulty walking on Mar2021. The physician felt it was important to call and report this since she has heard of people throwing clots with vaccines. The physician did not know if the events were related to the vaccine. The patient was in the hospital for 7 days (as reported). She was now able to walk with some assistance. The patient's son thought that the patient was okay when he went to see her. Since the reporter was a physician, she knew which questions to ask the patient and saw that she still had some problems. The patient could remember some things but has problems with recent memories on an unspecified date in 2021. She also has some confusion on an unspecified date in 2021, such as she thought she was in a boat hospital. The physician was uncertain if the vomiting and diarrhea was somewhat better. They were wondering if the patient has some food sensitivities (2021). The patient also had carotid studies done in the hospital on Mar2021 and they were fine. They believed that patient's potassium dropped, and she went into A-Fib and threw clots on Mar2021. The clots must have come from the heart valves since the carotid studies were fine and if the clots had been in her legs, they would've gone to her lungs not to her brain. The patient underwent lab tests and procedures which included CT scan: negative, potassium: low, carotid studies: fine, MRI of the head: multiple small emboli in brain, all on Mar2021; and potassium: low, height: 5 foot 2 inches (shrunk), and weight: lost, all on an unspecified date. The events diarrhea, vomiting, weakness and fall required emergency room visit. The outcome of the event difficult walking was recovering and unknown for all other events. The events diarrhea, vomiting, weakness, fall, potassium low, cerebral embolism, thrombosis, and ""potassium dropped and she went into A-Fib and threw clots"" were assessed as serious due to hospitalization and being life threatening; and the events aphasia and gait disturbance were assessed as serious due to hospitalization. All other events were non-serious. The patient was hospitalized from 27Mar2021 until 02Apr2021. Information about the lot/batch number has been requested."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1205576-1	I was diagnosed with a blot clot in my right leg.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1205919-1	"Approximately 7-10 days or so after receiving the first dose of the pfizer Covid vaccine I started to develop this red, nodule-like, painful ""rash"" on both of my forearms. Not only the left arm where the vaccine was administered. This did not go away after a couple of days, and I was instructed by a physician to being taking Benadryl. This appeared to alleviate the symptoms (no visible red nodules anymore) 4-5 days prior to my second dose, so I stopped the medication and presumed it was done. Then, within 4-5 hours of receiving my second Pfizer covid vaccine, I broke out into the same red, painful, nodule-like ""rash"" along both of my forearms. This time was worse than the previous reaction. I was instructed to take Allegra at the time by a physician, which I did for approximately 4 weeks. This seemed to get rid of the symptoms I was experiencing, and I could not visualize the nodules on my forearms anymore. However, after approximately 2 weeks, I noticed a painful and swollen vein in my right forearm, opposite arm to the one I received the vaccine in. I did not think much of it at the time, since I had no other associated symptoms. Then, over the next couple of weeks, it appeared to become more swollen and painful. Also, I saw another swollen vein near my knee. On April 6, I had labs drawn and CBC, CMP, TSH, and ESR came back normal, but the CRP was elevated. Thus, I was instructed to begin aspirin 325mg, which I took for 4 days beginning April 8th and ending April 12th. On April 13th, an official ultrasound confirmed a DVT in my right popliteal vein, a superficial thrombus in my proximal great saphenous vein, and a superficial thrombus in my right forearm where the pain originated from weeks prior. The initial presumption at this time is that the clots are vaccine-related, but could be linked to an underlying blood disorder that was triggered by the vaccine. Blood hyper coagulation labs have not yet come back. I was initiated on Eliquis starting April 12th for the DVT."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1205989-1	Blood clots in leg and both lungs; shortness of breath. Shortness of breath for 1 week before seeking medical attention on March 7th. Admitted to hospital through emergency room on March 8th. Hospitalized from March 8-14. Put on Coumadin. Sent home with supplemental oxygen.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1206421-1	Blood clot in right calf. Noticed pain and knot in calf on Sunday 3/14/2021. It worsened over the next few days. Went to Urgent care on 3/18 and they sent us straight to ER for Ultrasound. In hospital they found it was a clot identified as deep vein thrombosis, and started a treatment of blood thinner and pain medication.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1207618-1	Blood clot left calf
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1207683-1	Blood clot in left leg; first noticed on March 17; swelling and pain bad by March 19, called advice nurse who told me it sounded like a blood clot and told me to go to emergency room.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1208759-1	Blood Clots in both little toes. Eliquis 5 mg, Gabapentin 300 mg, Tramadol 50 mg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1209115-1	On April 6 a.m. developed fewer-like symptoms with muscle pain typical foe fewer with these symptoms steadily worsening by evening. Night shower at appr. 9 p.m. caused a feeling of spinal nerve pinch followed by acute pain during certain movements and breathing. Following day (Wednesday) spent in bed taking Ibuprophene thinking is just a nerve pinch. Called virtual appointment tele doctor who prescribed Katorolac which I took from Wednesday night to Thursday morning per prescription with no effects. On Thursday evening called tele doctor again and got Ibuprophene 800 mg prescription. It helped for 20-30 min only. Same evening coughed some blood. On Friday kept taking Ibuprophene for very temporary relieve. On Saturday April 10 was admitted to hospital and was hospitalized with blood cloth diagnosis for 4 days.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1209140-1	Pain in calf, swelling in lower leg (left leg), weakness/numbing in that leg-led to blood clot diagnosis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1209216-1	Blood Clot-Bad swelling in right arm. Arm tingling, numbness and discolored. Arm is swollen from top of arm to fingers. Notice first sign on Monday morning (04/12/2021) when i awoke from bed around 7:00am.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1209217-1	4/9/21-30 minutes to 1 hour after receiving 2nd dose of vaccine patient had bloody nose, clots, with bleeding sores on tongue and side of mouth. Began to have bruises on arms, feet and legs. Blood in urine and bruises on back. Presented to emergency department on 4/11/21. CT brain revealed a right sided posterior temporal hemorrhage. Patient was emergently treated with platelets, DDAVP, factor VII, Decadron, IVIG. 4/12/21, patient denies headache, changes in vision, double vision, change in sensation, including burning, pinprick, numbness, no focal or motor weakness. No change in bowel or bladder control. No sensory changes. Patient states she is now back to her normal state of health, save for the bruising described.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1209247-1	Severe Vascular Thrombosis (vomit, loss of balance, numbness in arms and legs, lost of speech, impaired vision) had to be hospitalized for 4 days, and had some brain damage from the thrombosis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210248-1	Patient states that 3 days after she received the Pfizer vaccine she started a mentrual cycle. When she came back to the pharmacy three weeks later she stated she was still having her menstrual cycle, and the bleeding was very irregular, sometimes clotty. She said she had just finished her menstrual cycle before receiving the first dose so shouldn't have had a cycle, and that also she was always very regular with her cycles prior to this.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210346-1	Hemorrhage like bleeding, soaked up 4 maxi pads within a 30 minute span. Expelled Blood clots the size of apples. Went to ER. Blood loss and clots resembled the look of a miscarriage, but pregancy test, can back negative. Multiply tests were ran. This carried on from 1:30 to 8pm, when fentaNYL and omipaque was given for pain. Blood Pressure was also high 154/119 and temp 99.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210395-1	I got my second Pfizer covid shot on 2/17/21. Since then I have ahad 2 large blood clots form in my lower right leg. I had one in the same place 24 years ago.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210605-1	I had the 1st shot on Wednesday March 10th. On Sunday the 14th, just before going to bed at about 11pm, I went to pee. The urine was very dark brown. Didn't appear to have any red for blood. The next day I felt like water was being retained and it felt that I couldn't empty the bladder and pee was coming out slower than usual. Late Monday the 15th before bed I peed another dark brown urine and a red blood clot came out. At that point I became very concerned and sent to see the doctor on the 16th. Referred to a kidney specialist for March 31st. Between March 14th to March 29th , somedays I felt like I was retaining water, most days felt like I couldn't empty my bladder with slow flow. With the recent news of the concerns of blood clots with vaccines, I felt that I should report this.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210894-1	Blood clot in lower right leg (calf), resulting in pulmonary embolisms and use of blood thinners. Due to fibroids, this caused excessive bleeding leading to anemic condition and the need for several blood and iron transfusions. Solution was to perform hysterectomy to eliminate fibroids and bleeding to be able to continue use of blood thinners for blood clot condition.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1210966-1	My mother was found unconscious on the kitchen floor. She was taken to the hospital. She had different test including a Cardiac stress test and it was found to be okay. However, my mother developed a blood clot in her left leg. She still been under medical observation in the hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1211350-1	Early on Sunday morning, April 4th, patient woke up with swollen right hand with immense pain. On Monday, April 5th an ultrasound confirmed a blood clot in her right forearm.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1211991-1	Clots Ongoing
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1213306-1	blood clotting in legs, lungs, resulting in hospitalization. Second clotting event resulting in stroke and hospitalization, ongoing treatment and evaluation
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1214146-1	March 14 2021 - 2 days after first vaccine shot, leg pain, 1-3 days later diagnosed with superficial blood clot in left leg via ultrasound, treated with baby aspirin for 3 days and heat pad, symptoms slowly lessened till 2nd dose of vaccine on April 2nd, after second dose symptoms worsened again, both legs still fatigue very quickly to this day (April 15 2021) when I'm sitting down on chairs, seats etc. April 8 2021, 12:30pm - heart pain for 30 minutes, felt very weak, similar to light heart attack, ER visit later that day couldn't find any explanation

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1215649-1	Patient advised she took COVID-19 Vaccine and formed blood clots in leg; This is a solicited report from a non-Pfizer sponsored program: -CW2891702, received from a contactable consumer, based on information received by Pfizer from manufacturer (manufacturer control number: BMS-2021-032881), license party for apixaban. This 86-year-old female patient was involved in a patient support program. The patient (patient ID: (ID)) received APIXABAN. The report describes a case of THROMBOSIS (Patient advised she took COVID-19 Vaccine and formed blood clots in leg). Co-suspect products included Covid-19 Vaccine for an unknown indication. On an unknown date, the patient started APIXABAN (unknown route), (unspecified dose and frequency). On an unknown date, the patient experienced THROMBOSIS (seriousness criterion medically significant). The action taken with APIXABAN (Unknown) was unknown. At the time of the report, THROMBOSIS outcome was unknown. For APIXABAN (Unknown), the reporter did not provide any causality assessments. This report was received from consumer. The patient received therapy with apixaban and COVID-19 Vaccine for unknown indication. Follow-up is unable to be performed. Medical evaluation comment: This patient had blood clots in the leg after apixaban therapy. Based on the limited information regarding medical history, event details, treatment details, it cannot be ascertained that the suspect drug contributed to the reported event. The reporter's assessment of the causal relationship of the event Thrombosis leg with the suspect products was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality assessment. Amgen's assessment: the event, Thrombosis leg, was assessed as unrelated to apixaban. No follow-up attempts are possible; information about batch number cannot be obtained.; Sender's Comments: As there is limited information in the case provided, the causal association between the event of thrombosis and the suspect vaccine BNT162B2 cannot be excluded. Based on known mechanism of action of the drug, the event of thrombosis is assessed as not related to Apixaban. The case will be reassessed once new information is available. The impact of this report on the benefit/risk profile of the Pfizer drug is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1215846-1	Severe abdominal pain,nausea,emergency hospital visit,atrial fibrillation,thrombosis,left.kidney infarct.admittance to local hospital for 5 days and ongoing follow up testing and multiple specialist visits for months
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1215952-1	Began with a severe cough 02/27/21 lasting for days....possibly causing blood clots in the lungs.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1217054-1	Painful cramps, heavier period on first day with clots. Not my usual period, heavier.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1217866-1	Systemic: Blood Disorder (diagnosed by MD)-Severe, Additional Details: Patient hospitalized for blood clot (called pharmacy @ 835pm on 4/15/21). Asked by hospital to contact pharmacy. Patient called from personal phone number.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1218142-1	Nausea. Heavy bleeding including clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1218741-1	Severe Shortness of Breath 10 days after vaccine. Discomfort in the right lower chest that radiates to the back.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1219609-1	symptoms pain in left arm continue April 6, 2021 every day went to see doctor on April 14, 2021 was prescribe a inflammation and muscle relaxer. Left arm continue to hurt the next day with finger tips on left hand begin to tingle, I call my doctor he sent me to have a ultra sound of my left arm and do blood work. upper extremity duplex ultrasound
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1219629-1	EXTENSIVE BLOOD CLOT ,DVT AND PE
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1219841-1	50 hours after receiving vax shortness of breath and found blood clotting in lungs and legs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1220036-1	I had intense stomach pains, muscle pains for 4 days. Could not get out of bed. Also had a fever. I thought these were regular side effects and eventually got better. Then, about a week later I started getting nose bleeds with blood clots coming out of my nose. These lasted a few more days. This never happened to me before, and must have been a side effect from the vaccine

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1220442-1	Developed a 7mm Blood Clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1220494-1	I started being really short of breath and felt pain/inflammation in lower body. I remember telling my brother I hurt everywhere below my waist; was just home from work where on my feet unless I sit down to rest. It all culminated with increased breathlessness and went to emergency at hospital at 8:30 pm on April 22, 2021. Diagnosis was MANY pulmonary embolisms, some large. I have taken week long road trips for the past several years with hours and hours at the wheel with no problems. It just seems like a real strange coincidence of the vaccine and subsequent symptoms.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1220690-1	On the evening of March 6, 2021, patient tried to exercise (consistent with regular routine) and couldn't do more than 15 minutes. He was very winded. On March 7, we skipped exercise. On March 8, patient tried to do simple stretching exercise, and couldn't do it. He got winded and shaky. On March 9, he was seen at an urgent care where he was told he had an inverted T wave on his EKG and a very fast heart rate (125 resting pulse). He was told to not exercise and to make an appointment with a cardiologist. On March 9, I called the urgent care clinic to ask questions about monitoring his blood pressure and when to seek more medical care. I spoke with Dr, who prescribed Atenolol. Around 4:30 pm on
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1221024-1	Extreme bruising at allergy shot site Nose bleed and blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1221080-1	Lower left leg pain, swelling and redness 3 days after 2nd dose of Pfizer Covid vaccine. Ultrasound confirmed superficial blood clot from ankle to knee.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1221447-1	Started having headache, chills and body aches on 4/1/2021. On 4/2/2021 I had major fatigue. On 4/4/2021 my right leg began to swell terribly and then on 4/6/2021 it was discovered that I had one big and many small blood clots in my right leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224177-1	Blood clot blocking blood flow to brain - 1st episode: (3/12/21) stabilized, minor limited movement left side - 2nd episode: (3/24/21) no blood flow to brain, death (maintained on life support for organ donation)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224239-1	Had blood clot in left leg. Blood in eyes. Sinus sore throat short breath congested headache vision problems. Tested positive for strep one month after receiving one vaccine been taking antibiotics for 6 days but I'm getting worse.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224262-1	strange pain on her left collar bone; going to see someone just in case it's a blood clot; lump on her collar bone, left collar bone; does hurt a little bit; This is a spontaneous report from a contactable consumer or other non hcp (patient herself). A 19-years-old female patient received bnt162b2 (BNT162B2, Formulation: Solution for injection), dose 1 intramuscularly administered in Arm Left (like the shoulder area) on 28Mar2021 09:00 (Lot Number: ER8734 and Expiry date: unknown) as single dose for covid-19 immunisation. The patient medical history was not reported. There were no concomitant medications. No additional Vaccines administered on same date of the Pfizer suspect. On 30Mar2021, the patient experienced going to see someone just in case it's a blood clot, lump on her collar bone, left collar bone, does hurt a little bit and on unspecified date strange pain on her left collar bone. 19-year-old female got the first Pfizer Covid vaccine at 9 am on Sunday, March 28 and this morning, she woke up with a strange pain on her left collar bone and there was a lump there, states when she woke up, she noticed a pain, checked in the mirror, she saw on the left side she saw a bump and checked on the right side and didn't see anything and that's when she got worried. Caller states she was just calling to report it, she's worried that it's a blood clot or something else. Why was the patient taking Pfizer BioNTech COVID Vaccine (Verbatim): she wanted to be vaccinated to protect herself and her community. Caller states she would not like to provide her last name at this time. However, provided her last name initial. She also reports she would not like to provide her address, height, or weight right now. She states she used to volunteer at a hospital. She does not see expiration or NDC number on card she received when she got her vaccine. Caller states she does know that one of her family members has the Factor 5. She is not sure if she has it but would like to be tested for it. The outcome of the event going to see someone just in case it's a blood clot was recovered, lump on her collar bone, left collar bone and does hurt a little bit was not recovered and strange pain on her left collar bone was unknown. Information on batch/Lot number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Lower chest partial atelectasis or scarring in the lung bases bilateral; Partial lung collapse (both upper and lower parts collapsed); Lower chest partial atelectasis or scarring in the lung bases bilateral; Beneath this area was a red circle where blood clotted or something; Wheezing; Shortness of breath; Swelling of the face; underneath the arm where the injection site was, a big red circle was noted and it was swollen/under her nose and left elbow were red; underneath the arm where the injection site was, a big red circle was noted and it was swollen; I would get hot and cold; I was sweating; Pain in her chest, stomach, back and head; Pain in her chest, stomach, back and head; Pain in her chest, stomach, back and head; Pain in her chest, stomach, back and head; Fatigue; Pain in left elbow; goes to bathroom and bowel was not emptying the way it should; rib cage is going on the side of body noticed this thumping; Her lips were red; Throat was red; Been sick and just laying around; Her eyes were red; Cramp in left arm where site of injection; Sharp pains were hitting her in her eye, thigh, chest, head, and the arm; Sharp pains were hitting her in her eye, thigh, chest, head, and the arm; This is a spontaneous report received from contactable consumer (patient). A 70-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via unspecified route of administration on 15Mar2021 (at the age of 70 years) at a single dose in the left arm for COVID-19 immunization. Medical history included lung issues and allergic asthma. Family history included brothers and sisters who died of lung issues and third sibling sister died many years ago due to allergic asthma. The patient previously took flu vaccine for immunization and experienced severe pain throughout her whole body. On 15Mar2021, patient received the first dose of Pfizer COVID-19 vaccine. Later on, that day when she got home after receiving the vaccine, underneath the arm where the injection site was, a big red circle was noted and it was swollen, beneath this area was a red circle where blood clotted or something, she would get hot and cold and take her clothes off because she was sweating but did not have fever, there was pain in her chest, stomach, back and head, wheezing, shortness of breath, fatigue, swelling of the face, pain in left elbow. She goes to bathroom and bowel was not emptying the way it should. She also noted that her rib cage is going on the side of body noticed this thumping, lips and throat were red. She's been sick and just laying around, her eyes, under her nose and left elbow were red, had cramp in left arm where site of injection, and sharp pains were hitting her in her eye, thigh, chest, head, and the arm. Patient clarified that these symptoms began on 15Mar2021, the same day she received the Pfizer COVID-19 vaccine. On 23Mar2021, patient went to the ER and the findings were lower chest partial atelectasis or scarring in the lung bases bilateral and partial lung collapse (both upper and lower parts collapsed). At the hospital her temperature was 98.9 F on 23Mar2021, but 97.1F - 97.7F is her normal temperature. She went to the Pulmonary doctor also, but was getting a second opinion because she knew it could be life threatening. The pulmonary doctor made light of it saying they were just creases, like creases in pants. Patient said crease was a far cry from partial lung collapse on both bases and the Pulmonary doctor also said that it is in the upper part of the lung also. This scared her and made her think nobody cared. She did not know if she should provide the information because the doctor did not seem to believe that that her symptoms were coming from receiving the Pfizer COVID-19 vaccine. She guessed she should provide the information because Pfizer does research. She wanted to know how long do the side effects last and if she should take the second dose. The outcome of the events underneath the arm where the injection site was, a big red circle was noted and it was swollen/under her nose and left elbow were red, pain in her chest, stomach, back and head, wheezing, shortness of breath, fatigue, swelling of the face, pain in left elbow, goes to bathroom and bowel was not emptying the way it should, rib cage is going on the side of body noticed this thumping, lips were red, throat was red, been sick and just laying around, eyes were red was not recovered; while unknown for the other events. Information on the lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224338-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224392-1	Developed a blood clot behind the left knee; This is a spontaneous report from a contactable consumer. A 72-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 1 via an unspecified route of administration, administered in Arm Left on 18Jan2021 16:00 (Batch/Lot Number: EL3246) as SINGLE DOSE, dose 2 via an unspecified route of administration, administered in Arm Left on 17Feb2021 (Batch/Lot Number: EN6200) as SINGLE DOSE for covid-19 immunisation. Medical history included Known allergies: Feathers. Concomitant medication included cetirizine hydrochloride (ZYRTEC [CETIRIZINE HYDROCHLORIDE]) taken for an unspecified indication, start and stop date were not reported. On 22Mar2021 18:00, the patient Developed a blood clot behind the left Knee. Ae resulted in : [Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care]. The patient received treatment for the event: Blood Thinner Meds and Compression Stocking. The outcome of the event was not recovered. The patient did not have covid prior vaccination, and the patient was not COVID tested post vaccination.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"hoped she wasn't having a heart attack; Elevated Pulse/high pulse/pulse was 165 beats per minute/climbed up to 195; Drenched in sweat/sweating/sweating like a dog in the middle of the night; Wasn't feeling well; Tightness in chest; Cough; Sinuses congested; Sore; Puffy, bubbled up arms, inflamed; Puffy, bubbled up arms, inflamed; Feverish/felt hotter than usual; Really, really tired/wiped out tired; so sleepy; Sore Arm/her arm started to hurt; Nauseous; Sore Throat/throat hurts; eyes half closed/eyes closing; Puffy Face; blood clot; This is a spontaneous report from a contactable consumer (patient). A 72-year-old female patient received the first dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number ER8733, expiry date: unknown), via an unspecified route of administration, on 29Mar2021 11:45, as single dose, for COVID-19 immunisation. Medical history included ongoing latex allergy (developed welts where the straps met the skin) diagnosed 20-30 years ago; ongoing dye allergy (developed warm soreness to leg, tightness in the chest, could not breathe) diagnosed many years ago; sulphite allergy (diagnosed about 1975-1980; cannot eat processed foods, bleached foods, frozen pre-made foods; tries to eat only single item foods; last anaphylactic reaction after eating olive oil at a restaurant because it had sulphites in it; Mother was possibly allergic to sulphites; Son also allergic to sulphites); and blood pressure that runs low. The patient previously took epinephrine and experienced epinephrine allergy; ciprofloxacin (CIPRO) and experienced Cipro Allergy (eyes closed, got puffy, couldn't breathe); amoxicillin for abscessed tooth and had a reaction because of the sulphites in it; epinephrine (EPIPEN) and experienced epinephrine allergy (does not use it anymore); plantago ovata (METAMUCIL) which she then found out the orange one had sulphites. Historical vaccine included flu shot for immunization and she experienced 104-degree temperature and was sick as a dog and she went to the emergency room (she had flu shot maybe 3 times). Ongoing concomitant medication included multivitamins. The patient got her first dose of BNT162B2 on 29Mar2021 at 11:45. On the same day, once she got home, she became really, really tired/wiped out tired; was so sleepy; experienced sore arm/her arm started to hurt; was nauseous; had sore throat/throat hurt; eyes half closed/eyes closing and puffy face. On 30Mar2021, the patient experienced elevated pulse/her pulse got too high/pulse was 165 beats per minute/climbed up to 195. Her heart rate was 194 at some point for a couple of minutes then stayed at 165 for a long time. She hoped she wasn't having a heart attack. She had tightness in her chest, wasn't feeling good and was coughing. She stayed in her bed to wait out her symptoms. These lasted for 10 minutes (as reported). She started meditating hoping her heart rate would decrease and it did after about 15-20 minutes. The patient also felt feverish/felt hotter than usual. Before she took the vaccine on 29Mar2021, her temperature was at 96. She stated that her temperature usually runs low as 95-06 degrees. She also felt like her sinuses were congested; was sore; had puffy, bubbled up arms, inflamed. The patient hoped it was not a blood clot she developed. At the time of the report, the patient stated that her heart rate was 61 beats per minute, earlier it was in the 70s; and her temperature was at 96.3. Her second dose is scheduled on 19Apr2021 and mentioned that if these were her reactions with the first dose, she is questioning getting the second dose. The outcome of ""blood clot"", ""hoped she wasn't having a heart attack"", ""Wasn't feeling well"", ""Tightness in chest"", cough, and sinuses congested was unknown. The patient recovered from ""Drenched in sweat/sweating/sweating like a dog in the middle of the night"", and ""Elevated Pulse/high pulse/pulse was 165 beats per minute/climbed up to 195"" on 30Mar2021; and was recovering from ""Sore Arm/her arm started to hurt"", and ""eyes half closed/eyes closing"". The outcome of the remaining events was not recovered. Information on the lot/batch number has been obtained. Additional information is expected."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224396-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				tip of left ring finger turned white and numb/finger was completely white and the rest of her hands were blue and red and white and yellow; tip of left ring finger turned white and numb; her finger felt funny; Reynauds phenomenon; sore arm; fever; chills; headache; swollen lymph node in the armpit of the arm she got her Covid 19 vaccine in; swollen lymph node started with a sharp pain and it's very sore; left foot, it was pulsing and throbbing all night; feet were modeled red, blue; feet were modeled red, blue; very sick; needed her glasses trying to give her primary care doctor's phone number; has something like gout in her foot; The swollen lymph node started with a sharp pain and could be a blood clot as far as she knows; This is a spontaneous report from a contactable consumer (patient) via the medical information team. A 57-year-old female patient received first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot: EW0150; expiry: 31Jul2021), via an unspecified route of administration in left arm on 31Mar2021 01:30 as single dose for covid-19 immunisation. The vaccination facility type was a warehouse. The vaccine was not administered at a military facility. No previous immunization with the Pfizer vaccine considered as suspect. No vaccines administered on same date with the Pfizer vaccine considered as suspect. No prior vaccinations (within 4 weeks) and no following prior vaccinations. Medical history included covid-19 in Nov2020. The patient had no concomitant medications. The patient had the first dose of bnt162b2 on 31Mar2021 and 12 hours later (on 01Apr2021), sore arm, fever, chills and headache. Got a swollen lymph node in armpit, same arm as injection left side, today tip of left ring finger turned white and numb. She thinks it was Reynauds phenomenon. The patient called and told nurse in her doctor's office and they said if it was painful, it could be a pain clot. It was painful at first, that lymph node started 24 hours later with a sharp pain, and then she just assumed it could be a lymph node. Swollen lymph node started with a sharp pain and it was very sore. The nurse told the patient to call Pfizer. The patient was fine until the tip of her finger happened. The patient said If it was painful it could be a blood clot, would need an x ray and asked if did the shot cause this. The complete ring finger left hand, the patient looked it up and it said it could happen to her feet and she looked, and her feet were modeled red, blue it was bizarre. Never happened before. The patient was not happy about getting the vaccine, she already had covid in November of last year (2020). The patient wouldn't have called if it wasn't for her finger. Not just her feet, her hands got like this too. The patient didn't want to get the second shot. The patient asked if there was any information or mandate on timing information for the second dose, should she needed to postpone it. The patient asked if what was the efficacy after first dose as she felt like she was immune because she had the virus and now the vaccine. She was very hesitant to get the Covid 19 vaccine. She thinks her body created Covid 19 antibodies. A couple days ago she was very sick. The swollen lymph node started with a sharp pain and could be a blood clot as far as she knows (in 2021). Even putting on deodorant she can tell it's there, it was very sore and there's a lump. Today (on 01Apr2021), she noticed the tip of her finger got white it was numb. She looked it up online and she thinks it was Reynaud's phenomenon. She has a picture if Pfizer needed it. The finger was on the same arm that she received her vaccine in. It was the ring finger on left hand. Once it started happening her finger felt funny and felt numb. She put her finger under warm water. She reported that the white color was maybe a third of the way down towards her knuckle. She was asking if this was normal or has this been reported. She had called her doctor today and spoke with the nurse who told her to call the CDC and report her symptoms. She needed her glasses trying to give her primary care doctor's phone number. The patient wanted to add that she has something like gout in her foot and on that side too, the left foot, it was pulsing and throbbing all night. She's had this pain in her left foot, she thinks its gout because it was all the symptoms of gout and she had it for a while, it hurts off and on but that night it was hurting and throbbing. She has not been diagnosed with gout in that left foot. She still has a headache but today she felt good except for her finger turning white and going numb. The lymph node in her armpit hurts. Her doctor's nurse told her if its painful to the touch it could be a blood clot so she was nervous about that. The tip of her finger was completely white and the rest of her hands were blue and red and white and yellow. She reported that her finger was back to normal now. She didn't want to get too inoculated since she also had the Covid 19 virus. The events did not require visit to emergency room nor physician office. The outcome of the events was unknown for thrombosis, Raynaud's phenomenon, erythema, cyanosis, illness, visual impairment, and gout while for the remaining events was recovering.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1224608-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224636-1	Blood clots after the 2nd dose; Blood clot split and some went into side of lung; This is a spontaneous report from a contactable consumer (patient). A 73-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration in the right arm, on 14Feb2021 at 15:45 (Lot Number: EL3247) (at the age of 73-years-old) as a single dose for COVID-19 immunisation. The patient had no medical history, family history, or concomitant medications. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot number: E10142 (as reported)), in the right arm, on 24Jan2021 at 15:45 (at the age of 73-years-old) for COVID-19 immunisation. The patient did not receive any other vaccines within four weeks prior to the vaccination. The patient experienced blood clots after the 2nd dose and blood clot split and some went into side of lung in Feb2021, which required hospitalization from 27Feb2021 to 28Feb2021. The clinical course was reported as follows: The patient stated he did not know what date the blood clots occurred and can't tell when that happened, but it was maybe a few weeks after the second dose of the vaccine. He stated that one blood clot is in the right leg near the calves and the blood clot broke and split some and went up into the side of his lung as it was detected in a doppler on 27Feb2021. The patient was hospitalized from 27Feb2021 to 28Feb2021. The doppler was performed as soon as he went into the emergency room and they went through a series of tests with unknown results on 27Feb2021 and 28Feb2021. The patient stated that the right leg is bigger than the other leg and they told him it might stay that size and may not go down. The patient is now on apixaban (ELIQUIS) 5 mg for the rest of his life. In the beginning, they gave him an unspecified shot in his stomach, then after that he began taking apixaban at two pills in the morning and two pills at night, then they cut him down to 1 pill in the morning and 1 pill in the evening, and he started it right when he went into the hospital immediately. Therapeutic measures were taken as a result of the event as aforementioned. The clinical outcome of blood clots after the 2nd dose and blood clot split and some went into side of lung was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"This is a spontaneous report received from a contactable consumer (patient). A 54-year-old male patient (height: 193cm, weight: 99.79kg) received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, batch/lot number: EN6207) via an unspecified route of administration at right upper arm on 18Mar2021 18:00 (54-year-old at time of vaccination), at single dose, for COVID-19 immunization. The patient's medical history included blood pressure, teeth cleaned (Shortly before the shot, went to the dentist to have teeth cleaned. Directly from Dentist went to pharmacy. Went to pharmacy's Drive Thru to pick up a little antibiotic from the dentist. It was 4 capsules of Amoxicillin. He was supposed to take the Amoxicillin before the dentist office visit, but he forgot to do that. Therefore, soon as he left the dentist office he went to pharmacy to pick it up so he could take it. While he was there, was offered the COVID 19 vaccine and that it was the last shot). Concomitant medications included atenolol tablet from 2016 (been taking for 5 years) and ongoing for blood pressure, and ascorbic acid/ cyanocobalamin/ ergocalciferol/ nicotinamide/ pyridoxine hydrochloride/ retinol/ riboflavin/ thiamine mononitrate (ONE-A-DAY). No additional vaccines administered on same date. No history of all previously immunizations. No prior vaccincations wtihin 4 weeks. He got the shot Thursday and by Sunday, many different side effects were coming in. He has had side effects from the COVID VACCINE, but initially he does not want to file report, just thinks that everyone needs to get the shot. He does not have complaints. He is wondering how long the side effects will last, and when they go away. States that he is being treated as he has gone to the doctor's. Also states that he made an appointment with the doctor and almost went to the emergency room a couple of times. Right after the shot, he was getting in the car and his right leg started hurting (on 18Mar2021). It didn't go away for a couple of days. It went away but then noticed his shoes started feeling funny. He wasn't looking at his feet, put his shoes on they felt tight. Then he started to look at feet more. Below his knees, his legs are swollen. Swelling started within 2 to 3 days after the vaccine. The swelling is really severe when standing. Everybody at work has seen how sick he is and his legs. Swelling in the legs has him concerned, thinking maybe it's a blood clot. He was given medicine and would like to see it go away. He keeps his legs up. He has other side effects (all in Mar2021) and is having flu like symptoms. He basically has the flu. He has never had the flu shot. This is the first flu shot (He referred to the COVID vaccine as the flu shot as documented. However, confirmed suspect product as Pfizer covid vaccine) he has had that he knows of. He is questioning if he should get the second shot. He feels a little dizzy. Quick questions get him confused and that it comes and goes. Lungs felt full, gasping for air: not now but, since the doctor gave him antibiotics (unspecified injections, he was given 2 shots at the doctor's office, one in each arm), his lungs felt full. At times he was gasping for air. It went away but thought he was going to die. Now when he gets winded, it feels like he has a dry cough. Gets winded easily. The doctor ordered him an inhaler, Symbicort, but it was very expensive, so he did not get it at that time. He never picked it up because it was too much money. The fluid is not there anymore and he does not feel like he needs this as does not feel like it needs to be cleared. He is having wheezing. He can now take deeper breaths. Fatigue: initially, that he was feeling tired within 1 or 2 days, but he did not recall the next day feeling that way (later stated it was within 2-3 days he started to feel tired). He is still tired. Unable to clarify time frame of fatigue further. He feels bloated and full. Hasn't really eaten but feels real full and bloated. Hot and cold flashes started occurring about the same time as the other symptoms. He had these last evening. He does not get the flu very often and doesn't take the Flu shot. This is the closest thing he can remember having a pretty good episode with the flu. He was also given prescriptions for 2 new medications: Triamterene 37.5mg daily, for leg swelling; Clindamycin, 300mg, twice a day, an antibiotic. The odd thing is that it seems like his eyes are big, like really, really big, like they are really open. It is like big eye balls. States that he is just sick and his body is reacting to getting better. He has not been to work in the last 3 days. States that he is fine, just worried about work. Patient asked for personal advice about going back to work. The outcome of ""his right leg started hurting"" was recovered; outcome of ""below his knees, his legs are swollen/ swelling in the legs/ put his shoes on they felt tight"" and Fatigue/feeling tired was not recovered; outcome of other events was unknown."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224670-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224703-1	Some blood clot on both of her legs; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as a single dose for covid-19 immunization. The patient's medical history and concomitant medications were not reported. The patient experienced some blood clot on both of her legs on an unspecified date. It was reported that patient would have a procedure on Friday and then she was going to go for the vaccination on Saturday. they would like to known if there was any problem for her to have the vaccination done. The outcome of the event was unknown. No follow-up attempts are possible; information about batch number cannot be obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224898-1	Stroke due to a large blood clot to the brain; Stroke due to a large blood clot to the brain; Widespread blood clots in both of her arms; This is a spontaneous report from a contactable consumer (daughter). A 67-year-old female patient received the 2nd single dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration, in the left arm, on 17Mar2021 (at the age of 67 years old, not pregnant at time of vaccination) (Lot number was unavailable since unable to locate or read the details), for COVID-19 immunisation. The patient's medical history included ongoing atrial fibrillation, high blood pressure, and scoliosis. Concomitant medications included ongoing acetylsalicylic acid (ASPIRIN) for atrial fibrillation, naproxen, pregabalin, tramadol, and metoprolol. The patient never had a history of blood clots. There were no allergies to report. The patient had not received any other vaccines in the four weeks prior to receiving BNT162b2. The patient had received the first single dose of BNT162b2 on an unspecified date, in the left arm. The patient's daughter stated that her mother suffered a stroke due to a large blood clot to the brain on 25Mar2021 at 12:30 am, requiring hospitalization. During hospitalization, a surgery to remove blood clot from the brain was needed. Hospitalzation lasted 12 days. The reporter added that in the week prior to the reporting, the mother 'passed' (as reported) but had widespread blood clots in both of her arms. The patient required unspecified treatment due to the events and had not recovered at the time of the reporting. The events were considered also serious since life threatening and due to disability. The patient did not have COVID-19 prior to the vaccination and had not tested positive post vaccination. Information on lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224924-1	"I had emergency surgery to remove a femoral blood clot from my groin to the bottom of my left leg; This is a spontaneous report from a contactable consumer (patient). A 67-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in left arm on 17Mar2021 12:00 (Batch/Lot Number: EP6955) as single dose for covid-19 immunisation at a Pharmacy or Drug Store. Medical history included known allergies to sulfa drugs, latex, animal dander, dairy, wheat, mold, mildew, ragweed and other wildflowers; and clot once before in the same leg-left inner thigh in 1974. Concomitant medications included thyroid (ARMOUR THYROID) and apixaban (ELIQUIS). The patient previously had a pneumonia vaccine on 24Feb2021 in the left arm. The patient does not know if her first dose of Pfizer vaccine caused this, but on ""04Mar2010"" (as reported) she had an emergency surgery to remove a femoral blood clot from her groin to the bottom of her left leg which started on 03Apr2021 at 12 AM (as reported). The event resulted in emergency room/department or urgent care visit, hospitalization for 2 days and was considered life threatening illness (immediate risk of death from the event). She had a clot once before in the same leg-left inner thigh in 1974. The patient was on Eliquis 5 mg 2 pills 2 X day for two more days, then 1 pill in am and 1 pill in evening for rest of her life. The patient had the second dose of BNT162B2 (lot: ER8732) on 07Apr2021 in the left arm. The patient has no covid prior vaccination and was not tested for covid post vaccination. The event recovered with lasting effects."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224928-1	Blood clot DVT in left leg.; Blood clot DVT in left leg.; This is a spontaneous report from a contactable consumer. A 48-year-old non -pregnant female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), at the age of 48 years, via an unspecified route of administration, administered in the left arm on 01Apr2021 10:45 (Batch/Lot number was not reported) as a single dose for covid-19 immunization. The patient has no medical history. Patient has not had Covid prior to vaccination. The patient has no known allergies. The patient's concomitant medications were not reported. The patient experienced blood clot DVT in left leg on 02Apr2021 at 09:00. Treatment for the event includes blood thinners. It was reported that the AE resulted in: Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Life threatening illness (immediate risk of death from the event). The patient underwent lab tests and procedures which included nasal swab for covid test on 02Apr2021 which had a negative result. The outcome of the event was not recovered. Information about Lot/batch number is requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1225687-1	Hospitalized for what was determined with Guillain Barre -Miller Fischer Syndrome. Symptoms slowly started a couple weeks after the vaccine and eventually worsened to the point of not being able to open my mouth. Symptoms included facial paralysis, Muscle weakness, fatigue, vision issues, Neuropathy issues. I was transfused with multiple doses of Immunoglobulin and IV steroids. Still currently in Physical therapy I have lost 3 months of not being able to work. Also had a blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1225939-1	The adverse event occur the day after receiving the 1st dose of the Pfizer covid-19 vaccine. I received the 1st dose on Sunday March 28,2021 and start feeling pain on my right leg in the calf area. The area was hot and swollen and in pain, these symptoms started Monday March 29. 2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1225999-1	Repeated, extreme nosebleed with large clots. I was unable to stop it and had to go to the ER on 4/16. They had to cauterize the site of the bleed. I do have a history of mild nosebleeds but this was unusual. I do not take any blood thinning medications. The silver nitrate did stop the bleeding. The ER ductus was not concerned that this was vaccine related but I thought it was unusual enough to report.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1227158-1	found a deep blood clot.; my calf, foot, and knee swelled up by evening./It stayed swollen for a week; my calf, foot, and knee swelled up by evening./It stayed swollen for a week; This is a spontaneous report from a contactable consumer (patient). A 61-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection) (at 61-years of age), dose 1 via an unspecified route of administration, administered in arm right on 17Mar2021 12:00 (Batch/Lot number was not reported) as a single dose for COVID-19 immunisation. Medical history included blood clot after a back fusion about 16 years ago (2005), and had a knee replacement 3 months before the vaccine (Dec2020). The patient's concomitant medications were not reported. The patient previously took ANCEF [cefazolin sodium] and had allergies; and oxacillin which the patient had allergies and hepatitis. The patient was not diagnosed with Covid prior to vaccination nor was he tested for covid since vaccination. The patient did not receive any other vaccine within 4 weeks prior to Covid vaccine. It was reported that the day the patient received the first vaccine his calf, foot, and knee swelled up by evening. It stayed swollen for a week and he went to see his doctor. He did an ultrasound on his calf and found a deep blood clot. He was now on blood thinners. He mentioned that he had a knee replacement 3 months before the vaccine but it was progressing fine. Usually blood clots occur after surgery within a 2 week period. He had a blood clot after a back fusion about 16 years ago. He was reporting this because his doctor was not sure why he got the blood clot and it coincided with the vaccine. The patient underwent lab tests and procedures which included ultrasound scan: found a deep blood clot in Mar2021. The events started on 17Mar2021 18:30 with outcome of recovering. The events resulted to doctor or other healthcare professional office/clinic visit. Therapeutic measures were taken as a result the events with LOVANOX and PRADAXA. Follow-up attempts are completed. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1227231-1	cerebral venous sinus thrombosis; CT scan that revealed the clot; horrible headache after intercourse; vomiting; This is a spontaneous report from a contactable consumer (patient). A 29-years-old non-pregnant female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot Number: EP7534), second dose via an unspecified route of administration, administered in left arm on 25Mar2021 09:45 as single dose for COVID-19 immunisation. Patient was 29 year old at the time of vaccination. Medical history included asthma, gastrooesophageal reflux disease, peanut and treenut allergy, all from an unknown date. Patient had estrogen allergy. Patient was not diagnosed with COVID-19 Prior to or after the vaccination. Concomitant medications included cetirizine hydrochloride (ZYRTEC ALLERGY), medroxyprogesterone acetate (DEPO PROGESTIN), beclometasone dipropionate (QVAR), OMEPRAZOLE, SERTRALINE HYDROCHLORIDE, all taken on an unspecified date and for unknown indication. Patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Patient had her first dose of bnt162b2 on 04Mar20210 at 9:45 am in left arm for COVID-19 immunisation. On 28Mar2021 at 09:45 the patient was diagnosed with cerebral venous sinus thrombosis a few days after receiving the second dose of the vaccine. She had a horrible headache after intercourse, went to the ER after vomiting, and had a CT scan that revealed the clot. The patient was hospitalized for cerebral venous sinus thrombosis. The patient underwent lab tests and procedures which included computerised tomogram: revealed the clot. Patient received Heparin and Eliquis as treatment for the events. Outcome of the events was recovering. No follow-up attempts needed. No further information expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1227273-1	massive clots; Pulmonary Embolisms; Occluded Left Lung, Right Ventricle; Occluded Left Lung, Right Ventricle; This is a spontaneous report from a contactable consumer (parent). A 44-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not reported) via an unspecified route of administration administered in the right arm on 15Mar2021 13:45 as a single dose for COVID-19 immunisation. Medical history included multiple sclerosis. The patient is not pregnant at the time of vaccination. The patient has no known allergies and has not had COVID prior to vaccination. Concomitant medications included gabapentin, carbamazepine (TEGRETOL), fluoxetine, colecalciferol (VITAMIN D), fish oil, and baclofen. The patient did not receive any other vaccine in 4 weeks prior to the COVID vaccine. On 30Mar2021 06:00, the patient experienced Pulmonary Embolisms, Occluded Left Lung, Right Ventricle and had emergency open heart surgery to remove massive clots. He coded for 4 minutes and was in ICU for 9 days. The patient was hospitalized due to the events on an unspecified date for 12 days. The events resulted in Emergency room/department or urgent care, Hospitalization, Life threatening illness (immediate risk of death from the event), and Disability or permanent damage. Treatment for the events included open heart surgery. The patient underwent COVID test post vaccination via nasal swab on an unspecified date with unknown results. The outcome of the events was recovered with sequelae on an unspecified date. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1227958-1	Blood Clots; Pulmonary Embolisms; DVT (Deep vein thrombosis); This is a spontaneous report from a contactable consumer (patient). A 47-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: ENG207), dose 2 via an unspecified route of administration, administered in Arm Left on 25Mar2021 10:00 (at the age of 47years) as single dose for Covid-19 immunization. The patient received the dose 1 of BNT162B2 via an unspecified route of administration, administered in Arm Left on 04Mar2021 08:30 (Batch/Lot Number: ENG203) as single dose for Covid-19 immunization. The patient is not pregnant at the time of vaccination. Medical history included essential tremor, insulin resistance, and anxiety; all from an unknown date and unknown if ongoing. Concomitant medications included propranolol, metformin, bupropion, and citalopram; all taken for an unspecified indication, start and stop date were not reported. The patient was not diagnosed with COVID-19 prior to vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced blood clots, pulmonary embolisms, and a DVT (deep vein thrombosis) on 04Apr2021 (11:00AM). The patient was hospitalized due to the events for 4 days. The patient was treated with Heparin and Eliquis. The patient was tested for Covid-19 post vaccination via Nasal Swab on 03Apr2021 with Negative result. The outcome of events was recovering. The adverse events resulted in Emergency room/department or urgent care, Hospitalization, Life threatening illness (immediate risk of death from the event).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1227979-1	got clots all over the body; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient experienced got clots all over the body in Mar2021 and he passed away. Pfizer has not been reporting any effects of blood clot. The patient died on an unspecified date. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested.; Reported Cause(s) of Death: got clots all over the body
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1227980-1	Blood Clot/The Blood clot was in the legs and went to his lungs; Blood Clot/The Blood clot was in the legs and went to his lungs; fell; Leg pain; This is a spontaneous report from a contactable consumer. A 51-year-old male patient received his first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 27Mar2021 (Batch/Lot Number: EP6955) as single dose for covid-19 immunisation. Medical history included thrombosis from 2020 to an unknown date, it was Months ago in 2020, He was prescribed a medicine for blood clots, but since then he has had no issues, anxiety from an unknown date and unknown if ongoing. Her brother in law had anxiety about even getting the vaccine. The patient experienced leg pain on Mar2021 , pulmonary thrombosis, leg thrombosis and fall on an unspecified date. The patient died on 02Apr2021. An autopsy was not performed. The clinical course was the following: The Blood clot was in the legs and went to his lungs, his leg never got red or anything and it was a blood clot. He was having such bad leg pain, it's too bad, had there been some sort of warning, he would have thought to wait till next year. There was nothing the day he got vaccine, but that night his leg started hurting, his girlfriend said let me go get a cold wrap and she wrapped it, he's a farmer. He wakes up the next day and his leg is still hurting, the 3rd day it goes on, his leg still hurting and no one is thinking it a blood clot, he calls the doctor and tells the doctor his leg is hurting, they suggest ibuprofen for inflammation, the next day thought maybe it's better, by Friday he was making breakfast, fell and died. With his girlfriend she said what's going on, and he went straight to hospital, they said the clot in his leg went to the lung. The Blood clot when it was in the leg it never got hot, red, or anything like that. He had the vaccine in the morning and on that same day the leg pain started later that night. It Started out with Leg Pain, she does not know if it was right or left but it was just one leg.; Reported Cause(s) of Death: Thrombosis pulmonary; Thrombosis leg; Fall; Leg pain
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228293-1	Patient is a 46 yr/o female who presents with vaginal bleeding, started 3 to 4 days ago, states she is used 3-4 tampons this morning in the last 4 hours, states seems to be better currently, but waxes and wanes. She has had some clotting. Last. She believes was 1 week later than normal, around the first of this month. Denies sexual activity in the last 10 months, denies concern for STIs. No dysuria or flank pain. No fever or chills. Thought she could be anemic secondary to her fingers tingling earlier today. She does have an OB/GYN but has not seen them for a couple of years. Did have uro-Guynn surgery last year. Diffuse lower abdominal pain, fairly constant. No treatment prior to arrival. Denies weight gain or concern for pregnancy.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228364-1	On Monday April 12, my mother reported leg pain which she thought was related to a flare up of her knee (surgery 4 years ago). She described her pain as a strong sharp pain, that spread throughout her calf area. She reported it as increasing in pain and worsening daily. By Friday April 16th I asked her to report to Emergency Department. At which time she was told she had a blood clot in her right leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228406-1	A week after receiving the shot, I started experiencing severe pain in my left calf. On Friday, April 9, I was diagnosed with 2 blood clots in my left leg (Thrombus in the posterior tibial vein on the left.).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228508-1	"Pt was found in severe distress in the bathroom by his wife. He couldn't walk or stand, complained of being dizzy with slurred speech and was throwing up a lot. 911 was immediately called. The EMTs during their assessment said his BP was normal but when he was asked his name, he started saying the alphabet""ABC... XYZ"". He was transported by ambulance to ER. Cat scan showed he had a bleed on the brain. The doctors told us it was because he had very high blood pressure, however where the bleed was, they said he wouldn't need surgery and that the body would reabsorb it over time. He was admitted to the ICU and has been there ever since. Several days later, he started having difficulty breathing and they discovered several blood clots in his right leg that had broken off into his heart and lungs. He had emergency surgery to remove the clots in his heart (approximately 10) and was given heparin in small doses over an 8 hour period so that it wouldn't cause the bleed in his brain to increase. The doctor should fill in the rest."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228562-1	Period began a week earlier than usual and thus far as been very heavy with additional clotting. So far I am on day 3 of this period and had to change my cup 4 times throughout the night where I usually only need to change it once or twice. As I am very anemic this is concerning.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228712-1	Patient had blood clots in right leg and in each lung.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1228783-1	Patient received her 1st dose of Pfizer COVID vaccine on 02/26/2021. She noted some wheezing and slight shortness of breath and shared this information with her PCP at her regularly scheduled visit on 03/01/2021. PCP refilled inhaler at that time. Her shortness of breath progressively worsened, and she was seen by another provider as a walk-in at the office of her PCP on 03/13/2021. At that time, she was prescribed prednisone for 5 days, noting some improvement after a few days. She received her 2nd Pfizer COVID vaccine on 03/19/2021. The patient woke up on the morning of 03/20/21 with a swollen and painful leg and by 03/22/21 called PCP and was seen in the office. PCP ordered D-Dimer and LE ultrasound. Per the patient, the D-dimer was abnormal and the LE U/S showed a blood clot. PCP sent the patient to the ER, where a chest CT was ordered and showed a pulmonary emobolus. She was admitted overnight and started on Eliquis, and discharged the next day after seeing a hematologist.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1229122-1	Blood clots-1. 4/14/21 @ 1:35 pm on right side of mouth where the tooth was pulled. Tooth on the right was pulled on 4/9/21 @ 10:00 am. This is the side the abscess was on & 10 day amoxicillin was prescribed. 2. 4/17/21 @ 12:31 am woke up blood on pillow went to bathroom rinsed mouth out blood cloth came out just like the same as 4/14/21. Prior to vaccine no problems.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1229461-1	I had my first dose of Covid 19 Pfizer vaccine on the 5th of April, 2021 at 1pm. I had sore arm and was fatigued which was expected. However, the day I had my vaccine was also the day my menstrual cycle started and just to assure myself I informed the pharmacist and asked if that was okay and they said it shouldn?t be a concern. I was doing fine until toward the end of day 3. I was expecting my period to stop as it always does on 3rd day but I started bleeding heavier that day there after. It didn?t worry me much until I started discharging huge blood clots as big as an apple 2 times every after 30 mins from that moment. I rushed to the ER later that night(still bleeding heavily with blood clots every now and then) but I was not admitted since my vitals and the blood counts were normal at that moment. I went back home and hoped the bleeding would stop but again with on pain or fever what so ever I kept bleeding all day. This is my 5th day of bleeding. Then on Sunday morning, I still kept bleeding heavy and I was too weak that I went back to the ER again. This time , (day 6) I got admitted because I have had lost a lot of blood and I needed blood transfusion. My HB dropped from 11.3 to 6.7 and it went further down to 5 something. I got 3 units of blood transfusion in the ER. They did ultrasound, CT scans(with and without contrast) and still couldn?t figure out the cause of the bleeding. My uterus showed no problems but my body was not even responding to Tranexamic acid to stop the bleeding. So the doctor proceeded with an angiogram and did an emergency uterine artery Embolization to stop the blood supply to the uterus. The bleeding stopped there after and I started regaining my vitals. However, the cause of the bleeding still remains unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1229471-1	Patient had 1st Pfizer vaccine on 3/13/21. She states she went to ER on 3/19/21 and was found to have a blood clot behind her left knee (found by ultrasound). She was put on Eliquis. Patient had 2nd Pfizer vaccine on 4/3/21. Patient did not mention blood clot at time of 2nd vaccine. Patient was admitted to hospital on 4/13/21 for stroke. It was attributed to bleeding in brain and patient was taken off Eliquis and put on a new blood thinner. She will be discharged on 4/20/21 to rehab.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1229499-1	Multiple blood clots, bleeding, hematoma, bleeding events after heparin, swelling in lower body, gangrene in both feet, double below knee amputation, situation ongoing, still in hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1229685-1	On 04/01/2021 I had a random nose bleed and passed a clot the size of half of a bean.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1230273-1	Partially occlusive peroneal blood clot that developed the week after he received the second Pfizer COVID vaccine
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1230622-1	Had a blood clot and a stroke and blind in right eye, and some paralysis in face. Happened day after first vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1230832-1	"4/4 2021 - Red raised Rash at the injection site about the size of a quarter 4/5 2021 _ Itchy raised rash, sore to the touch about the size of a dollar piece 4/6 2021 - Itchy raised, hard and hot. sore to the touch about 3"" round 4/7 2021 - Itchy raised, hard, sore to the touch about 3 1/2"" round Called Dr. Was told to take Benadryl, Tylenol, compress If no improvement the next bay then I had to come in to the office. I started to see some improvement-not as red. over the next two day it continued to itch less, redness was fading, still itchy and hard. By the 11th they swelling was down to the size of a quarter and redness was gone but still itchy. 4/18 it was about the size of a pea still a little itchy but doesn't hurt and no redness. However, on 4/13 around 6:30 pm I had to be rushed to the hospital couldn't breath, vision and speech was declining . was admitted for blood clot that cut the oxygen to brain causing a mini stroke"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1230862-1	Blood in left leg. Hospitalized for 3 days. Now on Eliquist. Couldn't walk on left leg. Had to pull the cord and maintenance called the ambulance. Still having pain in left leg even after a cortisone shot still not better and they don't know why. Next appoint is on Thursday. There is another appoint at the pain clinic for her back pain tomorrow
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1231003-1	Blood clot left arm, arm swelling
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1231186-1	Approximately 11 days after receiving my first dose, I began feeling dull pain in my left calf. By Saturday, 4/17, the pain was severe enough to warrant a trip to the emergency room. After receiving an ultrasound of my leg, two blood clots were found in my left calf. One fully occluded a vein and another partially occluded a different vein.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1231390-1	After first shot some swelling in lower leg, after shot 2 severe swelling and blood clots both legs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1232173-1	Superficial venous thrombosis of left arm Start 2 days after shot, heat, ibuprofen
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1233356-1	Blood clot in left nasal cavity
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1233567-1	I was diagnosed with Covid-19 on 12/02/2020 which was six weeks before my first dose of the vaccine on 01/21/2021. I also tested positive on 12/16/2020. I started to have a lot of problems with my legs. I had a lot of pain in my right leg. I was also very short of breath. It was excruciating pain at the top of my right thigh. I have been a runner in the past so this in unusual for me. I have a blood clot that was found on a scan with contrast. I have seen an ENT, Neurologist, Cardiologist, Orthopedist and Hematologist. I am also a cancer survivor of Non-Hodgkin?s Lymphoma. I have been prescribed Eliquis and taking it for one week and the pain has improved.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1233682-1	Blood clots developed in leg following week on Thursday. Went for treatment Monday.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1234130-1	Hematuria. Severely thickened urinary bladder with blood clots in organ and urine . Please note 1st Doze of Covid-19 vaccine was on 03/25/2021 and 2nd doze was on 04/15/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1234219-1	On Friday 4/16/21 evening, I had my left tonsil spontaneously rupture without any provocation. I was not eating or drinking but felt like I had something in my throat. I coughed it into a tissue, and it was a blood clot the size of a nickel. My tonsil had 2 places that it was oozing blood and continued to bleed for about 1.5 hours. It eventually tapered off and the redness/bruising disappeared over the next 48 hours. There was no pain, no warning - just a mouth full of blood. The tonsil that ruptured was on the same side as where I received the injection and I had swollen lymph nodes under my arm on that same side until Sunday 4/18. I can't help but think it was related to the vaccine as I had no other cold/hay fever/allergy symptoms of any kind to explain a bleeding tonsil.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1234443-1	After the first dose, mild dizziness and nausea followed by massive inflammation in joints and face, severe fatigue and loss of hearing in right ear. Hearing returned in 3 days, general malaise until a week before 2nd shot when she started having severe bloody noses and EXTREMELY heavy period. After the second dose, 45 minute nose bleed soaked through two wash cloths, so many blood clots it clogged the sink.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235719-1	left arm swelling started from 4/18, extended from upper arm to lower arm in the afternoon. went to Urgent care on 4/19 morning, the NP reviewed my blood report and referred me to do ultrasound and also ask me to go to ER if the swell extend. as the swell extend, i went to ER in the afternoon. blood test shows high D-dimer and then ultrasound finds blood clot in the vessel near left neck. diagonised to be Deep Vein Thrombosis. no family history of such problem. in the hospital, they gave me a shot of blood thinner and also prescript oral blood thinner to be take at home.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235740-1	I fainted; blood clots; light headed; I lost all color and my skin was wet and clammy; I lost all color and my skin was wet and clammy; This is a spontaneous report from a contactable consumer. A 60-year-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) lot number: EN6199, via an unspecified route of administration, administered in right arm on 04Mar2021 15:00 as single dose for Covid-19 immunisation. Medical history included known allergies to Atovaquone/proguanil. Concomitant medications were not reported. On 04Mar2021, approximately 10 minutes post vaccine, the patient fainted in the waiting area. She was told by a witness that she fainted two times. EMTs administered an IV and did a simple EKG. The patient lost all color and her skin was wet and clammy. EMTs were concerned about her heart and transported her to the ER where she had a full EKG and lab work for D-Dimer to consider blood clots. The patient has remained lightheaded for weeks after the vaccine. The outcome of the events was recovered with sequelae.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235767-1	small clot behind my left knee; left foot had swollen; This is a spontaneous report from a contactable consumer (patient). A 20-year-old female patient received BNT162B2 (lot number: ER2613) first dose on 20Mar2021 10:15 on left arm at single dose for COVID-19 immunisation. Medical history and concomitant medications were none. Patient is not pregnant. No other vaccine in four weeks. No Covid prior vaccination, No known allergies. She took the vaccine around 10:00am and around 6:00pm (also reported as 06:30 PM) she noticed that her left foot had swollen. She consulted with her general physician and he requested that patient did an Ultra Sound exam of her left leg. The result of the exam reported a small clot behind her left knee. Therefore, doctor suggested that she reported these events to Pfizer so an evaluation is made to identify if similar events have occurred with other patients. She was concerned if she should take the second shot on 17Apr2021. AE resulted in Doctor or other healthcare professional office/clinic visit. Patient received Eliquis 5mg (two times daily) as treatment. Patient had Nasal Swab on 01Apr2021 and tested negative. The outcome of the events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235798-1	Blood clots in legs where old injury was; This is a spontaneous report from a contactable consumer (patient). A 83-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), first dose via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. Medical history included patient had an old leg injury several times in the past, which was fine. The patient's concomitant medications were not reported. The patient experienced blood clots in legs where old injury was on an unspecified date with outcome of unknown. Received call from consumer who stated she had a speech impediment. She was calling about the Pfizer COVID vaccine. She had the 1st shot. She had an old leg injury several times in the past, which was fine. This time it had big blood clots that appeared since the vaccine. She was afraid to go for second shot. She would like some guidance about getting second shot because she was afraid to go get it. Unable to confirm if speech impediment was prior to receiving vaccine or after. No further details provided. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235823-1	DVT in right leg from calf to thigh; small blood clots; swollen right leg; This is a spontaneous report from a contactable consumer (patient). A 69-year-old male patient received bnt162b2 (BNT162B2; lot unknown: Not available/provided to reporter at the time of report completion), dose 1 via an unspecified route of administration, administered in Arm Left on 20Jan2021 14:00 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunization, at the age at vaccination of 68 years old. Medical history included high blood pressure from an unknown date and unknown if ongoing. No other vaccine in four weeks. Patient had no known allergies. Concomitant medications included losartan taken for an unspecified indication, start and stop date were not reported. The patient experienced dvt in right leg from calf to thigh on 25Jan2021 with outcome of not recovered, small blood clots on 25Jan2021 with outcome of not recovered, swollen right leg on 24Jan2021 10:00 AM with outcome of not recovered. Reported four days after the first vaccine, swollen right leg. Five days after the first vaccine, diagnosed with DVT in right leg from calf to thigh. Small blood clots. Events resulted in: Doctor or other healthcare professional office/clinic visit, Life threatening illness (immediate risk of death from the event). No Covid prior vaccination. No Covid tested post vaccination. Therapeutic measures were taken as a result of events included: Eliquis. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235824-1	blood clot; This is a spontaneous report from a contactable consumer (patient) from a Pfizer-sponsored program. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on an unspecified date (Lot number was not reported) at single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The patient experienced blood clot (life threatening) on an unspecified date. Event as reported: Patient had a blood clot due to our vaccine. The outcome of the event was unknown. Information about lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1235838-1	Extreme inflammatory response. In ER/hospitalized 5 days post vaccine; Had Angiogram, found clot in Obtuse Marginal; This is a spontaneous report from a contactable consumer (patient). A 54-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 21Mar2021 12:00 (Lot Number: EN6207), age at vaccination of 54-years-old, as single dose for covid-19 immunisation. Medical history included cardiac failure from an unknown date and unknown if ongoing, connective tissue disorder from an unknown date and unknown if ongoing. Patient was not pregnant. Concomitant medications included hydroxychloroquine sulfate (PLAQUENIL [HYDROXYCHLOROQUINE SULFATE]) taken for an unspecified indication, start and stop date were not reported; isosorbide mononitrate (IMDUR) taken for an unspecified indication, start and stop date were not reported; metoprolol (METOPROLOL) taken for an unspecified indication, start and stop date were not reported; rosuvastatin calcium (CRESTOR) taken for an unspecified indication, start and stop date were not reported; furosemide (FUROSEMIDE) taken for an unspecified indication, start and stop date were not reported. The patient previously took codeine and experienced hypersensitivity. The patient experienced extreme inflammatory reaction on 24Mar2021 at 04:00 AM and clot blood on 24Mar2021; events were serious as resulted in Emergency room/department or urgent care, hospitalization (for 5 days), life threatening illness (immediate risk of death from the event). The patient had extreme inflammatory response; in ER/hospitalized 5 days post vaccine. Angiogram was performed and clot in obtuse marginal was found; unable to remove clot, surgeon was able to break it apart and balloon the vessel open. It was unknown if patient had COVID prior vaccination. Patient had no other vaccine in four weeks. Sars-cov-2 test was negative on 15Apr2021. The patient was considered to be recovering from the events.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1237045-1	In the first week of April, patient developed bruise-like markings on her left upper arm where she received the vaccine. The following week she noticed swelling and discoloration of her left arm. She went to Urgent care on April 14th where bloodwork and an ultrasound came back normal. Over the weekend of April 17-18, the swelling got worse and she called her PCP on April 19th, who advised her to go to the Emergency Dept. At the ED, bloodwork, an ultrasound and CT of the chest was ordered and she was diagnosed with a blood clot. Acute deep vein thrombosis of axillary vein of left upper extremity. She was given a prescription for rivaroxaban.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1237463-1	Right ankle swelling upward to knee; diagnosed with blood clot per patient; treated with Eliquis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1237543-1	Patient developed palmar digital vein thrombosis in 4th digit of left hand
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1237755-1	pt says that he started getting a blood clot and his left hand wrist. He had this for 2 days. It disappeared and then the next day he discovered he had another blood clot on his right hand palm. He says it is a bump and he can see the blood clot under the skin. Pt will contact his PCP.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1238310-1	About 12+ hours after vaccination I began experiencing pain in the back of my right leg. On 4/14 my PCP sent me to the ER. After an ultrasound it was found that I had blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1238672-1	Pain in bilateral lower extremities that started after 1st dose of COVID vaccine. US completed on 4/16/21 showed superficial blood clot in right leg. Treatment included: warm compresses, taking ibuprofen 600 mg with food three times daily, and elevation of your right lower extremity. Outcome: unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1238999-1	I was having my menstrual cycle when I got my second vaccine. Approximately 24 hours after receiving my second COVID vaccine, my bleeding became very heavy and I passed a large amount of clots. This lasted approximately 8 hours then stopped. Previous periods have been light, as I had a tubal ligation and ablation several years ago. After this incident, I did not have a period for 2 months. My cycle has since resumed.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1239002-1	Venous Doppler on left leg, 2 blood clots found. Taking Xareto.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1239036-1	Blood clotting, hospitalized, emergency, paralyzed, thrombosis, body not responding, therapy, treatment.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1239294-1	I started having right leg swelling. I noticed a small red spot on my medial side of my knee. I had this for about a week and then the swelling went down. A few days later, I had redness, skin was warm to touch, and hardness developing where the red spot was. I went to urgent care and the NP sent me to the ER for a possible blood clot. I had a venous study done in the ER and it was determined I had a superficial blood clot and was started on ASA 325 mg daily x7 days. I followed up with a vascular doctor and it was determined the clot was worse and had clotted all the way up the vein into my thigh. It remains superficial in nature at this time. I was started on Xarelto 10 mg daily x1 month. I follow up with vascular in 1 month to make sure the clot is dissolving. I was instructed to Follow up with either the doc or go to the ER with worsening symptoms.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1239772-1	After an arterial catheterization, a blood clot developed in my forearm, about an hour later.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1240013-1	not feeling well and short of breath on 2/11/21, 2/12/21 more short of breath ambulance came. Went into PEA in ambulance with CPR. They gave TPA for suspected blood clot. He initially improved. He did not recover and died 4/4/21. He spent the entire time in hospital or TCU with complications. We brought him home 4/2 to die at home.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1242000-1	15 days after the 2nd shot i developed a blood clot in my left leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1242057-1	03/17: Got 2nd dose of Pfizer vaccine 03/23: Pain started in chest and leg 03/24: Pain forced her to go to ER where it was discovered that she had a clot in her lungs and leg. Now on Xarelto
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1243337-1	1st occurrence: After 1st vaccination on March 6th, 2021- intense pain in left leg, passed out, went to urgent care 2nd occurrence: After 2nd dose on March 27th, 2021 - on the 28th flu like symptoms- on the 29th intense leg pain, passed out, transported to hospital by ambulance, doppler revealed blood clot, blood work normal, placed on Eliquis I have since been to the hematologist and had a work up... blood tests are normal
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1243829-1	On 2/18/21, 10 days after receiving the vaccine, I donated blood as usual. About 30 minutes later I was light headed, had abdominal cramping pain, bowel emptying and possibly fainted. I was transported by EMTs to the ER and was later diagnosed with inflamitory bowel by a CT scan. I took antibiotics and was feeling fine. My gastroenterologist performed a colonoscopy on 3/24/21 and the findings were that I was healed and tissue samples were normal. 3/25/21 I had an annual physical and was told I was in good health. Later that afternoon I began feeling light headed upon standing. In the night I called my health insurance hotline because my blood oxygen meter was beeping continuously. The Dr. on call told me to go by ambulance to the ER. When sitting or lying down, I felt fine and all the tests at the ER for my heart and my vital signs were good. A few hours later, they did a hest Xray and then a CT scan of my lungs which showed a saddle embulism. I had also thought I had bruised my knee if and when I fainted on 2/18 and 1o hours later a blood lot was found by ultrasound of my left knee. I was given blood thinners in the ER and then in ICU where I spent the next night. I am now taking Eleris. I was also given pantoprozole for GERD coughing the next day. I continue to have periods of abdominal ramping/pain, light headedness in waves and pressure in my head. I have disontinued the stomach medication with some feeling better with the lightheadedness.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1243860-1	Changes in menstrual cycles including heavy bleeding, large blood clots, longer cycles, and spotting between periods. This has continued for four months since receiving vaccines (I?ve completed the series now).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1244088-1	Blood clots and related severe pain
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1244382-1	Blood clot causing heart attack
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1244800-1	Superficial blood clot in right leg two large hot swollen spots

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1245200-1	I've have an IUD for 3 years and I irregularly very lightly spot for a day or two every few months. I was vaccinated Monday afternoon and by Tuesday night I started bleeding. To put it into perspective I generally use 1 panty liner a day of the usually 2 maybe 2 days I lightly spot. But I have been using 6 or so panty liners a day since then and it's Thursday night. So this is highly irregular for my regularity. I aldo expelled blood clots which hasn't happened since I had the IUD placed. I thought it should be reported.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1245238-1	The back of my left leg hurting all of a sudden on 04/15/2021. The pain on the leg is getting worse each day. Went to the ER on 04/20/2021. Per the doctor no swelling and no redness. I said something is really wrong please check. They did blood work and everything came back good. They did a sonogram and that's where they saw a Blood Clot. Was given Eliquis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1245306-1	I got a blood clot, venous, in my upper right leg. Started on or about March 3rd, 2021
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1246807-1	Blood clot in left leg and phonomia
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1247916-1	Pulmonary embolism, and Left Atrial thrombus Unclear which covid vaccine he got
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1248248-1	One week after injection woke up with a blood clot in left calf. No history of clots. No injury or muscle strain, etc. Just happened out of no where.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1248373-1	hurting warm leg, blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1249768-1	I passed a number of large blood clots (the size of a walnut) from my body on the two days after I received the shot. The day that I got the shot coincided with the day that I got my period, so I was already menstruating when the clots passed from my body. The clots were much larger than normal, and much more frequent. I passed almost a half dozen large clots. By the third day, my menstruation flow started to subside (the first two days tend to be heavier days for me). I did not pass any large clots on the third day going forward.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1250586-1	Severe pain in left leg, swelling, hot Sever pain in upper body-left side only--from jaw, to shoulder, to arm to side, front and back, all the way to the hip CT scan at hospital--pulmonary embolisms in both lungs, blood clot diagnosed in left leg at vascular surgeon's office before going to the ER Now on XARELTO for life.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1250960-1	In mid March, I was very physically active and had no physical limitations. I had no underlying conditions except for a slightly elevated homocysteine level diagnosed a year prior. I have never smoked in my life or has any prior lung issues. After the two doses I developed difficulty in breathing, shortness of breath in any minor activity, which began about two weeks after the second dose, and pains in my right side.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251077-1	Severe menstrual bleeding with clotting causing severe anemia. Hemoglobin level 9. Received first dose of Iron transfusion through IV yesterday on 4/23/2012
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251492-1	I delivered one baby in 1999 and one in 2002. Two miscarriages in between. I woke up 4/6/21 with a strange pain in my left leg. It felt circulatory, not muscular or skeletal. Swelling behind knee. I went to ER for ultrasound and they found a blood clot. Cat scan revealed pulmonary embolism. The nurses at the hospital confirmed that this was happening frequently. They said it wasn?t rare. My PA at the clinic told me it was 1 in 5000 like me. Why doesn?t the hospital report? I?m on bed rest for 10 days, compression stockings and Eliquis to thin my blood. I have halted all HRT, but have been told by my dr that it?s clearly not related since i e been on them for over 6 years and never had a blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251504-1	I have an IUD (I've had it for 3 years - no trouble) for controlling my fibroids and have not had a period or any bleeding for 3 years. 3 days after my first dose of the Pfizer vaccine, I experienced very heavy bleeding which was a bright red color. The next day, (4 days after my first dose), I experienced very strong (the strongest I've ever felt) cramping and a very large piece of uterine lining expelled from my body. Again, very bright red color. I just received my 2nd dose on 4/22/21 and am waiting to see if I have bleeding and/or cramping again.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251629-1	On March 2nd woke up with right calf pain. Thought I had a charley horse in the middle of the night and thought this was the soarness you get afterward. Because I had been on Crestor for approximately 5 weeks I thought that the crestor may be causing muscle pain. After 2 days of calf pain which was subsiding by then I visited Dr. who ordered a right ultrasound and I had 2 blood clots in the right leg (one in calf and one behind the knee). Due to the fact that I did not meet any standards for getting blood clots I was put on blood thinners (Eliquis) and told to see a hematologist.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251715-1	Blood clot in the brain, stroke, and seizure on 27th of March. Heparin drip, Keppra as a seizure medication, blood thinner medication. No movement right arm and hand, speech difficulties, impairment of concentration.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251780-1	4/16/21 Second Dose received. 4/17 and 4/18 noticed light swelling in ankles. No other effects. 4/19 - Emailed my Cardiologist about swollen ankles. Didn?t think about the vaccine as a potential cause. He scheduled an Echocardiogram for May. 4/20 to 4/22 - Edema is ankles and lower legs with pain on bottom left. Tried heat and elevation with no relief. 4/23 - Called Advice Nurse with recommendation to go to ER. ER determined a blood clot in left leg at the left posterior tibial vein measured at 6.3cm.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1251846-1	Blood clot in left lower extremity, requiring TpA
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1254584-1	3 days after the first dose I developed a blood clot in my right calf, never had one before, no trauma, i am very active; This is a spontaneous report from a contactable consumer (patient). A 59-year-old male patient received first received first dose of bnt162b2 (Pfizer, Formulation: Solution for injection, Lot Number: EP 7534b) via an unspecified route of administration, administered in left arm on 13Mar2021 11:00 at a single dose for covid-19 immunization. Medical history and known allergies were none. Concomitant medication included rivaroxaban (XARELTO). The patient developed a blood clot in his right calf 3 days after the first dose, never had one before, no trauma, he was very active on 16Mar2021 07:00. No covid vaccination was taken prior. No covid tested post vaccination. Blood thinner for the clot was given as the treatment. The outcome of the event was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1254628-1	Blood and clots came out of nose and mouth, first vaccine.; Blood and clots came out of nose and mouth, first vaccine.; Blood and clots came out of nose and mouth, first vaccine.; Face felt tight, first vaccine.; Mouth and nose dry, first vaccine.; Mouth and nose dry, first vaccine.; Sore arm, first vaccine.; This is a spontaneous report received from a contactable consumer(patient). A 73-year-old female patient received first dose of BNT162b2 (Pfizer-BioNTech COVID-19 Vaccine), at the age of 73-years-old, via an unspecified route of administration, administered in Arm Left (in the back of the arm) on 20Jan2021 (maybe at 11:10 am) (Lot Number: EL3246; Expiration Date: Apr2021) as single dose for covid-19 immunisation. Medical history included ongoing Postural vertigo maybe 10 years ago for a day. No family medical history relevant to AE. The patient's concomitant medications were not reported. Patient got the Shingles vaccine years ago and the next day, she had a sore arm and tingles, up and down her arm. She didn't feel food for 7 days and felt achy. It was the first Shingles shot. No prior Vaccinations (within 4 weeks). The patient experienced blood and clots came out of nose and mouth on 21Jan2021 (the same time that she woke up) with outcome of recovered on 21Jan2021. She blew her nose and leaned over and noticed a drop of blood on the rug and grabbed a Kleenex. She blew her nose, easy, and all of a sudden blood and clots came out of her nose and mouth. It lasted for about 8 minutes. Sore arm on 20Jan2021 (started in the evening) with outcome of recovered on 21Jan2021. Face felt tight on 21Jan2021 (around 8:30 or 9:00 am) with outcome of recovered on 21Jan2021. Mouth and nose dry on 21Jan2021 (in the morning) with outcome of recovered on 21Jan2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1254765-1	blood clot in her right ankle; varicose vein in her right ankle and it started to hurt; varicose vein in her right ankle and it started to hurt; ankle on her right side started swelling; This is a spontaneous report from a contactable consumer (patient). A 66-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on arm left on 26Feb2021 14:00 (Batch/Lot Number: EN6198) as SINGLE DOSE for covid-19 immunisation. The patient medical history was not reported. The patient's concomitant medications were not reported. The patient previously took the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EM9809 and expiry date: 30Jun2021) on left arm on 05Feb2021 for covid-19 immunisation. The patient experienced blood clot in her right ankle, varicose vein in her right ankle and it started to hurt, ankle on her right side started swelling, all on an unknown date. All events caused patient physician office visit. Treatment received for all events. It is reported that she is calling about the Covid 19 vaccine. She wanted to report that she developed a blood clot in her right ankle. She went to the doctor yesterday on 06Apr2021. She would like to add she has no prior history of any type of blood issues. The doctor she saw thought that the blood clot and the Covid 19 vaccine could be related but would not commit to saying so directly. She started having issues about 6 days after the second Covid 19 vaccine. She doesn't exactly remember when, maybe the 04Mar2021 or 05Mar2021. She ignored the issues attributing them to over exercising. She reached a point where she couldn't ignore the issues anymore and went to the doctor. Caller clarifies she has a varicose vein in her right ankle and it started to hurt then her ankle on her right side started swelling. She only payed attention to her ankle when it got worse. The last 3 weeks have been awful. She would like it known that she does not lead a sedimentary lifestyle. She rides her bike 60 miles a week and walks 10 miles a week. She's always moving. This is unusual for her. The doctor she saw diagnosed her with a blood clot and he did a sonogram in his office. She has been prescribed Xarelto 15mg tablets. Take 2 tablets twice daily for the first 21 days then on the 22nd day start taking one 20mg tablet once daily. The patient underwent lab tests and procedures which included sonogram: unknown result on 06Apr2021. The outcome of the all events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Blood clots in right leg below knee; he passed out from pain; pain in his right knee; he couldn't bend his knee at all; knee was swollen; couldn't walk/limping around; his knee more stiff; This is a spontaneous report from a contactable consumer (patient). A 53-year-old male patient received the first dose of BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, lot number: EN6206), via an unspecified route of administration, administered in the right arm on 21Mar2021 at 10:00 as a single dose for COVID-19 immunization. Medical history included ongoing Factor V Leiden heterozygous, allergies to mushrooms from an unknown date and unknown if ongoing and a torn meniscus in his right knee and he had surgery 3 to 4 years ago. The patient's concomitant medications were not reported. The patient previously took Cephalexin and Percocet and experienced allergies. The patient did not have COVID prior to vaccination and has not been tested post vaccination. On 22Mar2021 at night around 2:30AM, the patient woke up and started walking to the bathroom and he passed out from pain. He had pain in his right knee. He thought he had torn his knee again. He had surgery 3 or 4 years ago for a torn meniscus in his right knee and he thought he had torn his meniscus again. He passed out, his wife found him and he went back to bed. In the morning, his knee was still very very sore, he couldn't bend his knee at all. He took an Advil between 6:30AM and 8:00AM. He hadn't taken any medications prior to the COVID-19 vaccine. His knee was swollen but the pain was manageable. He couldn't walk, he was on crutches for about 24 hours then the swelling started coming down and he put a brace on his knee. He could then walk without the crutches but it was still painful. The next day he had pain in the back of his knee. He thought the pain was caused because he had put the brace on and had been using the crutches. Like he had made his knee more stiff from not using it. He was limping around and the pain was bearable. At that point, the pain was maybe a 4 on a scale of 1-10 where his pain had been a 9 on a scale of 1-10. He took another Advil and he realized the pain in the back of his knee wasn't going away. He went to see his family doctor and his doctor was able to get him in that same day for an ultrasound. The ultrasound showed 2 blood clots behind his right knee. The patient had blood clots in right leg below knee. The week of the report, he got an appointment with a hematologist. The hematologist reviewed his case and the doctor isn't 100% sure the blood clots were from the Pfizer COVID-19 vaccine but the doctor isn't ruling it out either because it happened within 24 hours of receiving his first COVID-19 dose. He hadn't done anything strenuous before getting the COVID-19 vaccine, nothing that he thinks would have caused the blood clots. He would like to add, what he thinks is relevant, is that he is positive for Factor V. He reports he has one copy and Factor V only impacts about 3% of the population. The hematologist thought that he might be prone to developing blood clots. The hematologist reports that he has had patients who have had the COVID-19 virus who have developed blood clots. Maybe it's possible that the COVID-19 antibodies were trying to build up his immune system to COVID-19 and it created something similar to having the COVID-19 virus. If he is predisposed to blood clots then the COVID-19 vaccine may have precipitated the blood clots. His second COVID-19 vaccine is scheduled for Sunday, 11Apr2021. He had talked to both his family doctor and his hematologist and they both thought he should be ok in getting the second COVID-19 vaccine. He was prescribed blood thinners. If the second COVID-19 vaccine caused more blood clots the blood thinners would help him. He was asking if there is any additional information or guidance on receiving the second COVID-19 vaccine after his reaction. The pharmacist told him that the COVID-19 vaccine trials were all done in the left arm. He received his vaccine in his right arm and his right knee was affected. If he gets the second COVID-19 vaccine he's getting it in his left arm. He was prescribed Eliquis 5mg take one tablet twice daily. The outcome of the event blood clots in right leg below knee was recovering while the outcome of the other events was unknown.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1254796-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255233-1	infection; Abnormal nose bleed with blood cloths the first day; Abnormal nose bleed with blood cloths the first day/6 more nose bleed that it was taking me to control up to 30 min; This is a spontaneous report from a contactable other HCP (patient). A 75-year-old female patient (not pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EN6207), via an unspecified route of administration on 15Mar2021 10:45 (at 75 years old, not pregnant), single dose for covid-19 immunisation. Medical history included Type II diabetes, Hypertension, GERD and hypercholesterilymia, sulfa allergy, all from an unknown date. Concomitant medications included verapamil; rosuvastatin calcium (CRESTOR); metformin; omeprazole, unspecified blood thinners. The patient previously took tetracycline for known allergies: Tethracycline. Patient experienced abnormal nose bleed with blood cloths the first day (reported as 30Mar2021 0130). Five days later in two days had 6 more nose bleed that it was taking me to control up to 30 min. She went to ER in the morning and had a CBC having results normal with normal levels of platelets. She was referred to PCP. She saw a NP next day and gave me referral to ENT specialist. Using an scope could not find anything and ordered to have CT Scan. Results show negative for polips, tumors and show only a shadow by eye socket which MD believe it was an infection that was leading to the bleed. Prescribe a very small amount of Amoxicilline 125 mg, twice a day for 7 days plus a saline gel. Patient had the last nose bleed at midnight. For 3 days did not have any bleeding until today, at 10:15 am, with heavy bleeding which took me 15 min. to control. She has no history of nose bleed, taking blood thinners, no head traumas either. No other vaccine in four weeks. No Covid prior vaccination. No Covid tested post vaccination. The outcome of the event infection was unknown, other events was not recovered.; Sender's Comments: Information provided was so limited to prevent a meaningful and definite medical assessment for the events. A causal relationship cannot be completely excluded for BNT162B2 injection and development of Thrombosis and Epistaxis, only based on a plausible chronological sequence. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255241-1	blood clot in lessor saphenous vein left leg; This is a spontaneous report from a contactable consumer (patient). A 68-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 17Mar2021 at 15:15 (lot number EN6207) as single dose for covid-19 immunization, administered at the hospital. The patient received the first dose on 24Feb2021 at 15:15 in the right arm (lot number EN6198) for covid-19 immunisation. The patient did not receive other vaccines in four weeks. Medical history included type 2 diabetes, deep vein thrombosis (DVT), and pulmonary embolism (PE). The patient has not had COVID prior to vaccination and not tested for COVID post vaccination. The patient's concomitant medications were not reported. The patient experienced blood clot in lessor saphenous vein left leg on 27Mar2021 at 18:00 with outcome of recovering. The patient's hematologist put the patient on Eliquis 5mg 2 BID and at the time of report, this medicine has started to dissolve the clot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255308-1	she thought she had a blood clot; maybe it is MS (multiple sclerosis); possible septal infarct; weakness in her left arm; losing her balance; vaccine was given up high on her arm, not 2 fingers down. She thought they had hit bone; less range of movement in her left foot; trouble walking; like someone who had a stroke; numbness; legs felt heavy climbing stairs; immediate pain when needle went all the way in and worse when vaccine went in; pain in shoulder joint; This is a spontaneous report from a contactable consumer (patient). A 49-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: Ep6955), dose 1 via an unspecified route of administration, administered in Arm Left on 25Mar2021 18:00 (at the age of 49years) as single dose for Covid-19 immunization. The patient is not pregnant at the time of vaccination. Medical history included anemia that is treated, living kidney donor, gastric bypass in 1999, gallbladder removal, hysterectomy, intussusception x2, migraines, high heart rate, and smoking. Concomitant medications included omeprazole (OMEPRAZOLE), amfetamine aspartate, amfetamine sulfate, dexamfetamine saccharate, dexamfetamine sulfate (ADDERALL), and albuterol [salbutamol] (ALBUTEROL [SALBUTAMOL]); all taken for an unspecified indication, start and stop date were not reported. The patient previously received Flu vaccine and Tetanus vaccine. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient has not been tested for COVID-19 since the vaccination. She got the shot on 25Mar2021 on Thursday evening at 1800. She is 49 years old and has had all the flu shots and tetanus shots. She has never had one hurt so bad. She stated that the vaccine was given up high on her arm, not 2 fingers down. She thought they had hit bone. It was reported that the patient experienced immediate pain when needle went all the way in and worse when vaccine went in. The whole time it hurt really bad and her arm hurt severely for 2 days from 25Mar2021. She still has pain in shoulder joint. That aside wasn't a big deal, it went away even though the joint is sore. The following Monday 29Mar2021 in the evening, her legs felt heavy and she had trouble going up the stairs. Then she had complete numbness of her left leg, from her hip or butt crack to the tip of her toes. It was on the same side she had the shot. It was further reported that the patient's legs felt heavy climbing stairs then she woke Thursday morning with her left leg numb. She couldn't feel herself starting to have a bowel movement because the numbness went all the way up to that area. She went to the ER (emergency room) on Saturday night (03Apr2021)

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				area. She went to the ER (emergency room) on Saturday night (03Apr2021) because she was afraid of a blood clot. An ultrasound was done on her left leg, CT of head and neck, and blood work with nothing found. She has numbness and less range of movement in her left foot with trouble walking and have had no improvement with calf being numb. It was further clarified that when she was Easter shopping, there was numbness in her foot was causing her to drag her foot around the store. It was like someone who had a stroke, with slight impairment on one side. Her foot slaps the floor. She doesn't have control of it, she can't flex her foot up and down at all and she has little movement in her toes, even now. When she went to the emergency room, she thought she had a blood clot. She is a kidney donor and is healthy over all except for anemia that gives her tingly toes since she was 30, off and on. She never had a body part felt so numb, other than when she sits on her leg wrong. They did a bunch of tests there. They did a doppler, CT scan, and an EKG (unspecified date) which was abnormal. Her EKG has never been abnormal before. It said possible septal infarct undetermined time. The doppler showed no deep or superficial blockages noted from her groin down to her toes/ankle. The doppler was to check for blood clots. They referred her to her primary care provider who said that she will do a neurological exam. She has weakness in her left arm as well. The weakness in her left arm was only told by the neuro touch test. The patient further clarified that she doesn't have less range of motion, it is really just pain (arm). She can still move her arm all the way. Her left side is weaker or less sensitive as far as the neuro test. She wasn't aware she had less feeling on her arm until the nurse said she did. Her arm pain is nothing significant, it can hurt when moving her arm. She had no leg pain or anything like that until later in the week. She still had symptoms of losing her balance. Getting in and out of the chair she almost fell down. She didn't realize the numbness. Monday, she felt heaviness going up the stairs. For 2 days she had the worst pain ever in her arm. She had the tetanus shot and didn't have anything like that. It went away, when she raises her arm off from the side, she still has pain in the joint. She has migraines, it is nothing that is causing difficulty. She is more concerned something else happened. Patient thought maybe it is MS (multiple sclerosis) and asked if it is peripheral artery disease. She did smoke half a pack every other day. She is 49 but didn't start smoking till she was 36. It is a terrible habit and her doctor is aware. She also had her good and bad cholesterol which was just over 100. She is just reporting it because it happened on only the left side and she is a healthy person. She never had high blood pressure and never been treated for it, but she has had high heart rate before. It is not something that has ever caused any issue. She doesn't have something like peripheral artery disease, when she (software company name withheld) that is said plaque in the arteries. The doppler showed no signs of blockage superficial or deep on that side. She doesn't have an appointment until 29Apr2021. She doesn't know what to do to get more urgent care. The pain in her arm was better after the first 2 days. It is like a tooth ache, a dull ache. It is not all the time, just when moving her arm. Her legs were worse to where Thursday, she woke, and she was numb from the top of her leg to the tip of her toe. She can still lift her leg, but when she walks the tip of her second toe which is longer will drag on the ground. When she walks it is hard to hold her shoe on the left foot or put her shoe on. She can't maneuver her foot. When her foot hits the ground, the ball area of her foot smacks the floor because she doesn't have control over it. It is laboring to walk and is hard. She was kind of limping and pulling that leg on the left side. Her laboratory tests were BMP, CBC auto diff, CBC with diff, CT, angiogram of the head and neck, EKG 12 lead, and they did a venous doppler of the left lower extremity. It all came back good except the EKG which says septal infarct undetermined date, which has never come up on any tests before. The events resulted in Emergency room/department or urgent care, Disability or permanent damage. There was no treatment received for the adverse event. The patient has not been tested for COVID-19 since the vaccination. The outcome of events was not recovered.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255342-1	"blood clot; Right leg swelled and achy; Right leg swelled and achy; This is a spontaneous report from a contactable nurse (patient). A 67-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: EP7534),dose 1 via an unspecified route of administration, administered in Arm Left on 16Mar2021 (at the age of 67years) as single dose for Covid-19 immunization. Medical history included ongoing Beta thalassemias. The patient's concomitant medications were not reported. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient stated she has never had a problem with vaccinations, and she gets the flu vaccine every year. The patient reported that after the first injection coincidentally she did develop a blood clot in her lower leg. She thinks that it was due to a plane ride, but she has never had a blood clot before the injection. She added that she had the first dose on 16Mar2021 and developed a clot in a week. She further clarified that she got her first injection on 16Mar2021 and realized when she went on a flight, which is not that far she developed that clot and was having problems shortly after flight home on 24Mar2021 where her leg swelled up and she is on blood thinners since she went to the emergency room. She added that she developed the clot sometime between that flight and 02Apr2021. She had a doppler study on an unspecified date and it showed a blood clot. She stated that the outcome was she does not have the blood clot (as reported) but she will be on blood thinners and she needs to follow up on her scan to see if the clot issue is resolved but she will not have a doppler scan for another month or two and the blood thinner she is taking is Xarelto. She also stated that her right leg swelled up and it felt like a log. She reported it felt achy and swollen quite a bit from knee down and her leg was achy. It started between 24Mar2021 and by time she went to the emergency room she was not getting better. She had her symptoms for a week and toward the end of the week she thought she had a blood clot since it was not getting better it just got worse. She added that right now it is not swollen but she has been supine, and she does believe that it is better. The outcome of events was recovering. (As reported ""outcome is unknown but pretty sure it has improved because of the blood thinners""./She stated she does not think it has resolved because it takes time for clots to resolve./She added that right now it is not swollen but she has been supine, and she does believe that it is better.) The reporter stated for the seriousness criteria for event blood clot: she believes as a nurse that any blood clot is extremely dangerous, and she was not hospitalized but obviously a blood clot is serious, and she could have ended up with pulmonary embolism. The seriousness for events right leg swelled and achy was reported as medically disabling. Causality assessment: Patient stated for the blood clot she has doubts if it was related to the vaccine but is it something that should be known about; for the leg swelling and achy she has doubts but it could have been the plane ride.; Sender's Comments: Based on the current available information, the reported events are most likely related to an intercurrent or underlying condition which is unlikely related to the suspected drug. The plane ride may provide an explanation for the events. The case will be reassessed if additional information becomes available."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255460-1	blood clot on the left side of neck; This is a spontaneous report from a contactable HCP (patient). The patient was a Respiratory Therapist. A 60-year-old male patient received his secondo dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot Number: EN6206), via an unspecified route of administration on 03Apr2021 (at the age of 60-years-old) as single dose for COVID-19 immunisation. The patient took the first dose of BNT162b2 (lot number EN6199) on an unspecified date for COVID-19 immunisation. Medical history included high cholesterol and hypertension from an unknown date and unknown if ongoing. Concomitant medications included nebivolol hydrochloride (BYSTOLIC) taken for hypertension, start and stop date were not reported; rosuvastatin taken for blood cholesterol increased, start and stop date were not reported. The patient experienced blood clot on the left side of on an unspecified date in Apr2021. The event was treated with blood thinner. Patient thought it was called Xarelto because they called him on 07Apr2021. He was not sure how to spell that. Treatment was started on 07Apr2021. Consumer stated he first started having symptoms on the night of the injection. On an unspecified date in Apr2021 the patient did the ultrasound and that's when they found blood clot and he just got the results back. Recently, may be 2 weeks before reporting, patient had routine blood work and It was normal. It was nothing abnormal. The outcome of event was unknown.; Sender's Comments: Based on the current available information and the plausible drug-event temporal association, a possible contributory role of the suspect product BNT162B2 to the development of event Clot blood cannot be excluded. The case will be reassessed if additional information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255462-1	Her husband got a heart attack a week after the first dose; there are blood clots on his legs; This is a spontaneous report from a Pfizer-sponsored program. A male patient of an unspecified age received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 24Mar2021 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The reporter stated that her husband got a heart attack a week after the first dose and there are blood clots on his legs on an unspecified date. Outcome of the events was unknown. Information on the Lot/Batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255463-1	Developed like a blood clot, black and blue in the arm; Developed like a blood clot, black and blue in the arm; This is a spontaneous report from a non-contactable consumer (patient). A patient of unspecified age and gender received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot Number and Expiration Date were not reported), via an unspecified route of administration, on an unspecified date, at a single dose, for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient previously took the first dose of BNT162B2 for COVID-19 immunization. The patient was calling about the side effects of Pfizer COVID-19 vaccine and was wondering. The patient got the first and second vaccine of Pfizer for COVID-19. After the second dose of vaccine on an unspecified date, the patient developed like a blood clot, black and blue in the arm. The patient wanted to ask if this is a rare side effects or common. The outcome of the events was unknown. No follow-up attempts are possible; information about batch number cannot be obtained. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255545-1	"I am bleeding heavily because I am getting blood clots and stuff like that; I am bleeding heavily because I am getting blood clots and stuff like that; Right side of face, my tongue inside is very swollen and irritated; Lot of pain in my joints and in my bones; Right side of face, my tongue inside is very swollen and irritated; Right side of face, my tongue inside is very swollen and irritated; Lot of pain in my joints and in my bones; This is a spontaneous report from a contactable consumer (patient) A patient of unspecified age and gender received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Lot number FN6200) as SINGLE DOSE for covid-19 immunisation; axitinib (INLYTA), via an unspecified route of administration from an unspecified date (Batch/Lot number was not reported) to an unspecified date, at 2 DF, 1x/day (two pills a day) for renal cancer; pembrolizumab (KEYTRUDA), route of administration, start and stop date, batch/lot number and dose were not reported for renal cancer; zoledronic acid (ZOMETA), route of administration, start and stop date, batch/lot number and dose were not reported for bone disorder . The patient received the first dose of BNT162B2 on unknown date (Lot number CL9265). The patient medical history was not reported. The patient's concomitant medications were not reported. The patient stated ""I got a clear cell renal cancer. And I am taking a combination Inlyta okay which is your drug and Keytruda and I also have also taken Zometa that is for the bones and I have had two Pfizer Vaccines (Unspecified Vaccines) for it.I am getting terrible side effects from either the Inlyta or the Keytruda. I am bleeding very heavily and the right side of face, my tongue inside is very swollen and irritated. I am getting a lot of pain in my joints and in my bones. I am bleeding heavily because I am getting blood clots and stuff like that. I just wonder if there is anything that I have been taking Inlyta two pills a day and I have taken probably 80 pills from the start. So, if there is anything to mitigate what's going on. The action taken with axitinib , and pzoledronic acid embolizumab was unknown."".The outcome of the event was unknown"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255628-1	died yesterday due to blood clots; This is a spontaneous report from a contactable consumer. A 55-year-old female patient (mother) received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 28Mar2021 (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and the patient's concomitant medications were not reported. The patient experienced blood clots and died due to the event. It was not reported if an autopsy was performed. Information on the lot/batch number has been requested. ; Reported Cause(s) of Death: died yesterday due to blood clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255683-1	painful little clot in her front leg; pain/ache; shortness of breath like wheezing; shortness of breath like wheezing; This is a spontaneous report from a contactable consumer via a Pfizer sponsored program. A female patient (consumer) of an unspecified age received first dose of BNT162B2 (Pfizer COVID-19 Vaccine, Formulation: Solution for injection, Lot number and expiration dates were not reported), via an unspecified route of administration on 26Mar2021, at single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient reported that she had the first dose 26Mar2021 and scheduled for the second dose on 21Apr2021, which was more than 21 days. Caller read the news about the adverse events with (Other company vaccine). They experienced pain/ache and shortness of breath like wheezing on an unspecified date in 2021. They also had a painful little clot in her front leg on an unspecified date in 2021. They wants to know if she can take the second dose. The outcome of the events was unknown. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255692-1	Blood Clot; This is a spontaneous report from a contactable nurse. A 60-year-old female patient (no pregnant) received her second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number: EN6208), intramuscular at the age 60-year-old at arm right on 11Mar2021 at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient previously received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EN6200), intramuscular at the 60-year-old at right arm on 19Feb2021 for COVID-19 immunisation and experienced pneumonia. There was no other vaccine in four weeks. Facility type vaccine was reported as public health clinic/administration facility. The patient experienced blood clot on Mar2021 after 2nd. The patient was hospitalized for event for 5 days. Adverse event resulted in doctor or other healthcare professional office/clinic visit, emergency room/department or urgent care, hospitalization, life threatening illness. The patient received the treatment for event. The outcome of event was unknown.; Sender's Comments: Based on information available, a possible contribution role of the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) to the onset of event clot blood cannot be completely excluded. The case will be reassessed should additional information become available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics committees and Investigators, as appropriate.,Linked Report(s) : US-PFIZER INC-2021422806 same patient/product, different dose and event
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255698-1	NSTEMI; Pulmonary Embolism; left upper arm thrombosis; This is a spontaneous report from a non-contactable other healthcare professional (patient). A 48-year-old male patient received the first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE), on 22Mar2021 (at the age of 48-years-old) at single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. No other vaccine received in four weeks. On 15Apr21, the patient presented to hospital with complaints of dyspnea, chest pain and arm swelling. The patient was found to have left upper arm thrombosis, pulmonary embolism, and NSTEMI. The adverse events started on 15Apr2021 and resulted in emergency room/department or urgent care visit, hospitalization on 15Apr2021 and life-threatening illness (immediate risk of death from the events). An unspecified treatment was received in response to the events. The outcome of the events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255700-1	Felt something in my leg like a cramp / did a scan of my leg and found the clot; dizzy; nearly passed out; This is a spontaneous report from a contactable consumer (patient). A 54-year-old male consumer received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 26Mar2021 at 16:30 at single dose in right arm for COVID-19 immunisation at the age of 54-year-old. Lot number was ER8732. Medical history included Human papilloma virus (HPV) positive tonsil cancer, ankylosing spondylitis, tinnitus, neuropathy, decreased memory from chemo and radiation. The patient did not have Covid prior to vaccination. Concomitant medications were unknown. The patient felt something in leg like a cramp for a week since 06Apr2021 before ER visit. On 06Apr2021, the patient got dizzy at home nearly passed out, and he went to ER. On 14Apr2021, Doctor did a scan of leg and found the clot. The patient was hospitalized due to the events and they were considered life-threatening. The patient was treated with heparin and ELIQUIS. On 14Apr2021, Rapid Covid test was negative. The patient was recovering from the events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255710-1	"DVT; multiple bilateral pulmonary embolisms/blood clots in both lungs after being sent to the ER; one blood clot still in my leg/ blood clots in both lungs; This is a spontaneous report from a contactable Other HCP. A 41-year-old female reported that she received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number BL9269), intramuscular into the left arm on 08Feb2021 at single dose for COVID-19 immunization. No COVID prior vaccination. Medical history included type 1 diabetes mellitus, Hashimoto's Disease. Concomitant medication(s) included metformin (METFORMIN) and levothyroxine (LEVOTHYROXINE) . The patient had received the first dose of BNT162B2 on 19Jan2021. About a week before the second dose, she self-diagnosed with a calf injury due to extreme pain that felt like a constant cramp that lasted until her second dose. After the second dose, she was hospitalized 5 days (16Feb2021-20Feb2021) with multiple bilateral pulmonary embolisms/blood clots in both lungs after being sent to the ER by a nurse practitioner, and told that her previously self diagnosed leg injury was really DVT. She actually left the hospital with one blood clot still in her leg.Hospital's Cardiac ICU by physician's transport because her case was too to be treated at (Withheld). The patient underwent lab tests and procedures which included antiphospholipid antibodies (unknown date): unknown results (won't be in until mid-Jun because it requires testing 12 weeks apart)-tested negative for genetic conditions that could have caused the clots , had a negative COVID nasal swab test on 16Feb2021.The events DVT and pulmonary embolism resulted in Emergency Room visti /Phisician office visit .The event ""DVT"" was treated with Heparin, still on blood thinner. The outcome of the event DVT was recovering.; Sender's Comments: Based on vaccine-event chronological association causality between reported events and BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) cannot be excluded. . The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.,Linked Report(s) : US-PFIZER INC-2021427569 same patient, and drug different dose"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255715-1	This is a spontaneous report from a contactable consumer. This 81-year-old female consumer (patient) reported that she received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number unknown) at single dose for COVID-19 immunisation on 13Jan2021. Relevant history included mild dementia, high cholesterol. Relevant concomitant drugs included memantine 10MG,donepezil 5MG,atorvastatin, cetirizine hydrochloride (ZYRTEC). The patient was not pregnant. No known allergies. On 30Jan2021, the patient fell and became unconscious after getting up from bed to use bathroom, was able to get up and go back to bed. Next day, 31Jan2021, she told her daughter at 2pm what had happened and daughter took her to urgent care. Urgent care was performed EKG and called ambulance as EKG suggested suspicious activity. She was coughing with blood. Cat scan of the lung revealed acute saddle pulmonary embolism with acute cor pulmonale and a large blood clot in the calf. On 02Feb201 then placed her on blood thinner and discharged her. As soon as arrived home, she began to suffer a massive gastro intestinal bleed and was rushed back to the hospital. On 04Feb2021, she began to have heart issues and suffered a mild heart attack and underwent a procedure to have a stent inserted. She developed the shingles the next day (05Feb2021). The events assessed as serious due to Hospitalization, Life threatening illness (immediate risk of death from the event), Disability or permanent damage. The patient visited Emergency room/department or urgent care. The outcome of events was resolved with sequel. Treatment therapy involved. The patient was not diagnosed with COVID prior vaccination. The patient had Nasal Swab/Fast Test/PCR (for COVID) in Apr2021 with negative result. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255731-1	Hot a fever then full blown covid; Hot a fever then full blown covid; blood clot; organ failure; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer reported that a male patient of an unspecified age received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. Previously on an unknown date, the patient received the first dose of BNT162B2 vaccine. On an unspecified date, the patient experienced hot a fever then full blown COVID, blood clot and organ failure leading to patient death on an unknown date. It was not reported if an autopsy was performed. No follow-up attempts are possible. Information about lot/batch number cannot be obtained. No further information is expected; Reported Cause(s) of Death: Drug ineffective; Covid-19; Blood clot; Organ failure
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255926-1	Blood clots, Hospitalization, Blood thinners
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1256737-1	Died blood clot in brain

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1256823-1	Started 10 minute after the shot I was nauseous with a headache watery eyes , nose I had an asthma attack, the nurse I seen took me to a curtain room and gave me 6 pumps of albuterol my bp was dropping at first and I was extremely weak and I took 2 more puffs and my BP went up now and the nurse wanted me to go to the ER and I got up and almost fell I was so dizzy I was given Benadryl at the hospital and received chest X-rays and was watched over and I was continuing to take Benadryl and albuterol around the clock, I was itchy all over and I?ve had to take my albuterol more often then usual now even though feeling better my heat rate is all over the place and and my period is extremely off as well clotting, tissue passing as the vaccine is now itchy again and swollen and hurting after 2 days as of your symptoms were gone
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1257053-1	Got first pfizer vaccine 4/7. Started to feel pain in right calf 4/9. Went to emergency room Hospital, 4/14 where after an ultrasound was diagnosed with blood clot in saphenous vein. Repeated ultrasound per primary care recommendation 4/21, no changes in size were noticed. Currently taking Advil.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1257058-1	April 13, 2021 at approximately 9am I started passing blood clots, April 16,2021 blood clots stopped and I have not had any more. It felt as though I was having a period , except the blood clots were coming from my rectom.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1257145-1	Fever. Chills, hallucinations, renal failure, ischemic colitis, gi bleed, blood clot,, dehydration
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1257393-1	5 days after I had the second dose of the Pfizer Covid vaccine, I had blood clots that caused several strokes. I have normal cholesterol, normal blood pressure, normal A1C and other diabetes measures, and my blood was tested for multiple things to identify what caused the clotting. Every single test came back within normal limits. The ONLY thing that changed was that vaccine. And it was 5 days after the second dose. The doctors cannot figure out why I had a stroke.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1258788-1	Blood clots leading to TIAS and strokes
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1258918-1	Client received vaccine Wednesday morning. Client started to feel weak the next afternoon. Got out of vehicle and almost fell due to leg weakness and had an abnormal gate. After 20 minutes had passed, symptoms got better. A couple hours later that day it happened again, but this time with arm weakness and hand fell at side. Client then went to ER. Had CT scan and MRI which showed client had a small clot causing a small stroke. Client had no risk factors for a stroke. Not a smoker, no high blood pressure, and good cholesterol. Doctors were unaware if the stroke was correlated with vaccine or not. Primary care physician is running labs to see if there is a clotting disorder.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1258979-1	On April 17th, approximately 2 weeks after receiving the second shot, I woke up to use the bathroom at which point I passed an extreme amount of blood from my rectum along with what appeared to be remnants of blood clots. I took myself to the hospital who confirmed that hemorrhaging and blood clots had formed behind the site of my old ostomy scar inside of my intestines. I had no other signs of ulcerative colitis when scoped and no active flare, only this clotting that arose specifically after this second vaccine dose.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1259025-1	1 week after injection I developed superficial blood clots in my Left Calf, that still have not resolved. Left arm and shoulder pain and headaches.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1260536-1	Heavier period than normal, bigger blood clots expelled

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261769-1	Approximately 2 weeks after the vaccine, inexperienced deep leg pain. 3 days later I went to the hospital and was diagnosed with a blood clot/DVT.; Approximately 2 weeks after the vaccine, inexperienced deep leg pain. 3 days later I went to the hospital and was diagnosed with a blood clot/DVT.; Approximately 2 weeks after the vaccine, inexperienced deep leg pain. 3 days later I went to the hospital and was diagnosed with a blood clot/DVT.; This is a spontaneous report from a contactable consumer. A 46-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 1 via an unspecified route of administration, administered in Arm Left on 25Mar2021 13:00 (Batch/Lot Number: ER2613) as SINGLE DOSE for covid-19 immunisation. Medical history included none. The patient's concomitant medications were not reported. Approximately 2 weeks after the vaccine, inexperienced deep leg pain. 3 days later, the patient went to the hospital and was diagnosed with a blood clot/DVT. AE resulted in: [Emergency room/department or urgent care]. The patient received treatment for the events: Vitals taken, blood work, prescribed blood thinner. The patient underwent lab tests and procedures which included blood test: unknown result on an unspecified date ae treatment= Vitals taken, blood work , investigation: blood clot/dvt on 08Apr2021 3 days later I went to the hospital and was diagnosed with a blood clot/DVT, investigation: unknown result on an unspecified date ae treatment= Vitals taken, blood work. The outcome of the event was recovering. The patient does not have COVID prior vaccination, and was not COVID tested post vaccination. The patient has no known allergies.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261778-1	Swelling underneath my left arm pit which may be a possible blood clot; Swelling underneath my left arm pit which may be a possible blood clot; This is a spontaneous report from a contactable consumer (patient). A 26-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number and expiration date were not reported) on the left arm on 08Apr2021 (16:15) as a single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. On 09Apr2021 (05:45), the patient had swelling underneath my left armpit which may be a possible blood clot. The patient did not receive any treatment for the reported events. The outcome of the events was not recovered. The patient did not have COVID-19 prior to vaccination, and had not been tested post-vaccination. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261799-1	I was diagnosed with a blood clot in my right leg; This is a spontaneous report from a contactable consumer (patient). A 49-year-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in right arm on 03Mar2021 15:45 (Lot Number: EN6202), at the age of 49-years at vaccination, as SINGLE DOSE for covid-19 immunisation. Medical history included none. The patient is not pregnant at the time of vaccination. The patient received dose 1 of the vaccine on 10Feb2021, lot number: EM9810, on 10Feb2021 03:45 PM at right arm. The patient did not have COVID-19 prior vaccination. The patient's concomitant medications were not reported. On 04Mar2021 12:00 AM, patient was diagnosed with a blood clot in her right leg. The event required emergency room visit. The patient was put on Xarelto, a blood thinner for clots. The patient has not been tested for COVID-19 post vaccination. The outcome of the event was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261801-1	Heavy menstrual bleeding with clots; Heavy menstrual bleeding with clots; This is a spontaneous report from a contactable nurse (patient). A 33-year-old female patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; lot number and expiration date unspecified), via an unspecified route of administration, administered in left arm on 08Apr2021 13:00 as single dose for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient was not pregnant and had no other vaccine in 4 weeks. On 08Apr2021 13:30, the patient experienced heavy menstrual bleeding with clots. There was no therapy for the events. The patient had no COVID prior vaccination and was not COVID tested post vaccination. Outcome of events was recovered on an unspecified date. Information about Lot/Batch number is requested.; Sender's Comments: Based on the available information the reported events were attributed to an underlying or an intercurrent medical condition and it is assessed as unrelated to the suspect drug bnt162b2. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"blood clot in my lungs/Multiple blood clots; Pains; This is a spontaneous report from a contactable consumer. This 69-year-old female consumer reported for herself that: Patient characteristics: Weight (kg): 102.97 Height (cm): 165 Sex: Female Relevant medical history and concurrent conditions: Structured information (Patient episode name): Hypertension Patient Medical comments: Verbatim: Hypertension Reaction(s)/Event(s): Reaction/event as reported by primary source: Blood clot in my lungs within the first 2 weeks after the second shot Reaction(s)/Event(s): Reaction/event as reported by primary source: Hospitalization Reaction/event in MedDRA terminology (LLT): Hospitalization Reaction/event MedDRA term (PT): Hospitalisation Reaction(s)/Event(s): Reaction/event as reported by primary source: Pains Results of tests and procedures for investigation of the patient: Test: CTA scan More information available (Y/N): No Drug(s) Information: Characterization of drug role: Suspect Proprietary medicinal product name: Covid-19 Vaccine Batch/lot number: ER8732 Date of start of drug: 30Mar2021 Action(s) taken with drug: Unknown Drug(s) Information: Characterization of drug role: Concomitant Proprietary medicinal product name: Lisinopril HCTZ Dosage text: 20-25 mg Tablet Indication for use in the case: Hypertension Narrative case summary and further information: Case narrative: Selected Report Type: Initial Patient Ethnicity: (Ethnicity withheld) Is the patient also the reporter? Yes Reporter type: Consumer or other non-health professional Reporter telephone: (Phone no withheld) Primary / Prescribing Healthcare Professional Info Dates for Blood clot in my lungs within the first 2 weeks after the second shot.: (From: Unspecified To: Unspecified) Dates for Hospitalization: (From: Unspecified To: Unspecified) Dates for Pains: (From: Unspecified To: Unspecified) Is Covid-19 Vaccine a Pfizer product? Yes Covid-19 Vaccine manufacturer: Unspecified Dates for Covid-19 Vaccine: (Start: 30Mar2021 Stop: Unspecified) NDC number of Covid-19 Vaccine: Unknown UPC number of Covid-19 Vaccine: Unknown Expiry Date of Covid-19 Vaccine: 31Jul2021 Other Products: Yes Dates for Concomitant Products Lisinopril HCTZ: (Start: Unspecified Stop: Unspecified) Patient History: Yes Patient history: Hypertension (From: Unspecified To: Unspecified) Investigation Assessment: Yes Investigation: CTA scan (Date: Unspecified, Result / Units:) Additional Context: Consumer stated, ""I could not follow on the parts. So may be should have done certainly but I am not technically astute. So, I wanted to report that I just had the second Covid-19 test and as a experience in emergency room discovery of blood clot in my lungs within the first 2 weeks after that second shot. So, I wanted to report that. Is this the place I do that?"" When paraphrased, consumer stated, ""Multiple blood clots."" Start date of event (Multiple blood clots.): Consumer stated, ""10Apr2021. I should probably say that the pains were 09Apr. I went in the emergency room on the 10th. So, I am not sure which day."" Treatment :Consumer stated, ""Yes, I went To the emergency room and was kept in the hospital and then I saw my own Doctor just today who is going to help me try to determine what is going to happen next ? I was released by the hospital."" Details of hospitalization: Duration of hospitalization: 24 hrs. (overnight) Date of Admission:10Apr2021 Date of Discharge:11Apr2021 Lab work: Consumer stated, ""They did lab. work while in the emergency room and in the hospital. They did Chest CTA scan."" This is a spontaneous report from a contactable consumer. A 69-year-old female patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 30Mar2021 (Batch/Lot Number: ER8732; Expiration Date: 31Jul2021) as single dose for covid-19 immunisation at age of vaccination 69-year-old. Medical history included hypertension from an unknown date. Concomitant medication included hydrochlorothiazide/lisinopril (LISINOPRIL HCTZ) taken for hypertension, start and stop date were not reported. The patient experienced blood clot in my lungs/multiple blood clots (thrombosis) (hospitalization) on 10Apr2021 with outcome of unknown , pains (pain) (hospitalization) on 09Apr2021 with outcome of unknown. The patient was hospitalized for blood clot in my lungs/multiple blood clots (thrombosis) from 10Apr2021 to 11Apr2021. The patient was hospitalized for pains (pain) from 10Apr2021 to 11Apr2021. The patient underwent lab tests and procedures which included computerised tomogram: unknown results on , laboratory test: unknown results on . The action taken in response to the event(s) for bnt162b2 was not applicable. Therapeutic measures were taken as a result of blood clot in my lungs/multiple blood clots (thrombosis), pains (pain)."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1261806-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1261824-1	swelling in her right foot and right calf and pain/blood clot was diagnosed in her right calf under her knee; This is a spontaneous report from a contactable consumer (patient). A 52-year-old female patient received the second dose of bnt162b2, via an unspecified route of administration, administered in left arm on 31Mar2021 (Lot Number: ER8732) as single dose for covid-19 immunisation. There were no medical history and concomitant medications. The patient previously received the first dose of bnt162b2, at the age of 52-year-old, via an unspecified route of administration, administered in left arm on 10Mar2021 11:15(Lot Number: EN6206) as single dose for covid-19 immunization. Started experiencing swelling in her right foot and right calf and pain 3 days following 2nd injection (03Apr2021). Went to the doctor and blood clot was diagnosed in her right calf under my knee 9 days following 2nd injection (09Apr2021). Event resulted in Doctor or other healthcare professional office/clinic visit, Life threatening illness (immediate risk of death from the event). The treatment for event was 21 Days Xarelto BID then Pradaxa for 90 days. The patient was not pregnant. The outcome of the event was recovering.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261830-1	Blood clot in leg; This is a spontaneous report from a non-contactable consumer reported for himself. A 42-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via unspecified route of administration on 01Apr2021 02:00 PM at 42-year-old at single dose for COVID-19 immunisation. Patient had known allergies (unspecified) and other medical history (unspecified). There was no covid prior vaccination. There is no other vaccine in four weeks. Other medications in two weeks included antibiotics. Patient experienced adverse event: blood clot in leg on 03Apr2021 02:00 AM. The adverse event resulted in: Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Life threatening illness (immediate risk of death from the event). Patient received treatment for events included blood thinners. Patient had covid tested post vaccination on 21Apr2021: Nasal Swab: Negative. Patient was not recovered. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1261831-1	Blood Clots both legs; had other vaccine same date at 2nd dose in the left arm; had other vaccine same date at 2nd dose in the left arm; This is a spontaneous report from a contactable Consumer (patient). A 69-year-old male patient received the 1st dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL9269) via an unspecified route of administration in the left arm on 15Apr2021 12:00 PM at single dose for COVID-19 immunization. Medical history was unknown. The patient had no known allergies. The patient had no covid prior vaccination, no covid tested post vaccination. The patient had no other vaccine in four weeks or other medications in two weeks. Concomitant medications were none. The patient had other vaccine same date at 2nd dose in the left arm. The patient experienced blood clots both legs on 15Apr2021 12:15 PM. The patient had extensive dvt IVC and both legs as treatment for event. Outcome of the event blood clots both legs was not recovered. AE resulted in: Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Hospitalization (duration: 5 days), Life threatening illness (immediate risk of death from the event).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1262578-1	Chills, Sweats, Cramps night of vaccine and SOB
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1262687-1	Acute non occlusive thrombus of common femoral and iliac vein in left leg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1263346-1	Patient began very heavy menstrual cycle, 2 weeks earlier than expected, 6 days after receiving second Pfizer COVID-19 vaccine. She is bleeding through maxi pads every 15-30 minutes, passing smaller than quarter-sized clots, with associated severe menstrual cramping, when she previously had no history of this.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1263521-1	Symptom: Very sharp pain at top of stomach followed by very low heart rate and brief loss of consciousness after 1 hour Diagnosis: Partial thombus (blood clot) of celiac axis which extended into the common hepatic artery and splenic artery Treatment: Blood thinners Timeline: Stabilized farily quickly and then pain subsided after 2 days Course of treatment: Monitor for any additional symptoms through additional CT scans
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1263560-1	Per patient's Daughter the patient developed blood clots in his small intestine on 4/13/2021 and died 4/22/21 with physicians unable to explain how or why he developed these clots while on apixaban. She requested his case be reported and reviewed in hopes it helps make the vaccines safer if that was what caused the clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1263908-1	On April 24, 2021, a vein in my arm swole up like a blood clot, the a cephalic vein. I started experiencing nerve burning and tingling in hands, arms and shoulders. The blood clot was in an area that I had damaged years ago and is now clogged. Today is 4/27/2021 and I still have the blood clot and tingling, sometimes in my face also, burning in the shoulder blades, very similar to symptoms of brachial neuritis. I am scheduled for my second shot on 5/4/2021, but thinking I may not get the shot since I am having so much trouble. I have an appointment schedule with a vascular doctor to have an arterial dopler on 5/6/2021, and have the vascular doctor advise.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265579-1	Blood clots in left leg

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265810-1	sweating; felt cold/intense cold even in his bones; He was told by the professional health care that he had small clots in his blood; death cause: Medication; arm started to sore; Doctor identified he had DVT; her husband during that night was not able to sleep; He started having fever; This is a spontaneous report from a contactable consumer (patient's spouse). A 61-year-old male patient received bnt162b2 (BNT162B2), via an unspecified route of administration on 21Mar2021 09:00 (Batch/Lot number was not reported) as single dose f(at the age of 61-year-old) or COVID-19 immunisation. Medical history included dialysis, diabetes mellitus, known allergies: A7, Penicillin, Aspirin, Iodine, Povidone, Pepcid, dyes, iodine allergy. The patient's concomitant medications were not reported. The patient previously took Aspirin, povidone and pepcid ac and experienced drug hypersensitivity with all. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient experienced death cause: medication on 18Apr2021, he was told by the professional health care that he had small clots in his blood in Mar2021, felt cold/intense cold even in his bones on 21Mar2021, sweating on 26Mar2021, her husband during that night was not able to sleep on 21Mar2021, he started having fever on 21Mar2021, arm started to sore and DVT on 29Mar2021. The patient was hospitalized for he had small clots in his blood, felt cold/intense cold even in his bones, sweating for 27 days. The event DVT was medically significant. The course of events was as follows: After getting the vaccine in 21Mar2021 her husband during that night was not able to sleep. He started having fever and felt cold. Days later he continued with the symptoms. On 26Mar2021 after vaccination he had dialysis same day in the afternoon. When arriving home the person notifies symptoms of intense cold even in his bones and then he started to sweat excessively on 26Mar2021 (Friday). The reporter decided to take her husband to the emergency room on 28Mar2021 (Sunday) where he had a general checkup. He was told by the professional health care that he had small clots in his blood. After some time he had health complications where they had suggested to amputate some of the limbs because of this, the reporter alleges those complications were due to the vaccine. On Monday 29Mar2021 same symptoms reappeared and he was admitted to Hospital. Had a blood test and notified to health professional that blood presents small clots. His arm started to sore severely after the sample. Doctor identified he had DVT. Doctor decided to proceed with various medications. Patient received treatment and he was injected: Percose, Morphine, Benadryl, Triphetarin for blood clot reduction. The patient underwent lab tests and procedures which included blood test: blood clot in Mar2021, Sars-cov-2 Nasal swab test: negative in Mar2021 post vaccination. The patient died on 18Apr2021. It was not reported if an autopsy was performed. The outcome of death cause: medication was fatal, of the other events was unknown. Information about the lot/batch number has been requested.; Reported Cause(s) of Death: death cause: Medication
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265958-1	Developed blood clot in right leg (behind knee); This is a spontaneous report from a contactable consumer (patient). A 34-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date not provided), via an unspecified route of administration, administered in Arm Left first dose on 26Feb2021 11:15 at single dose for covid-19 immunisation. The patient's medical history included Developed blood clots after twin c section. The patient has been off blood thinners for 2.5 years. The patient was not pregnant at time of vaccination. The patient's concomitant medications included venlafaxine hydrochloride (EFFEXOR) and vitamin d [vitamin d nos]. On 17Mar2021, the patient Developed blood clot in right leg (behind knee). The patient was put on blood thinners as treatment. The events cause the patient emergency room visit and physician office visit. The outcome of the event was recovering. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265982-1	Blood clot in her lower abdominal area; had the worst pain through her side that ran down her left leg and to her ankle; had the worst pain through her side that ran down her left leg and to her ankle; nerves are all tangled up; This is a spontaneous report from a contactable consumer (patient). A 76-year-old female patient received the second dose of bnt162b2 (BNT162B2, Solution for injection, Lot Number: EN6207), via an unspecified route of administration, administered in Arm Right on 17Mar2021 13:50 (at 76-years-old) as single dose for COVID-19 immunisation. The vaccine was administered at a hospital. It was not administered at a military facility. Medical history included hip replacement four years ago in 2017 (First right then 2 months later left) and knee replacement six years ago in 2015 (both 2 months apart). There were no concomitant medications. The patient previously received the first dose of bnt162b2 (BNT162B2, Solution for injection, Lot Number: EN6203), administered in Left Arm on 25Feb2021 (at 76-years-old) for COVID-19 immunisation and experienced bad headaches. The patient received the second dose of vaccine on 17Mar2021. On 29Mar2021, she was not hurting and went to brunch with friends. Afterwards (on the same date, 29Mar2021), she was sitting and got the worst pain through her side that ran down her left leg and to her ankle. She drove herself home but was in such misery. She went to the emergency room the next day, on 30Mar2021, wherein she had a CT (computerized tomography) scan that showed a blood clot in her lower abdominal area. She was taking pain medication every 6 hours, or she could hardly stand it. On 08Apr2021, she went back to the emergency room because she could hardly stand it. She was kept in observation status but was not admitted. She was scheduled to have another CT scan on 06May2021. She is not sure what they will do because the nerves are wrapped around it. She also mentioned that her doctor wants to do a biopsy. The pain was also running down to her ankle. She stated that she has had both of her hips replaced in the past. She added that her doctor will do an MRI (magnetic resonance imaging). He will put in a drain if it has not shrunk. The doctor can't do anything because the nerves are all tangled up (2021). As corrective treatment, the patient was taking pain medication for the blood clot and pain through her side that ran down her left leg and to her ankle. Outcome of nerves are all tangled up was unknown, while outcome of the other events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265986-1	Doppler showed a blood clot; Pain behind her cast; Really bad case of diarrhea; Swelling in her knee; Leg was swelling/right shoe was tight; Lower extremity venous ultrasound right calf vein acute deep vein thrombosis of the right lower extremity involving the popliteal vein; It was tender on her right leg on the left side of the tibia/It is very tender to the touch; Red mark on her leg with a little nodule; Red mark on her leg with a little nodule; Calf pain; This is a spontaneous report from a contactable consumer. A 62-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number: EP7533; Expiration date was not reported) on the right arm on 30Mar2021 as a single dose, with route of administration unspecified, for COVID-19 immunization at the clinic. Medical history included pain and headache. Concomitant medications included paracetamol (TYLENOL ARTHRITIS) for pain; hydrocodone for pain; amitriptyline for headache; and ongoing paracetamol (TYLENOL EXTRA-STRENGTH) for pain. The patient had previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number: EN6208; Expiration date was not reported) on the right arm on 11Mar2021 (when the patient was 62 years old) for COVID-19 immunization. On 07Apr2021, the patient's doppler showed a blood clot. On 30Mar2021, the patient had a really bad case of diarrhea; swelling in her knee; and leg was swelling/right shoe was tight. On 02Apr2021, the patient had pain behind her cast. On an unspecified date in 2021, the patient's lower extremity venous ultrasound showed right calf vein acute deep vein thrombosis of the right lower extremity involving the popliteal vein; was tender on the right leg on the left side of the tibia that was very tender to the touch; had a red mark on her leg with a little nodule; and had calf pain. The events had resulted into an emergency room visit and physician's office visit. The patient had received treatment for the events, 'blood clot', 'lower extremity venous ultrasound right calf vein acute deep vein thrombosis of the right lower extremity involving the popliteal vein', 'leg was swelling/right shoe was tight', 'pain behind her cast' and 'tender on her right leg on the left side of the tibia that was very tender to the touch'. The outcome of the events was recovered on 30Mar2021 for 'really bad case of diarrhea'; was recovering for 'leg was swelling/right shoe was tight', 'pain behind her cast' and 'red mark on her leg with a little nodule'; and was unknown for all the other events.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1265987-1	4 days later minor blood clot in my finger (not sure it is drug related); Bruise at site of injection; This is a spontaneous report from a non-contactable consumer (patient). A 35-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 09Apr2021 (Batch/Lot number was not reported) as SINGLE DOSE for COVID-19 immunization. Medical history and concomitant medications were not reported. On 13Apr2021, the patient had bruise at site of injection and 4 days later (17Apr2021) minor blood clot in her finger was noted (not sure it was drug related). The patient had no COVID-19 prior vaccination and not tested post vaccination. The patient requires doctor or other healthcare professional office/clinic visit for the events and unknown if treatment was received. The patient had not yet recovered from the events. No follow up attempts are possible. information about lot/batch number cannot be obtained

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1266022-1	Blood clots; headache; nervousness; anxiety; This is a spontaneous report from a non-contactable consumer (patient). A 34-year-old female patient not pregnant received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 03Apr2021 09:00 (Lot Number: ER8733) as single dose (at the age of 34-years-old) for covid-19 immunisation. Medical history included allergies (known allergies: yes). Concomitant medications in two weeks prior to the vaccination included vitamin B complex (B-COMPLEX), magnesium (MAGNESIUM), chlorophyll (CHLOROPHYLL), vitamin D NOS (VITAMIN D NOS). Patient did not receive other vaccine in four weeks prior to the COVID vaccine. It was unknown if patient had COVID prior vaccination. Patient was not tested for Covid post vaccination. On 03Apr2021 patient experienced blood clots, headache, nervousness and anxiety. Events resulted in Doctor or other healthcare professional office/clinic visit. No treatment was required. Outcome of the events was unknown. No follow-up attempts are possible. No further information is expected.; Sender's Comments: Based on the available information, the known safety profile and the temporal association of BNT162B2 administration to the event of Thrombosis, a possible contribution of the drug to the event cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1266035-1	have developed 2 massive blood clots; This is a spontaneous report from a contactable consumer (patient). A patient of unknown gender and age received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) in Feb2021 at single dose for COVID-19 immunisation. Medical history and concomitant medications were unknown. Historical vaccine included first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) in Jan2021 for COVID-19 immunisation. Since then, the patient had developed 2 massive blood clots. The patient did not have a history of blood clots. The outcome of the event was unknown. There was no doubt this was related. No follow-up attempts are needed; information about lot/batch number cannot be obtained.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1266038-1	Subsequently - a doctor discovered numerous blood clots in that leg; then a blood clot in one of her lungs; her left leg (she was inoculated on the left side) swelled up; it was painful; She is being treated but the lymphadenopathy has not subsided in her leg; she is now unable to walk without a cane; This is a spontaneous report from a contactable consumer. A 69-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number not reported) on an unspecified date in 2021 at single dose (inoculated on the left side) for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient received the vaccine few weeks ago, in 2021, and within 3 hours, she recounted her left leg (she was inoculated on the left side) swelled up dramatically it was painful. Subsequently a doctor discovered numerous blood clots in that leg, then a blood clot in one of her lungs. She is being treated but the lymphadenopathy has not subsided in her leg and she was now unable to walk without a cane. The outcome of lymphadenopathy and unable to walk without a cane events was not recovered; outcome of the other events was unknown.Reporter asked if there were reports of blood clots that have been validated as a result of the Pfizer shot. Information on the lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1266039-1	big heart blood clot; This is a spontaneous report from a contactable consumer. A 66-year-old male patient received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 21Mar2021 12:00 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation (Age at vaccination: 66 years). Medical history was reported as none. There were no concomitant medications. The patient experienced big heart blood clot on 23Mar2021 14:00. Therapeutic measures were taken as a result of big heart blood clot. Outcome of the event was not recovered. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1266045-1	15cm Blood clot in left leg - lower calf.; Superficial vein prominence; This is a spontaneous report from a contactable consumer (patient herself). This 50-year-old female patient (not pregnant) received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number ER8732), via an unknown route in the left arm, on 27Mar2021 at 14:45 (at the age of 50-year-old) at single dose for COVID-19 immunisation, administered at hospital. Relevant medical history included Factor V Leiden and varicose veins. The patient did not have allergies. The patient did not have COVID-19 before vaccination. Relevant concomitant medications included acetylsalicylic acid (ASPIRIN), levothyroxine sodium and Women's multivitamin (unspecified). On 07Apr2021, the patient presented 15 cm blood clot in the left leg-lower calf and superficial vein prominence. The events required a physician office visit. An ultrasound was done but results were not provided. Therapeutic measures taken as result of the events included enoxaparin sodium (LOVENOX) injections for 14 days. Post-vaccination COVID test was not performed. The patient was recovering from the events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266051-1	blood clot; lymphadenopathy; unable to walk without a cane; her left leg swelled up dramatically; painful leg; This is a spontaneous report from a contactable consumer. A 69-year-old female patient received her first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date (at the age of 69-years-old) as single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. It was reported that patient received Pfizer shot a few weeks before reporting. Within 3 hours her left leg (she was inoculated on the left side) swelled up dramatically - - it was painful. Subsequently - a doctor discovered numerous blood clots in that leg; then a blood clot in one of her lungs. She was being treated but the lymphadenopathy has not subsided in her leg and she was now unable to walk without a cane. Lymphadenopathy was not resolved; the outcome of the other events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266057-1	strokes signs; Two blood clots in the veins in legs; This is a spontaneous report from a contactable consumer. A 64-year-old female consumer (patient) reported that she received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) (Lot number EN6199/EN6207 (as reported)) intramuscular into the left arm on 10Mar2021 (at the age of 64-years-old) at single dose for COVID-19 immunization. She received the second dose of BNT162B2 dose on 31Mar2021 (lot number unknown). Medical history included high blood pressure and GERD. Concomitant drugs included Toprol 25mg, oral , twice a day for blood pressure stated ''in Feb some time she don't know'' and Protonix 40mg, oral twice a day for years for GERD. The consumer reported that 'she got her first vaccine and she has had the strokes signs''. Further she reported she got her first vaccine dose, then she had leg pain, she went to see the doctor and ordered a doppler. She was put on blood thinners and three days later she got two blood clots in the veins of her legs, right. On 20Mar2021, her leg pain went pretty severe since I had the blood clot, when I got the second shot, it was still there. On 27Mar2021 she started treatment receiving Eliquis 5mg, twice a day for the blood clot and she did take a Physical therapy. When she got the second vaccine dose on 31Mar2021 she was on blood thinners. At the time of reporting the outcome of the event was reported as 'hasn't gotten better, it's the same not better not worse.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266075-1	Multiple blood clots in the left leg had to be submitted to the Emergency Room at the hospital for immediate treatment; This is a spontaneous report from a non-contactable consumer. A 40-years-old male patient received 1 dose of bnt162b2 ((PFIZER-BIONTECH COVID-19 VACCINE) lot number: ER2613 via an unspecified route of administration, in arm left on 12Apr2021 at the age of 40 years as SINGLE DOSE for covid-19 immunization. The patient medical history was not reported. There were no concomitant medications. The patient experienced multiple blood clots in the left leg and he had to be submitted to the emergency room at the hospital for immediate treatment on 16Apr2021 12:00 with outcome of recovering. The patient was hospitalized for 2 days and was treated with unspecified treatment. Covid test post vaccination: Nasal Swab on 20Apr2021 result Negative No follow-up attempts are possible. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266340-1	1/28/2021-swollen lymph nodes and pain to left side of chest and injection site. This pain continued and I went to Dr. on 4/06/2021. Dr. treated me with Toradol and Tylenol#3. Schedule me to have a mammogram and ultrasound on 4/20/2021. This is when it was discovered I had a clot in left arm. Currently being treated with blood thinners(Eliquis).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266361-1	"April 22, 2021 morning my auntie got her first dose of Pfizer vaccine. She happily shared here vaccine experience with family member and relative. April 23 morning, she told us she has severe abdominal pain, has the need to have a bowel movement, but she was defecated. She also told us that she vomit couple times already, then she has no more energy to talk. She got into the hospital and she was announced death right after midnight at 12:04 April 25th. Reason of death given is ""septic shock secondary to ischemic bowel & intra-abdominal sepsis"". But it is the blood clots issue that is causing it. I was told that after my auntie got into hospital the very next day after Pfizer vaccine, hospital later found out that my auntie has blood blockage issue that is causing blood couldn't flow to her intestines. The acid in her blood was very high, 6+. After more than 6 hours without blood to her intestines, her intestines tissues start dying and turned black color. Doctor found out about this at the night of April 23, 2021 nighttime. April 22, 2021 morning is the day my auntie got her Pfizer covid vaccine. There is a very high chances of the Pfizer vaccine is causing the blood clots that lead to blood blockages."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266724-1	First day of period marginally heavier, accompanied with 2 large blood clots. First day of period is always light & clots are not typically found until day 3+ of menstrual cycle. Will continue to monitor menstrual cycle at home.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266827-1	Stroke caused by bloodclot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266861-1	4-16-21 @ 2PM SHE HAD A 'SHADOW IN HER RT EYE'. DROVE TO ER. DX WITH CLOT TO RT EYE AND LT OCCIPITAL AREA OF BRIAN. NO SURGERY. WAS GIVEN PLAVIX. HAS APPT WITH EYE DOCTOR ON MAY 3RD. SAW A NEUROLOGIST AND PCP. SHE IS DOING BETTER NOW AND RT EYE HAS 'CLEARED UP'. WILL NOT BE GETTING SECOND DOSE.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266996-1	I started to have weakness on my left side around 9 hours after receiving the vaccine. I went to the hospital and was told I had a stroke in the right side of my brain from a clot in my brain, which they took the clot out using a catheter through my leg. I am currently undergoing rehabilitation at a rehab hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1267117-1	DVT blood clot formed in left leg, intense pain started almost 14 days after vaccine. No history of any health concerns with blood clots, healthy 35 year old male who works out and diets. No long drives or sedentary lifestyle.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1267155-1	Patient has second Pfizer covid vaccine on 4/12/21, patient had chemotherapy treatment infuse through L portacath on 4/14/21. Patient called office with complaints of arm pain on 4/26/21. Ultrasound on 4/26 was read as: partial, nonocclusive thrombus within superficial left cephalic vein at the antecubital, midportion. Patient was initiated on oral anticoagulation therapy
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1267360-1	After the first vaccine: Leg pain, foot pain, chest pain. Four days after vaccine, the chest pain was so bad that my dad went to ER and had a Pulmonary Embolism . He was hospitalized two nights. After the second vaccine: right leg pain, weakness in both legs. Three days after the second dose, my dad had a stroke.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1268364-1	Stroke from blood clot. .Surgical thrombectomy . After initial full paralysis on left side, not adverse effects.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1268967-1	First period after vaccine was very. very heavy! I was sent to the ER due to big clots and significant blood loss. Period started a week early on 4/22 and is still going on. Usually done within 5 days and not nearly as heavy with clots like this. ER suggested I report a possible correlation between that and the vaccine as they have seen it with other patients as well. ER doctor medicated with birth control hormones to slow down bleeding.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269502-1	Caller states she was not due for her monthly menstrual period for another week, so she found, passing the clot, odd.; Blot clot/Very big blood clot, that passed out of body; Fever, of 102; Sore arm; This is a spontaneous report from a contactable Other HCP (patient). A 35-years-old female patient (no pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Right on 29Mar2021 09:00 (Lot Number: ER2613) as single dose for covid-19 immunisation. Medical history included ongoing asthma , contraception. Concomitant medication included ethinylestradiol, norgestimate (SPRINTEC) taken for contraception from an unspecified start date and ongoing. The patient previously took bactrim and experienced drug hypersensitivity. The blood clot only was for one day on 31Mar2021, and it was one blood clot. The patient went to the bathroom and it was passed, one time. The patient was not due for her monthly menstrual period for another week, so she found, passing the clot, odd. The patient menstrual period did happen on time and it was normal. The patient was not due for her menstrual period for another week, and she takes birth control. The patient contacted her Gynocologist, and it was confirmed that she was not pregnant, and states her menstrual period happened on time. The clot did not happen again. The patient had Fever, of 102 on 30Mar2021, the temperature resolved in a day. The patient took Aleve, drank a lot of water and napped, reclarifies, and the temperature improved. The patient experienced sore arm on 30Mar2021. The outcome of event Clot blood was recovered on 31Mar2021; outcome of event Fever was recovered on 30Mar2021; outcome of event Sore arm was recovered on 01Apr2021, outcome of other event was unknown. No other vaccine in four weeks; No covid prior vaccination; No covid tested post vaccination.; Sender's Comments: Based on the temporal relationship, A possible contributory role of the suspect product to the development of Thrombosis and Menstrual Disorder cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269503-1	Blood clots in colon; 2nd dose 07Jan2021; This is a spontaneous report from a contactable other hcp (patient). A 61-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19), second dose at the age of 61-years-old via an unspecified route of administration, administered in arm left on 07Jan2021 (Batch/Lot Number: Ek5730) as single dose for covid-19 immunisation. The patient medical history was not reported. The patient has no known allergies. Concomitant medication included unspecified medication. The patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), administered in the left arm on 22Dec2020 (lot number: Ek5730) at the age of 61-years-old for covid-19 immunisation. The patient experienced blood clots in colon on 14Jan2021. It was noted that the patient received the second dose on 07Jan2021. The patient was hospitalized for blood clots in colon for 30 days. The patient underwent lab tests and procedures which included sars-cov-2 test: negative on 21Jan2021. Therapeutic measures were taken as a result of blood clots in colon which included surgery. The patient was recovering from the event.; Sender's Comments: Based on event-vaccine chronological association a causal relationship between event blood clots in colon and BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), cannot be completely excluded. . The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269505-1	asked if this was a blood clot; a big black spot next to her knee, and if she touched it, it hurt; a big black spot next to her knee, and if she touched it, it hurt; This is a spontaneous report from a contactable consumer (patient). A (age: 70; unit: unspecified) female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; solution for injection) via an unspecified route of administration on 12Apr2021 as a single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. On 12Apr2021 the patient experienced a big black spot next to her knee, and if she touched it, it hurt. The patient called to report that she received the first dose of the COVID-19 vaccine yesterday 12Apr2021 and she was supposed to receive the second dose on 03May2021. When she got home yesterday afternoon, she ended up with a big black spot next to her knee, and if she touched it, it hurt. It was a little larger than a quarter coin, about the size of a silver dollar. She stated it was next to the inner part of her knee. The patient asked if this was a blood clot. She stated that she was worried with all the information about blood clots and the COVD-19 vaccines. She stated that she did not know if it was a blood clot but asked if it was a blood clot. The clinical outcomes of the events a big black spot next to her knee, and if she touched it, it hurt were both unknown. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269525-1	atrial fibrillation; atrial fibrillation; splenic infarction; extreme left upper abdominal pain; large portion of his spleen unusable; blood clots causing that portion of his spleen to die; blood clots causing that portion of his spleen to die; This is a spontaneous report from a contactable consumer (patient). A 68-year-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot: EN6206), via an unspecified route of administration in right arm on 23Mar2021 09:00 (at the age of 68-years-old) as single dose for covid-19 immunisation. The vaccination facility type was a health department. Medical history included atrial fibrillation (under control with medicine). The patient's concomitant medications were not reported. The patient previously took ciprofloxacin hydrochloride (CIPRO) and experienced allergies: Cipro. The patient had no COVID prior vaccination and was not COVID tested post vaccination. No other vaccine in four weeks. It was reported that within 13 days of the injunction (on 06Apr2021 at 04:00), the patient had three atrial fibrillation events on three separate days between 01:00 and 03:00 while sleeping and on day 13 was admitted to the hospital with extreme left upper abdominal pain they diagnosed with splenic infarction and rendering a large portion of his spleen unusable from blood clots causing that portion of his spleen to die. Now the patient have to take blood thinners the rest of his life. The events resulted in emergency room/department or urgent care, hospitalization for 1 day, disability or permanent damage. The events treatment included several pain shots administered then given enoxaparin sodium (LOVENOX). The outcome of the events was recovered with sequel.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269527-1	multiple pulmonary embolisms/Pulmonary embolism and subsegmental pulmonary embolus; felt sluggish directly; feeling discomfort in my lungs; pain in my lung/chest area; pain in my lung/chest area/right-sided chest pain under the rib; difficult and extremely painful to breathe/significant shortness of breath/shortness of breath due to extreme pain; blood clots; This is a spontaneous report from a contactable consumer (patient) and physician. A 60-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number EP7534 and expiration date not provided), via an unspecified route of administration, administered in Arm Left second dose on 07Apr2021 17:00 at single dose for covid-19 immunisation. The patient medical history was not reported. The patient was not pregnant at time of vaccination. The patient's concomitant medications included cetirizine, diphenhydramine, and phenylephrine. The patient historical vaccine includes bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number EN6203), right arm first dose on 10Mar2021 for COVID-19 Immunization. On 07Apr2021 21:30, the patient felt sluggish directly after receiving the second dose. The patient began feeling discomfort in my lungs about 5 hours after the dose. The pain in my lung/chest area grew worse throughout the night to the point where it was difficult and extremely painful to breathe. The patient went to urgent care in the morning, and they ran an EKG, urine analysis, and chest X-ray: all with unknown results. They were concerned it could be something worse along with potential pneumonia, so they sent me to the emergency room. In the emergency room, I received a CT scan where they found multiple pulmonary embolisms (1 large one in my right and multiple in my left lung). The patient stated that she had just had my yearly physical the previous week (Apr2021), with no signs of any health issues. With that, the doctors believed my condition to be related to my 2nd dose, so I was admitted to the hospital. I had to stayed for 2 nights. I am now on blood thinners, other meds (pain medications), and require oxygen during sleep (O2 levels drop too low). I have been referred to lung and hematology specialists. Also, reported that patient experiencing significant shortness of breath and right-sided chest pain under the rib. This persisted for some time until she presented to the emergency department found to have segmental and subsegmental pulmonary embolus. Additionally, the patient experienced blood clots, chest pain, and shortness of breath due to extreme pain. Hypercoagulable work-up pending. The patient underwent lab tests and procedures which included COVID-19 virus test (nasal swab) with result of negative. The outcome of the events was recovering.; Sender's Comments: Based on the information available, a possible contributory role of the suspect BNT162B2 cannot be excluded for the reported events occurred in a plausible temporal relationship. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269535-1	blood clot under her eye; bright red bruise in the corner of the eye to the middle of the eye underneath; feeling crummy; she was down; headaches; couldn't move; arm hurt all the way below her arm pit into her breast; arm hurt all the way below her arm pit into her breast; arm hurt all the way below her arm pit into her breast; got very sick; This is a spontaneous report from a contactable consumer (patient). A 59-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: ER8737), dose 1 via an unspecified route of administration, administered in Arm Left on 05Apr2021 08:30 (at the age of 59years) as single dose for Covid-19 immunization. Medical history included ongoing rheumatoid arthritis and osteoarthritis (not bad enough that she takes treatment, but she is diagnosed; she was diagnosed way back in 2006). Concomitant products: patient stated she takes some medications (not further specified). The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient reported that she had her first dose of the Pfizer Covid-19 vaccine on 05Apr2021 and got very sick. She was down for two days and after day 10 (as reported) she has a blood clot under her eye. She would like to know what would happen if does not get a second dose. She consulted if there is any talk on whether or not people should receive an additional dose of the Pfizer Covid-19 vaccine next year. The patient further reported that on 06Apr2021 she woke up feeling crummy right away and a couple of hours into that day she couldn't lift her arm, literally couldn't lift arm for 6 inches and stated it went on for 2 days. She further stated that her arm hurt all the way below her arm pit into her breast. She stated that it all ended on 08Apr2021 and she went back to work on 08Apr2021. The patient added that she was down for two days, had headaches, and stated she never knew what people meant when they said they got hit by train. She doesn't believe she had a fever and couldn't move. The patient further stated that she got pretty darn sick and this morning 14Apr2021, she woke up with a blood clot under her eye. She is an esthetician and is pretty good with skin. She stated that she doesn't know how she could get this overnight unless she injured herself in her sleep, but she is a light sleeper. She does not know if it is a clot. She also stated that she called a medical esthetician who told her to take a baby aspirin and cold compress. The patient clarified that she didn't feel so good and today she feels better. She doesn't know that it is a blood clot, it is not raised and not hard, it looks like she has a bright red bruise in the corner of the eye to the middle of the eye underneath, not in the eye. She wants to make it clear she doesn't know if it is a blood clot. She feels like if she hurt herself, that would be the reason, but it would be painful, and it is not. She added that the bruise is about the same. Therapeutic measures were taken as a result of the events (except for feeling down and couldn't move). The outcome of events thrombosis and contusion was not recovered; the outcome of other events was recovered on 08Apr2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269617-1	"Got worse and the leg swelled and was also having foot swelling because blood flow wasn't there; sore arm; blood clot; pain behind left knee; This is a spontaneous report from a contactable consumer (patient's wife). An 81-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration, administered in right shoulder on 23Feb2021 (Batch/Lot Number: EN6202) (at the age of 81-years-old) as single dose for covid-19 immunisation. Medical history included diabetes mellitus, stage 3 renal failure (was not bad, was in stage 5 couple of years ago, was a month or so away from full failure, was in the hospital and had dialysis, was able to get reversed with medication and diet), high blood pressure, and blood cholesterol increased/high cholesterol. The reporter stated that with regards to patient's height, he shrunk a little. Patient had no prior vaccinations within 4 weeks, didn't get flu shots. Concomitant medications include about 12 other unspecified drugs, takes statins, has high blood pressure and high cholesterol. Patient started experiencing pain behind his left knee about 6 or 7 days later in Mar2021 (also reported as 04 or 05Apr2021, pending clarification). By that Sunday, it had gotten worse. The leg swelled and was also having foot swelling because blood flow wasn't there. There was no TIA's no stroke, no heart racing, no sweating, was all pain. Got leg up and kept elevated until they could get to the doctor and get imaging. Patient went Monday 08Mar2021 to see the doctor. Patient was brought to the doctor and was then sent to the imaging place. Radiology read the imaging around 5:15pm on the Monday, 08Mar2021. The radiologist said there was 2.5"" blood clot behind the left knee. He had never had a blood clot before. Patient was then sent to the Emergency Room at (Name) Hospital, in (Name) where they live. Was kept there several hours. Did a lot more blood tests. He was immediately put on 10mg ELIQUIS 10mg, once in the morning, and once at night. He is now taking 5mg Eliquis in the morning, and 5mg at night. He has not had any of these problems before the Pfizer shot. The reporter knew that not a lot had been reported. It seemed it should be safe. The correlation and timing having the shot on 23Feb2021 and all of a sudden a week and a half later this happened. It was 10 days later. There was nothing they can do about it. Just continuing with the ELIQUIS, trying to dissolve the blood clot, will be staying on it for 4 to 6 months. It was a pretty big blood clot and were worried about it breaking up and traveling. The patient had no other symptoms, had a sore arm, that was it. Patient going back to (Name) 01Jun2021 and he will have another CT scan or ultra sound when they get back. The reporter did not want to need to get the leg amputated. Patient was only in the ER, was not admitted to the hospital. Patient was concerned because of his age. If it doesn't dissolve, they know there will be a lot more trouble. The event of pain left knee, blood clot and sore arm required a visit to the emergency room and physician's office. The patient received the second dose of BNT162B2 on 01Apr2021 (lot number: ER8730) in the right arm. The outcome of events was unknown."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269707-1	left leg deepvein thrombosis; blood clot; This is a spontaneous report from a contactable consumer (patient). A 73-year-old male patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: EL9269), via an unspecified route of administration, administered in Arm Right on 12Feb2021 19:30 (7:30 pm) as single dose for COVID-19 immunization in a hospital. Medical history included high blood pressure and high cholesterol both from an unknown date. No known allergies. The patient took an unspecified medication. No other vaccines in four weeks. No COVID-19 prior to vaccination. The patient received the first dose of bnt162b2 (lot number: EL9261) on 22Jan2021 07:15 am on the right arm for COVID-19 immunization. On 25Feb2021 21:00 (9:00 pm), the patient experienced left leg deep vein thrombosis and blood clot which required a visit to the emergency room and subsequent hospital admission. The patient was given Eliquis as treatment. The patient was tested for COVID-19 post vaccination via nasal swab on 08Apr2021 with a negative result. The outcome of the events was reported as recovering.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269731-1	Clotting; Leg got worse/ leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis; Leg got worse/ leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis; Leg got worse/ leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis; Leg got worse/ leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis; Leg got worse/ leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis; This is a spontaneous report from a contactable consumer (patient). A 37-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE lot/batch number and expiration date not reported) via an unspecified route of administration on 10Mar2021 at a single dose for COVID-19 immunisation. Medical history included allergies and asthma. Concomitant medication included levocetirizine dihydrochloride (XYZAL) taken for allergies. The patient previously received first dose of BNT162B2 on 17Feb2021 for COVID-19 immunisation; and experienced thrombophlebitis and leg swelled up and got very sore, painful, and red (lot number: EL9266 at 04:00 PM). About 2 weeks after the first dose, patient's leg swelled up and got very sore, painful, and red. Doctors advised that it was likely thrombophlebitis. She received the second dose on 10Mar, and her leg got worse a few days later. She visited the emergency room and was sent to a specialist who put her on blood thinners for clotting. Patient stated that the only significant change or occurrence in her life was that she received the vaccine. The events resulted to a doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. Prior to vaccination, the patient was not diagnosed with COVID-19. Post vaccination, the patient has not been tested for COVID-19. No known allergies. Patient was not pregnant. Outcome of the event clotting was unknown, while not recovered for the other events. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269733-1	blood clot; swelling and pain in right leg; swelling and pain in right leg; right arm hurting tremendously; This is a spontaneous report from a contactable Nurse. A 52-years-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Right on 01Apr2021 13:45; at the age of 52-years-old, (Batch/Lot Number: ER8734) as SINGLE DOSE for covid-19 immunisation. Medical history included ongoing hypertension, ongoing memory impairment (She states that she was diagnosed either last year or the year before that she can't remember. She states that it is ongoing, but it is well controlled). Historical vaccine included vaccine to tetanus (About 3 or 4 years ago). There were no concomitant medications. The patient experienced blood clot (thrombosis) (disability) on 13Apr2021 with outcome of not recovered, swelling and pain in right leg (disability, medically significant) on 09Apr2021 with outcome of not recovered, right arm hurting tremendously (disability) on 01Apr2021 with outcome of recovered on 04Apr2021. The patient underwent lab tests and procedures which included ultrasound scan: positive on 13Apr2021 had a blood clot. Therapeutic measures were taken as a result of blood clot (thrombosis) included Xarelto and due to pain in extremity the patient received ibuprofen (7 ibuprofen every day); Sender's Comments: Based on the information available and a close temporal association, a possible contributory role of the suspect BNT162B2 cannot be excluded for the reported events of Thrombosis. The case will be reassessed once new information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269745-1	after 3 years of not having her period, she suddenly started bleeding heavily and having menstrual cramps. The bleeding was very clotted and seemed somewhat abnormal; after 3 years of not having her period, she suddenly started bleeding heavily and having menstrual cramps. The bleeding was very clotted and seemed somewhat abnormal; after 3 years of not having her period, she suddenly started bleeding heavily and having menstrual cramps. The bleeding was very clotted and seemed somewhat abnormal; This is a spontaneous report from a non-contactable consumer. A 20-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation (Age at vaccination: 20 years). The patient's medical history and concomitant medications were not reported. On an unspecified date, the patient stated that after 3 years of not having her period, she suddenly started bleeding heavily and having menstrual cramps. The bleeding was very clotted and seemed somewhat abnormal. The patient underwent lab tests and procedures which included nasal swab: negative on an unspecified date. The patient did not have COVID prior vaccination. The patient was not pregnant. Outcome of the events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269746-1	Regular pain feeling; Headache; Muscle pain; Calf feels tight, stiff, it feels like dead weight; Calf feels tight, stiff, it feels like dead weight; Pain running up the right arm up to the shoulder/Calf pain; He asked if someone can tell him if it is a blood clot in the back of his calf; This is a spontaneous report from a contactable consumer (patient). A 46-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in left arm (shoulder) on 12Mar2021 10:07 (Lot Number: EN6202) as single dose for covid-19 immunization. There were no medical history and concomitant medications. The patient has no prior vaccinations within 4 weeks. The patient was not sick at time of vaccination. The patient reported that on 12Mar2021, he has been feeling the regular pain feeling, a headache, muscle pain before, it comes and goes. However, he said that the last few days (2021), there had been, by his calf muscle in the back there, they've been tightening, like stiffening up on him when he's sleeping. His calf feels tight, stiff, it feels like dead weight. It wakes him up at times. He also reported pain running up the right arm up to the shoulder on unknown date in 2021. He was just curious in terms of what that might be. He asked if someone can tell him if it is a blood clot in the back of his calf. The headache went away after 2-3 days. He also reported that his eyes also hurt a little bit and was the same time frame as the headache (onset date reported as 12Apr2021: after the second dose, pending clarification). He stated that those things were just mild, nothing major. He did not take anything for these. He confirmed he no longer had muscle pain. He just has pain behind the calf and right arm up to his shoulder. His calf pain, started to feel tight, started within the last 2- 3 weeks (2021). The arm pain in right arm up to shoulder started maybe within 2 days. The patient received the second dose on 02Apr2021. Outcome of the events muscle pain and headache was recovered in Mar2021, events pain running up the right arm up to the shoulder/Calf pain was not recovered, and outcome of other events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269761-1	blood clot; feels like there is a worm in between her ankle and knee, crawling around her leg; This is a spontaneous report from a contactable consumer (patient reported for herself) received from a Pfizer sponsored program. A female patient of an unspecified age received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for injection, Lot Number: Unknown) via an unspecified route of administration on 15Apr2021 as single dose for covid-19 immunization. The patient's medical history and concomitant medications were not reported. She felt like there was a worm in between her ankle and knee, crawling around her leg. She thought that it might be a blood clot and it might crawl going to her brain. She said she's been up all night and she put an elastic bandage on her leg so that it won't crawl going to her brain. The outcome of the event was reported as unknown. Follow up attempts are needed. Information about lot/batch number has been requested

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269769-1	Blood clots; Major blood loss; Blurred vision; Heart palpitations; Shortness of breath; Extreme fatigue; Dizziness; This is a spontaneous report from a contactable other hcp (patient). This 37-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in arm left on 29Mar2021 12:00 (Lot number was not reported) as single dose (at 37 years old) for COVID-19 immunisation. Medical history none. There were no concomitant medications. On 31Mar2021 01:00, the patient experienced blood clots, major blood loss, blurred vision, heart palpitations, shortness of breath, extreme fatigue, dizziness. The outcome of the events was unknown. Therapeutic measures were taken as a result of the events included blood transfusion, ongoing tests, ultrasounds. The events required a visit to the emergency room and a visit to the physician's office. Seriousness criteria of the events was reported as serious due to disability and life threatening. Prior to vaccination, the patient was not diagnosed with COVID-19 and since the vaccination, the patient had not been tested for COVID-19. No allergies. The patient was not pregnant at the time of vaccination. The information on the Lot / Batch number has been requested.; Sender's Comments: Based on the limited information currently available, a possible association of the suspect drug administration with the reported events cannot be completely excluded, due to a plausible temporal relationship. This case will be reassessed when additional information becomes available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1270155-1	Patient had 3 different clots in her left leg and a small-moderate PE for the first time ever, 2 weeks after #2 Pfizer vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1271348-1	The day after the vaccine I experienced brain fog, fatigue, and sore arm. I also got my period. I have a history of painful periods but after an endometrial ablation over 10 years ago I have had normal pain. The period after the vaccine was incredibly painful and the consistency of the discharge was completely different. Pale pink clots that almost looked like uterine lining instead of clots I'm used to seeing. I want to report this because I've talked to other women who've had something similar but I didn't see anything about it on the CDC website.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1271486-1	Patient had vaccine on 4/23/21 and developed calf pain on 4/26/21. No other symptoms. No other triggers known at that time for a DVT. US did confirm a small blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1271804-1	The day after receiving the first dose of the Pfizer COVID-19 vaccine, the patient began to experience heavier than normal menstrual blood loss and blood clots. The patient returned in 21 days and received her 2nd dose of the vaccine and began to experience heavy clots and blood loss again along with headaches, paleness, and weakness. The patient saw a physician and was given medroxyprogesterone to decrease the bleeding on 4/15/2021. On 4/28/2021 the patient reported to an ER and was given a transfusion due to low iron levels.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1271962-1	On April 5th, 2021 I went to the E.D. , for shortness of breath and severe right shoulder pain. While there they got a chest x-ray, ran labs, and did a chest CT. The CT revealed that I had one blood clot in my left lung and two blood clots in my right lung, as well as slight pleural effusion in the right lower lobe. They put me on Xarelto for the next 3-6 months and monthly follow up visits with my primary care provider. I was also given Hydrocodone as need. I experienced extreme pain with breathing and sort of movement for a week and half but currently only have occasional shoulder and back pain usually at the end of the day.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1272256-1	1.5 weeks after my first shot I started my period 1 day early. Everything was fine the first day but by the afternoon of day 2 I was bleeding extremely heavy and several very large blood clots past in a matter of minutes. I called consulting nurse who told me to go to the ER immediately. I then went to the ER for treatment. They did some tests and sent me home and told me to watch my symptoms closely for the next few days. This was last night and I have no change today. I'm still bleeding very heavily and passing large clots. I have never in my life experienced a period of this nature.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1272549-1	Heavy menstrual cycle came the day I got the Covid vaccine, despite not having period while on IUD for many months before this. Clotting in period, cramping.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1272707-1	Day 3 cramping in uterus, uteran contractions Day 4 spotting, intense pain and reddish pink discharge Day 5 bloody discharge turned red and then it turned brown and continued that way for a week and then the next week it started clotting and the bleeding started again and continues to this day.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1273067-1	Blood clot in right leg.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1274088-1	Pulmonary embolism, pea, heart arrest x2. 0100 4/14, some up with difficulty breathing. O2 sats low 80's and tachycardia low 100's. 4/15, says in mid 90's yacht at mid 120' s. 4/15 increased diffficulty breathing. Went to ed. Troponin in 800's. Saddle pe discovered on ct. Clots in rt leg found also. 4/16 is removed pe, ir decided to let clots in let reabsorb. 4/17, ready to d/c, when acute onset of respiratory distress occured. Went into PEA x 's 2. Back to ir to remove pe and place filter.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1274232-1	"Leg started swelling 4/22 and became painful. Saw Dr. on Monday 4/26 and had venous ultrasound which showed a ""blood clot"" was started on xarelto"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1274501-1	Swollen left leg noticed on 3-19-2021 as a result of blood clots identified by ultra sound on 03-31-2021
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1274793-1	My left leg started hurting from my foot all the way up to the top of my thy was swollen painful.. the morning of 4/21/21
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1275514-1	9 days after receiving the first dose of the Pfizer vaccine, the patient developed superficial thrombophebititis in her right lower extremity that extended despite compresses, elevation, and aspirin use. A 2nd ER visit showed clot above and below her knee and in her greater saphenous vein which is a risk for deep venous thromboembolism and she was started on rivaroxaban (anticoagulation).
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1275842-1	After 2 days of receiving my first COVID vaccine, I started my menstrual cycle. My cycle was very heavy with large clots. It lasted four weeks. Very unusual, I never had a menstrual cycle like this before.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1277344-1	headache; nausea; light headed/ dizziness; unsteady gait; vomiting; chills; sweats; superficial blood clot; pain to back of Rt knee; bruising; swelling; This is a spontaneous report from a contactable Nurse. A 57-year-old non-pregnant female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscularly, administered in Arm right on 31Mar2021 (Batch/Lot Number: EW0150) (at age of 57-year-old) as single dose for COVID-19 immunisation. Medical history included osteoarthritis, and migraines. The patient's concomitant medications were not reported. The patient previously took the first dose of BNT162B2, intramuscularly, administered in left Arm on 10Mar2021 (at age of 57-year-old) (Lot number: EN6207) for COVID-19 immunisation. It was unknown if other vaccine in four weeks. It was unknown if COVID prior vaccination. The patient experienced onset of pain to back of Rt knee, bruising, and swelling on 06Apr2021. She called doctor on 09Apr2021, provider diagnosed her with superficial blood clot. Treating with applying warm compress to area. Client had onset of headache, dizziness, nausea, light headed, unsteady gait, vomiting, chills and sweats on 10Apr2021. Patient called doctor on 14Apr2021, they recommended covid-19 testing, results are negative on 12Apr2021. Symptoms ongoing since 14Apr2021. The outcome of events was not resolved.; Sender's Comments: Based on plausible temporal association, a causal association between the reported event thrombosis and suspect drug bnt162b2 cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278155-1	Stroke caused by blood clot. Spent 6 days in medical facility as a result. Unknown if vaccine and blood clot are related - but it seemed wise to report for further investigation

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278466-1	stroke caused by a blood clot; stroke caused by a blood clot; right arm went completely numb; She is tired; shortness of breath; her eyes were blurry; her head felt like somebody put a spike in it on the right side; sore arm; This is a spontaneous report from a contactable consumer. A 77-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 06Feb2021 13:15 (Lot Number: EM9809) (at age of 77-year-old) as single dose for COVID-19 immunisation. Medical history included colitis ulcerative from Oct2020 (dropped weight, she dropped from 127 pounds to 109 pounds) , cardiac disorder, atrial fibrillation, and osteoporosis, all were ongoing. Concomitant medications included carvedilol taken for cardiac disorder, atrial fibrillation from an unspecified start date and ongoing; cyanocobalamin (VITAMIN B12) taken for cardiac disorder from 2018 (reported as has taking it close to 3 years) and ongoing; denosumab (PROLIA) taken for osteoporosis from an unspecified start date and ongoing; ubidecarenone (Q10) taken for cardiac disorder from unknown start date and ongoing; calcium, colecalciferol (CALCIUM + D3) taken for osteoporosis from 2020 (reported as a year ago) and ongoing; influenza vaccine (FLU VACCINE VII) taken for an unspecified indication, start and stop date were not reported, and vitamins NOS (MULTIVITAMIN) from an unspecified start date and ongoing, the patient started taking it when she was a senior in high school and stated because she is not the best eater in the world. There was no prior vaccinations (within 4 weeks). AE(s) following prior vaccinations: The patient stated arm is sore with flu shot and probably did when she was little when she got the loaded shot, the mumps one. She wouldn't remember that because she was little. The patient experienced stroke caused by a blood clot on 20Feb202114:30 with seriousness criteria hospitalization, sore arm on 07Feb2021, and right arm went completely numb, tired, shortness of breath, her eyes were blurry, her head felt like somebody put a spike in it on the right side on an unspecified date. The event stroke caused by a blood clot result in emergency room visit.The patient was hospitalized for stroke caused by a blood clot from 23Feb2021 to 24Feb2021. The patient reported that she received the first dose and she is thankful she got it on 06Feb2021 and the only side effect she had was a sore arm. Stated on 20Feb2021 she had a stroke caused by a blood clot obviously, she ended up in the hospital. She didn't think at the time, doesn't fault the physicians that took care of her. She didn't think it was related and her eldest son thought it was related. The patient stated they thought it was a mild stroke and after they did the C scan the neurosurgeon, while she was lying in the hospital bed, looked at her assistant and said he couldn't believe that she was talking and walking. The patient stated part of it was ignorance on her part, everyone thought it was the left side that numbs, but hers was the right side. Stated she didn't fall, she dropped something and went down to get it and her right arm went completely numb and she couldn't get back up and she tried to push with legs and couldn't get up. Stated all of a sudden her eyes were blurry and then she never gets headaches and her head felt like somebody put a spike in it on the right side, stated it is supposed to affect the left side. stated she has a heart problem. The patient stated she had a sore arm for about a week, lasted for a few days. stated she gets a sore arm with the flu shot. stated when she would roll over to lay on her left side and roll over on that arm it was enough to wake her up. stated it lasted about 5 days. She couldn't figure out why it was lasting so long. Stated the second thing that happened was that she went to her son's home that night for dinner, hadn't told him about it and while she was eating dinner she asked am if she leaning to the left and he said yes, she was and that her words were being slurred and only lasted for about 10 mins. Stated she didn't call and ignorance is unreal with this stuff. She went to urgent care the next day and they didn't call the ambulance. When she went to the doctor on 23 Feb2021 she was sent immediately to Hospital so she could go through the ER and that's when the neurosurgeon became involved. Stated right now her balance isn't quite right but she is driving and she was taking care of herself and they didn't send anyone home to help her. She was not having any problem talking. Stated she was not going to get into this and will beat it and get back to normal. She was tired a lot, which they said it's normal. Stated she was a real dynamite. Stated it is improving slowly, she was extremely tired and not taking her long walk like she was used to and has shortness of breath, she was not trying to over do it. Stated she always gets a sore arm with the flu shot. Stated her pain level is lower than most people. The patient stated the sore arm started late evening but the next day was really hurting and the main thing was rolling over. The patient stated she didn't take anything for her sore arm, she won't even take an aspirin because some of these Advil medications will mix with the heart medications. Stated in the hospital they were giving her medications she had through an IV in her arm but doesn't know what was going through that. Stated they put her on a blood thinner and they were testing the blood thinner through the IV. She was so miserable, she just wanted to go home and get a good nights sleep. Stated they took all kinds of test, checked her oxygen and had to have a certain number before she could go home. The patient received treatment for event stroke caused by a blood clot and not received treatment for sore arm . The outcome of events stroke caused by a blood clot was resolving, of event sore arm was resolved on 12Feb2021, of other events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				problem with her cholesterol; pain on the top of my head and my hands hurt/headache; pain on the top of my head and my hands hurt; Nauseous; my hands and feet started going numb; they itch they burn; feet are painful; uncomfortable; blood clot/a clot or something in her legs; from the knees down I was having a lot pain/leg pain felt like it is really deep in the vein; she thought she was having a stroke; cramps; burning from my calf to my feet; pain in her neck then to her back; pain in her neck then to her back; different temperatures like hot and cold make it bad and irritating; can't sleep well because of her symptoms; Swelling in her Feet; can't think right; This is a spontaneous report from a contactable consumer. A 43-years-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), via an unspecified route of administration in the right arm in the morning of 01Apr2021 (Batch/Lot Number: ER8727) as a single dose for COVID-19 immunization. Medical history included ongoing complex regional pain syndrome, ongoing fibromyalgia, ongoing arthritis - all diagnosed 20 years ago. Her mother has lupus and rheumatoid arthritis and she can't have the vaccine, states she doesn't know if something could be genetic. Low potassium in the past and stated that she has a lot of allergies to medications, but she consulted with the pharmacist and he said it would be okay. There were no concomitant medications. No additional vaccine was administered on the same date of BNT162B2 and no prior vaccination within four weeks. The patient was having weird symptoms and she tried looking online and in different groups to find out if anyone has similar symptoms, but she hasn't been able to find anything. She began by saying she has health issues; she has Fibromyalgia and Complex Regional Pain Syndrome. She doesn't know if studies were performed on patients with those disorders that took the vaccine. It's been about 3 weeks since getting the vaccine (Apr2021) and she started having symptoms that she thought could be a blood clot. From the knees down, she was having a lot of pain, cramps, burning from her calf to her feet. She would keep rubbing them and massaging them. It was excruciating pain. She was having symptoms and she did go see her Nurse Practitioner on 14Apr2021 because she thought she was having a stroke or a clot or something in her legs or that she was having a problem with her cholesterol. The nurse thought maybe it was her electrolytes, because she had low potassium in the past, but everything came back normal. Her feet were painful but not as bad in Apr2021. She felt like her condition but it's flaring up like 200% it's like over a 10 on the pain scale. She can't think right, she can't work, and was very uncomfortable in Apr2021. The leg pain felt like it is really deep in the vein, it doesn't feel like it is superficial, and she can feel stuff or liquid flowing. Her pain alternates from her head to her neck to her feet and it is very uncomfortable. Her hands and feet started to go numb, not completely numb but like when they fall asleep from sitting on them and then hit something and it hurts. Her arms were burning like crazy; they itch, they burn in Apr2021. Different temperatures like hot and cold make it painful, bad and irritating, and she can't sleep well because of her symptoms. She doesn't know the exact dates, but it has been about a week, but it is changing. It started with cramps in her legs from her knees down. That is where she had injuries and she thought it was her cholesterol. From her knees down, she has cramps and pain and it is very uncomfortable, she felt a really deep pain. She has pain in her neck then to her back, it started in her lower extremities and it is going up. It has stayed the same and she has to keep taking Tylenol to control the pain. Her symptoms got worse and she started taking Tylenol and that helped and then on 15Apr2021, her hands and feet started going numb and she had a headache and pain on the top portion of her head (reported as brain) and her hands hurt, and she was nauseous. She hasn't heard back from the nurse practitioner, so she decided not to go and get it done because she felt it is from the vaccine. She also has swelling of feet in Apr2021, and now she just woke up an hour ago and she slept somewhat okay, but she had to take Tylenol. Now that she is standing, it felt like her legs are starting to hurt again. Treatment included Tylenol, which helps a little and she has to keep taking it. If she doesn't take Tylenol, she can't even function. The events from the knees down I was having a lot pain/leg pain felt like it is really deep in the vein, cramps, my hands and feet started going numb, they itch they burn, feet are painful, uncomfortable, pain on the top of my head and my hands hurt/headache, swelling in her feet, nauseous, pain in her neck then to her back, different temperatures like hot and cold make it bad and irritating had not resolved, while the event can't sleep well because of her symptoms was resolving, and the outcome rest of the events was unknown.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	[1278527-1]	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"swollen joints; her back condition and may have ""pinched a nerve or something; describes it as the worst pain; the leg pain that she is experiencing and would like to know if it is possible that it could be related to a blood clot; leg pain; also has some tingling on the skin; It feels like a Charley's horse; had a horrible flu; Her every muscles and every joint hurt, and her head hurt; Fever; body aches; chills; joint pain; Her every muscles and every joint hurt, and her head hurt; This is a spontaneous report from a contactable consumer (patient). A 53-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in left arm on 07Apr2021 at 14:00 (Batch/Lot number was not unknown; Expiration Date: 31Jul2021), at age of 53 year-old, as 0.3 mL single, for Covid-19 immunisation. Medical history included liver disease from 2016 and ongoing. There were no concomitant medications. The patient previously took the first dose on 17Mar2021 for Covid-19 immunization and experienced fever. The patient received her second dose of the Pfizer BioNTech COVID-19 vaccine on 07Apr2021 and reports having side effects after. She is specifically worried about the leg pain that she is experiencing and would like to know if it is possible that it could be related to a blood clot and if she needs to be worried about it. She stated that with the ""swollen joints"" and everything from the vaccine, it is possible that it worsened her back condition and may have ""pinched a nerve or something."" She describes it as the worst pain she's ever felt in her life and states it ""comes out of the blue"" so she does not have time to prepare for it. She likens it to a ""Charley horse"" but a ""really bad one"" that goes away and has no pain afterwards. She states she is scheduled for an MRI next week of her lower back to investigate. The patient also mentions that she thinks she had COVID-19 at the time of her vaccine because ""when I got the shot, I had a super strong response."" On 07Apr2021, the patient had body aches, chills, she was on the couch for quite a while. The joint pain was incredible, even in her toes. She felt like she had a horrible flu. Her every muscles and every joint hurt, and her head hurt. She was on the couch for like 4 days. On day 3 or 4 she developed intermittent pain in the leg and shin on 08Apr2021. It feels like someone is drilling a jackhammer in it. It feels like a Charley's horse, but not, more like it's in the bone. She didn't associate it with the vaccine at first, even though she had never had this pain before until she got the vaccine. She said that she has had some back problems and scheduled a MRI to see if she had pinched nerve or something. On 08Apr2021, she also has some tingling on the skin. She read about side effects of blood clots and doesn't know if this leg pain could be related to a blood clot. She assumed it could be from her back. She read that with blood clots there is usually an ache and discoloration, which she does not have now. She also read that intermittent Charley's horse could have something to do with blood clots. She has pins and needles on the skin. She also had a headache after the vaccine, but she does not have it now. She said that the pain in her leg that shoots on and bangs in her tibia hurts when bending over, lifting her leg to her chest, squatting to pick up something, and going up and down the stairs. She said that the pain is horrible and it is worse than leg pain, it is worse than a charley horse, it is the worst pain she has ever felt in her life. She cannot stop screaming when it happens. States with the joint pain that she had, there may be a nerve that's being pressed on her back. She said that with all of the information being reported in the news about blood clots, she asked is that something she needs to be concerned about. She said that she did a virtual visit with her doctor and notified her of the leg pain and asked the doctor to schedule her for an MRI because of the pain. She stated her hemoglobin and hematocrit, MCV are high. She also said that the clotting is affecting her red blood cells. She ran a fever with the first vaccine, but she had a high fever between 103-104 degrees Fahrenheit after the second vaccine that stopped on 12Apr2021 or 13Apr2021. Her high fever lasted for about 3 days. She had a low grade fever that lasted for about a week. Her body aches resolved completely on 12Apr2021 or 13Apr2021. Her joint pain is in her back and elbows. She also had lot of joint pain in her shoulder and arm and radiates to her neck on the left side. She still has some muscle pain in her leg, but overall her muscle pain is gone. For treatment she used Ibuprofen 600 mg by mouth and fluids. The day before her shot, her son called and told her he tested positive for covid. She got a PCR test immediately on 07Apr2021 and the next day she was told it was negative. She thinks she might have tested too soon. She also says her response to it was so strong, she must have been getting covid and didn't know it. She might have tested negative because she had the first shot already. She wasn't worried about exposing others and went to get her second shot, but she got really sick after that and it kicked her butt pretty hard. The outcome of the events for leg pain and tingling on skin was not recovered, fever, body aches and headache recovered in Apr2021, chills on 11Apr2021, muscle hurt/pain on 12Apr2011, joint pain was recovering, and outcome of other events was unknown."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1278543-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278714-1	A little over a week after the First vaccination, I developed two blood clots in my leg; This is a spontaneous report from a contactable consumer (patient). A 67-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: unknown), dose 1 via an unspecified route of administration, administered in the left arm on 18Mar2021 as SINGLE DOSE, for covid-19 immunization (at the age of 67 years-old). The patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: unknown), dose 2 via an unspecified route of administration, administered in the left arm on 14Apr2021 as SINGLE DOSE. The patient was not pregnant at the time of vaccination. The patient did not receive any other vaccine within 4 weeks prior to the vaccine. Medical history included peripheral arterial occlusive disease from an unknown date and unknown if ongoing and bad circulation. There were no known allergies. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient did not receive any concomitant medications. The patient developed two blood clots in her leg on 28Mar2021 a little over a week after the first vaccination. There was no treatment provided for the blood clots. The outcome of the blood clots was not recovered. It was also reported that since the vaccination, the patient had not been tested for COVID-19. No follow-up attempts are possible; information about lot/batch number cannot been obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278723-1	three blood clots in right leg; This is a spontaneous report from a contactable consumer. A 45-year-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: ER2613, Expiry date not reported), via an unspecified route of administration, administered in the left arm on 25Mar2021 15:00 (age at vaccination was 45 years) as single dose for COVID-19 immunization. Medical history included drug hypersensitivity (sulfonamide allergy) from an unknown date and unknown if ongoing, and seasonal allergy from an unknown date and unknown if ongoing. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient was vaccinated at a clinic. The patient was not diagnosed with COVID-19 prior to vaccination. Since the vaccination, the patient has not been tested for COVID-19. Concomitant medications included fluticasone propionate taken for an unspecified indication, start and stop date were not reported; olopatadine hydrochloride (PATADAY) taken for seasonal allergy, start and stop date were not reported. The patient experienced three blood clots in right leg on 09Apr2021 20:00 with outcome of recovering. Therapeutic measures were taken as a result of three blood clots in right leg (thrombosis) that included blood thinners (Eliquis). The event was reported as serious, medically significant. No follow-up attempts are possible. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278730-1	big bump on arm; rash/red blotch on arm; nose bleed, blood clots coming out of nose; nose bleed, blood clots coming out of nose; This is a spontaneous report from a contactable consumer (patient). A 52-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), intramuscular, administered in the left arm on 05Apr2021 (Batch/Lot Number: EW0151) as single dose, for covid-19 immunization. Medical history included ongoing sjogren's syndrome and ongoing lupus. Concomitant medications included levothyroxine sodium (SYNTHROID) taken for an unspecified indication, start and stop date were not reported. 07Apr2021, she started having a nose bleed. She explains she wasn't doing anything in particular, just talking on the phone, and blood just started pouring out of her nose. Blood clots started coming out as well. She ended up calling 911 and they sent an ambulance. The EMS helped her stop the bleeding. She didn't go to the ER or anything. However, she is getting implants done for her upper teeth so she wanted to double check and make sure nothing was pressing against her nasal cavity because the nose bleeds were so off/random. The doctor said there was no pushing. The doctor completed a panoramic exam/x-ray and confirmed there was nothing pushing. The doctor did suggest she go to the ENT. She made an appointment with an ENT doctor for the following day, 08Apr2021, and the ENT put a camera down. She confirms she explained to them she just had the vaccine and she was googling information. She was told to not google anything and to let them check it out. The caller states she explained to the ENT the reason she was there was because she was concerned, she is not a nose bleeder and nose bleeds never happened to her. Once blood clots started coming out that way, she was terrified. They didn't see anything alarming, but they wanted to make sure and do further research, so basically they planned to do a CT/CAT scan. She was scheduled for a CT/CAT scan on Monday, 12Apr2021, but she was allergic to the contrast and was sent home because they were not prepared for pre-medication. She is waiting for the doctor to schedule another appointment so she can go back. She felt she needed to report these nose bleeds because she has been seeing Johnson & Johnson had severe blood clots, but she took Pfizer's vaccine, and was also getting blood clots. Her nose bleeds just happened right after both vaccines. On an unspecified date, she had a little bump that went away. She had this big bump on her arm and a big rash developed. At first, the bump was the size of a quarter and then grew double in size. The bump finally went down, but the rash she can still see. The rash hasn't grown, it's just there- there is just a red blotch on her arm. She has not had another nose bleed since the one that occurred on 07Apr2021. She just noticed when she had her first dose on 15Mar2021, the nose bleed happened within that week also. The outcome of events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278778-1	calf feels tight, stiff, it feels like dead weight/started within the last 2- 3 weeks; calf feels tight,; pain running up the right arm up to the shoulder; pain running up the right arm up to the shoulder/his calf pain, started to feel tight, started within the last 2- 3 weeks; He asks if someone can tell him if it is a blood clot in the back of his calf; his eyes also hurt a little bit; This is a spontaneous report from a contactable consumer (patient). A 46-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), second dose at the age of 46-years-old via an unspecified route of administration, administered in arm left on 02Apr2021 (Batch/Lot Number: ER8734) as single dose for covid-19 immunisation. There was no medical history reported. There were no concomitant medications. The patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 12Mar2021 (lot number: EN6202) at the age of 46-years-old for covid-19 immunisation and experienced If it is a blood clot in the back of his calf, Regular pain feeling, Headache, Muscle pain, Calf feels tight, stiff, it feels like dead weight, Pain running up the right arm up to the shoulder and Calf pain, started to feel tight. The patient reported that he has been feeling the regular pain feeling, a headache, muscle pain before, it comes and goes on 12Mar2021. However, he says that the last few days there had been, by his calf muscle in the back there, they've been tightening, like stiffening up on him when he's sleeping on an unspecified date. It wakes him up at times. He also reports pain running up the right arm up to the shoulder on an unspecified date. He says he is just curious in terms of what that might be. He asks if someone can tell him if it is a blood clot in the back of his calf on an unspecified date. The headache went away after 2-3 days. He says his eyes also hurt a little bit on 12Apr2021 and was the same time frame as the headache. He says those things were just mild, nothing major. He did not take anything for it. He confirmed he no longer has muscle pain. He just has pain behind the calf and right arm up to his shoulder. He says his calf pain, started to feel tight, started within the last 2- 3 weeks. The arm pain in right arm up to shoulder started maybe within the last 2 days. The outcome of the events was unknown. Follow-up attempts completed. No further information expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278908-1	A Blood clot was found; Cellulitis; Migraine started within 24 hours; left calf pain/Leg increased in swelling and burning pain; severe swelling and burning sensation; severe swelling and burning sensation; Leg increased in swelling and burning pain; Phlebitis; This is a spontaneous report from a contactable consumer (patient). A 50-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: EN6199/EN5318), via an unspecified route of administration administered in left arm at doctor's office/urgent care facility on 12Mar2021 15:00 as SINGLE DOSE for COVID-19 immunisation. The patient was not pregnant at the time of vaccination. The patient was not COVID tested post vaccination. The patient had no other vaccines in four weeks. Medical history included high blood pressure and sleep apnea. Concomitant medications included amoxicillin, valsartan, amlodipine, and vitamin d [vitamin d nos]. The patient previously took doxycycline and experienced allergies. Historical vaccine includes first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) on an unspecified date for COVID-19 immunisation. On 05Apr2021 at 03:00 PM, the patient experienced a blood clot was found, cellulitis, migraine started within 24 hours, left calf pain/leg increased in swelling and burning pain, severe swelling and burning sensation, leg increased in swelling and burning pain, and phlebitis. It was further reported that migraine started within 24 hours. 48 hour left calf pain that got increasingly worse leading to a severe swelling and burning sensation. The patient went to right time urgent care on Friday, 09Apr2021. The patient was diagnosed with cellulitis. It was also reported that pain increased swelling spread and went to emergency room (ER) on 10Apr2021 for ultrasound of leg. No clots and was advised if worsened to come back. The patient's leg increased in swelling and burning pain. The patient went to urgent care patient first on 11Apr2021 and then the doctor on Monday, 12Apr2021 where she was sent for a STAT ultrasound due to major pain and swelling. A blood clot was found. Phlebitis was diagnosis. The patient was sent to a vascular surgeon on 13Apr2021 and vein specialist on 15Apr2021. The patient was given Xarelto blood thinners and started taking them on 15Apr2021. All of this began 48 hours after taking the second vaccination (pending clarification) and has the patient worried. The events resulted to doctor or other healthcare professional office/clinic visit and emergency room/department or urgent care. Therapeutic measures were taken as a result of all events. The patient had not yet recovered from all events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278972-1	"find out if his side effects are normal or if he maybe has a ""blood clot.""; the shoulder and neck on the side where he received his injection are swollen and painful; the shoulder and neck on the side where he received his injection are swollen and painful; the shoulder and neck on the side where he received his injection are swollen and painful; the shoulder and neck on the side where he received his injection are swollen and painful; across his collar bone is painful; my lips, nodes and top of my right my trap area is swollen; my lips, nodes and top of my right my trap area is swollen; I was told you might swollen on arm where shot went it or under your armpit not on top on my neck; This is a spontaneous report from a contactable consumer (patient). A male patient of an unspecified age received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 17Apr2021 (lot number and expiry date not reported) as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. Historical vaccine included first dose of bnt162b2 on an unspecified date for COVID-19 immunization. The patient stated that he received his second dose of the Pfizer COVID-19 vaccine on Saturday 17Apr2021. He stated that he called Pfizer yesterday, Sunday 18Apr2021 and reported his side effects, but was not able to have his questions answered. He was calling back to find out if his side effects were normal or if he maybe had a ""blood clot."" The patient stated that the shoulder and neck on the side where he received his injection were swollen and painful. The patient stated that across his collar bone was painful. He stated that the swelling and pain had improved slightly since yesterday. He stated that other than these side effects he feels great. He added, ""I just got my second shot yesterday. No symptoms showed up except my lips nodes and top of my, right my trap area is swollen. So I want to make sure, is that normal? Because I was told you might swollen on arm where shot went it or under your armpit not on top on my neck."" Clinical outcome of blood clot was unknown, while for the other events was recovering. Information on the lot/batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1279039-1	Large blood Clotting after each dose during menstral cycle.Bled for more than 2 weeks.; Large blood Clotting after each dose during menstral cycle.Bled for more than 2 weeks.; This is a spontaneous report from a contactable consumer (patient). A 51-year-old female patient (not pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), second dose via an unspecified route of administration, administered in Arm Right on 19Mar2021 04:15 (at 51 years old, not pregnant), single dose for covid-19 immunisation. Medical history included stroke, high blood pressure, diabetes mellitus, all from an unknown date and unknown if ongoing. The patient's concomitant medications was unspecified (mentioned other medications in two weeks: yes but prescribed). The patient had her first dose of BNT162B2 vaccine on 26Feb2021 04:15 at 51 years old, for COVID-19 immunisation on left arm. The patient experienced large blood Clotting after each dose during menstral cycle. Bled for more than 2 weeks. All on 27Mar2021 12:00. No other vaccine in four weeks. No covid prior vaccination. Not covid tested post vaccination. No known allergies. No treatment for the events. The outcome of the events was unknown. Information about lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1279063-1	Thrombocytopenia; He had clot in the legs and in the lungs; He had clot in the legs and in the lungs; Low platelet count; This is a spontaneous report from a contactable Other-HCP. A 67-years-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation (Age at vaccination: 67 years). Medical history included diabetes. Concomitant medication(s) included atorvastatin (LIPITOR [ATORVASTATIN]); colecalciferol (VITAMIN D [COLECALCIFEROL]); lisinopril; insulin taken for diabetes mellitus. The patient experienced thrombocytopenia, he had clot in the legs and in the lungs and have a low platelet count on an unspecified date. The patient was hospitalized for the events on an unspecified date. The patient underwent lab tests and procedures which included CBC: unknown result on 08Apr2021, Heparin associate antibody: unknown result, comprehensive metabolic panel (CMP): unknown result and platelet count: low on an unspecified date. Therapeutic measures were taken as a result of thrombosis included Heparin drip and Argatroban. Outcome of the events was unknown. Information about lot/batch number has been requested.; Sender's Comments: The reported events are considered unrelated to BNT162B2 vaccine, being rather intercurrent occurrences. Clots in the legs and in the lungs were likely favored by the mentioned diabetes and by a possible hyperlipidemia (the patient was taking atorvastatin) in a setting of possible arterial hypertension (the patient was taking lisinopril). The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1279190-1	two blood clots in his leg; This is a spontaneous report from a contactable consumer reporting on behalf of the patient. A 57-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), first dose on 08Feb2021 (Lot Number: EM9810, unknown expiration) and second dose on 02Mar2021 (Lot Number: ENU202, unknown expiration; pending clarification), both received at the age of 57 years old via unspecified route of administration as a single dose for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The reporter called to report that the patient had a Pfizer vaccine, received second COVID vaccine on 02Mar2021, and couples of weeks ago, a week longer that the patient was diagnosed with two blood clots in his leg. The patient got both of COVID-19 vaccine. Outcome of the event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1279205-1	I got a blood clot in my varicose vein in my left leg; This is a spontaneous report from a contactable consumer(patient). A patient of unspecified age and gender received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number was not reported), on an unspecified date at single dose for covid-19 immunisation. The patient medical history included varicose vein in the left leg. Concomitant medications were not reported. On an unspecified date in Apr2021, the patient developed a blood clot in the varicose vein in the left leg. Patient clarified that after talking to the doctor patient thought may had the adverse reaction to the first vaccine. Patient was supposed to get Pfizer Vaccine at the day of reporting but when talk to doctor and the urgent care where was supposed to be getting it, rescheduled it for 04May2021 because 3 days before (in Apr2021) patient got a blood clot in varicose vein in left leg so, they didn't want to go and get the vaccine and rescheduled it for 04May2021. Patient asked if he/she should even get the second vaccine.The final outcome of the event was unknown. Information on the lot/batch number has been requested
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1279276-1	"big knot there and now its black and blue and it's also like ball/could be a blood clot on her hand; Headache; sickness; felt like little vomit; not having a very good appetite; feeling like sinusy; feeling little nauseas; bruise on the top of her hand and then that bruise move to left part of her upper hand; This is a spontaneous report from a contactable consumer (patient) reported that a 53-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: ER8727), via an unspecified route of administration on 08Apr2021 (at the age of 53-years-old) as a single dose for covid-19 immunisation. The patient medical history included cardiovascular disease. She has had open heart surgery, a double bypass when she was 44. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 18Mar2021 (at the age of 53-years-old) for covid-19 immunisation, and experienced face got swollen and got red and she think she got wet nose right there swelling a little bit same time her face swollen. Additional medical history included anxiety and depression. Concomitant medications included acetylsalicylic acid, ascorbic acid (ASPIRIN [ACETYLSALICYLIC ACID;ASCORBIC ACID]; atorvastatin; midodrine; metoprolol; clopidogrel; ranolazine; meclozine hydrochloride (MECLIZINE HCL); hydrocortisone; glyceryl trinitrate (NITROGLYCERIN); zinc sulfate (ZINC SULPHATE); gabapentin; colecalciferol (VITAMIN D [COLECALCIFEROL]); and paracetamol (ACETAMINOPHEN); all were taken for an unspecified indication, start and stop date were not reported. Additional concomitant medications included duloxetine taken for depression and anxiety; and alprazolam (XANAX) taken for anxiety; both start and stop date were not reported. On 13Apr2021 5 days after the second vaccination, the patient experienced headache every single day with sickness and felt like little vomit and not having a very good appetite at all. She was feeling ""sinusy"" but she don't have any sinus issues and then she was feeling little nauseas and then her right hand for some reason she got bruise on the top of her hand and then that bruise move to left part of her upper hand and gotten a big knot there and now its black and blue and it's also like ball; her husband and she think that could be a blood clot on her hand. The outcome of the events was unknown."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1281374-1	Woke up in the middle of the night dizzy on the evening of 3/11/21,started vomiting blood, syncope, when woke up had altered mental state, when reoriented he was able to call EMS, taken to hospital where he was diagnosed with an abdominal clot size of a grapefruit
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1281770-1	PT PRESENTED TO ER WITH COMPLAINTS OF CHEST PAIN. REPORTED RECEIVING COVID VACCINE 5 AND 2 EEKS PRIOR. IT WAS NOTED SHE HAD ELEVATED TROPONIN LEVELS, NSTEMI. PT TRANSFERRED TO ANOTHER MEDICAL CENTER FOR HIGHER LEVEL OF CARE/CARDIOLOGY. HEART CHAT PERFORMED. THE DISTAL ONE FOURTH OF THE LAD WAS OCCLUDED AND APPEARED TO BE FROM THROMBUS. OTHER FINDINGS ALSO NOTED . PT DID NOT REPORT PRIOR HISTORY OF CORONAY HEART DISEASE OR MI.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1281855-1	Blood clotting in the left leg, Use ultrasound to identify and confirm clotting locations in the left calf leg. Cramp like pain in the left calf with blue-ish/purple color in the ankle and left foot. Foot and leg was cold to the touch with extraordinarily strong migraines. Person leads healthy lifestyle and is physically and athletically fit. I was given Gemtesa to treat the clots, 150mg x2 a day for 7 days then to 75mgx2 a day for an unknown period. First dose on 3/24/ 2021 Pfizer Lot#EN6205 Second Dose ON 04/14/2021 LOT#EW0161 10:30AM.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1281892-1	Blood clotting in the left leg, Use ultrasound to identify and confirm clotting locations in the left calf leg. Cramp like pain in the left calf with blue-ish/purple color in the ankle and left foot. Foot and leg was cold to the touch with extraordinarily strong migraines. Person leads healthy lifestyle and is physically and athletically fit. I was given Gemtesa to treat the clots, 150mg x2 a day for 7 days then to 75mgx2 a day for an unknown period. First dose on 3/24/ 2021 Pfizer Lot#EN6205 Second Dose ON 04/14/2021 LOT#EW0161 10:30AM
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1282796-1	Very heavy menstrual flow during the first cycle after vaccine, noticed small clots. This is a first time ever for me. Also, longer menses than usual by a few days.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1283034-1	Headache stomach issues and arm pain Thursday chest pain and shortness of breathe went to the er had blood clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1283043-1	On April 12, shortly after I got my second Pfizer shot, I started my period. It was very early by several weeks, and only a week since my previous period ended, which is why I made note of it and have kept a record. For the first week it was fairly light and mostly spotting. After a week, it progressed and became more like a normal period in terms of the amount of blood I was seeing. By April 23rd I started noticing a lot more blood clots in my period than usual--ordinarily I'll see one or two each time I get my period, but there were about five days in a row when I noticed multiple blood clots. The clots stopped around April 28th, but as of today (May 3rd) my period is still ongoing, making it exactly three weeks long, and more unless it suddenly stops today. For the most part the blood has been red and I haven't noticed any other unusual discharge. I have noticed the past few days that the blood is more brown, like old blood. I have not experienced bad cramps or stomach upset during this very long period--I had the same level of menstrual cramping/stomach upset that I am accustomed to over the first couple of days, but nothing since then. I have felt bloated and tired the entire time, however. I have also experienced a really low mood for these few weeks, though I can't be certain if this is worse than I usually experience on my period, because it may just be because it's lasting so long that it feels worse.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1283062-1	heavy blood in stool and small blood clot passed with stool. blood with stool lasted for approximately one week.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284761-1	blood clot at upper right chest underneath collar bone; Swollen right arm and fingers; tingling and numbness in arm; numbness in arm; headaches; This is a spontaneous report from a contactable consumer (patient). A 56-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) dose 1 via an unspecified route of administration, administered in Arm Left on 08Apr2021 04:00 (Lot Number: EWO153) as SINGLE DOSE for covid-19 immunisation. Medical history included Implantable cardioverter defibrillator insertion from 2005. Concomitant medication in two weeks included verapamil 120 mg. No other vaccine in four weeks. No Covid prior vaccination. No Covid tested post vaccination. No Known allergies. On 12Apr2021, the patient experienced blood clot at upper right chest underneath collar bone. Swollen right arm and fingers with tingling and numbness in arm. Also, suffering from occasionally headaches. The patient was relatively heathy with no know blood clot issues. Events resulted in Doctor or other healthcare professional office or clinic visit. Unknown if treatment was received for events. The outcome of events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284805-1	more than usual vaginal bleeding/with very dark blood with clots.; menstrual cramps; sharp pelvic pain; more than usual vaginal bleeding; more than usual vaginal bleeding; This is a spontaneous report from a contactable consumer (patient). A 40-year-old female non-pregnant patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 05Apr2021 10:30 (Batch/Lot Number: ER8734) as single dose for covid-19 immunisation. Facility type Vaccine: Other. Medical history included known allergies: Sulfa and Folate deficiency. The patient's concomitant medications were not reported. The patient previously received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot number: EN6206) on 15Mar2021 10:30 AM in left arm for covid-19 immunisation and experienced vaginal bleeding/blood clots at mid-cycle on 15Mar2021 08:00 PM. If other vaccine in four weeks: No. Other medications in two weeks: No. If Covid prior vaccination: No. If Covid tested post vaccination: No. The patient experienced more than usual vaginal bleeding/with very dark blood with clots, menstrual cramps, sharp pelvic pain, all on Apr2021 with outcome of not recovered. Clinical course: After the first dose, the patient had vaginal bleeding/blood clots at mid-cycle. The same day after the 2nd dose, she started feeling menstrual cramps and sharp pelvic pain. After taking paracetamol (TYLENOL) and feeling a bit of relief, she stood up and started having more than usual vaginal bleeding at the level that it reached the floor as if it was a hemorrhage with very dark blood with clots. This lasted for 1 minute. This lasted for 5 days and 7 days later, she started bleeding again with dark blood clots for a period of 3 days. The events resulted in: Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. No treatment received for events more than usual vaginal bleeding/with very dark blood with clots.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284832-1	Big Blood Clot in left leg; This is a spontaneous report from a non-contactable consumer (Patient). A 80-years-old female patient (Non-pregnant) received bnt162b2 (Pfizer-Biontech COVID-19 vaccine, Solution for injection), via an unspecified route of administration on 28Mar2021 11:00 as single dose for COVID-19 immunisation. The patient medical history included allergies to Demerol. The patient's concomitant medications were not reported. On 17Apr2021 21:30 patient experienced big blood clot in left leg. Patient was hospitalized for 2 days. The patient underwent lab tests and procedures which included Nasal swab test results negative on 16Apr2021. Treatment received for the adverse event was blood thinners. The outcome of event was Not recovered. No follow-up attempts are possible; information about lot/batch number cannot been obtained
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284856-1	5-6 Blood clots, blood clots were present; heart attack; This is a spontaneous report from a contactable consumer reporting for himself. A 54-year-old male patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 29Mar2021 12:00 on left arm at single dose for COVID-19 immunization. Facility type vaccine was at Pharmacy or Drug Store. Medical history included high blood pressure, high cholesterol. Concomitant medications included atorvastatin (LIPITOR), amlodipine besilate (NORVASC). Historical Vaccine included first dose of BNT162B2 on 08Mar2021 12:00 on left arm for COVID-19 immunization. The patient experienced 5-6 Blood clots, had heart attack and blood clots were present on 14Apr2021 13:00. The events were resulted in Emergency Room Visit, Hospitalization, Life threatening illness (immediate risk of death from the event). Treatment was received for the events included Stent. The outcome of the events were resolving. Information on Lot/Batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284864-1	blood clot in her leg; blood clot in her leg; Cardiac arrest; Heart attack; This is a spontaneous report from a contactable consumer (patient's daughter). A 96-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 2 via an unspecified route of administration on 29Jan2021 (Lot Number: EL9265) as SINGLE DOSE for COVID-19 immunization. Medical history included blood clots in her legs from an unknown date and unknown if ongoing , diabetec, bone infection, surgery and anemic; all from an unknown date and unknown if ongoing; and a family history of gangrene from an unknown date and unknown if ongoing of her mother. Concomitant medications included apixaban (ELIQUIS) taken as blood thinner and furosemide (FUROSEMIDE) taken for an unspecified indication; both start and stop date were not reported. The patient previously had BNT162B2 (Lot Number: EL1283) dose 1 on 08Jan2021 for COVID-19 immunization. The facility where the most recent COVID-19 vaccine was administered was in the military facility. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. On 15Mar2021, the patient died due to a heart attack and cardiac arrest. The patient developed a blood clot in her leg and had to have her leg taken off. The date of surgery was 05Feb2021, not early Mar2021 like was originally stated at the hospital. The patient was admitted either on 01Feb2021 or 02Feb2021 and discharged on 23Feb2021. They had a bunch of bad weather and then they put the patient in the nursing facility for about a month and when she was brought home she had a heart attack and died due to cardiac arrest. She doesn't know about the blood clot and this being related to the COVID vaccines but the patient had surgery a week after she had her second shot. The heart attack was on 15Mar2021. The patient had a history of blood clots in her legs before and she had problems with that so that might of made it worse but she doesn't know for sure. The patient died on 15Mar2021. An autopsy was not performed. The outcome of the event blood clot in her leg was unknown. No follow-up attempts are possible; information about lot/batch number has been obtained.; Reported Cause(s) of Death: heart attack; cardiac arrest
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1284975-1	My mother Received vaccine on 3/18/21. Approximately the week of 4/4/21 she started to experience shortness of breath and got progressively worse. On 4/9/2021 she was admitted to hospital and diagnosed with extensive clot burden within the interlobal pulmonary arteries bilaterally as well as upper lobar branches. Venous Doppler was a performed on 4/10/21 blood clots found in both lower legs Please note that I attempted several times to report this to Pfizer, I was disconnected once and was on hold for over hour with the same automated message replaying that there were 6 callers ahead of me. There was no option to leave a message for a return call to share what I feel is important safety information.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1285007-1	Diagnosed with CML few months ago and taking Dasatinib daily with great response and normalized WBCs. Had second dose of Pfizer vaccine 2/2. Blood work on 2-5 WBCs 150K and 2/6 bilateral blood clots in legs which then became pulmonary blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1285475-1	Blood clots, organs shut down, death.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1285649-1	ON 4/28 felt intense pain in legs and blurry eyes, slowly went away after taking advil, came back at night, on thursday 4/29 pain continued and had shortness of breath decided to go to urgent care made appt for 4/30 813AM, at Urgent Care symptoms pointed to blood clots was told to go to local imaging company to get sonogram, sonogram showed blood clots in leg told to go to emergency room, went to Hospital and was admitted, confirmed blood clots and put on blood thinner neprin IV, released 5/2 at 1PM, at home now on blood thinner pills 10mg Eliquis

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1285758-1	Menstrual cycle heavy blood clots - my next cycle after the first dose of covid vaccine which I received on 4/14, my period flow was comprised of 2 inch blood clots which I passed for an hour on 4/27. I haven?t passed period blood clots this large since I was 12 years old so I wanted to report it since I happened to get these unusual blood clots the cycle after I got the first dose. I am getting the second dose on 5/5 so I will see if the same results take place. I have reported this to my gynecologist but am waiting on her response.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1285922-1	Pulmonary embolism also blood clot in leg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1286071-1	Pt reported painful blue-purple lump on palmar surface of R middle finger at crease of the PIP joint, onset 2-3 days after first dose of the Pfizer vaccine, diagnosed by this provider as a palmar digital vein thrombosis. No other precipitating factors such as trauma or exposure to intense vibrations to the finger or other causative factors could be identified. Pt without any history of clotting disorders. Pt healthy, not taking any medications. Pt reported that symptoms were improving at the time of her clinic presentation on 4/13/21. Of note, pt reported her husband, who received the vaccine the same day, had very similar symptoms on the palmar surface of his 5th finger onset a few days after this patient.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1286437-1	Pfizer-BioNTech COVID-19 Vaccine EUA: nine and eleven days after vaccination during clinic visit patient reports left axillary swelling radiating down left arm. Subsequently developed numbness and tingling to left hand/fingers seven days after vaccination. Arthralgias, joint swelling, mylagias, and weakness also noted. After first visit patient prescribed methylprednisolone. Two days later patient returned to clinic with improvement in swelling but reports tremors and three new bumps on arm, palpitations, and weakness/numbness of 4th and 5th digits on left hand proximally. Patient referred to neurology and physical therapy with diagnosis of ulnar neuropathy.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1286446-1	Bilateral pulmonary clots, bilateral lower extremity clots, right heart strain, atrial fibrillation, right atrial clot, heart attack
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1286717-1	Blood clot causing the need for emergency surgery
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1286817-1	Received first dose on 1/4/21 and second dose on 1/25/21. 4/28/21 reported having leg pain and was seen by primary care where it was determined he had a blood clot and was placed on blood thinners.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1287018-1	The next day after I received the Pfizer vaccine I started feeling sick and lightheaded it had a headache. I think nothing of it and then my menstrual cycle started early and it started off being heavier than usual. Then I notice when I went to wipe myself or use the bathroom blood clots was falling out. Being that I work in the medical field I know like if is not bigger than 1/4 there's nothing to really worry about. So as the day went on and the following days I noticed that the blood clots kept on getting bigger and bigger and coming faster and faster. At this point I started getting concerned, that's when I decided to go to the hospital. At the emergency room at Hospital they really didn't do nothing for me but gave me an ultrasound and gave me some birth control told me to follow up with my OBGYN. But I already made an appointment with OBGYN cuz I just feel like if something was going on and I can't get in to see them till June 30th. They did give me some birth control and tell me your help with the bleeding and for me to take some ferrous sulfate. They said they'd never heard of this reaction before. So I went on May 1st to get my second vaccine. Before I received it, I talked to doctor there and they told me it would be fine for me to get it. Now my symptoms and reactions are starting all over. I'm so concerned about what's going on with my body. I just need somebody to explain to me what's going on. I cannot afford another hospital ER bill again.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1287041-1	Blood Clot Deep Vein Thrombosis
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1287662-1	A few days after the vaccine started feeling uncomfortable in the left leg. It did not get better with rest and time so went to the ER and an ultrasound and found a clot from the groin to the ankle. He had no injury to preclude this clot. He was put on an anticoagulant (Xarelto) and standing and walking is very painful. He has a follow up appointment set with his PCP 5/6/21 He was told he might have to be on this medication for months maybe longer.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1288353-1	"may be blood clot; Something happened with my knee and it has just got worse; I was thinking maybe it is a pulled muscle or something but it is very bad now; This is a spontaneous report received from a contactable consumer (patient). An 83-year-old female patient (5 feet 1 inches, about 120 pounds) received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot number: EN6198, patient was not sure) via an unspecified route of administration at left arm on 22Feb2021 (83-year-old at time of vaccination), at single dose; the second dose of BNT162B2 (lot number: unknown) via an unspecified route of administration on 15Mar2021 (83-year-old at time of vaccination), at single dose, for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. Patient stated she has got both the shots of Pfizer (Covid-19 vaccine) and at the end of the first one or it is like eighteen days, just before the second shot, something happened with her knee and it has just got worse (from 12Mar2021). She did not connect that with the Pfizer shot and I was thinking maybe it is a pulled muscle or something but it is very bad now. She would like to know what she can do. Also, she is thinking may be blood clot at this point and she would like to have it checked. She thinks that she should be able to contact somebody to actually get some physical test on this. She tried heat and cold alternate and Absorbine Jr. and Aspirin. It varied with the days, she thinks she has taken Aspirin about twice a day and it is a tablet and it was 325 mg and she took two of those, orally. Therapeutic measures were taken as a result of the event and included ""heat and cold alternate and Absorbine Jr. and Aspirin"". The outcome of the event was not recovered. Information on the lot/batch number has been requested."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1288355-1	"Blood clot in lungs and leg; Trouble breathing/Shortness of breath, but not as bad; didn't feel right; Chest tightness; Slightly ill; Sore arm, in left arm where the shot was; Slightly tired; This is a spontaneous report from a contactable consumer. An 81-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in the left arm on 10Feb2021 (Batch/Lot Number: EN6201) as SINGLE DOSE for covid-19 immunization. Medical history included Type 2 Diabetes from 2004 and ongoing, High Blood Pressure from 2000 and ongoing (reporter stated she was 81; it's not very high, usually when she goes to the doctor). Concomitant medications not specified but reporter mentioned that everything she has been on she has been on long time; she takes blood pressure medication because sometimes it is high. She has not had started any new medication prior to starting vaccines or during both vaccines. The patient previously received flu shot (September). The patient received her first vaccine dose of bnt162b2 (Lot # EL9262, Expiry date UNKNOWN), on 20Jan2021, at 1:30-1:45, in the left arm and experienced sore arm and slightly tired. The reporter stated that after her second vaccine on 10Feb2021, she had a sore arm, (left, where shot was) and was slightly tired from 10Feb2021 until 12Feb2021. She confirmed she noticed this started the same day, 10Feb2021. On 31Mar2021, she felt slightly ill and had some slightly trouble breathing. She saw her doctor who did some tests. The next day 01Apr2021, her tests came back and the doctor told her to go to the hospital. While in the emergency room she had a CT of the lungs that showed blood clots in both lungs. She was admitted to the hospital where on 02Apr2021 they did a ultrasound of her legs. She was told she had blood clots in her right leg. She stated she had no symptoms of blood clots in her legs like redness, sore, swelling, or hot. She was asked if she had fallen suddenly, had an accident, or was sitting for a long time, traveled a long distances. On 02Apr2021 she had a Echocardiogram and was told her heart was ok. Caller stated she was given Lovenox shots in the hospital. She did not have Lot # or Expiry date on the Lovenox. She stated on 02Apr2021 after having two Lovenox shots that day, she met with a Vascular Surgeon who started her on Eliquis that evening. She stated she started taking Eliquis that evening. She stated she took 2 tablets, twice a day, for a week. Now she takes one tablet, twice a day, ongoing. The Eliquis tablet says 5mg. She stated she was discharged on 03Apr2021, that evening. Patient went back to the emergency room on 10Apr2021 after speaking with a nurse at (university name withheld) because she had tightness in the chest but the shortness of breath, her breathing was not as bad. She just didn't felt right. She stated she was not admitted. Patient stated she saw a NP (name withheld) with (university name withheld) Pulmonary the day of 31Mar2021. When discussing the tests performed that day, Chest X-ray, D-Dimer, and Pro INT, CH asked caller the results from them. She did not provide results, just stated she had looked up her results in ""EPIC mychart"" while getting the result information from the office. Where she was instructed to go to the hospital. She went to the Clinic of the Vascular Surgeon at (Hospital withheld) on Wednesday 14Apr2021 and was feeling better. Her second appointment with them will be in July. Where they will do another ultrasound of her legs and talk about the medication. She has an appointment with her primary on 19Apr2021. The outcome of events Trouble breathing (recovered on 14Apr2021), Shortness of breath, but not as bad (recovered on 12Apr2021), Sore arm, in left arm where the shot was (recovered on 12Feb2021), Slightly tired (recovered on 12Feb2021) Chest tightness (recovered on 12Apr2021), and didn't feel right was recovered. The outcome of event Blood clot in lungs and leg was unknown."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288373-1	vaginal bleeding/blood clots at mid-cycle; vaginal bleeding/blood clots at mid-cycle; vaginal bleeding/blood clots at mid-cycle; This is a spontaneous report from a contactable consumer. A 40-year-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) lot number: EN6206, via an unspecified route of administration, administered in left arm on 15Mar2021 10:30 as single dose for Covid-19 immunisation. Medical history included folate deficiency and allergies to sulfa. There were no concomitant medications. After the first dose on 15Mar2021, the patient had vaginal bleeding/blood clots at mid-cycle. No treatment information was provided. The outcome of the events was not recovered.; Sender's Comments: Based on temporal association reported events causal relationship with BNT162B2 cannot be excluded. Case to be re-assessed upon receipt of new information. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288388-1	Developed two blood clots in my left calf; This is a spontaneous report from a contactable consumer, the patient. An 18-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Lot unknown, first dose) solution for injection intramuscular in the right arm on 05Apr2021 at 10:00 (at the age of 18-years-old) as a single dose for COVID-19 vaccination. Medical history included ADD (attention deficit hyperactivity disorder). Concomitant medication included mirtazapine (AVANZA). The patient had no known allergies. The patient did not receive any other vaccine within 4 weeks prior to the vaccine. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient developed two blood clots in my left calf on 13Apr2021 at 14:00 which resulted in emergency room/department or urgent care visit. Treatment for the event blood clots included rivaroxaban (XARELTO). The outcome of the event blood clots was recovering. The patient was not tested for Covid post vaccination. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288390-1	"passing blood clots/having a total of 4 of them she has passed/noticing one when she went to the bathroom to urinate; she went into the bathroom and wiped, without actually going to the bathroom, she noticed the blood clot(s); felt like she needed to go to the bathroom/felt weird like something needed to come out; headache; dizziness; hot and cold sweats; This is a spontaneous report from a contactable consumer (who is also the patient). A female patient of an unspecified age received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiry date not reported), via an unspecified route of administration, on 16Apr2021, as single dose, for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. In Apr2021, after the first vaccination, the patient experienced ""some of the usual side effects"" described further as headache, dizziness, and hot and cold sweats. On 18Apr2021, 2 days after the first vaccination, the patient experienced passing blood clots having a total of 4 of them she has passed. She reported noticing one when she went to the bathroom to urinate. For the other blood clots, she reported it felt like she needed to go to the bathroom further described as ""felt weird like something needed to come out."" When she went into the bathroom and wiped, without actually going to the bathroom, she noticed the blood clot(s). The outcome of the events was unknown. No follow-up attempts are needed; information about lot/batch number cannot be obtained."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288411-1	sustain cramps in her right leg; limited range of motion; not sure if there was a nerve that has been damaged or early stage of thrombosis; not sure if there was a nerve that has been damaged or early stage of thrombosis; severe arm pain; This is a spontaneous report from a contactable consumer reporting for herself. A 56-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE) (lot number/expiration date: not provided), via an unspecified route of administration, on 12Apr2021 at 09:30 (at the age of 56 years old) as a single dose in the right arm for COVID-19 IMMUNIZATION. Relevant medical history and concomitant medication was none. The patient did not have any known allergies. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient was not pregnant at the time of vaccination. On 18Apr2021 at 18:00, the patient experienced severe arm pain that was described as arm pain was too painful and have been woken up out of her sleep and she has limited range of motion and she was not sure if there was a nerve that has been damaged or early stage of thrombosis. On 20Apr2021, the patient experienced sustain cramps in her right leg. The patient did not receive treatment for the events. The outcome of the events was not recovered. Since the vaccination, the patient had not been tested for COVID-19. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow-up.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288420-1	clots; I received the pfizer vaccine and within 2 weeks had bruising; low platelets; This is a spontaneous report from a contactable consumer, the patient. This patient of unspecified age and gender received BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: UNKNOWN) via unspecified route on unspecified date as a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient reported that he/she received the vaccine and within 2 weeks had bruising, low platelets and clots (unspecified date). Lab data included platelets: low on unspecified date. The clinical outcomes of bruising, low platelets and clots were unknown. The lot number for the vaccine, BNT162b2, was not provided and will be requested during follow-up.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				BNT162B2, was not provided and will be requested during follow up.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288423-1	"Blood clots in her nose; Nose bleed; her arms are both numb on her; Couldn't move them; She feels lethargic; This is a spontaneous report from a contactable consumer (patient herself) via a medical information team. A female patient of an unspecified age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection), via an unspecified route of administration on an unspecified date as a single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. It was reported that, enquired have people died from the vaccine or not. Enquired what do you know about the news in the media about reports of death in nursing home elderly patients. Response: Pfizer and BioNTech are aware of reported deaths following administration of Pfizer-BioNTech COVID-19 vaccine . We are working with the (withheld) to gather all the relevant information. (Withheld) Authorities have prioritized the immunization of residents in Nursing Homes, most of whom are very elderly with underlying medical conditions and some which are terminally ill. (Withheld) confirm the number of incidents so far was not alarming, and in line with expectations. All reported deaths would be thoroughly evaluated by (Withheld) to determine if these incidents are related to the vaccine. (Withheld) Health Authorities have now changed its recommendation in relation to vaccination of the terminally ill (Clinical Frailty Scale 8 or higher). Our immediate thoughts are with the bereaved families. The Global Advisory Committee on Vaccine Safety (GACVS) COVID-19 Vaccine Safety subcommittee met virtually on Tuesday, 19Jan2021, to review available information and data on deaths reported in frail, elderly individuals who had received the Pfizer BioNTech COVID-19 mRNA vaccine, BNT162b2 (hereafter, BNT162b2). Experts invited from the (Withheld) and the (Withheld) provided an overview of deaths reported in (Withheld) and in the WHO global database (VigiBase) following vaccination with BNT162b2."" Based on a careful scientific review of the information made available, the subcommittee came to the following conclusions: The current reports do not suggest any unexpected or untoward increase in fatalities in frail, elderly individuals or any unusual characteristics of adverse events following administration of BNT162b2. Reports are in line with the expected, all-cause mortality rates and causes of death in the sub-population of frail, elderly individuals, and the available information does not confirm a contributory role for the vaccine in the reported fatal events. In view of this, the committee considers that the benefit-risk balance of BNT162b2 remains favourable in the elderly, and does not suggest any revision, at present, to the recommendations around the safety of this vaccine."" Enquired what was the efficacy after one dose. Caller had a terrible experience after her first dose of the Pfizer BioNTech COVID19 vaccine. She had blood clots in her nose, every time she blew her nose, blood clots would come out. She never had a nose bleed before. She also added that her arms are both numb on her. She couldn't move them, she feels lethargic. She's supposed to get her second dose next Friday. Enquired do we have any information regarding blood clots or not. Caller was upset and disconnected the call before some of the answers were given. Caller also asked the following questions: Has there been deaths reported from the vaccine or not, what was the efficacy after 1 dose, any guidance on getting the second dose after having side effects after the first dose. Response: Proposed response : CONS-Blood clots. Caller said ""this was ridiculous"" when I tried providing her with a long website with more information. She feels like we are withholding information from the public. She thinks the deaths should have been on the media. Caller hung up. Enquired do you have any guidance on getting the second dose after having side effects with the first dose. I had a side effect after the first dose of the vaccine. Should I get the second dose or not. Caller disconnected before agent could provide response. Response: Proposed response: As noted in the Fact Sheet for Recipients, you should not get the Pfizer-BioNTech COVID-19 Vaccine if you had a severe allergic reaction after a previous dose of the vaccine or if you had a severe allergic reaction to any ingredient of the vaccine. The decision to receive the second dose for any other reason could not be made by Pfizer. We refer you to speak to your healthcare provider about the risks of the vaccine compared to the risks of potentially not being fully protected against COVID-19 infection. Your healthcare provider knows your health situation and has access to information that can better help inform this decision. The outcome of the events was unknown. No follow-up attempts are needed. Information about lot/batch number cannot be obtained."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288429-1	"I went back to the hospital and couple days later they told me that actually I have Deep vein thrombosis now in my legs; my Crohn's had caused really bad flare up/I have been in the hospital for 5 days now with the clot and Crohn's Flare; I also have superficial clot in that left leg that was the pain began on the edges/blood clot; the arm is much better now. You know that's not really sore; I just was too tired/extreme fatigue; really bad like pain; Shortness of breath; This is a spontaneous report received from a contactable consumer (patient). A 54-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 2 via an unspecified route of administration, administered in Arm Right on 29Mar2021 (Batch/Lot Number: EN6201) as SINGLE DOSE for covid-19 immunisation. Medical history included multiple sclerosis from an unknown date and unknown if ongoing, ileostomy from an unknown date and unknown if ongoing, back surgery from an unknown date and unknown if ongoing, Crohn's disease from an unknown date and unknown if ongoing , blood clot in arm back from 2014 to an unknown date. Historical vaccine included BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 1 on 08Mar2021 and experienced terrible diarrhea and neuralgia. Concomitant medication(s) included gabapentin (GABAPENTIN) taken for an unspecified indication, start and stop date were not reported; valaciclovir hydrochloride (VALTREX) taken for an unspecified indication, start and stop date were not reported; diazepam (DIAZEPAM) taken for an unspecified indication, start and stop date were not reported; (ZOFRAN) taken for an unspecified indication, start and stop date were not reported; oxycodone (OXYCODONE) taken for an unspecified indication, start and stop date were not reported; lansoprazole (PREVACID) taken for an unspecified indication, start and stop date were not reported. After second dose on 29th (29Mar2021) immediately he was really bad like pain and had shortness of breath. The patient stated that she had a back surgery and so she always felt bad, so she didn't think anything of it. But then she ended up into ER after the second dose and discovered that her Crohn's had caused really bad flare up. She mentioned her foot was looking strange but she was just too tired and left the ER. She then went back to the hospital and couple days later they told her that she actually had deep vein thrombosis now in her legs. She also had superficial clot in that left leg where the pain began on the edges. The patient reported that the vaccine was basically started the Crohn's flare up. The patient also stated that ""the arm was much better now and not really sore"". The patient was hospitalized for DVT from unknown date and for Crohn's flare up for 5 days. The patient underwent lab tests and procedures which included endoscopy, sigmoidoscopy, ultrasound scan: all on unknown date with unknown result. The patient was treated with Entyvio and on blood thinners now for a blood clot. The outcome of pain in extremity was recovering while others was unknown."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288432-1	"My grandma got a PFIZER COVID VACCINE and has now experienced a blood clot that caused a stroke.; My grandma got a PFIZER COVID VACCINE and has now experienced a blood clot that caused a stroke.; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Lot number unknown), via an unspecified route of administration on an unspecified date as a single dose for COVID-19 immunization. The patient's medical history was not reported. The patient's concomitant medications were not reported. The reporter stated that ""my grandma got a pfizer covid vaccine and has now experienced a blood clot that caused a stroke."" The patient experienced a cerebrovascular accident on an unspecified date with outcome of unknown and thrombosis on an unspecified date with outcome of unknown. No further information was provided. Information on the Lot/Batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288442-1	minor blood clot on his leg; This is a spontaneous report from a Pfizer-sponsored program received from a contactable consumer (patient). A male patient of unspecified age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number not reported), on 03Apr2021 at single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. It was reported that patient had the first dose on 03Apr2021 and he was scheduled on Saturday 24Apr2021 to get the second dose of the vaccine. However, the patient had a minor blood clot on his leg, and he was taking blood thinners. Patient asked if it was okay to get the vaccine while talking to the blood thinners (as reported). The outcome of the event was unknown. Information about lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288824-1	Pfizer BioNTech-Covid-19 vaccine EUA--4/10/2021 started having abdominal pain radiating to back, left chest pain and it progressively got worse. 4/13/21 called PCP to make appointment for 4/14/21, ended up going to ER on 4/14/21 they did EKG, blood work ultrasound of abdomen, X-ray and CT scan. They discovered the clot in portal vein to the liver. ER prescribed blood thinners, states she went back on the 15th because pain was not improving. She saw Dr., hematology, on 4/19/21 she thinks the clot is from the COVID vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288839-1	Systemic: Blood Disorder (diagnosed by MD)-Severe, Additional Details: Patient passed away on 4/23/21 and had blood clots throughout body. Patient did not have any other symptoms that were reported. Patients family requested a VAERS report be submitted.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1288911-1	Period lasting over 14 days (still on going). Multiple pads filled in a 2 hour window. A piece of uterine lining came out of body, approximately 2 inches long and an inch across.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1289494-1	Patient was hospitalized with lower extremity DVT and bilateral pulmonary emboli. He is young and previously very healthy and has no risk factors for venous thromboembolism.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1290246-1	1/7/2021 vaccination 1/8/2021 bruise/pain R lower leg, hard warm to touch. Painful and sore. Progressively got worse; red, bluer. 1/10 or 1/11 ER . Ultra sound on leg. 10cm clot. Put me on aspirin, warm compresses. Had to increase aspirin up to 2x day. It eventually cleared. Referred to surgeon. Found some other veins their inspecting further. *have not had any more clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1290888-1	Blood Clot discovered in my leg my Wednesday April 27. Incredible amounts of pain. By May 5 pain caused by unknown bruise on foot on same leg.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1290956-1	After the vaccination in the morning, Sa-04-17-2021, have been in Urgent Care in the afternoon for an ongoing kidney stones issue that had occurred earlier this week. Felt tightness and pain in both lower legs which continued the next day, and it became worse. Mentioned this issue to the primary physician the following Monday 04-19-2021, but predominance had been given to the Kidney stones issue. On 04-26-2021 mentioned again the pain predominantly in the left calf to the primary physician. The same day, blood clots were found in both lower legs by Urgent Care. Had received information for Deep Vein Thrombosis by physician who prescribed Lovenox (enoxaparin) injections and advised to report the findings to V-safe. Continued narrative: Being informed by Urgent Care on 04-30-2021 that blood clots were not only in both lower legs, but also in both upper legs, now, after complaining about more pain. Ultrasound checks. Being advised to move and rest moderately. Received again information about Deep Vein Thrombosis. Advised to consider to skip the second Pfizer vaccination. Being on sick leave since 04-13-2021. Had switched from Lovenox to Pradaxa per prescription starting from Sa-05-01-2021. Urgent Care, 05-03-2021 conducted another ultrasound check. Another blood clot had developed. Again advised to consider to skip the second Pfizer vaccination, on SA-05-08-2021, to avoid more complications and risks. Under continued observation. Being alerted that this condition can lead to permanent damage and can turn life-threatening. Continued sick leave, considering medical leave. Concerned about the decision to take the second Pfizer vaccination, on Sa-05-08-2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291116-1	severe pain in the leg, it got worse, checked for clot, pain was from inner thigh to calf, it was swollen; severe pain in the leg, it got worse, checked for clot, pain was from inner thigh to calf, it was swollen; severe pain in the leg, it got worse, checked for clot, pain was from inner thigh to calf, it was swollen; heart palpitation for 6 hours; muscle pain; light headiness; chills; This is a spontaneous report from a contactable consumer. This consumer (patient) reported for herself that a 65-years-old female patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 09Mar2021 at single dose for COVID-19 immunization. Medical history included she had COVID-19 in the past and thyroid. Concomitant medications included she was also taking medications for the thyroid. Patient stated that she had already reported her symptoms with Pfizer COVID-19 vaccine she received on 09Mar2021 and a week after she experienced severe pain in the leg, it got worse, checked for clot, pain was from inner thigh to calf, it was swollen, had heart palpitation for 6 hours, light headiness, chills, she was wearing a heart monitor now, she still had the muscle pain. Patient wanted to know if with second dose she would have more sever side effects, should she take or not take the second dose. She was scheduled to take it this Thursday. Patient also wanted to know how she would know if she was allergic to the vaccine/ingredient. Outcome of the event severe pain in the leg was not recovered while of remain was unknown. No follow-up attempts are needed; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291138-1	The pain I feel, Could be due to a blood clot?; pain on my arm; felt numbness in my arm; now I feel it on the inside, like if it was in the bone.; This is a spontaneous report from a contactable consumer (patient). A 42-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), via an unspecified route of administration in the arm on an unspecified date (Batch/Lot number was not reported) as a single dose for COVID-19 immunization. Medical history was not reported. Concomitant medication included paracetamol (TYLENOL) taken for an unspecified indication, start and stop date were not reported. The patient received the Pfizer Covid 19 vaccine on Friday, since then she felt pain on her arm, it got stronger in the afternoon, she couldn't lift anything or move that arm. On Saturday, the pain diminished. On the day of the report, the patient still feel pain, but now is like an internal ache, like if it is in the bone. She had trouble getting dressed, specially putting on t-shirts, when she goes to sleep, she cannot lay on that arm. It was reportedly very painful, and she was feeling as if it has moved to the other arm. On the day of the report, she applied heat on her arm and took Tylenol on Friday and Saturday. On Friday, she couldn't do anything, she felt numbness in her arm. Initially the pain was external, now she feels it on the inside, like if it was in the bone. The patient also asked if the pain she feel could be due to a blood clot. The event pain on my arm resolved on an unknown date in 2021, while the outcome of the rest of the events was unknown. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291184-1	"This is a spontaneous report from a contactable consumer (patient's fiance). A 46-year-old male patient received the first dose bnt162b2 (BNT162B2), via an unspecified route of administration, administered in Arm Left on 18Apr2021 11:20 (Lot Number: EW0162) (received at the age of 46-years-old) as SINGLE DOSE for COVID-19 immunisation. Medical history included right knee pain. Concomitant medications were not reported. The patient received the Pfizer 1st shot 18Apr2021 11:20 am. On 19Apr2021, at midnight, he had low grade fever then swelling right leg. On the day of the report, 20Apr2021, he had swelling to both legs. The patient had taken ibuprofen as treatment beside propping legs up. The reporter wanted more information on the swelling and if it's related to taking ibuprofen. It was further reported that the patient experienced severe swelling from the knees down to the calves on both legs on 19Apr2021. He also experienced night sweats, fever, lightheadedness and dizziness on 19Apr2021. They want to know if there are any reports regarding the swelling side effects and if it is from the vaccine. The first dose of the vaccine was given 2 days ago (18Apr2021) and the second dose is due on 09May2021. The patient got his first shot on 18Apr2021 about 11:20am. After about 15 hours he developed swelling in one leg, she later clarified this to be his right leg, and now reports that both of his legs are swollen at the time of report (20Apr2021). She noted that patient told her his legs are not painful. She doesn't know if it's a blood clot or if they needed to go to the hospital. She also wanted to know if there is information about this as a side effects or if other people are having this too. She then reported that the patient started experiencing a fever, profuse sweating, and was spacey about 12 hours after the vaccine, a little after midnight 19Apr2021. She doesn't know if the dehydration made it worse. His fever was low-grade 99.8 degrees and lasted 4 and a half to 5 hours and has now resolved. When asked if patient received any treatment for fever, caller said it was so recent, and that was listed as a common side effect that goes away. Reported the patient drank fluids and she kept an eye on him. Leg swelling was about 15 hours after the vaccine, early morning 19Apr2021. She noted that the swelling in the right leg got bigger and didn't go down, but the left leg did. But today the left was more swollen and more prominent today. She further clarified that the patient takes ibuprofen for pre-existing right knee pain. Caller also reported that the patient propped his legs to alleviate the swelling, it has improved some but the swelling did not go away. The adverse events did not require a visit to the physician office or emergency room. Outcome of the event fever was resolved on 19Apr2021 (lasted 4 and a half to 5 hours); outcome of the event ""severe swelling from the knees down to the calves on both legs"" was not resolved; outcome of the remaining events was unknown."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				took 2 generic Whole Foods 365 brand Acetaminophen Expiration Date: Jun2020 in Apr2021; ear crystals; Nausea; she wanted to make sure her not feeling well was normal; She had a really bad headache. Clarifying the headache started 1-2 hours after she got the first COVID-19 Vaccine shot; her (left) arm was really sore after she had the COVID-19 Vaccine, but she had expected that; She was still achy in her muscles, but not everywhere. She said her muscles ached from her shoulders up only.; she had a scratchy (sore) throat; she felt tired; she had a little bit of congestion; after she received the COVID-19 Vaccine that she had some clotting, and menstrual irregularities; she had a swollen lymph node in her left arm pit for the last couple days; after she received the COVID-19 Vaccine that she had some clotting, and menstrual irregularities; she had a pressure inside of her head/Her head doesn't feel well. She said she does not quite have a headache, but her head does not feel normal.; This is a spontaneous report from a contactable consumer (patient). A 48-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: EW0170), dose 1 single dose via an unspecified route of administration in the left arm on 16Apr2021 between 3:53 PM-3:55 PM (at the age of 48 years-old) for COVID-19 vaccination. Medical history included hypermobility syndrome from an unknown date and unknown if ongoing, tends to get headaches with her hypermobility issues, and those headaches felt like pain in her head. Reported she was on the hypermobility spectrum with tension headaches, vertigo positional from an unknown date and unknown if ongoing, 3-4 weeks prior to getting the COVID-19 Vaccine, she tried to take a half of a beta blocker pill to treat the benign paroxysmal positional vertigo (BPPV). She said the BPPV had cleared, and she felt better, vestibular migraine from an unknown date and unknown if ongoing had some residual dizziness from the vestibular migraines, Ehlers-Danlos syndrome from an unknown date and unknown if ongoing. She said she did not have a technical diagnosis for Ehlers-Danlos Syndrome (hypermobility) yet, and finally sees a geneticist next month. She said she was on the weaker end of the Ehlers-Danlos hypermobility spectrum disorder, and did not have a definitive diagnosis. She didn't drink alcohol, exercised every day and slept enough. The patient did not receive any previous immunization with the Pfizer vaccine. The patient did not receive any other vaccine within 4 weeks prior to the vaccine. Concomitant medications included cyanocobalamin (VITAMIN B-12) taken for an unspecified indication, start and stop date were not reported and ergocalciferol (VIT D) taken for an unspecified indication, start and stop date were not reported. Cyanocobalamin and ergocalciferol were discontinued a week before she got the COVID-19 vaccine to avoid any issues.The patient experienced she had a little bit of congestion on Apr2021, she wanted to make sure her not feeling well was normal on 16Apr2021, she had a really bad headache, clarifying the headache started 1-2 hours after she got the first covid-19 vaccine shot on 16Apr2021, her (left) arm was really sore after she had the covid-19 vaccine, but she had expected that on 16Apr2021, she had a swollen lymph node in her left arm pit for the last couple days on Apr2021, after she received the covid-19 vaccine that she had some clotting, and menstrual irregularities on Apr2021, she had a pressure inside of her head/her head doesn't feel well, she said she does not quite have a headache, but her head does not feel normal on Apr2021, she was still achy in her muscles, but not everywhere. she said her muscles ached from her shoulders up onlyon 16Apr2021, she felt off/felt a little run down on Apr2021, she had a scratchy (sore) throat on Apr2021, she felt tired on Apr2021, nausea on 16Apr2021, her (left) arm was really sore on an unspecified date and swollen lymph node in her left arm pit on an unspecified date. She said it was not possible that her head pressure, and her head feeling off had anything to do with the BPPV or the vestibular migraines. She said her ear crystals had fell out since then, and what she was experiencing now must be due to the COVID-19 Vaccine. She had no fever. She only had nausea for the first 24 hours after getting the COVID-19 Vaccine. She said she had no vomiting. She said her headache had been constant since she had the COVID-19 Vaccine on 16Apr2021. The headache started 1-2 hours after she got the first COVID-19 Vaccine shot. The events did not require an emergency room or physician's office visit. The patient underwent lab tests and procedures which included blood test and urine analysis on an unknown date in Mar2021 or Apr2021: came back normal . Therapeutic measures were taken as a result of she had a really bad headache. clarifying the headache started 1-2 hours after she got the first covid-19 vaccine shot (headache). She took 2 generic Whole Foods 365 brand Acetaminophen 500mg (1000mg total): NDC Number: 42681-7076-1, Lot Number: 8HE1354B, & Expiration Date: Jun2020 in Apr2021. The events did not require an emergency room or physician's office visit. The clinical outcome of not feeling well, headache, arm was really sore, swollen lymph node, had some clotting and menstrual irregularities was recovering and not recovered for pressure inside of her head, felt off, scratchy (sore) throat , congestion, felt tired, hadn't felt 110%, recovered for nausea and unknown for achy in her muscles and ear crystals.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1291204-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291268-1	This is a spontaneous report from a contactable consumer (patient). A 59-year-old male patient received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: ER8732) at the age of 59-years-old, via an unspecified route of administration in left arm on 22Mar2021 at 15:00 at single dose for COVID-19 immunisation. The patient's medical history was reported as none. He has no known allergies and has COVID prior to vaccination. Concomitant medications included metoprolol, rosuvastatin, oxybutynin, and tamsulosin. There were no other vaccines in four weeks and no additional vaccines administered on the same date of the COVID-19 vaccine. The patient reported that he was having some wild effects from the vaccine and did not know what to do. On 26Mar2021 at 15:00, he was having neuropathy like effects. His hands, arms, legs, and feet were experiencing pain and burning sensations, they were burning like nerve damage; he also has a little burning sensation in his nose and cheeks. He has never had this before. The events resulted in doctor or other healthcare professional office/clinic visit and emergency room/department or urgent care. He was also having some weird sensations in his calf muscles, these sensations started within the past week and a half (Apr2021) which prompted physician office visit. The physician placed him on gabapentin 100 mg to start from 12Apr2021, but it was not enough, it was titrated up to 300 mg, three times a day and he is scheduled to see specialist soon. The patient mentioned that this vaccine has ruined his health. He also went to an urgent care center because his leg was bothering him, it was swelling on an unspecified date in 2021; he thought he had a blood clot; they did some kind of test where they looked at the leg and did not find anything. He didn't have a particular question. He would just like to talk to someone to see if this was going on with anyone else, if it will go away, or if this was permanent problem he will have. The patient was not tested for COVID post vaccination. He received his second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EW0161), via an unspecified route of administration in left arm on 13Apr2021 at 15:00 at single dose for COVID-19 immunisation. The outcome of the events was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291930-1	Immediate chest pain (tightening) , then arm pain while in store/pharmacy Next Day began spotting (menstrual bleeding) and lasted for 2.5 days Chest pain continued and has not stopped for last 10 days, intermittent throughout the day - tigheting of left side and right side of chest, radiates to theback Sunday following Monday vaccination (6 days later) got a blood clot on palm of hand, lasted for 4 days that started bleeding out and blood spread to front of hand like a birthmark as it bleed out
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1292074-1	I started my period on Tuesday (which has always been incredibly regular and predictable for 5+ years while being on birth control). I have been experiencing heavier bleeding and more clotting than average. My cramps are also worse. This never happens to me, and it only makes sense due to the vaccine. I would like there to be some organization that looks into this because I have not changed anything in my life for this to be happening.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1292206-1	Extreme blood clots in lungs and legs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1292432-1	Significant change in menses. Typically cycles once every three months with continuous birth control use. Menses in January (after first vaccination) was significantly heavier with passage of clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1292987-1	Injection site tenderness, flu-like symptoms including headache, low grade fever, muscle and joint aches, chills, making work impossible for two days. Intensely painful and heavy menstrual period a week later, including passing palm-sized blood clots and intense pain that did not subside with otc pain killers, TENS therapy or heat. Patient does not have a history of painful menstrual periods.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1293193-1	Patient received her first Pfizer vaccine on April 8, 2021. On April 24, her home health care nurse noticed swelling of the left calf. Pt is wheelchair dependent and has little sensation below the waist due to MS. She never had any pain. She was seen in the office on April 27 with significant swelling and discoloration of her left leg. A venous duplex scan revealed a large clot burden from the groin to the ankle. She was started on Eliquis and evaluated by vascular surgery. She required surgical embolectomy which was preformed 5/5/21.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1293871-1	Blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1293888-1	BLOOD CLOTS IN BOTH ARMS AND BRUISERS BACKS ON BOTH LEGS SOME WHEEZING

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294515-1	"After receiving the first dose on 4/1/21 Lot# ER8733, I had light spotting for 2 days. Second dose given 04/23/21 and by 04/28/2021 I started bleeding again And by the second day I was soaking through pads and huge clots were coming out. This has never happened to me before. After 7 days of bleeding heavily I woke up on May 5th at 4:00 a.m. with intense abdominal cramping The only way I can describe was labor. I felt back labor, pelvic pressure, frequent urination and intense pain. I reached out to my gynecologist And he immediately told me to remove my estrogen patch and later that morning he started me on progesterone treatment for 10 days to stop the bleeding. After 7 hours of pain on May 5th, I went to the emergency department and an ultrasound revealed a blood clot. From ultrasound results ""there is a complex area in the area of the cervix measuring 5.7 x 3.6 x 2.6 cm."" After several hours I was released from the emergency room taking toradol for pain. That evening I passed a huge blood clot! I'm very grateful it was where it was and I could expel it. But there's no doubt the second vaccine caused this issue that has never happened to me before. I still get my periods though irregular and very light. So this event was highly unusual and very painful"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294576-1	First shot 12/23, 2nd 1/12/21. A week after #2 my menstrual cycles (which have been clockwork all of my life-I have 4 children and track it monthly) became much more frequent and heavy with excessive clotting. I thought perhaps I was going though early menopause (44) but then realized correlation with vaccine. My mood changed, I feel sad all of the time, I am dreading recommendation of additional shots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294661-1	"Blood clots on my legs; tired; This is a spontaneous report from a contactable consumer (the patient). A 30-year-old patient of an unspecified gender received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Lot number ER8737), via an unspecified route of administration, on 07Apr2021 at 12:30 (at the age of 30 years old) as a single dose in he left arm for COVID-19 immunization. Medical history included COVID-19 from an unknown date and unknown if ongoing. The patient has no known allergies. The patient's concomitant medications were not reported. The patient had received no other vaccines within fourteen days prior to vaccination. The patient experienced ""blood clots on my legs"" on 07Apr2021 at 17:00 with outcome of unknown, and ""feeling drained and tired for two weeks now"" begining on 07Apr2021 at 17:00 with outcome of unknown. The patient has not been tested for COVID-19 since the vaccination ."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294665-1	three smallest toes on both feet (left/right) feel swollen; discomfort in wearing shoes and sleeping at night; balance is affected; nerve damage; discoloration; blood clots; This is a spontaneous report from a contactable consumer (patient). A 57-years-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; lot number and expiry date: unknown), via an unspecified route of administration, administered in Arm Left on 06Mar2021 11:00 as single dose for COVID-19 immunization. The patient had no medical history and known allergies. Patient did not have COVID prior vaccination. Patient was not tested for COVID post vaccination. Concomitant medications were not reported. It was reported that patient had three smallest toes on both feet (left/right) feel swollen. He had discomfort in wearing shoes and sleeping at night. He had movement of the toes but balance is affected. Just really troubling as nothing like this was affecting him before. He had been told it was COVID toe. He first thought it is was nerve damage. Could be blood clots small based on reading information. He never got sick. He was attempting to set a doctors appointment. There is a small amount of discoloration. He was hopeful with time this will get better. Events started on 09Mar2021 14:00 with outcome of not recovered. No treatment received. Events resulted in Doctor or other healthcare professional office/clinic visit, Disability or permanent damage. Events were reported as serious due to disability. Patient had second dose of bnt162b2 (lot no.: ER2613) on 27Mar2021 on Left Arm. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				blood clot; anemia; knee down felt tingly/like her legs were falling asleep; knee down felt tingly/like her legs were falling asleep; could not sleep; She was feeling uncomfortable constantly/she did not feel well; Shortness of breath; Her arm started hurting/was just at the injection site; This is a spontaneous report from a contactable consumer. This consumer (patient) reported for herself that a 36-year-old female patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EP7533) at the age of 36-years, via an unspecified route of administration in left arm on 04Apr2021 12:15 at single dose for COVID-19 immunisation. Medical history included had a pregnancy (a while ago, not ongoing). No other products. The patient stated that she experienced shortness of breath on 08Apr2021 (4 days after receiving the first dose), it got so severe on day 7 that she went to the ER on 11Apr2021. She started getting progressively more shortness of breath. Twice, she called the nurse line for her doctor and was advised to go to the emergency room. Added she could not catch her breath, and it was worse at night, could not sleep, so she went to the emergency room, she was in emergency room for 2 and a half hours and then went home. She wanted to know if the second dose could cause issues or if she should get it, added that the paperwork said that people should not get the vaccine if they received an adverse reaction to the first dose or any vaccination. She did not know if this was normal for it to get progressively worse and 2 to 14 days later. Patient wanted to know if the shortness of breath was something that can get worse, if this was a reaction to the vaccine even though it happened so many days later. Patient added it was the only thing she could think of and that her heart rate, oxygen levels, and everything was good minus she felt like she could not catch her breath. The doctor in the ER gave her anti-anxiety medicine, something with an L, but the doctor gave her that and it really seemed to help. It was kind of like, she did not know, it made her feel calm enough to fall asleep. Caller added that she has that and an salbutamol (ALBUTEROL) inhaler. She was not sure whether progressively worsening shortness of breath had gotten any better or was still going on, added that yesterday was the day she did not need either treatment medication. The day before, she took just the anxiety medicine and not the albuterol. Yesterday, she took none. During the day on 11Apr2021 (Sunday), she had called because she was feeling uncomfortable constantly. She has two children and all day she did not feel well. She told her husband she did not feel well at all. Sunday when she went to lay down, she could not sleep. Patient reported that as her shortness of breath subsided, her arm started hurting again, thought this began on 19Apr2021 and was not sure, it went away completely and was just at the injection site, probably lasting half a day or so and then it went away, clarified as left arm, same as injection. Caller added that 15Apr2021 was after the ER visit, was that she was standing up doing things and from her knee down felt tingly. Stated that knee down felt tingly on 15Apr2021, which was after her ER visit occurred, she was standing up and doing things. She had not been doing things for long periods but her knee down felt tingly. Added not like numb, but like her legs were falling asleep. She was preparing dinner and moving around and felt tingling in both legs. This probably lasted about three hours. She just kind of took it easy and it started to subside and then it was gone. While mentioned investigations, patient stated the ER did an EKG, where they put stickers and monitor her heart. They did a chest x-ray, and they took blood. She believed they checked for blood clot tests and anemia. Patient did not know any results but stated the doctor came into the room and told her everything was clear and nothing came back with anything on it. She stated the date was either the night of 11Apr2021 or the morning of 12Apr2021 for the tests. Outcome of the events shortness of breath and arm hurting was recovered/resolved in Apr2021; of the event knee down felt tingly/like her legs were falling asleep was recovered/resolved on 15Apr2021; while of remain was unknown. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1294675-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294744-1	urinating blood; might be a UTI; arm pain; fatigue; blood clots; he started bleeding after the first shot; This is a spontaneous report from a contactable consumer, the patient. This 63-year-old male patient reported that he received the first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EL3249) on 18Feb2021 (also reported as 17Feb2021) (at the age of 63-years-old) via an unspecified route as a single dose for COVID-19 immunization. Medical history included enlarged prostate, urinary health issues (gets up in the middle of the night), and has glasses as he had cataract surgery last year (2020) so he lost his close-up vision. Concomitant medications were not reported. On 18Feb2021, the patient got his first shot. Three weeks later, he had an episode of urinating blood and blood clots profusely. The patient stated the amount of blood and blood clots was scary at best. The patient reached out to his daughter who is a nurse who advised him to go to the Emergency Room (ER). He reached out to the nurse hotline as well, which recommended that he go to the emergency room that day as well. He went to the ER (it was a Sunday). They did a blood test, a sonogram or some kind of sonogram, and a urine sample. They also flushed the patient's bladder out with a catheter. This emergency room visit determined that it might be a urinary tract infection (UTI) and started him on antibiotics. The patient confirmed he was not admitted to the hospital. The patient then made an appointment with his doctor because that was on a weekend. He saw his doctor who referred him to a urologist. The patient also reported that he did have the normal arm pain and fatigue the second day or the day after the shot. The patient reported 11Mar2021 or 12Mar2021 was when he started bleeding after the first shot, he was unsure of the exact date, and stated he went to the hospital on 14Mar2021. The patient reported he is going to the doctor today (21Apr2021), to see the urologist but the urologist is going to do a scope in his bladder because that is the only thing that they cannot see. The clinical outcome of events urinating blood, blood clots, UTI, bleeding, arm pain, and fatigue was unknown.; Sender's Comments: Linked Report(s) :PFIZER INC-2021471072 same patient/reporter, different AE/2nd dose
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294746-1	blood clots; urinating blood; arm pain; fatigue; he started bleeding from the second one; This is a spontaneous report from a contactable consumer, the patient. This 63-year-old male patient reported that he received the second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EN6206) on 17Mar2021 (at the age of 63-years-old) via unspecified route as a single dose for COVID-19 immunization. Medical history included enlarged prostate, urinary health issues (gets up in the middle of the night), has glasses as he had cataract surgery last year (2020) so he lost his close-up vision. Concomitant medications were not reported. The patient previously received the first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Batch/lot number: EL3249) on 18Feb2021 (also reported as 17Feb2021) (at the age of 63-years-old) via unspecified route as a single dose for COVID-19 immunization and experienced urinating blood and blood clots profusely, UTI, bleeding, arm pain, and fatigue. The patient received his second shot on 17Mar2021 and almost to the date three weeks later, had the same incident with urinating blood and passing blood clots. He waited an extra day because he could not go to the bathroom and then he went to the ER that same day so that they could flush him out. He also had an appointment on that same day with his urologist. A CT scan was done at the ER and then in the afternoon he went to the doctor. The doctor did a catheter flush. The patient was unclogged but he was passing blood. The patient also reported that he did have the normal arm pain and fatigue the second day or the day after the shot. He has no other adverse reactions at this time. The patient reported 11Apr2021 or 12Apr2021 was when he started bleeding from the second one (shot). He went to the hospital on 14Apr2021. The patient confirmed he was not admitted. The patient reported he is going to the doctor today (21Apr2021) to see the urologist but the urologist is going to do a scope in his bladder because that is the only thing that they cannot see. The clinical outcome of the events urinating blood, blood clots, bleeding, arm pain, and fatigue was unknown. The patient stated he does not know if this is an issue, however, it coincided with both of his shots. The patient was calling to see if this is something that has been seen or if it is in his head.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021470922 same patient/reporter, different AE/first dose
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1295866-1	Swollen red left arm. Diagnosed with blood clots in all his veins in left arm and spent 4 days in hospital on blood thinners and had 2 surgeries to remove them
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1296602-1	Shortness of breath upon exertion.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1296762-1	Had an additional mensural cycle with heavy clotting and cramping.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1296804-1	4/28/2021-Lower left leg hurting; 4/29/2021-can't put weight on leg; ER visit with doppler to confirm blot clot in calf; ER gave Norco 5-325 mg for pain; fever of 101; 4/30/2021-still can't put weight on leg and fever of 101.8; 5/1/2021-woke up with no pain; clot still there; 5/3/2021-saw primary doctor for ER follow-up on clot; primary doctor confirmed there is a small clot in deep vein and could take up to 6 months for it to dissolve on its

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1297346-1	Tinitus in left ear as of 4/13/21. And disgnosed with a blod clot in right calf symptom of clot felt on 4/17/21
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1297853-1	Clotting and breakthrough bleeding for 4 weeks following 1st vaccination, then swelling of lymph nodes 48 hrs. following 2nd vaccination.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299097-1	blood clot; This is a spontaneous report received from a contactable pharmacist from a Pfizer-sponsored program. A 70-year-old male patient received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date were not provided), via an unspecified route of administration on an unspecified date at single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The pharmacist was asking if there have been any reports of blood clots associated with the COVID-19 vaccine as the patient developed blood clot 1.5 weeks after receiving his first Pfizer COVID vaccine on an unspecified date. The outcome of the event was unknown. Information about batch/lot number has been requested.; Sender's Comments: Based on the information currently available, a possible contributory role of the suspect vaccine BNT162B2 or comirnarty in triggering the onset of Blood clots cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299103-1	Blood clot in the calf; Swelling/pain behind the knee; Swelling/pain behind the knee; A 62-years-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 02Apr2021 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation. The patient's medical history was not reported. Concomitant medication included vitamin d3 (VITAMIN D3). The patient experienced blood clot in the calf and swelling/pain behind the knee on 10Apr2021. The events resulted in emergency room visit. The patient underwent lab tests and procedures which included nasal swab: negative on 17Apr2021. Therapeutic measures were taken as a result of thrombosis, joint swelling and arthralgia included anticoagulation medication. The patient did not have COVID prior vaccination. Outcome of the event was recovering. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299104-1	Two large hot swollen areas on right leg Went to ER ultrasound detected superficial clot; This is a spontaneous report from a contactable consumer (patient). This 46-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in arm right on 19Apr2021 12:15 (Batch/Lot Number: EWO164) as a single dose for COVID-19 immunization. The patient's medical history was not reported. Patient is not pregnant at the time of vaccination. Concomitant medications included fluticasone propionate (FLONASE), fexofenadine hydrochloride (ALLEGRA), and ibuprofen. The patient previously took tobramycin and Wellbutrin and experienced allergies for both. Patient received the first dose of bnt162b2 for COVID-19 immunization on an unspecified date (lot number unknown). Patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient was not diagnosed with COVID-19 prior to vaccination. Patient has not been tested for COVID-19 since the vaccination. On 20Apr2021 18:30, patient had two large hot swollen areas on right leg. She went to the ER. Ultrasound detected superficial clot. AE resulted in emergency room/department or urgent care. Patient has not recovered from the event. Treatment included antibiotics and ibuprofen.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299150-1	He completely lost mobility on the left side of his body; Stroke; They found they clot, but could not extract it from what he understood; This is a spontaneous report from a contactable consumer. A 46-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on 31Mar2021 (Batch/Lot number was not reported) as single dose (at the age of 46 years old) for COVID-19 immunisation. The patient has no medical history. Concomitant medications included aspirin [acetylsalicylic acid] (ASPIRIN [ACETYLSALICYLIC ACID]) taken for an unspecified indication, start and stop date were not reported. Since his stoke, he is doing better. He completely lost mobility on the left side of his body. His mobility on the left side of his body is slowly coming back. His job is in jeopardy. The doctors told him that there is no guarantee that he will regain full mobility. They did not give him a patient record card. If they did give him one, he left it at the house. He has been in the hospital ever since his stroke. They put him on blood thinners after his stroke. Whenever he was in the ambulance, he did not have high blood pressure. They were able to catch his stroke quick. They found they clot, but could not extract it from what he understood. Adverse events reqiored Emergency Room. The patient was hospitalized on 01Apr2021. The outcome of clot was unknown and other events was recovering. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299152-1	period is extremely heavier than it's ever been, I have large blood clots; period is extremely heavier than it's ever been, I have large blood clots; cramping; irritability; premenstrual symptoms like cramping and irritability were the worst they've ever been; This is a spontaneous report from a contactable consumer, reporting for herself. A 30-year-old non-pregnant female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 2 via an unspecified route of administration, administered in the right arm on 15Apr2021 (Batch/Lot Number: Er8729) as a single dose (at the age of 30 years old) for covid-19 immunization. The patient medical history was not reported. Concomitant medication(s) included ethinylestradiol, levonorgestrel (SEASONIQUE) taken for an unspecified indication, start and stop date were not reported; spironolactone taken for an unspecified indication, start and stop date were not reported; magnesium taken for an unspecified indication, start and stop date were not reported; polycarbophil calcium (FIBER) taken for an unspecified indication, start and stop date were not reported, and an unspecified multivitamin taken for an unspecified indication, start and stop date were not reported. Historical vaccine information included BNT162B2, dose 1 via an unspecified route of administration, administered in the right arm on 25Mar2021 at 12pm (Batch/Lot Number: Er8730) as a single dose (at the age of 29 years old) for covid-19 immunization. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient experienced period is extremely heavier than it's ever been, she has large blood clots on 21Apr2021 at 08:00 with outcome of not recovered, cramping on 21Apr2021 at 08:00 with outcome of not recovered, irritability on 21Apr2021 at 08:00 with outcome of not recovered, and premenstrual symptoms like cramping and irritability were the worst they've ever been on 21Apr2021 at 08:00 with outcome of not recovered. The patient underwent lab tests and procedures which included sars-cov-2 test (nasal swab): negative on 31Mar2021. Treatment was not received for the events.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299184-1	"Right knee swelled and felt like it dislocated; Right knee swelled and felt like it dislocated; Pain in groin; Felt like a muscle cramp; Blood clot as well as fluid on that knee/blood clot right leg; Muscle tear; the pain went to his knee; Pain in the arm opposite that he received the injection; his other leg ""got messed up""; Blood clot as well as fluid on that knee; Knee was stiff as a board; He could not walk on either of his knees for a couple of days; Inflammation in the calf; His leg was stiff; This is a spontaneous report from a contactable consumer (patient). A 67-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: ER8730), via an unspecified route of administration, administered in left arm on 30Mar2021 10:45 as single dose for COVID-19 immunization. Medical history included knee surgery, dislocated shoulder (he has dislocated that shoulder in the past several times but had no recent issues), carpal tunnel syndrome and, rheumatoid arthritis, all from an unknown date and unknown if ongoing. There were no concomitant medications. In Mar2021, the patient experienced pain in the arm opposite where he received the injection. He reported he like had a muscle tear that it has an intense excruciating pain. He took three aspirin and put a heating pad on it that didn't touch it. He then reported that the pain went to his knee and it ""got all blow up"" and stated that that knee had to be drained. He then stated that the calf on his other leg ""got messed up"" and was found to have a blood clot as well as fluid on that knee. He reported that it was as stiff as a board and that he could not walk on either of his knees for a couple of days. He reported that he still has problems with the one knee that ""blew up"" and had to be drained but stated that the inflammation in the calf where blood clot was has receded and is back to normal. He stated that he had to visit the emergency room twice and see an orthopedic surgeon for these issues and has been prescribed Xarelto to take to treat the blood clot. In the morning his leg was stiff. It was also reported that the patient's right knee swelled and felt like it dislocated on 16Apr2021, blood clot on right leg on 12Apr2021, and pain in groin on 13Apr2021. The pain in his groin went down, but his calf swelled to twice the size and it felt like a horse because he was favoring it. It felt like a muscle cramp. The orthopedic saw him for his left knee and he was seen in the Emergency Room for his right and they called his primary doctor. He tried to work and paid the price with his left knee. He has been taking Tylenol and icing it. The last two days (unspecified date) he has been babying it, the weather has been not so great, so he is laying low and give it a rest, but knows he is supposed to get up and move around so he has been doing that and elevating it. He would like to know if, based on these symptoms he experienced after the vaccine, he should get the second dose of the vaccine, which he states he is supposed to get on Tuesday. The outcome of the events ""Right knee swelled and felt like it dislocated"", and ""Pain in groin"" was recovering, and for other events was unknown."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299195-1	Blood clot, inner right upper leg; Chills; stomach upset; nausea; headache; This is a spontaneous report from a contactable consumer (patient). A 55-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 01Apr2021 11:30 (Batch/Lot Number: Er8735) as single dose for covid-19 immunisation. Medical history included covid-19 prior to vaccination. The patient has no known allergies. The patient's concomitant medications were not reported. The patient has no other vaccine in four weeks. The patient experienced blood clot in inner right upper leg, chills, stomach upset, nausea, and headache on 01Apr2021 at 20:00. The events were not treated. The patient was not tested for covid post vaccination. The patient had the second dose on 22Apr2021 at 11:30 AM in the left arm (lot: Er8731). The outcome of the events was recovered with sequel. No follow-up attempts are possible. No further information is expected.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299197-1	I had an ultrasound done on my leg where a blood clot was found. I had a chest CT performed where several clots were found; Two days after my vaccine I began having severe pain in my right leg in the calf area and pain in my chest when I took a deep breath or coughed. Over the weekend it got worse and worse, on Tuesday th; Pain in my chest; This is a spontaneous report from a contactable consumer (patient). A 51-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration, administered in left arm on 14Apr2021 13:00 (Batch/Lot Number: Ew0153) (at the age of 51 years old) as single dose for COVID-19 immunization. The patient medical history was not reported. Concomitant medications included duloxetine; trazodone; pregabalin (LYRICA); oxycodone hydrochloride, oxycodone terephthalate, paracetamol (PERCOCET [OXYCODONE HYDROCHLORIDE;OXYCODONE TEREPHTHALATE;PARACETAMOL]); and meloxicam, all taken for an unspecified indication, start and stop date were not reported. The patient previously took vancomycin and daptomycin and experienced drug allergy to these medications. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. On 16Apr2021 09:00, two days after the vaccine, the patient began having severe pain in his right leg in the calf area and pain in his chest when he took a deep breath or coughed. Over the weekend, it got worse and worse, on Tuesday 20Apr2021, the patient had an ultrasound done on his leg where a blood clot was found. He had a chest computerized tomography (CT) performed where several clots were found. The event resulted in doctor or other healthcare professional office/clinic visit. The treatment included blood thinners. No COVID prior vaccination and not COVID tested post vaccination. The case was assessed as non-serious by the reporter. The outcome of the event was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299222-1	This is a spontaneous report from a contactable consumer (patient). An 83-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on 03Feb2021 (Batch/Lot number was not reported) as a single dose (at the age of 83-years old) for COVID-19 immunisation. Medical history included the patient had known allergies (unspecified) and unspecified chronic health conditions from an unknown date. Prior to the vaccination, the patient was not diagnosed with COVID-19. The patient was not pregnant. The patient received unspecified concomitant medications. The patient did not receive any other vaccine within 4 weeks prior to the vaccine. On 15Feb2021 at 13:00 the patient experienced getting out of breath, three blood clots/clots were in my leg heart and lung, and cough. The events resulted in emergency room visit and the patient was hospitalized (from an unknown date) for the events getting out of breath, three blood clots/clots were in my leg heart and lung, and cough for 6 days. The clinical course is as follows: 2 or 3 weeks after the patient got the vaccine she started getting out of breath when she walked. Each day she got worse. Then the patient was taken to the hospital where they discovered the patient had three blood clots. The clots were in the patient's leg, heart and lung. The patient stated the Doctors do not understand where the clots came from. The patient was in the hospital 6 days. The patient stated they are still having breathing problem and still on oxygen as needed and they still cough at night. The patient asked if there have been any other reports of this happening to other people who got the vaccine as the patient never had blood clots before this. The patient underwent lab tests and procedures which included COVID test (nasal swab): negative on 15Mar2021. Therapeutic measures were taken as a result of getting out of breath, three blood clots/clots were in my leg heart and lung, and cough and included blood thinners and oxygen therapy and complete bed rest. The clinical outcome of the events getting out of breath, three blood clots/clots were in my leg heart and lung, and cough was not recovered. The batch/lot numbers for the vaccine, PFIZER-BIONTECH COVID-19 MRNA VACCINE, were not provided and will be requested during follow up.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299242-1	This is a spontaneous report from a contactable consumer(Patient). A 83-years-old non pregnant female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection Formulation, Batch/Lot Number: EN6200), via an unspecified route of administration on Arm left on 17Feb2021 10:00 AM as single dose for covid-19 immunisation. Medical history included colitis ulcerative. Concomitant medications included mesalamine (MESALAMINE). Patient had no past drug history. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. post Covid vaccine patient did not test for Covid. The Device Date given was 25Apr2021. The patient received treatment for raised blood thinner as Elequis 5mg. On 20Feb2021 at 10:00 (Three days after injection), the patient suffered severe pain in the right leg. She went to ER next day and blood clot was discovered behind right knee and later she subsequently had been seen by hematologist. The outcome of the events was not recovered. Follow-up attempts are required. Further information was expected.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299252-1	"blood clot it is from my hip all the way down to my left leg to my ankle; blood clot but it is the big one from my hip all the way down to my foot; foot going down the swelling went up to my leg/foot and my leg is swelled my toes all over up to my leg swelled; This is a spontaneous report from a contactable consumer (patient). An 85-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection; Batch/Lot Number: EW0162), via an unspecified route of administration, administered in left upper arm on 10Apr2021 at the age of 85-years-old as single dose for COVID-19 immunization. The patient's medical history included ongoing arthritis and ongoing chronic obstructive pulmonary disease (COPD). The patient reported that she was not taking anything for arthritis, just some Tylenol once in a while over the counter (it was not prescription). The patient was taking an inhaler for COPD. The patient was taking unspecified concomitant medications. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection; Batch/Lot Number: ER2613), administered in left upper arm on 20Mar2021 at the age of 85-years-old for COVID-19 immunization and experienced bladder infection and viral infection. The patient got the first Pfizer vaccine on 20Mar2021 and was alright for a few days afterwards. Then all of a sudden, she started getting infections on 28Mar2021, 8 days after she had the first vaccine. She got a real bad bladder infection then she went to the doctor and she got medication for that. She was with bladder infection then she came down with the viral infection. The patient reported that she has not even been going anywhere because of the virus and she has been staying home. She just goes to the grocery store and clinic to the doctor, that was it. The patient had an antibiotic for the bladder infection. Sulfamethoxazole was the name of the medication that was for bladder infection. They didn't give the medication for the viral infection. The patient reported that it was just a viral infection, it was not the Covid. Since then, she had her second dose of the vaccine on 10Apr2021 and from an unspecified date in 2021, her foot started, her foot swelled up and she didn't know. She takes lot of retention pills, so she thought maybe it was that. So she took her pills and everything and washed and start getting but the next day instead of the foot going down, the swelling went up to her leg and then the next day it went all away up to her hip. The swelling on the left side from the foot all away up to the hip. She went to urgent care on 21Apr2021 (reported as ""21st"") and saw a doctor and she was tested and she had blood clots. Regarding lab test, the patient reported she had two infections and then she got the blood clot that was from her hip all the way down to her left leg to her ankle. She could not figure out that they called the test and what they tested. But they had a lab work done then one has a test, veins, and everything. She can't remember. The patient reported that her foot and her leg was swelled, her toes all over up to her leg swelled. She had a fat foot and a fat leg just one side. She would say since she has started taking medicines for that, she took the first dose at the evening of 21st (21Apr2021), the day she went to the urgent care, and she would say it was not any worse. The medication was supposed to break up the blood clot, but it was the big one from her hip all the way down to her foot. Just a little bit better. The outcome of the events was recovering."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299281-1	bad pain in the leg, near the calf/ confirmed the patient had blood clot in the leg; hurting in the arm; chills; This is a spontaneous report from a contactable consumer (patient). A 54-year-old patient of an unspecified gender received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), via an unspecified route of administration, administered in the right arm on 15Apr2021 (batch/lot number and expiration date were unknown) as a single dose for COVID-19 immunization. The patient had no medical history. The patient was taking other unspecified medications. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 19Mar2021 at the age of 54 years old for COVID-19 immunization. The patient reported having the normal side effect like the hurting in the arm, chills, and stuff on an unspecified date, but after that the patient got a really bad pain in the leg, near the calf and left it go for a week maybe, thinking it would go away, but didn't go away, so the patient went in to the emergency room and the emergency room work on it. That's what they did something to come up wrong with one of the blood test, so then they did an ultrasound and they confirmed the patient had blood clot in the leg, for which the patient was currently on a blood thinner. The patient didn't know if this was going to fix the problem or this was not going to fix the problem. The patient knew that some of the other vaccine caused blood clots, but the patient's doctor won't give any kind of pain medicine, and all they do was to give paracetamol (TYLENOL). They did start the patient on a pill called apixaban (ELIQUIS) taken at 25 mg tablet in the morning and 25 mg in the evening, and this was the thing the patient had never had; anything like this had never happened with the patient before, since the patient had been healthy in all his/her life, so the patient called to ask if there would be any relief as the patient didn't want this. Before the patient went into the emergency room, the patient had taken TYLENOL as treatment for all the events. The patient underwent lab tests and procedures on an unspecified date, which included blood test and chest x-ray with unknown results and an ultrasound, which revealed blood clot in the leg. The outcome of the events was unknown. Information on the batch/lot number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299293-1	bloodclots in my leg and lungs; bloodclots in my leg and lungs; This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: unknown), single dose, dose 2 via an unspecified route of administration on 05Apr2021 for COVID-19 immunization. The patient's medical history was not reported. The patient's concomitant medications were not reported. The patient experienced blood clots in the leg and lungs in Apr2021 within days of receiving the second dose. The clinical outcome of blood clots in the leg and lungs were unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299332-1	Heart Attack due to blood clot in RCA. Occurred on 4/27/2021 at 11 pm; Heart Attack due to blood clot in RCA. Occurred on 4/27/2021 at 11 pm; This is a spontaneous report from a contactable consumer. A 57-year-old male patient received first dose of BNT162B2 (BNT162B2) via an unspecified route of administration, administered in Arm Right on 29Mar2021 03:00 (Batch/Lot Number: ER8733) as SINGLE DOSE for covid-19 immunization. The patient's medical history was not reported. Concomitant medications included pitavastatin calcium taken for an unspecified indication, start and stop date were not reported. The patient experienced heart attack due to blood clot in rca. occurred on 27Apr2021 at 11 pm. The patient was hospitalized for the reported events and was Life threatening (immediate risk of death from the event). The patient underwent lab tests and procedures which included sars-cov-2 test: negative on 27Apr2021 Nasal Swab. Therapeutic measures were taken as a result of the events which includes Emergency Heart Cath and Stent. The outcome of the events was recovered with sequel. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299351-1	My wife is having side effects that have to do with the formation of blood clots in the reproductive system.; This is a spontaneous report from a contactable consumer (consumer's wife). A female patient of an unspecified age received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Formulation: solution for injection, Batch/Lot Number and Expiration Date were not reported), via an unspecified route of administration on an unspecified date as single dose for covid-19 immunisation. The patient medical history was not reported. The patient's concomitant medications were not reported. The reporter has been reporting on COVID. His wife and he has been vaccinated. His wife was having side effects that have to do with the formation of blood clots in the reproductive system. Consulted in different forums in two countries there are many women reporting the same. The outcome of event was unknown on an unspecified date. Information on lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299360-1	This is a spontaneous report from a contactable consumer (patient). A 40-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 26Apr2021 08:15 (Batch/Lot Number: EW0171) (at age 40 years old) as single dose for covid-19 immunisation. Medical history included high blood pressure, high cholesterol, carpel tunnel, absent vas deferens, all from unspecified dates. No COVID prior to vaccination. He underwent Covid test post vaccination (nasal swab) on 29Apr2021 with negative result. The patient's concomitant medications included unspecified prescribed and store-bought medications. No other vaccines in four weeks. On 28Apr2021 22:00, the patient experienced chest pains and shortness of breath testing of EKGs, echo, CT scan, and cardiac cath show- NStemi heart attack showing from damage to heart caused by blood clot. No problems with arteries or existing with heart. Also, lower lobe of left lung shows collapsed. The events resulted in doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. The events were considered serious: hospitalization (2 days), life threatening illness (immediate risk of death from the event), disability or permanent damage. Treatment included blood thinners- apixaban (ELIQUIS) and metoprolol. The patient was recovering from the events.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1299361-1	Blood clot on the cerebellum that caused a stroke; Blood clot on the cerebellum that caused a stroke; This is a spontaneous report from a contactable consumer (patient). A 29-year-old female patient (not pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: EW0167), first dose via an unspecified route of administration, administered in Arm Left on 30Apr2021 11:30 (at 29 years old,(not pregnant), single dose for covid-19 immunisation. The patient's medical history was not reported. The patient's concomitant medication included an unspecified vitamin. The patient previously took ibuprofen and experienced Ibuprofen drug allergy. On 02May2021 08:30, patient experienced a Blood clot on the cerebellum that caused a stroke. The outcome of the events was recovering. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The adverse event resulted in visit to Emergency room/department or urgent care. Treatment received for the adverse event was Blood clot meds. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has been tested for COVID-19 (02May2021), negative for swab test. Events caused 3 days of hospitalization. Information on Lot/Batch number was available. Additional information has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1301127-1	Menstrual irregularities. Light spotting for at least one full week (never experienced). Heavy bleeding and clotting (currently happening). This is day 16 of this. I have experienced heavy bleeding in the past but it only lasts 2-3 days at the most (usually only 2 days). Clotting is heavy unlike anything I've ever experienced. Minor soreness of nipples shortly after 1st vaccine, which was constant for 4 weeks but has since stopped (this has never happened).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1301289-1	Severe joint pain, fevers, severe headaches, cough, severe fatigue, the first week. Second & 3rd week -Positive D dimer,low o2 sat.s. high fevers and body/joint pain. 4-6 week-Hemoptysis, Systemic inflammattory response syndrome, sepsis, acute bronchitis, uveitis and vertigo. Info below is for answer 21 1 admissions to Hospital was 4 days 2nd admission to Another Hospital. 6 day stay Most of the treatment was antibiotics, oral steroids, steroid drops for my eyes and breathing treatments to open my airway.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1302270-1	2 days after 2nd shot, noticed left underarm area was enlarged. Felt Lump. Date of shot: 4/10/2021. Date noticed lump: 4/12/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1302470-1	DVT, LLE pain, occlusive thrombus
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1302479-1	Blood Clots in left leg and both sides of lungs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1303074-1	1st dose in series on 03-18-2021; 2nd dose in series on 04-08-2021 2 days after 2nd dose, he developed severe pain in both legs. He was taken to the hospital where clots were diagnosed in both legs and groin area. A previously undiagnosed lung nodule was found and he was given a cancer diagnosis. He was discharged home on apixaban (Eliquis) and comfort measures 2 days after admission. Over the following weeks, he had a permanent lung drain placed. He died on May 5th or 6th at home. No autopsy performed.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1303441-1	Prolonged heavy menstrual cycle with excessive blood clot starting on 3/22/21 till present 5/10/21. Despite intervention by ob/gyn such as performing pelvis ultrasound and performing a uterine biopsy with negative result for cancer as well as treating me with medroxyprogesterone for a course of 15 days that I completed as prescribed . After the medroxyprogesterone, I have to be placed 3 days ago on birth control pill, the menstrual bleeding continues . It?s been almost for 50 days. Need help!
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1304151-1	Patient was found at home Saturday afternoon passed out. Patient was rushed to the hospital where it was discovered he had a blood clot. He is going to hospitalized for a few days per patients mother.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1304179-1	Cerebellar thrombotic stroke 1/28 manifested by sudden onset dizziness and vomiting. Symptoms not improved with fluids and rest. Misdiagnosed 1/29 in ER as vertigo. Not diagnosed until 1/30/2021 upon readmission to ER. Treatment: sub-occipital craniotomy, steroids, and 3 days ICU with ICP monitoring. Outcomes: chronic left cerebellar atrophy with minimal clinical symptoms
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1304182-1	My left leg and ankle became very, very swollen and sore. I experienced severe pain in my left thigh, left knee and left calf whenever I sat, walked, went up and down stairs, while sleeping and I was limited in what activities I was able to do without pain. I went to the doctor on 3/29/2021 (9 days after my first shot) and the doctor sent me for an ultrasound of the left leg and an X-ray of the left ankle..
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1304689-1	Blood clot

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1305699-1	swelling behind the knees and swelling of the veins; it had to do with blood clot warnings / it was some kind of clotting because of the position of the areas of swelling.; she got like three balls on the right arm; pain in her left arm from the armpit to the elbow, it was a line of pain going down the arm; the lymph nodes in left armpit got swollen; swelling behind the knees and swelling of the veins/ she had swelling under her armpit and swelling on her right arm below her elbow; This is a spontaneous report received from a contactable consumer. A 48-years-old female patient received bnt162b2 (BNT162B2), dose 1 via an unspecified route of administration, administered in Arm Left on 12Apr2021 10:30 (Batch/Lot Number: ER8729) as SINGLE DOSE for covid-19 immunisation. Medical history included covid-19 from Nov2020 to an unknown date. The patient's concomitant medications were not reported. The patient experienced it had to do with blood clot warnings / it was some kind of clotting because of the position of the areas of swelling on 15Apr2021, swelling behind the knees and swelling of the veins, swelling behind the knees and swelling of the veins/ she had swelling under her armpit and swelling on her right arm below her elbow on 12Apr2021, the lymph nodes in left armpit got swollen on 13Apr2021, pain in her left arm from the armpit to the elbow, it was a line of pain going down the arm on 14Apr2021, she got like three balls on the right arm on 15Apr2021. The patient underwent lab tests and procedures which included ultrasound scan: unknown results.She was calling because she got her shot on 12Apr2021 and then she ended up having swelling behind her knee, then she had swelling under her armpit and swelling on her right arm below her elbow. She stated that they weren't lymph nodes, and she went to the ER (EmergencyRoom) had to check and see if they were blood clots. She stated that it turns out that her veins were swollen underneath her arm. She said that was what happened after the shot and that was her first shot. She states so that's her report on what happened. She stated she got the shot in the morning so her symptoms at 6 or 7 pm that evening. She stated she went to the doctor first and then went to the hospital. She went to the doctor on the 16Apr2021 and the hospital on 16Apr2021. She stated that the swelling behind her knee went away on 20Apr2021.Tuesday, 13Apr2021 the lymph nodes in left armpit got swollen. On Wednesday, 14Apr2021, there was a pain in her left arm from the armpit to the elbow, it was a line of pain going down the arm. On Thursday, 15Apr2021 she states she got like three balls on the right arm, on her elbow on the right side. She went to the doctor on Friday 16Apr2021 she went to the ER (emergency room) because they couldn't figure out if it was some kind of clotting because of the position of the areas of swelling. She got an ultrasounds for them to check her blood flow. She stated that the pain in her left arm was going down but didn't resolve until the 22Apr2021. For the little balls on her arm, the MD (Medicne Doctor) gave her a medication for the swelling of her veins but she forgot the name and she doesn't have the paper with her. She states the balls of fluid went away between the 20Apr2021 and 22Apr2021. Everything starting going away that week.She states it had to do with blood clot warnings, and swelling under her skin had to do with the veins. Outcome of with blood clot warnings was unknown, like three balls on the right arm and pain in her left arm from the armpit to the elbow was recovered, remaining events was recovering.Therapeutic measures were taken as a result of swelling behind the knees and swelling of the veins/ she had swelling under her armpit and swelling on her right arm below her elbow, she got like three balls on the right arm.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1305754-1	Blood clot in right calf; This is a spontaneous report received from contactable consumer for herself. A 42-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) dose 2 via an unspecified route of administration, administered in Arm Left on 11Apr2021 17:15 (Batch/Lot Number: EW0151) as 2nd dose, single for covid-19 immunisation. No pregnancy at time of vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient medical history allergy to penicillin. The patient's concomitant medications were not reported. Historical Vaccine included the first dose of BNT162B2 on 21Mar2021 05:15 PM in Left Arm (Batch/lot number: Ep6955) as 1st dose, single for COVID-19 Immunization. The patient experienced blood clot in right calf on 15Apr2021 03:00 AM. The adverse event result in doctor or other healthcare professional office/clinic visit and Emergency room/department or urgent care. Therapeutic measures were taken as a result of blood clot in right calf included Blood thinners. No Hospitalization Prolonged. Prior to vaccination, the patient did not diagnose with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. Outcome of the events was recovering.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1305769-1	she thinks may be blood clot or a muscle but not really sure; passed out; very anxious and nervous; her heart is beating fast and she is nervous.; burning sensation in her ear; tiredness; headache; heaviness on her back and chest on the left side; heaviness on her back and chest on the left side; no appetite; nauseous; diarrhea; throbbing is almost like a band-aid is too tight. If she rubbed her hand across her thigh she could feel crushing and throbbing; her heart is beating fast and she is nervous.; sharp pain in her thigh/pinching sensation on inner thigh; sick; throbbing; muscle pain; weird sensation; throbbing sensation to the left side of her belly button/It was like at her stomach; fever of 100.3F; had a feeling of heat on her neck, face, and ears; redness and itchiness at the injection site; redness and itchiness at the injection site; the injection was large, red, hot, and itchy; the injection was large/ the swelling went down; dizzy; This is a spontaneous report from a contactable consumer reported herself. A 56-years-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 22Apr2021 (Batch/Lot number was not reported) as 1st dose, single for covid-19 immunisation. Medical history included disability, obese, covid-19 when she

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				immunisation. Medical history included disability, obese, covid-19 when she was sick with COVID last year, she has never been that sick. she thinks she made it through because she owns a nebulizer and can say her lungs have never felt like they were full of fluid. If anything, they felt heavy. It was heavy in her back and chest. She also had a fever. She guesses she would be considered obese. Concomitant medications included paracetamol (TYLENOL). The patient received the first dose of the Pfizer COVID vaccine last week and she has experienced several side effects and is very anxious and nervous. She experienced redness and itchiness at the injection site, pinching sensation on inner thigh that she thinks may be blood clot or a muscle but not really sure, fever of 100.3F from 24Apr2021, burning sensation in her ear, tiredness, headache, heaviness on her back and chest on the left side, no appetite, nauseous, diarrhea, had a feeling of heat on her neck, face, and ears, throbbing sensation to the left side of her belly button from 26Apr2021 and she felt dizzy from 22Apr2021. The patient stated there was a bad connection and she was having difficulty hearing. The patient was a little nervous. Maybe this was a psychosomatic issue and she was thinking of things too much. She was sick for 5 weeks. She was never in the hospital or intubated. The site where she had the injection was large, red, hot, and itchy. It was from 22Apr2021 (Thursday) for the next maybe 3 days. Now the color is diminishing, the color is going down more like to pink and smaller and it not itching anymore. Once in a while she got a pinching sensation on her inner thigh ever since she had the injection. In the last couple days, she has been feeling a that a little more. She wanted to make sure it was not a symptom of a blood clot. She doesn't feel short of breath. She was nervous talking the call handler so that is why she is talking fast. She doesn't want to feel so sick like she did last year when she had it in March or April. She wanted to know that it's not a symptom of a blood clot. It was a throbbing pinching pain that goes and doesn't come back for a little bit, but it might come back a little bit later. It happened when she woke up. She has no swelling in her legs or thighs or anything else. She says she would not be talking this fast, but her heart was beating fast and she was nervous. She tended to be an anxious and nervous person. The patient never had blood clot issues. She wanted to make sure with everything that she has heard about Pfizer she hasn't heard of anyone having blood clot issues. She had to read to calm herself down a bit. It is a sharp pain with throbbing that comes and goes. She didn't read where it could be a blot clot, she just wanted to know. She is horrible at running to the hospital when she feels something because she gets so nervous and she thinks knowing is better because she's a realist. It takes her a minute to go to the hospital to figure it out. If she continues to feel like this she and her daughter will go to the emergency room. It is 5 blocks away. It is a small little hospital and they will be able to ease her mind. It could be her nerves getting to her too. Her fever got up to 100.3 and that was only on Saturday, 48 hours after the vaccine. Once she took Tylenol it was gone and didn't return again. She took her temperature; it was a beeping one. She always says she would take like 2 out of 3, and every time she took it went up. She felt a weird sensation of burning in her ears, the same as when she first had Covid, feeling like a fever. It brought her back to a bad place and it freaked her out. She is on disability and has people go to the store for her. She did that for 3 to 4 months. She is not agoraphobic, but she hadn't gone out until she got the vaccine. It took her all day to get back home because she loves the outdoors. It is kind of sad, and she knows she is not the only one who experienced that. She was able to actually stay at home, and when she went out, she wanted to stay. She still has her second dose to take, not everyone in has this and she doesn't want to feel that sick again. Eventually she will perish away one day, from God knows what but a slow death of not being able to breathe she cannot picture for herself or anybody else. She tries not to watch the news, it was depressing, because it is all over the news, she tries to stay in the loop. She was tired and had a headache. She took Tylenol and slept for an hour or hour and half. When she woke up she felt heaviness on her back and chest on the left side, and as she moved around more it was on her back and less on her chest. It was the same as she felt when she had Covid. She didn't have any aches or pains. The other thing was she had no appetite for the next 3 days. Every time she woke up, she felt extremely nauseous but had nothing to vomit. For the next 3 days or 72 hours, she had diarrhea. If she ate anything, she was in the bathroom. On 24Apr2021 (Saturday) after 48 hours she started to feel the heat in her neck and face and ears like right on her ears. She was talking to her daughter and her daughter said these are normal. Where she got the injection was itchy and a little more red. She told herself not to scratch it in her sleep, she is one of those people if something itches, she scratches. What she would do is rub her arm up and down on the site. It was almost the size of an 8 ounce tomato sauce can or maybe the size of a baseball. Then the third day it got redder. In her sleep she was rubbing her arm because it got itchy. Today the swelling went down, the color is like a light pink, and the size is also minimized. The fever lasted that night. She took her temperature again the next day and it was 97.9. That is usually what it is when she wakes up. She is not just tired; she is exhausted and feels like she has to go to sleep. Her daughter came with her granddaughter who is a baby, and she didn't know how to tell them. She was happy to see them but so exhausted. She worries about the sharp pain in her thigh. The other thing is she started getting pain in the middle of her thigh, her inner thigh. A sharp pain with like a throbbing is almost like a band-aid is too tight. If she rubbed her hand across her thigh, she could feel crushing and throbbing. She can feel it from touching the area. She had this before like when she is stretching. It started maybe 3 months ago and went away but then after she took the injection and heard about thing, she worried about a blood clot with Johnson & Johnson, not that she would know because she has never had a blood clot. It could be psychosomatic and all working in her head. It started the last 2 days when she woke up, she felt fine and as she was lying in bed, she started feeling a

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				she woke up, she felt fine and as she was lying in bed, she started feeling a sharp throbbing pain that eventually goes away. She has no swelling of her legs or thighs, and her daughter said it looks fine. She is semi-terrified of hearing bad news. It takes her longer to go to the hospital. If she starts feeling worse, she will be there in no time. She does not want to wait until it is worse, but she does not want to spook herself into going. She suffers from anxiety. She definitely had the fever not like she was just tired it was like she was exhausted. She got up early and would take a little nap because she was doing a bunch of stuff. That hasn't happened, but that's her. It will be 2300 when she goes to sleep. She was doing a bunch of stuff and felt like she just needed to take a nap. Maybe it is part of that too, but usually she doesn't feel tired like this. After her daughter left, she read something on her phone and passed out. When she woke up, she saw she actually dropped her phone. Her thigh is now not a pinch but more a little bit of a muscle pain. Now it is not like the inner thigh, it is more behind the thigh now, but not below the knee, right behind the thigh, closest to the knee but not on the knee. It is the same thing as if the middle had gone behind her thigh a bit. The little pillow on her chair is not helping anything either. It feels like it maybe muscular, its deep feeling. It feels like it could be muscle, if she rubs it, she can feel it. When asked has it improved persisted or worsened, caller states it is there, but it is dull. She can say there is pain, but it is a little duller just because it is more in the muscle. It is hard to explain, it was not the same throbbing pain but there is a dull pain, it goes around the chunkier part of her thigh, like the inner thigh. When she rubs it there is pain but a dull pain. When she rubs it, it feels better but she can still feel the site where it hurts. When she rubs it, it is still located right in the lower middle going toward her knee but not at the knee, maybe 3 to 4 inches away from the fold of her knee, almost at the center of her thigh almost. Almost like a dull muscle ache, there is a bit of throbbing with it. She will go to the doctor if something weird happens and will call back when they have an idea of what happened to her. This will sound crazy. The next thing she felt was a light throbbing where her belly button is but more to the left side, but not at the belly button. It was like at her stomach, not right at the belly button but right on the same line as her belly button but more at her waist. It was a small throbbing started today about an hour ago. She is hoping it is gas. Her fever went away pretty fast. She was on the phone and took her temperature to tell what it was. The woman said to take some Tylenol. It went from 99.9 to 100 to 100.3. She thought oh no. That was the highest she saw. Then eventually it started going down to 99.4, and then by morning it was back to normal. She only took Tylenol the one time. She is thinking maybe it is from being nervous and causing her body to heat up. She doesn't know she is just thinking. She thinks the fever went away the same day within an hour to 2 hours. She also wasn't drinking enough water. When asked if her fever has recovered completely or with last effects, caller states that is fine now, her temperature today was 99.4 without the fan on, she says she thinks that was pretty normal. When she first wakes up it is 97.9 usually. There was also a 98.4. The redness and itching at the injection site started 3 hours after the injection. She took her 2 sons with her. She took it and then her son took it. They said to wait 15 minutes. As soon as she stepped out, she looked up and felt so dizzy. It looks like when in summer and it seems to be boiling hot, that is how it got for her. Things got blurry. It was not like she was going to black out, but she had to close her eyes because the room was going to spin. She knocked back on the door and took a seat, so she sat down because she was not feeling right. When she gets nervous, she chews gum as a coping mechanism. Her son got some gum for her. She felt better but kind of nervous waiting for 15 minutes to go by, what if she felt dizzy again. She let 20 minutes go by and she got up and was fine. She wanted to be out of there and in the cool air outside. That is the first reaction that she had. She is one of those people that feel so nervous she is sure that happens that they feel like, oh they are so nervous, and they read so much, and it is in our faces everyday no matter what. Maybe it had nothing to do with the injection, maybe more that she was so nervous, but it was a weird sensation after. She knows when people first started getting the COVID vaccine, people were just like falling to the ground, she was like that would be terrible. She went with high expectations and was also nervous. Her heart was beating, and she could hear her own heartbeat. She still feels tired and like she can go take a nap now. She asks if that is normal. She would rather talk to Medical Information now. She is over this statistics part of it. She doesn't feel a release, she is just giving her information and she doesn't feel any better. Outcome of the event throbbing sensation to the left side of her belly button/It was like at her stomach was not recovered; of the event dizzy was recovered; of the events the injection was large, red, hot, and itchy/ the swelling went down, tiredness was recovering; of the other events was unknown. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1305784-1	Thrombosis; Vericose veins; Swollen, bulged nerves around left hand palm and fingers/The same symptoms noticed on right foot; Nerves are in black darkened color and the palm fingers, knuckle areas; the palm, fingers, knuckle areas had moderate pain. The same symptoms noticed on right foot.; the palm, fingers, knuckle areas had moderate pain; sensitive to touch; This is a spontaneous report from a contactable consumer(patient). The 36-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in arm left on 24Apr2021 14:45 (Lot Number: EN6205) as first dose, single for covid-19 immunization. Medical history included none. He had not known allergies to food or medications, no illlness history. Concomitant medications included minerals nos, vitamins nos (CENTRUM A TO ZINC) adutls vitamin, miconazole nitrate (LOTRIMIN AF ANTIFUNGAL) topical cream. The patient experienced one hour after first dose of vaccine shot at left arm shoulder, he saw swollen, bulged nerves around left hand palm and fingers. Nerves were in black darkened color and the palm, fingers, knuckle areas had moderate pain. The same symptoms noticed on right foot. The affected areas on both palm, fingers, foot area have dark patch areas with moderate pain. After 32 hours, he saw swollen nerve back of left leg thigh area with deep pain. It was sensitive to touch. It looked like thrombosis or vericose veins not sure about it. He reached out to doctor office and they said to wait for another day to see if he can recover. whereas he still have the darkened black nerve with pain in the affected areas. Events occurred on 24Apr2021 16:00 which was resulted in doctor or other healthcare professional office/clinic visit. There was unknown if treatment received. Outcome of events was not recovered. There was no covid prior vaccination and post vaccination.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1306670-1	On 3/23/21 pt had SOB so went to went to the Hospital ER. She had lab work, CT, Chest Xray, ECG 12 lead. DX is pneumonia of right lower lobe of the lung. She was given Augmentin and released after 7 hours and told to FU w/ PCP. On 3/31/21 pt went to see her PCP due to having SOB. She was given a Covid Test which was negative. She was sent home and told to breath deeply and hopefully this symptom would go away. On 4/18/21 she went back to the Hospital ER for SOB. Pt was admitted with acute DVT and other blood clots as well as pleural effusion of the right lung. They did US on legs and Chest X-Rays. They found fluid around her right lung which they drained. She was prescribed Eliquis. Pt was discharged on 4/21/2021 and was told to FU w/ PCP. On 4/24/21 pt went back to the Hospital ER, because she had a bruising/blood clots in the groin area. She had 3 blood clots on the left side and one on the right side. They told her the bruising was from the Eliquis and they did a US. She was told she had the same blood clots that she had during her hospital stay and they were right where they were before. She was discharged. On 4/29/2021 she went back to her PCP due to SOB. She told her PCP about the DVT that was bothering her. She was sent to another Hospital. She had a stat Chest X-Ray- results were that her lungs were stable with inflammation around the right lung. She was told to FU w/ PCP in July 2021 and go back to ER if she had another episode w/ SOB.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1307066-1	Blood clot in right calf and part of the thigh
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1307192-1	On April 8, 2021, I went to Hospital with symptoms of blood clots in my right leg. I was diagnosed with Superficial thrombophlebitis - lower extremity
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1307481-1	Patient is a 48-year-old male with patient history significant for CAD, CHF, ICD, HTN, hyperlipidemia, and smoking who presented to the ED yesterday morning with 1 day history of chest pain. Patient described the CP as moderate aching in the substernal area that did not radiate. The pain was intermittent, with each episode lasting 1-2 minutes. The CP was associated with LUE numbness and SOB on exertion. The chest pain and LUE numbness have now resolved. Patient also reports a three week history of RUE swelling. The swelling gradually worsened and is now associated with pain and proximal RUE bruising. RUE pain is rated 5/10. In the ED, patient was found to have NSTEMI and small right lower lobe segmental pulmonary artery embolus. Venous Doppler was positive for occlusive thrombosis in the R subclavian, basilic, and cephalic veins. Patient was started on IV heparin. Patient is scheduled for cardiac cath today. Hematology is consulted regarding PE and RUE DVT. Patient denies personal or family history of blood clots or recent travel. Reports receiving COVID vaccine in LUE approximately three weeks ago. Also admits to being more sedentary recently due to being unemployed. Denies nausea, vomiting, bleeding, bowel changes, fever, or chills. Significant PE findings: RUE edema, firm and tender to palpation, ecchymosis on proximal RUE, distal sensation and active ROM intact
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1308494-1	Swelling, cramping, redness, pain, and warmth in left leg. He went to Urgent Care. The ultra sound lady said she saw a small clot. The PA he saw reported the radiologist did not diagnose a clot. The PA cautioned from prescribing any blood thinners for fear of interacting with DM medications (metformin and lisinopril).
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309527-1	have some clots; Neuropathy; autoimmune disorder; Fell and hit elbow and 3 ribs on left side; lost 6 (lbs) pounds/Within 2 months, he has lost 47 lbs and eats all the time and is still losing weight; falling; felt super strong; dizziness; failed all the tests for motor skills for his arms and legs; Fell and hit elbow and 3 ribs on left side; Fell and hit head; Sensory nerves are all gone; Fatigue; scabs; delusional; tore tendons in between ribs; They are

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	tender(ribs); hurting all over; Cannot walk or lift anything; chest pain; his **Adverse Event Description**
				arm was sore a little; Left arm injection site pain; Disoriented; This is a spontaneous report from a contactable consumer (patient). A 49-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 1 intramuscular, administered in left arm on 03Feb2021 (batch/lot number and expiration date not reported) at the age of 49 years old, as 1ST DOSE, SINGLE for COVID-19 immunisation. Medical history included COVID-19 from Dec2020 to an unknown date, vascular disease, bilateral stents placement, cataracts since he was born. Concomitant medication included gabapentin taken for an unspecified indication from an unspecified start date and ongoing. On 04Feb2021, the patient experienced left arm injection site pain. The patient experienced dizziness, falling, and couldn't go to work on unspecified date. He was treated by his (MD) physician, was given a CT scan, echocardiogram, and found to have some clots. He stated he has a history of vascular disease and had bilateral stents placed a year ago. He was placed back on an aspirin/Plavix regimen and referred to a neurologist because he has no motor skills anymore. He stated he failed all the tests for motor skills for his arms and legs. He was advised not to take the second dose of the vaccine and was diagnosed with neuropathy and autoimmune disorder (caller does not specify which disorder). He reported being positive for Covid-19 in Dec2020. He wanted to know what else he can do for his symptoms and is offering to be tested by Pfizer for treatment. The symptoms come in really hard seven days later. He actually had to go to the (ER) emergency and they worked on him and a whole string of doctors. He went to the emergency room and something was going on and wanted him to go to a family practitioner. They sent him to vascular, neurological and now they want to send him to a research center. It was a whole week after receiving vaccine before he began having symptoms. With the neuropathy and autoimmune, he cannot work like he did. He can hardly walk. Unspecified autoimmune was diagnosed last week. He can't remember the name of it. They told him not to take second one because the first one did what it did to him. It hurt a little and it was quick. The next day (04Feb2021), he said his arm was sore a little. He is only taking 4 medications and they looked it up. He did not think they were relevant. He was disoriented and fell and hit his head a couple of times. They ran an MRI on his head trying to figure that part out. That was done last week. He doesn't know the date and said it may have been on Friday. Results were good. The date he was disoriented was Wednesday 03Feb2021, when he took the shot. He has been disoriented and has been that way ever since. They kept him for 15 minutes to make sure no one passed out. He has been disoriented since then. He could not think clearly, and he could not do his job. He tore tendons in between ribs. He first thought he actually broke his ribs, but did not. There were two different falls. He cannot feel feet or anything. His motor skills are OK but when they did neuropathy test, they put needles in and shocked him to make his motor fire. His reflexes were really bad. He fell and hit elbow to keep from falling on face and twisted going down hitting 3-4 ribs on left side. It has been over 3 weeks now for his ribs. They are tender but they are not like the initial blow. It feels like they are healing. He fell and hit head a week ago. He was unsure of the exact date. He did not go to the emergency room with this one because the bed caught him. He fell and hit his head. All he has left is scabs. He cannot walk or lift anything (11Feb2021). That next Wednesday, he got ready to go to work and was so delusional and could not go in. He slept from 5:30 all that day and all that night. Then Thursday, he gets up and goes to work and goes slower, just to see what happened on Wednesday. He had fatigue and was disoriented. He did not provide start date or outcome for fatigue. He walked in and said either call an ambulance or he would go home. He went home and that was the last day of work. He then stated he laid in the car for 30 minutes and he said he is going to the emergency room. He went straight there that day. He had stents put in lower part of his body a year before. The neuropathy and the autoimmune kicked in 7 days later after the shot. The reason he went to the ER is because he thought it was his stent. He had bilateral, 4 of them. He was having chest pain and hurting all over on 11Feb2021. He thought he was having a heart attack. He was seen in the emergency room and released. It depends on how far he goes, how the chest pain worsens. Right now, he is ok, but when he gets up and moves around, it does the same thing. Within 2 months, he has lost 47 lbs and eats all the time and is still losing weight. He was about 200 something and was kind of built and now is skin and bones. He lost 6 (lbs) pounds within a week and is eating like crazy. Sensory nerves are all gone. If hit with a baseball bat and he would not feel it. It came on slowly almost that he did not know it was happening. That is why they want to send him to Privacy for a research center and he does not want to go. He never had this happen before. ER or physician's office required: 1st date was on Thursday, 11Feb2021, when he went to Privacy. He was having chest pain and hurting all over. He thought he was having a heart attack. He was released. The patient stated he wanted to change the information he had provided since he was not in his right mind since he had taken the wrong medication dose than the doctor had given him. He stated the medication dosage he wanted to change was for Amiripyin. He stated he wanted to say he told the representative the symptoms he had experienced was enhanced which they were yesterday because he was trying to fight everything and stay awake. COVID 19 vaccine: he stated the doctor did not want him to take the second dose of the vaccine. He stated one of the symptoms he had experienced after the vaccine that he wanted to change the outcome for is the neuropathy. He stated he is back to where he used to be and yesterday it was bad he could hardly walk or talk. He stated also the other symptoms he wanted to change the outcome for is the autoimmune. The patient has not recovered from neuropathy, autoimmune disorder, disoriented, can't walk or lift anything, chest pain, and sensory nerves all gone; while outcome of the

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				remaining events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309550-1	leg formed a blood clot above knee; Severe pain; Numbing in arm and leg; This is a spontaneous report from a contactable other healthcare professional (patient). A 52-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 23Mar2021 10:00 (Batch/Lot Number: unknown) as 1ST DOSE, SINGLE for Covid-19 immunization. The patient was not pregnant at the time of vaccination. Medical history included complications from bladder mesh and mitral valve prolapse. The patient's concomitant medications included Vitamins (unspecified). The patient was not diagnosed with COVID-19 prior to vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. On 24Mar2021 06:00, the patient experienced severe pain followed by numbing in arm and leg, and leg formed a blood clot above knee. The patient tried to make doctor appointment but 40 days out. The patient took aspirin daily for clot and will see doctor in 2 weeks. No treatment was received for severe pain and numbing in arm and leg. The patient has not been tested for COVID-19 since the vaccination. The outcome of events severe pain and numbing in arm and leg was recovering; outcome of event leg formed a blood clot above knee was recovered on an unspecified date. Information on the lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309553-1	blood clot in one of his leg; leg became red; This is a spontaneous report from a contactable consumer (patient). A 73-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in left arm on 23Apr2021 (at the age of 73 years old) (Lot Number: EW0169, unknown expiration) as 1st dose, single for COVID-19 immunization. Medical history and concomitant medications were reported as none. The patient received his first dose of COVID-19 vaccine on 23Apr2021. In the night of 24Apr2021, his leg became red and he spoke to his doctor. The experienced a blood clot in one of his leg on 25Apr2021. The patient received Eliquis for the events to be taken for 45 days. He would like to consult if he should receive the second dose after the side effect from the first dose. Patient would like to know if there have been reports of blood clots after receiving the COVID-19 vaccine. The patient is scheduled to receive his second dose on 14May2021. The patient underwent lab tests and procedures which included blood test and ultrasound on leg: unknown results on 26Apr2021. Outcome of the events was unknown. No follow-up attempts are possible. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309554-1	Calf muscle pain since 1 week. Clots upto 1 inch near the knee area. Observed on both legs; Calf muscle pain since 1 week. Clots upto 1 inch near the knee area. Observed on both legs; This is a spontaneous report from a contactable consumer. A 29-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), dose 2 via an unspecified route of administration, administered in Arm Left on 21Apr2021 16:00 (Batch/Lot number was not reported) as 2ND DOSE, SINGLE for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient received BNT162B2 for COVID-19 immunization on 31Mar2021 (Lot unknown=True, Lot unknown reason=Unable to locate or read the details, Administration date=31Mar2021, Administration time=02:00 PM, Dose number=1, Vaccine location=Left arm, No reaction.) On 24Apr2021, Calf muscle pain since 1 week. Clots upto 1 inch near the knee area. Observed on both legs. The patient did not receive treatment for the events. The outcome of the events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309557-1	"blood clot in her left calf; This is a spontaneous report from a contactable consumer, the patient. A female patient of unspecified age received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration on 24Feb2021 as a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot: EN6203) on 02Feb2021 for COVID-19 immunization. On an unspecified date, the patient experienced pain and on 15Mar2021 it was confirmed that the patient had a blood clot in her left calf via ultrasound performed on 15Mar2021 with positive results. The patient was treated for the blood clot. The clinical outcome of ""blood clot in her left calf"" was unknown. Information on the lot/batch number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1309578-1	To check if it was blood clot; Intermittent long-lasting palpitations; Lymph discomfort in one leg; Severe pain of bones in chest; Joint pains; Severe shooting intermittent head pain; Cold sweat; Felt like she was going to faint; COVID arm popped out; Severe nerve pain behind ear same side as vaccine; Exacerbated back pain; Bad menstrual cramps; This is spontaneous report from contactable consumer (patient). A non-pregnant 62-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot Number: ER8727), via an unspecified route of administration in left arm, on 31Mar2021 at 01:00 PM (13:00), as first dose, single, for COVID-19 immunization. The patient's medical history included fibromyalgia, complex regional pain syndrome (CRPS), lymphedema (previously resolved), mild asthma, osteoporosis, herniated discs, allergies, and menopause. The patient's concomitant medications included unspecified vitamins. The patient had no other vaccine in four weeks (however, it was also reported that other vaccine was administered in same date, 31Mar2021; pending clarification). The patient was not diagnosed with COVID-19 prior vaccination. On 03Apr2021 at 02:00 AM, the patient experienced intermittent long-lasting palpitations, lymph discomfort in one leg for which they went to check if it was blood clot. On 03Apr2021 at 02:00 AM, the patient also had severe pain of bones in chest, joint pains, severe shooting intermittent head pain, cold sweat, felt like she was going to faint, COVID arm popped out (nite 8; as reported), severe nerve pain behind ear same side as vaccine, exacerbated back pain, and bad menstrual cramps for 2 mornings 11 years post-menopause. The adverse events resulted in doctor or other healthcare professional office or clinic visit. No treatment was received for the adverse events. The events were considered non-serious. The patient has not been tested for COVID-19 post-vaccination. The patient had not recovered from the events. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1309626-1	part of my left kidney was no longer receiving blood; blood clot; placed on stroke protocols.; I developed severe abdominal pain in my left side during the evening; lack of appetite; bouts of severe discomfort; This is a spontaneous report from a contactable consumer (patient). A 40-years-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Batch/Lot Number: EW0167) via an unspecified route of administration, administered in Arm Left on 28Apr2021 12:15 (40-years-old) as 2nd dose, single for COVID-19 immunization. The patient medical history was not reported. Vaccination Facility Type was clinic. Concomitant medications included ibuprofen (ADVIL); desloratadine (CLARINEX). The patient previously took first dose of BNT162B2 (lot number=EN6208) on 07Apr2021 01:15 PM (40-years-old) on Left arm for COVID-19 immunization. It was reported that the patient developed severe abdominal pain in my left side during the evening of Wednesday, 28Apr2021 19:00. The pain appeared sporadically throughout the evening and continued through Friday, April 30. During this time, he experienced a lack of appetite and bouts of severe discomfort. Tried treating with GasX and Tylenol. Duration of Hospitalization reported as 3 days. On Saturday, 01May2021, starting around 10:30 a.m., the pain in his left-side abdominal area intensified and did not diminish. After 6 hours, he visited the ER at hospital. After being admitted to the ER, a series of blood, urine and imaging tests were performed on me. The doctor on duty reported that a CT scan showed that part of my left kidney was no longer receiving blood (disability). He was placed on Heparin and admitted to the hospital. He was diagnosed with a blood clot and placed on stroke protocols. The adverse event result in Emergency room/department or urgent care. Treatment received for the adverse events was Heparin, fluids, blood work, urine analysis. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. The outcome of the events abdominal pain, lack of appetite, and bouts of severe discomfort was recovering; while other events was unknown. Events reported as serious due to hospitalization and life threatening.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1310312-1	I felt tired the evening of the shot. The next morning I had fever and chills, body aches everywhere, severe headache, extreme fatigue, and very upset stomach. By the second day following the vaccination I have a very bloated stomach and extreme pain. These symptoms continued with the stomach and headache becoming worse. Entered the Emergency Room the evening of May 7th and admitted into the Hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1310568-1	Shortness of breath
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1310985-1	Heavy vaginal bleeding with clots, cramping in uterus and tenderness in this area prior to onset of bleeding. Vaginal bleeding lasted for 6 days. Please note that this was not like a regular menstrual period and I was not due for my period. I also bled after my first Pfizer vaccine but it was not alarmingly heavy like it was after this second dose. I was fatigued during the bleeding and after. The fatigue was resolved two weeks after I received the vaccine. One other strange side effect was that three days after I received the second vaccine my left hand was slow to respond when I needed to type on the computer. This lasted for several days.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1311022-1	Heavy vaginal bleeding with clots, cramping in uterus and tenderness in this area prior to onset of bleeding. Vaginal bleeding lasted for 6 days. Please note that this was not like a regular menstrual period and I was not due for my period. I also bled after my first Pfizer vaccine but it was not alarmingly heavy like it was after this second dose. After this second vaccine dose, I was fatigued during the bleeding and after. The fatigue was resolved two weeks after I received the second vaccine. One other strange side effect was that three days after I received the second vaccine my left hand was slow to respond when I needed to type on the computer. This lasted for several days.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1311675-1	My period flow have constantly been pretty normal. But this cycle I am have extremely heavy flowing and a lot of blood clots. I normally never have these.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1311962-1	Irregular, prolonged, heavy menstration with clots lasting greater than 7 days.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313014-1	Blood clot in left leg; This is a spontaneous report from a contactable consumer (patient). A 47-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), on 25Jan2021 10:30 (lot: EL3302) as 1ST DOSE, SINGLE then on 17Feb2021 10:30 (lot: EL9264) as 2ND DOSE, SINGLE; both via an unspecified route of administration in left arm (at the age of 46-years-old) for covid-19 immunisation. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Since the vaccination, the patient was not tested for COVID-19. Medical history included covid-19 (prior to vaccination, diagnosed with COVID) and no past drug event. The patient's concomitant medications were not reported. The reported event was blood clot in left leg on 18Apr2021 at 22:00. The event resulted in doctor or other healthcare professional office/clinic visit and Emergency room/department or urgent care. Treatment received for the adverse event was an unspecified medication. The outcome of the event was not recovered. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313019-1	blood clot; This is a spontaneous report from a contactable consumer. A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as unknown, single for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. It was reported that this patient who is a physician, experienced blood clot on an unspecified date. The outcome of the event was unknown. Information on batch number has been requested
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313030-1	two blood clots in his right calf; It is painful; This is a spontaneous report from a contactable nurse. A 73-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 1 via an unspecified route of administration, administered in arm left at 10 am on 16Mar2021 (Batch/Lot number was not reported) as 1st dose, single; dose 2 via an unspecified route of administration, administered in arm left at 10 am on 08Apr2021 (Batch/Lot number was not reported) as 2nd dose, single, both for covid-19 immunisation. Medical history included ongoing varicose vein (for years, barely noticeable). No history of all previous immunization with the Pfizer vaccine considered as suspect. No additional vaccines administered on same date of the Pfizer suspect. No prior vaccinations received within 4 weeks. The patient's concomitant medications were not reported. The patient experienced two blood clots in his right calf on 27Apr2021, it is painful. The event 'two blood clots in his right calf' was reported as serious and seriousness criteria were medically significant and disability. He said it is better today once he started apixaban (ELIQUIS). He did not know if that was physical or mental. The adverse event required a visit to physician office. The patient underwent lab tests and procedures which included sonogram on leg: unknown results. The outcome of events was recovering.; Sender's Comments: A causal association between BNT162B2 and the reported events cannot be excluded based on a compatible temporal relation. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313038-1	"possible thrombosis reaction; Unexpected bruising near injection site.; Bruise was purple and red splotches and has now progressed to mostly yellow/discoloration; Bruise was purple and red splotches and has now progressed to mostly yellow/discoloration; minimal pain/stiffness following the shot; minimal pain/stiffness following the shot; This is a spontaneous report from a contactable consumer (patient). A 53-year-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), at the age of 53-year-old via an unspecified route of administration, administered in Arm Left on 20Apr2021 11:15 (Lot Number: EW0153) at 1st dose, single for covid-19 immunisation. The patient medical history was not reported. The patient received no other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. Concomitant medication included famotidine (PEPCID AC). The patient previously took flu shot in Oct2020, prilosec [omeprazole magnesium] for seasonal allergy and experienced drug hypersensitivity. The patient experienced unexpected bruising near injection site. Bruise was the size of the bandaid, approximately 1"" x 2"". Bruising had persisted for more than a week now. The patient had never experienced bruising like this with other vaccines. He would like to rule out possible thrombosis reaction before second shot. He did not have bruising at any other locations other than the injection site. Bruise was purple and red splotches and has now progressed to mostly yellow. He believed the discoloration was a reaction to the vaccine and not the result of the needle. The shot itself was painful but patient felt minimal pain/stiffness following the shot. The event start date was on 20Apr2021. No treatment was received. The outcome was unknown."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313046-1	began having severe cramping and heavy bleeding with clots the next day; Started menstrual cycle a week early a day after getting second dose; began having severe cramping; heavy bleeding with clots the next day; This is a spontaneous report from a contactable consumer (patient). A 36-year-old female patient received bnt162b2 (BNT162B2, Pfizer COVID 19), dose 2 via an unspecified route of administration, administered in left arm on 26Apr2021 08:45 (Lot Number: unknown) as single dose for covid-19 immunization. Medical history included Anemia, ulcerative colitis, intracranial hypertension. None allergies reported. Concomitant medications included iron taken for anaemia. The patient received dose 1 on 01Apr2021 07:45 in right arm for covid-19 immunization. The patient started menstrual cycle a week early a day after getting second dose. The patient began having severe cramping and heavy bleeding with clots the next day 27Apr2021. No treatment received. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, patient was not diagnosed with COVID-19, Since the vaccination, patient has not been tested for COVID-19. The outcome was recovering. Follow up attempts are needed. Further information has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313048-1	Blood clotting with spotting and pain; Irregular period; Pain in lower left side of abdomen; Blood clotting with spotting and pain; This is a spontaneous report from a contactable consumer (patient). A 29-year-old non-pregnant female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on 24Mar2021 10:00 (Batch/Lot number was not reported) as single dose for covid-19 immunization. Facility where the most recent COVID-19 vaccine was administered was Other. Medical history included none. The patient's concomitant medications were none. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine or any other medications the patient received within 2 weeks of vaccination. The patient previously received the first dose on 05Mar2021 08:00 as single dose for covid-19 immunization. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. The patient experienced irregular period after second dose, pain in lower left side of abdomen and blood clotting with spotting and pain on 16Apr2021 12:00. The outcome of the events was unknown. No treatment received for the events. Information related lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313062-1	passing blood clots from her rectum; blood clots; deathly ill; the lymph nodes underneath the arm swell up / She swelled up on her right side; kind of in the breast/chest area, up a little higher than the breast and it was very uncomfortable; This is a spontaneous report received a contactable consumer, the patient. A 68-year-old elderly female received the first dose of intramuscular BNT162b2 (solution for injection; Lot ER8737 and expiry information not provided) as a single dose in the left arm on 08Apr2021 at 11:00 (at 68-years-old) for COVID-19 immunisation. Relevant medical history included ongoing back problems for which she takes medication. Concomitant medications included unspecified medication for back problems. The patient denied any other vaccines within four weeks prior to this vaccine. The patient reported that she generally gets the flu shot every year and has never had any adverse reactions. The patient also mentioned that she previously tramadol (for an unspecified indication) in the past but has not taken it for years. The patient reported that she received her first dose on 08Apr2021 at (Privacy). The patient explained that within six hours she was deathly ill. The patient reported that stayed very ill for quite a few days. The patient stated when she gets up in the morning, she gets up early and she always goes to the bathroom. The patient stated that on 13Apr2021, when she went to the bathroom, she started passing blood clots from her rectum which went on until 16Apr2021. When describing the blood clots, the patient stated 'envision sunny side up eggs, between six to eight of them, and in between that was all this webbing holding the clots together'. The patient stated while she was urinating, the clots were just pouring out of her rectum. The patient clarified that there was not blood in her stool. The patient explained that she took the opportunity to get some plastic gloves and look through her stool for blood and there was not any. The patient stated that her underwear and the toilet paper was stained. The patient also stated that she was reading that whichever arm you get the vaccine, the lymph nodes underneath the arm swell up. The patient explained that her lymph nodes did not swell up underneath her left arm where she got the shot. She swelled up on her right side. She clarifies she was swollen underneath her arm, kind of in the breast area and chest area, up a little higher than the breast. It was very uncomfortable. The patient denied any visits to the emergency room or doctor's office for the events. The outcome of the event passing blood clots from her rectum was recovered on 16Apr2021. The outcome of the events deathly ill, the lymph nodes underneath the arm swell up / She swelled up on her right side, and was very uncomfortable was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313900-1	Altered mental status, confusion, subdural hematoma. Was lifeflighted to Clinic and remained there about 5 weeks, developed a stage IV pressure ulcer on buttock. Diagnosed with multi organ failure. Went to a SNF for rehab after being discharged. Wife said then patient received 2nd dose of vaccine about 2 weeks ago and is currently at Hosptial in town for a GI bleed, passing frank blood and clots. He was admitted 5/9/21. Patient is still confused and lethargic. Before the vaccine, he was taking care of his farm and home. Now he doesn?t even know his wife or what is going on.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1314360-1	Patient stated she developed clots during hospital admission with the complaint of chest pain. Patient stated at hospital admission, they discovered blood clot in her right leg and right lung. Patient was treated with oxygen and medication. Patient was inpatient for two nights and sent home lovenox injections for one week and continuing with Eliquis. Patient is currently on Eliquis for unknown duration. Patient will see cardiologist in three weeks and reevaluate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1314427-1	2/4/2021 - Vaccine 2/26/2021 - Left Ankle surgery 4/9/2021 - Left leg swelling 4/22/2021 - Ultrasound found clots in left leg from groin to ankle

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316143-1	"they packed my nose to stop it from bleeding, clots were coming out, clots coming both ways,; nose bleed; my blood pressure went up; This is a spontaneous report from a contactable consumer (patient). A 58-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on 14Mar2021 10:00 as 2nd dose ,single for covid-19 immunisation. The patient medical history was not reported. Concomitant medication included acetylsalicylic acid (ASPIRIN BAYER). The patient previously received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) on 28Feb2021 for COVID-19 Immunization. The patient received the Second dose of Pfizer on 14Mar2021, states, ""On 17Apr2021, Saturday, nose bleed started, and my blood pressure went up. I Called ambulance took blood pressure and it was 210/107. I didn't go to hospital, nose bleed had stopped while they were there, I stayed in ambulance until blood pressure came down. Sunday I went to ER (emergency room) because my nose started bleeding again, it stopped Sunday, but they checked the nose, nothing was there. I stayed in the hotel, Monday morning I blood again, Tuesday I bled even more starting at 7am and it didn't stop. Went to the hospital to ER (emergency room), they packed my nose to stop it from bleeding, clots were coming out, clots coming both ways, they checked blood, not anemic, everything okay with blood. on 23rd they removed the pack from nose, ENT told me no vessels were broken, he said I was fine. That is when I received a phone call about someone else having this problem. A week later a class mate said her sister had the same reaction after getting the Pfizer vaccine."" She was calling about the Covid 19 vaccine. She received her second vaccine on 14Mar2021 and had no problems during that time. On 17Apr2021 she called an ambulance because she had a nosebleed. Her blood pressure was also high at that time around 210/107mmHg. Her nosebleed stopped with emergency management technicians and her blood pressure came back down. She did not go to the emergency room at this time. On 18Apr2021 at around 3:00AM, she had another nosebleed and she did go to the emergency department at about 10:00AM. She has never had a nosebleed before. When she was examined in the emergency room on 18Apr2021, she had no blood in her nose that the doctor could see. On 19Apr2021 around 7:00AM she had another nosebleed but the nosebleed stopped and she did not go back to the emergency room. On 20Apr2021, her nose started bleeding around 5:00AM and it didn t stop so she went back to the emergency room around 7:00AM and the doctor in the emergency room packed her left nostril and the blood was coming out of the right nostril and out of her mouth. She was bleeding for a few more minutes and then the bleeding stopped. Her blood pressure was fine. Her bloodwork was checked and everything was fine. She wasn t anemic or anything like that. She didn t have an bad effects from the nosebleeds. The emergency room told her to follow up with her primary care doctor on Thursday, 22Apr2021, but her primary care doctor no longer takes her insurance. She did go see an Ear, Nose, and Throat doctor on Friday, 23Apr2021. The Ear, Nose, and Throat doctor removed the nasal packing and examined the inside of her nose and told her that he could not see any broken vessels or any cause as to why she would be having nosebleeds. The patient's apartment is being renovated and she thought she might be having a reaction to the hotel she s staying in. She moved out of her apartment on 12Apr2021 and her nosebleeds started 17Apr2021. She has missed work because of her nosebleeds and she started back to work this week. When probing for her Covid 19 vaccine information she reports she does not have the vaccine record card with her. She received the second Covid 19 vaccine on 17Mar2021. NDC/Lot/Expiry not known. First Covid 19 vaccine date is in the suspect product field. She does not have the NDC/Lot/Expiry for the first date either. She reports she had a regular physical with her doctor in Nov2020. She received a tetanus vaccine at that visit. She reported she sometimes takes 2-3 regular Bayer Aspirin for minor pain. She took the Bayer Aspirin on Saturday, 17Apr2021, because she was going somewhere and she knew she would be sitting for a while at that place. She would like to report she experienced no discomfort, or aches or pains or headaches with her Covid 19 vaccine. There is no prior Vaccinations (within 4 weeks). The outcome of the events was unknown. Information about lot/batch number has been requested.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021476976 same reporter/drug/event, different patient"

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316152-1	clots were coming out; nose bleed; blood pressure went up; This is a spontaneous report from a contactable consumer (classmate's sister). This is the second of two reports. A 62-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; solution for injection) via an unspecified route of administration on an unspecified date (at an unspecified age) as a single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; solution for injection) via an unspecified route of administration on an unspecified date (at an unspecified age) as a single dose for COVID-19 immunization. On an unspecified date (almost a month after receiving the second COVID-19 vaccine) the patient experienced clots were coming out, nosebleed and blood pressure went up. The reporter called to report that she had received a phone call from an old classmate, who had a sister, who experienced the same reaction after getting the Pfizer vaccine. This patient experienced the same nosebleeds after the COVID-19 vaccine. This person's nosebleeds started almost a month after she received the second dose of the COVID-19 vaccine. That was when the reporter started trying to figure out where these nosebleeds were coming from. She did not have any further information on the person that she mentioned. The clinical outcomes of the events clots were coming out, nosebleed and blood pressure went up were unknown. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021476256 same reporter/drug/event, different patient
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316156-1	resulted into blood clots in his throat and left side of his body; This is a spontaneous report from a contactable consumer via a Pfizer sponsored program. A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), first dose on 22Jan2021 (Batch/Lot Number: EL1283) as 1st dose, single, second dose on 16Feb2021 (Batch/Lot Number: EL9264) as 2nd dose, single; both via an unspecified route of administration for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient reported that he ended up in the hospital after receiving both covid vaccines which resulted into blood clots in his throat and left side of his body from the waist down on an unspecified date. The patient was hospitalized due to the event from 25Apr2021 to 27Apr2021. The outcome of the event was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316157-1	they were concerned about blood clot; shaking uncontrollably; pouring sweat; sheet white with no color to skin at all; blood pressure 68/30; not particularly coherent; This is a spontaneous report received from a contactable consumer (patient's husband). A female patient of unspecified age received BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: UNKNOWN) via an unspecified route on an unspecified date as a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter stated his wife had adverse event to the COVID vaccine last week. He stated that they both got the vaccine. At 3 AM on unspecified date, the reporter woke up because his wife (patient) was shaking uncontrollably, pouring sweat, sheet white no color to skin at all, not particularly coherent and had blood pressure 68/30. They had to call an ambulance and she was brought to the hospital. They were able to get her blood pressure up and stabilized her after 8 hours. They were concerned about blood clot and did other scans. Lab data included blood pressure 68/30 (units unspecified) and scans with unknown results; both on unspecified date. The clinical outcomes of shaking uncontrollably, pouring sweat, sheet white no color to skin at all, not particularly coherent and blood clot were unknown, while of blood pressure 68/30 was recovering. The lot number for the vaccine, BNT162b2, was not provided and will be requested during follow up.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316160-1	left hand was edematous and cold; left hand was edematous and cold; left hand pain; blood clot; swollen submandibular gland that got bigger/could feel her mouth getting bigger; ill with something not related to the vaccine; This is a spontaneous report from a contactable nurse (patient) via Medical information team. A 67-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 1 intramuscular, administered in Arm Left on 03Mar2021 15:00 (Lot Number: EN6201) as 1ST DOSE, SINGLE for COVID-19 immunization. Medical history included stroke on 2007, COVID-19 from an unknown date and unknown if ongoing. Concomitant medications included clopidogrel bisulfate (PLAVIX) taken for stroke from 2007 to an unspecified stop date; and acetylsalicylic acid (ASPIRIN (E.C.)) taken for stroke from 2007 to an unspecified stop date. The next day on 04Mar2021, the patient had a swollen submandibular gland that got bigger. She could feel her mouth getting bigger. The patient was admitted into the hospital on 07Mar2021 and released on 09Mar2021. She was readmitted to the hospital on 11Mar2021 until 18Mar2021. The patient went back into the hospital on 21Mar2021 and released 20Apr2021. The gland was removed on 13Apr2021. She developed a blood clot. She went to the emergency room and was sent home on Xarelto. The patient mentioned that she received the 1st dose of the vaccine on 03Mar2021 and came down ill with something not related to the vaccine. She was hospitalized for 4 weeks and ended up missing her 2nd dose appointment on 24/Mar/2021. She would like to get her 2nd dose but she is now 5 weeks overdue for it. She mentioned the hospital is recommending that she restart her vaccination. She is looking for guidance from Pfizer. The facility where the most recent COVID-19 vaccine was administered was in the hospital. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. The patient underwent had CT and MRI of left arm/wrist which she does not have results to provide. Event relatedness to vaccine was unknown for events, swollen submandibular gland that got bigger, left hand was edematous and cold, left hand pain and blood clot. The outcome of the event blood clot swollen submandibular gland was recovered with sequel on unspecified date, the outcome of ill feeling was unknown and the outcome of the rest of the events was recovering.; Sender's Comments: The 67-year-old female patient had medical history of stroke. Based on information available, the reported thrombosis and other events were unlikely related to the vaccine of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE). This case will be reassessed should additional information become available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316174-1	"Blood clot in leg - first dose; This is a spontaneous report received from a contactable consumer, the patient. A 77-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot: CR2613), via an unspecified route of administration in the left arm on 30Mar2021 (at the age of 77-years-old) as a single dose for COVID-19 immunization. Medical history included broke femur of right leg on 09Nov2020 and hospitalized from 09Nov2020 until 15Nov2020 or 16Nov2020, surgery to repair broken right femur on 12Nov2020, rehabilitation following hospital discharge and got out of the rehab facility on 14Dec2020, been in either a walker or a wheelchair and still not fully recovered (ongoing); chronic obstructive pulmonary disease (COPD), asthma diagnosed at least 13-14 years ago, bronchitis, all ongoing from an unspecified date diagnosed at least 13-14 years ago. Concomitant medications were not reported. There were no other vaccinations within 4 weeks of the vaccine. On an unspecified date, the patient experienced a blood clot in the leg. The clinical course was follows: After he was discharged from rehab, the patient went to his own family doctor on 30Mar2021 because his ankle and leg were still swollen. The physician sent him to have a vascular done, on an unspecified date after 30Mar2021, with unknown results. The patient reported that the physician determined that he had blood clots in his leg. The patient was sent to the emergency room and started ""that stuff"" for about a week until he went to a regular vascular doctor and the physician determined what the patient should be doing. The patient reported that he was diagnosed with the blood clots after he got out of rehab. The patient received apixaban (ELIQUIS) from Apr2021 for blood clot in leg. The clinical outcome of blood clot in leg was unknown.; Sender's Comments: Linked Report(s) : PFIZER INC-2021478767 same patient, different dose/event"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1317146-1	Severe clots on right thigh rushed to urgent care and urgent care referred me to emergency Hospital. Was hospitalized for 5 nights with severe swelling on right thigh. The swelling has since abated but has still not subsided. Feel still very heavy on right thigh.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1318527-1	PATIENT WAS ADMITTED TO ICU ON 5-11-2021. CHIEF COMPLAINT BEING BLOOD CLOTS.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320025-1	Superficial blood clots in my right leg; This is a spontaneous report from a contactable consumer. A 55-years-old male patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in Arm Left on an unspecified date (Batch/Lot Number: EW0164) as SINGLE DOSE for covid-19 immunisation. Medical history included Alfa 1 Antitrypsin (hereditary), Sulfa allergies. Patient previously received the first dose of bnt162b2 on 27Mar2021 for covid-19 immunization (lot number=ER8734,vaccine location=Left arm). Patient was not diagnosed as COVID-19 prior vaccination. No covid tested post vaccination.The patient's concomitant medications were not reported. The patient experienced superficial blood clots in his right leg 5 days after his second COVID 19 vaccination shot on 22Apr2021 04:00 with outcome of unknown. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Treatment received for the adverse event included Rest and pain relievers. Case was reported as non-serious. Information on Lot/Batch number was available. Additional information has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320074-1	Second blood clot that traveled to her brain which caused her to have a stroke; Blood clot in her right leg; Weight Loss from 258 pounds, down to 208 pounds, back up to 214 pounds.; She has to learn how to walk again, she is in a wheelchair now; Having Problems Swallowing from the Stoke; Speech went down to 25% and now she is at 50%, because of the Stroke; high blood pressure; slept five days; Arm was Sore; Body was Sore; Diarrhea; Slept and was Totally Exhausted; This is a spontaneous report received from a contactable consumer (husband of the patient) and a contactable other healthcare professional. This consumer reported similar events with different doses of the same product; this case refers to dose 2; only this case is serious. A 61-year-old adult female received the second dose of BNT162B2 (solution for injection, Lot EL3246 expiry information not provided) as a single dose via an unspecified route on 30Jan2021 at 13:45 (at 61-years-old) for COVID-19 immunization. Relevant medical history included ongoing seizures (1964; began at 5 years old, minor seizures; under control in 1984, possibility configuration it is the right cocktail), and wound on her left leg (that was being treated by a wound doctor for the last 7 years). Concomitant medications included gabapentin; 300mg three times a day by mouth (6am, 2pm and bedtime) ongoing from 2014 for nerves; valproate semisodium (DIVALPROEX; 500mg two tablets in the morning and one tablet at 6pm ongoing from 2010 for seizures furosemide 20mg every 48 hours (odd days); States she takes it to reduce swelling (diuresis)on the leg wound leg; states he gives it to her on the odd day of the month from an unspecified date; metoprolol (50mg) one tablet twice a day (6am and 6pm), for Blood pressure management; ascorbic acid, zinc (VITAMIN C WITH ZINC; 500mg/50mg) once daily from an unspecified date for an unspecified indication; acetylsalicylic acid (ASPIRIN) 81mg once daily from an unspecified date for an unspecified indication; and pramipexole 0.5mg once daily at 6pm ongoing from an unspecified date for seizures. The patient previously received COVID-19 immunization with the first dose of BNT162B2 (solution for injection; Lot EL1284) on 09Jan2021 (at 61-years-old) and experienced arm was sore, slept for two to three-days, body was sore, and diarrhea. The consumer reported that on 30Jan2021, the patient experienced her arm was sore, her whole body was sore for about five days, she had diarrhea for five days, and she for slept five days and was totally exhausted. The consumer then reported that on 22Feb2021, she developed a blood clot in her right leg. The reporter explained that her leg swelled up and turned dark blue, almost black. The consumer reported that the patient was immediately rushed to the hospital. Caller states they did all sorts of tests and confirmed she had a blood clot in her right leg. States she was hospitalized from 22Feb2021 through 26Feb2021, so for five days. Caller states the leg went back to normal color, the swelling went down and she felt good. Caller states they came home on 26Feb2021, and she was up moving around, and everything was wonderful. On an unspecified date in Feb2021, around the time of the first blood clot, the patient was diagnosed with high blood pressure. Caller states on 12Mar2021 she had a second blood clot that traveled to her brain which caused her to have a stroke. Caller states when they got to the hospital they said she had a stroke from a blood clot that was somewhere in her system and it traveled to her brain. Caller states she was hospitalized from 12Mar2021 through 16Mar2021 when she was released from the hospital. Caller states she was too sick to go home so she was transferred to Rehab Center. She was there from 16Mar2021-27Apr2021. Caller states on 27Apr2021, she was transferred for rehab. Caller states at the center they only give rehab once a day for 30 minutes and he only works five days a week. Now she gets three hours of physical therapy a day and it is seven days a week. States they are working to get her to walk again and she is receiving speech therapy. States she has to learn how to walk again. Caller states her lowest weight was 208lbs, and they weighed her in yesterday and she was at 214lbs. Caller states with the stroke when she was at the hospital, they put her on baby food because she was having problems swallowing. While at rehab center she went from baby food to mechanical food. On 27Apr2021 she went to regular food. Speech because of the stroke went down to 25% and now she is at 50% and she is also getting one hour of speech therapy at hospital. Caller states they recognized it is important to get as much therapy as possible in the first 90 days, they call it a boot camp and patients are limited to 90 days to repair the damage. After 90 days, that is it. States that is why it was important to transfer her to hospital to heal and get back to normal. Caller states she is in a wheelchair and they are trying to get her to stand up. Caller states at the hospital they were able to get her to take 2 steps. At Hospital they were able to get her to do special training and he was able to get her to take 4 steps. Caller states he has no idea where she is with the walking right now. States she did not do anything on the 27Apr2021, because that was arrival date. Therapy started yesterday

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	on 28Apr2021 at Hospital. The consumer also reported that the patient's **Adverse Event Description**
				current weight is 214 pounds, from 258 when she took the shot and down to 208 pounds on an unspecified date in 2021. Treatment for the events included occupational, physical and speech therapies, as well as apixaban (ELIQUIS), rivaroxaban (XARELTO), and spironolactone 25mg once a day; takes in the evening at 6:00p.m. ongoing from 12Mar2021 for hypertension. The outcome of the events arm was sore, body was sore, diarrhea, and slept and was totally exhausted and slept five days was recovered on 04Feb2021. The outcome of the event blood clot in right leg was recovered on 26Feb2021. The outcome of the event second blood clot that traveled to her brain which caused her to have a stroke was recovered with sequelae on an unspecified date in 2021. The outcome of the event Having Problems Swallowing from the Stoke was recovered on 27Apr2021. The outcome of the events speech went down to 25%, back up to 50%, and weight loss from 258 pounds, down to 208 pounds, back up to 214 pounds was recovering. The outcome of the event high blood pressure was unknown.; Sender's Comments: Linked Report(s) : PFIZER INC-2021484454 PFIZER (Same reporter/patient/different doses/events)
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1320081-1	blood clots, Clots were going down the back of her throat; currently feels very weak; nose bleed; This is a spontaneous report from a non-contactable consumer. This report was received via a sales representative. A 70-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: unknown), single dose via an unspecified route of administration in Apr2021 for covid-19 immunization. The patient's medical history was not reported. The patient has been on icosapent ethyl (VESCEPA) and ibuprofen for years and currently feels very weak on 25Apr2021. Concomitant medications were unknown. The patient experienced nose bleed blood clots throughout the week on 25Apr2021. Clots were going down the back of her throat. The patient had to seek medical attention. and currently feels very weak on 25Apr2021, two weeks after vaccination. The outcome of nose bleed blood clots and weak was unknown. No follow-up attempts are possible. No further information expected.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320109-1	"Blood clots; Delirious; Oxygen was down to 80%; Sepsis; E. coli/sepsis with cultures of E. coli; C-diff; sick; Shaking uncontrollably; Achy; Extremely high fever/a high fever of 103 F; Chills; Aneurysm in his leg; Bleeding from somewhere; weak; missed his second dose of the Pfizer COVID-19 vaccine; Caller says her husband's kidney was failing too while he was in the hospital; This is a spontaneous report from a contactable consumer reported for her husband. A 69-years-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in arm on 11Mar2021 18:00 (Lot Number: EV6206) as single dose for covid-19 immunization. Medical history included Kidney transplant from 1998, same kidney all these years, on immune medication, polycystic kidney disease from 1998, the transplant saved him in 1998, blood pressure high went along with the polycystic kidney disease, had it for a while, Type 2 diabetes mellitus this was diagnosed probably about four years ago and immunocompromised. There were no concomitant medications. The patient had the first dose of the Pfizer vaccine on 11Mar2021, ""he felt okay first evening, Friday was okay, Saturday 13Mar2021 achy, Sunday running a high fever of 103 F, he started to get chills. Then Around 3am (Early Monday on the 15Mar2021) called the ambulance, oxygen was down to 80% he was shaking uncontrollably, temperature was still 103 F when they took it and even at the hospital was the same temperature. He was delirious on 15Mar2021. When he got to Emergency Room he was in critical section, had to put on BiPAP with 100% oxygen, When in the ambulance they tested for covid and was negative each time he was tested for covid he was negative. He was in the hospital from 15Mar2021 to 24Mar2021, in critical care unit. He Turned out having sepsis, E.coli, had to put him on strong antibiotics. He developed CDIFF on 15Mar2021, and an aneurysm in his leg in Mar2021 and needed a blood transfusion. They were trying to rule out blood clots and did sonogram and found it. His blood was going down, he was bleeding from somewhere and had to have a transfusion. They did a procedure on the leg to break up the aneurysm twice. On BiPAP for a few days and then did the medium one and then able to take off oxygen when he went home. He came home on antibiotics and tons of medication. Should he get the second vaccine? Is it related? Is it to late to get the second vaccine? following Sunday he had an extremely high fever and was shaking uncontrollably on 14Mar2021. He was told to get the shot by his provider. She had to call an ambulance, he almost died, and she didn't know if it is related to her husband getting the vaccine. They put her husband on a BiPAP with 100% O2 when he went to the hospital by ambulance and was admitted on 15Mar2021, which he was on the BiPAP a couple days then went on to the second level, then right before he came home he was weaned off of oxygen. They kept testing him for COVID every day, because the symptoms were similar but they tests kept coming up negative. On 15Mar2021, the patient had sepsis, with cultures of E. Coli, and very contagious C-diff, he was sick from the 15th to the 24th for all those things. She didn't know if the vaccine brought it on if it was brewing in him before that, or if it had to nothing to do with vaccine. He also had an aneurysm in his leg and had a medical procedure where they had to go in and take care of it to break up the aneurysm. The patient's kidney was failing too while he was in the hospital. The patient missed his second dose of the Pfizer COVID-19 vaccine due to his hospitalization from 15Mar2021 to 24Mar2021. She would like to know if it would still be recommended for the patient to get the second shot of his COVID-19 vaccination since he is past the deadline to get it, is it going to be too late for him to get the second dose to be fully vaccinated and effective to get it after waiting this long? His first dose was injected about 06:00PM in unknown arm. The patient had a fever Saturday evening, clarified to being 13Mar2021, which was higher the next day on 14Mar2021 and reached 103 degrees Fahrenheit, and he went to the hospital by ambulance at 03:00AM 15Mar2021. His fever started getting higher late Saturday to Sunday and he got progressively worse. The patient had had serious complications before related to his health problems, but she had never had to call an ambulance before this. The patient had chills on Saturday 13Mar2021 and then Sunday evening he was shaking and it wouldn't stop so she gave him Tylenol, then she called the ambulance Monday morning. The patient can't take anything more than Tylenol, and he never usually takes anything. The patient's kidney function was getting bad while he was in the hospital, his Creatinine went high and was 3.8, which is high for someone with a transplant. The patient was doing better, he was still very weak, he was under a doctor's care, he was on very strong antibiotics for a month, Vancomycin and other medications which also make him weak. The patient was on IV antibiotics at the hospital and then came home on oral. The 3.8 for the Creatinine was probably at the beginning when he went into the hospital. The outcome of the event Sepsis was Resolved with Sequel and the outcome of the other events was unknown."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320143-1	Heavy bleeding including passing a blood clot with my first cycle after my second dose.; Heavy bleeding including passing a blood clot with my first cycle after my second dose.; This is a spontaneous report from a contactable consumer (reporting for herself). A 37-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 25Mar2021 at 13:00 (Batch/Lot Number: ER2613) as 2ND DOSE, SINGLE for COVID-19 immunization. Medical history included hypothyroidism from an unknown date and unknown if ongoing. Concomitant medication included etonogestrel (NEXPLANON); levothyroxine sodium (LEVOTHYROXIN); cetirizine hydrochloride (ZYRTEC ALLERGY); fluticasone propionate (FLONASE ALLERGY RELIEF), all taken for an unspecified indication, start and stop date were not reported. The patient previously took first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Left Arm on 04Mar2021 at 13:00 (Batch/Lot number: EN6206) for COVID-19 immunization. On 28Apr2021 at 14:30, the patient experienced heavy bleeding including passing a blood clot with her first cycle after my second dose. The have NEVER bleed like this except for when she was having a miscarriage or postpartum after giving birth to her daughters. She usually can go all day with the same pad and soaked through a pad and her underwear was completely bloody in 3 hours (it was so bad I discarded the underwear). When she got on the toilet it kept gushing out and that was when she passed a clot. After that the bleeding slowed down. The outcome of the events was unknown. The patient did not received any other vaccines within 4 weeks prior to the COVID vaccine. The patient was not diagnosed with COVID-19 prior to vaccination and has not been tested for COVID-19 since the vaccination.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320149-1	Diagnosed with a blood clot; Diagnosed with stroke; not feeling well; wakes up at night; Numbness of left hand and mouth; Numbness of left hand and mouth; Constipation; severe headache, described as he felt his eyes were exploding out of his head.; Temperature of 100.2; Could not sleep; This is a spontaneous report from a contactable consumer (spouse reporting on behalf of husband). A 63-years-old male patient received the first dose bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration, administered in Arm Left on 23Mar2021 09:00 (Batch/Lot Number: ER2613) as SINGLE DOSE for covid-19 immunization. Medical history included ongoing rheumatoid arthritis started few years ago, ongoing back pain (for many years), ongoing pain (for many years) (patient is a farmer 'seeing another doctor' for back pain/pain, testis cancer from 1986 and ongoing. Concomitant medication(s) included adalimumab (HUMIRA) taken for rheumatoid arthritis from an unspecified start date ' a few years ago.' and ongoing; tramadol (TRAMADOL) taken for back pain from an unspecified start date ' for years' and ongoing. Vaccine was not Administered at Military Facility. No other vaccines were given within 4 weeks. No additional vaccines were administered on the same date of the Pfizer suspect product. The patient experienced diagnosed with a blood clot, diagnosed with stroke, numbness of left hand and mouth on 12Apr2021, severe headache, described as he felt his eyes were exploding out of his head, temperature of 100.2 (in the evening.) , could not sleep on 03Apr2021, constipation on 05Apr2021, not feeling well and wakes up at night on an unspecified date. The patient was hospitalized for stroke and blood clot from 12Apr2021 to 14Apr2021. The patient underwent lab tests and procedures which included body temperature increased: 100.2 on 03Apr2021; computerized tomogram: first cat scan, a blood clot was diagnosed on unspecified date., computerised tomogram head: unknown result on an unspecified date, electrocardiogram: unknown results (husband was not having a heart attack) on 12Apr2021 , laboratory test: unknown results an unknown date , sars-cov-2 test: negative on 12Apr2021, computerised tomogram: negative on an unspecified date. Treatment was given thrombosis, Stroke, Headache, Insomnia. On an unspecified date patient contacted his primary care doctor and was told to take some extra Motrin, twice a day but Motrin was not effective for his headache. The patient was not feeling well he went to his primary care doctor. Caller states her husband was at his doctor on 07Apr2021, and he went to radiology for a Cat Scan on an unspecified date and the results was negative. He had lab work drawn and a CT of the head. Her husband was given a medication to help him sleep, medication not specified. Caller mentioned she rescheduled her husband's second vaccine. On 12Apr2021, in the evening when he was getting out of the shower, he developed numbness in his left hand, and added later in the report, numbness of his mouth. Patient went to the emergency room, where they performed several tests, Multiple Cat Scans. Results of the first Cat Scan, a blood clot was diagnosed, and an EKG because they thought he may be having a heart attack but it was confirmed he was not having a heart attack.. When her husband was in the emergency room, they did a Covid test and it was negative. The patient was diagnosed with a blood clot and got Tissue plasminogen activator. At 7:33 pm, when he was given tissue plasminogen activator. After the treatment, the headache went away, but she is not sure if he has lasting effects. He was admitted to the Intensive Care Unit and was in the hospital for two days and diagnosed with a stroke, but he was able to walk out of the hospital. The patient still gets an ache in his head, and it comes and goes. Caller states it seems that these events were not coincidental, as they all happened after he got the vaccine. Since her husband has been home from the hospital, he is sleeping better but normally does not get a full night of sleep, and he usually wakes up at night and husband gets constipated when he takes Motrin. Caller mentioned that her husband is not fully constipated anymore, but he feels that his system is not back to normal. Caller assumed the stroke has cleared up, but is not sure. Caller rescheduled her husband's second vaccine, and she is not sure her husband should get the second vaccine. Caller states if these events were caused from the Covid vaccine, what will happen in the future? The clinical outcome of the events Clot blood, Stroke, malaise and wakes up at night was unknown; Numbness of left hand and mouth was recovered on 12Apr2021, Fever was recovered on 04Apr2021, while headache and Constipation was recovering. Follow up attempts are needed. Further information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320186-1	"Blood clot; He was peeing a lot of blood; all of the joints went wondering in one another that was swelling up and it just so painful. Rheumatoid Arthritis layered up; leg swells up twice the size of the other leg; This is a spontaneous report from a contactable consumer reporting for his father. A male patient of an unspecified age received BNT162b2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for Covid-19 immunisation. Medical history included rheumatoid arthritis. The patient's concomitant medications were not reported. The patient experienced blood clot on an unspecified date with outcome of unknown, he was peeing a lot of blood on an unspecified date with outcome of unknown, all of the joints went wondering in one another that was swelling up and it just so painful/ he is just swelling up, leg swells up twice the size of the other leg on an unspecified date with outcome of unknown. The patient underwent lab tests and procedures which included ultrasound scan: clot blood. The event thrombosis and hematuria were considered Important Medical Events. Course of the events: Reporter stated, ""My father got the Pfizer Covid vaccine couple of weeks ago and it caused I think Rheumatoid Arthritis layered up in his joints one of the times started swelling up. He got a blood clot, his legs swelled up like twice the size of the other leg and they did ultrasound on it and got a blood clot. They put him up on blood thinners (Unspecified Medication) now today he had an appointment with the doctor in the room and when he was there he went to the bathroom and then he was peeing a lot of blood. Doctor told him that It should be okay if stop taking the blood thinners and she cleared up and that's it and I am like freaking out that why would they not got him to the hospital. He went to the hospital because it's the other blood clot. People saying this is on the blood thinner. I don't know why is they rolling up a lot. Nobody wants to say its from the vaccine but it is happen in half an hour within the vaccine, its started."" Information about Lot number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320205-1	she has a blood clot; had a blood clot and it looked like dark red bruises on the left side of her chest; This is a spontaneous report from a Pfizer sponsored program COVAX US support from a contactable consumer (patient). A female patient of unknown age received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 27Apr2021 at single dose for COVID-19 immunisation. Medical history and concomitant medications were not reported. The patient previously received first dose of BNT162B2 on 06Apr2021. The patient had a blood clot and it looked like dark red bruises on the left side of her chest. She just noticed it after she received her 2nd dose on 27Apr2021. The outcome of the events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320208-1	There is nonocclusive thrombus in the proximal left femoral vein. There is occlusive thrombus from the mid femur vein to the popliteal and calf veins; This is a spontaneous report from a contactable consumer (patient). A 58-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection) via an unspecified route of administration, administered in left arm on 26Mar2021 at 12:00 PM (at age of 58 years old, Lot Number: ER8733) as a single dose for covid-19 immunization. Medical history included hypertension from an unknown date. Concomitant medications included amlodipine, zolpidem, lisinopril. The patient previous took the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection) via an unspecified route of administration, administered in left arm on 05Mar2021 at 01:00 PM (at age of 58 years old, Lot Number: EN6206) as a single dose for covid-19 immunization. It was reported the patient the left common femoral and great saphenous veins are patent. There was nonocclusive thrombus in the proximal left femoral vein. There was occlusive thrombus from the mid femur vein to the popliteal and calf veins on 22Apr2021 12:00. The patient was hospitalized for 3 days, it was life threatening illness. Treatment received IV heparin. The event result in doctor or other healthcare professional office/clinic visit and emergency room/department or urgent care. The patient underwent lab tests and procedures which included SARS-CoV-2 test (Nasal Swab): negative on 29Apr2021. The outcome of event was not recovered.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320242-1	"deathly ill after the vaccine; blood clots in stool; This is a spontaneous report from a contactable pharmacist. An adult female patient received COVID-19 vaccine (UNSPECIFIED TRADE NAME), via an unspecified route of administration on 06Apr2021 (Batch/Lot Number: Unknown) as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. Facility where the most recent COVID-19 vaccine was administered: Pharmacy or Drug Store. The patient experienced deathly ill after the vaccine and blood clots in stool on 16Apr2021 with outcome of recovering. Patient stated they had blood clots in stool 2 weeks after the vaccination but was deathly ill after the vaccine. The adverse event result in Doctor or other healthcare professional office/clinic visit. Information on the lot/batch number has been requested.; Sender's Comments: Based on the temporal relationship, the association between the events ""deathly ill"" and ""blood clots in stool"" with COVID-19 vaccine can not be fully excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320251-1	blood clot; heart attack; This is a spontaneous report from a contactable consumer. This consumer reported for his father. A male patient of an unspecified age received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for covid-19 immunisation . The patient medical history and concomitant medications were not reported. The patient experienced blood clot and had an heart attack on an unspecified date with outcome of unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320270-1	Patient reported clot developed in calf 1 week following first dose; Confusion; This is a spontaneous report from a contactable consumer (reporting for herself). A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 04Mar2021 (Batch/Lot number was not reported) as 1ST DOSE, SINGLE for COVID-19 immunization; and sarilumab (KEVZARA), via an unspecified route of administration from 05Mar2021 (Batch/Lot number was not reported) at an unknown dose and frequency for an unspecified indication. The patient medical history was not reported. Concomitant medication included enoxaparin sodium (LOVENOX [ENOXAPARIN SODIUM]) taken for an unspecified indication, start and stop date were not reported. In Mar2021, the patient reported that clot developed in calf 1 week following first dose of Pfizer covid-19 vaccine. Also, with a lot of confusion. She stated that she was driving and got lost in her 1 stoplight/1-mile small town. The outcome of the events was unknown. The action taken in response to the event for sarilumab was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320289-1	it is weird clot it is not that area where I got the shot; feeling like I was having heart pain; His left upper chest area has swelling and when he touches it he feels something hard. Pain when he touches; His left upper chest area has swelling and when he touches it he feels something hard. Pain when he touches; His left upper chest area has swelling and when he touches it he feels something hard. Pain when he touches; He feels pressure on his chest; It feels like pain on the heart. It's a sharp needle like pain in some spots; my left arm and also my back there is going to pain seems like needle anything from the back; my left arm and also my back there is going to pain seems like needle anything from the back; This is a spontaneous report from a contactable consumer (Patient) via a Pfizer sponsored program. A male patient of an unspecified age received second dose of BNT162B2 (solution for injection, Lot number: ER8735) via an unspecified route of administration on 25Mar2020, as 2nd dose, single for COVID-19 immunization. Medical history and concomitant medications were not reported. Patient received first dose of BNT162B2 (solution for injection, Lot number: CVS8601) via an unspecified route of administration on an unspecified date, as 1st dose, single for COVID-19 immunization. It was stated that everything was fine on the first day after second dose and then from the second and third day the patient had a pain which was weird clot, and it was not that area where he got the shot.It was like between shoulder and neck, left shoulder, and neck. A kind of in deeper area there was lot of bone and just like where neck was connected to shoulder but it was in front of right. Patient had pain when he touched there in like size of the corn was like he could feel something hard inside and then just started day prior to this report further on the evening and started having chest pain like whole that spot felt like he was having heart pain and his left arm and his back there was going to pain seemed like needle anything from the back. The pain was constant. It was not on chest, it was not like that would come and go, would say it was bearable thing, but it was constant and breath that goes inside his body it was still there when touched, it was still paining. Patient upper left chest area had swelling and when touched it, he felt something hard. It would pain when touched. Patient felt pressure on his chest. It feels like pain on the heart. It was a sharp needle like pain in some spots. Patient took aspirin on the day and a day prior to this report. The outcome of the event of all the events was unknown. Additional information about batch/lot has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320341-1	"I am bleeding heavily because I am getting blood clots and stuff like that; I am bleeding heavily because I am getting blood clots and stuff like that; Lot of pain in my joints and in my bones; Lot of pain in my joints and in my bones; Right side of face, my tongue inside is very swollen and irritated; Right side of face, my tongue inside is very swollen and irritated; Right side of face, my tongue inside is very swollen and irritated; Right side of face, my tongue inside is very swollen and irritated; This is a spontaneous report from two contactable consumers (one was the patient). A patient of unspecified age and gender received the second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot Number: FN6200), via an unspecified route of administration on an unspecified date as a single dose for COVID-19 immunisation; axitinib (INLYTA), via an unspecified route of administration from an unspecified date (Batch/Lot number was not reported) at 2 pills once a day for clear cell renal cancer; pembrolizumab (KEYTRUDA), route of administration, start and stop date, batch/lot number and dose were not reported for clear cell renal cancer; zoledronic acid (ZOMETA), route of administration, start and stop date, batch/lot number and dose were not reported for the bones. The patient received the first dose of BNT162B2 on unknown date (Lot Number: CL9265) for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient was getting terrible side effects from either the axitinib or the pembrolizumab. The patient was bleeding very heavily and the right side of face and tongue inside was very swollen and irritated. The patient was getting a lot of pain in the joints and bones. The patient was bleeding heavily because was getting blood clots and ""stuff like that"" (as reported). The actions taken with axitinib, pzoledronic acid, and embolizumab were unknown."" The outcome of the events was unknown. Follow up (23Apr2021): New information received from a contactable consumer includes reporter details. No Follow-up attempts are needed. No further information is expected. Follow-up (07May2021): This is a follow-up report to notify that the case 2021406450 and case 2021513212 are duplicates. All subsequent follow-up information will be reported under Manufacturer report number 2021513212."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320345-1	got acute deep vein thrombosis of right lower extremities; inflamation and blood clot in vein; inflamation and blood clot in vein; This is a spontaneous report from a contactable consumer (patient). A 53-years-old female patient reported for herself that received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Left on 18Apr2021 (Batch/Lot number was not reported) at the age of 53-years-old, as single dose for covid-19 immunisation. The patient had no medical history. The patient's concomitant medications were not reported. The patient reported: I got acute Deep Vein Thrombosis of right lower extremities, inflammation and blood clot in vein on 19Apr2021 01:00 AM. The patient was visited by physician in Emergency Room. The reported events were deep vein thrombosis, thrombosis and inflammation, all considered life-threatening. Clinical outcome of the events was not recovered. Lab tests: CBC with auto diff CBC/ ETC. Unspecified therapeutic measures were taken as a result of the events. The patient was not diagnosed with COVID-19 prior to vaccination and since the patient was not vaccinated. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320357-1	cough; chest hurts; clots on both legs and stomach; hemorrhage; Blood sugar decreased; Oxygen saturation low; Blood pressure low; Skin discoloration; vein on her right leg was torn; Difficulty breathing; Tiredness; Breast pain; blockage all over the heart; This is a solicited report from A contactable consumer (patient) based on the information received by Pfizer from AbbVie (Manufacturer Control No: 21K-163-3840719-00). A 73-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 intramuscular on 27Feb2021 (Batch/Lot number was not reported) as 2ND DOSE, SINGLE for COVID-19 immunization; adalimumab (HUMIRA), subcutaneous from 2005 (Batch/Lot number was not reported) to Feb2021, at 40 mg/0.4 ml, then subcutaneous from 21Apr2021 (Batch/Lot number was not reported) and ongoing, at 40 mg 1 in 2 week for moderate to severe rheumatoid arthritis psoriatic arthritis. Medical history included patient consumed a pack of cigarette a day from 1967 to 2019, heart attack from 2003 to 2003, abstains from alcohol, and high blood pressure. Concomitant medication included metoprolol taken for high blood pressure. The patient previously took metformin to manage blood sugar. On an unspecified date, the patient experienced cough, chest hurts, clots on both legs and stomach. In 2010, the patient experienced breast pain and blockage all over the heart. She first tried to do the balloon and had three stents on her heart due to blockage sometime in 2010. In Jan2021, the patient had bypass surgery as treatment heart blockage as well. On 18Feb2021, echocardiogram/ECHO showed blockage all over the heart again. In 2021, the patient had difficulty breathing and experienced tiredness easily. In Mar2021, she was told to have an outpatient diagnostic heart catheter procedure and they had to take her by ambulance to be admitted in a hospital, when she moved during the procedure, a vein on her right leg was torn and also had hemorrhage, blood sugar decreased, oxygen saturation low, blood pressure low, skin discoloration. She experienced a recurring pain in between the breast bone after the triple bypass surgery that was done last 09Mar2021 while in the hospital. She also had blood transfusions while hospitalized as treatment for hemorrhage. The patient was hospitalized for the events for 28 days (also reported as 4 weeks). The event vein on her right leg was torn was considered life threatening. The events breast pain, blockage all over the heart, difficulty breathing, tiredness, hemorrhage, blood sugar decreased, oxygen saturation low, blood pressure low, and skin discoloration were also considered medically significant. The patient underwent lab tests and procedures which included stress test and echocardiogram/ECHO with unknown result in Jan2021;

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				echocardiogram/ECHO with unknown result in Jan2021; echocardiogram/ECHO showed blockage all over the heart on 18Feb2021; blood sugar: decreased, blood pressure: low, and oxygen saturation: low, all in Mar2021. The action taken in response to the events for adalimumab was dose not changed. The outcome of vein on her right leg was torn was recovered in Apr2021; hemorrhage and skin discoloration were recovered in Mar2021; clots on both legs and stomach, difficulty breathing, tiredness, blood sugar decreased, oxygen saturation low, blood pressure low was recovering; breast pain, blockage all over the heart was recovered on an unspecified date, while other events was unknown. Causality: Drug : Humira 1.Vein on her right leg was torn Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 2.Clots on both legs and stomach Causality as per reporter (Drug/Vaccine) : Not Reported Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 3.Blockage all over the heart Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 4.Difficulty breathing (Dyspnoea) Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 5.Hemorrhage Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 6.Breast pain Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 7.Skin discoloration Causality as per reporter (Drug/ Dose Not Changed) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 8.Blood pressure low Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 9.Cough Causality as per reporter (Drug/Vaccine) : Not Reported Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 10.Oxygen saturation low Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 11.Blood sugar decreased Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 12.Chest pain Causality as per reporter (Drug/Vaccine) : Not Reported Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility 13.Tiredness Causality as per reporter (Drug/Vaccine) : Not Related Causality as per Mfr.(Drug/Vaccine) : No reasonable possibility Causality: Covid-19 Vaccine 1.Vein on her right leg was torn Causality as per reporter (Drug/Vaccine) : Not Related 2.Clots on both legs and stomach Causality as per reporter (Drug/Vaccine) : Not Reported 3.Difficulty breathing Causality as per reporter (Drug/Vaccine) : Not Related 4.Hemorrhage Causality as per reporter (Drug/Vaccine) : Not Related 5.Breast pain Causality as per reporter (Drug/Vaccine) : Not Related 6.Skin discoloration Causality as per reporter (Drug/Vaccine): Not Related 7.Blood pressure low Causality as per reporter (Drug/Vaccine) : Not Related 8.Cough Causality as per reporter (Drug/Vaccine) : Not Reported 9.Oxygen saturation low Causality as per reporter (Drug/Vaccine) : Not Related 10.Blood sugar decreased Causality as per reporter (Drug/Vaccine) : Not Related 11.Chest pain Causality as per reporter (Drug/Vaccine) : Not Reported 12.Tiredness Causality as per reporter (Drug/Vaccine) : Not Related 13.Blockage all over the heart Causality as per reporter (Drug/Vaccine) : Not Related Information on the lot/batch number has been requested.; Sender's Comments: Based on the current available information, the reported events are most likely related to an intercurrent or underlying condition which is not related to the suspected drug. The case will be reassessed if additional information becomes available.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1320793-1	After 1st shot; Patient developed multiple blood clots (lungs and legs) about 10-11 days afterwards. All tests for cause of clots were inconclusive. Patient received his second dose and with in 48 hours was hospitalized for appendicitis. If it was a reaction after just one shot, I would normally think it was a coincidence. With him have severe reactions after both shots, it seems very suspicious!
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1320860-1	Got my second Pfizer Covid shot Friday, by Monday I had a huge blood clot in my neck and also a huge one in my left arm. I am now permanently on a blood thinner because of this.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1321826-1	Blood Clot in my lower right leg. Which can be life threatening if it were to dislodge and travel to the lungs and or Heart. Treatment: 30 day starter pack Eliquis 5mg tabs blood thinner, Methylprednisone 4mg dospak 21s, Acetaminophen 325mg 2@-6hrs or as needed
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1322526-1	Transient moderate headaches and nose bleeds 3-7 days post-injection, including sneezing out blood clot Menstruation lasted longer than normal Mild fatigue 1-5 days post-injection
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1323193-1	Blood clots
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1323477-1	I started having shortness of breath and went to the emergency room. It was determined I had blood clots in both lungs and I was prescribed blood thinners for the next 3-6 months

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1324020-1	About ten days after the vaccine I started to have intense cramping and I lost a blood clot. My belly was tightening up and I started to feel downward pressure. I also had back pain. I was checked out at the hospital, and they did not find anything wrong. I was 23 weeks pregnant at the time. I have not had any more cramping. I am also breast feeding my toddler and she was cranky and out of sorts after each dose of the vaccine. She was really irritable and she was running a low grade fever. My daughter is 22 months old. Pregnancy-second
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1325380-1	On April 14, 2021 at 2:15 AM, I woke up with severe pain in my lower right calf/ankle. Unable to bear weight on right leg. Reduced to crawling around house on hands and knees. Went to former Primary Care Physician on 4//14/21, where I was treated for Sciatic Nerve flare up. Returned to Dr on 4/19/21 in severe severe pain. Dr referred me to hospital where I was again treated for Sciatic related symptoms. Symptoms worsened daily. On April 28, 2021, I crawled into Hospital, was admitted. Received injection in spine. Symptoms worsened. Changed Primary Care Physicians to Dr on 4/11/21.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1326142-1	"developed a thrombosis on the left arm; varicose veins on the arm; whole arm is swollen; veins are swollen; burning pain in left lung, and it goes up to the breast and lymph nodes/ discomfort on left side Lung; burning pain in left lung, and it goes up to the breast and lymph nodes, it is sensitive there; burning pain in left lung, and it goes up to the breast and lymph nodes, it is sensitive there; has tenderness near the wrist and elbow, clarifying the inside of the elbow; felt numbness in the left arm; hot flashes; This is a spontaneous report from a contactable consumer (patient) and physician. A 76-years-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), second dose via an unspecified route of administration, administered in left arm on 10Mar2021 (Batch/Lot number was not reported) as single dose for covid-19 immunisation. Medical history included heart conditions, he has had 2 heart attacks in the past, the first one was 21 years ago, and the second was 2 years ago. Concomitant medications included all his regular medications, nothing new; (also reported as No other products). Patient previously received BNT162B2 first dose on 13Feb2021 administered in left arm for COVID-19 immunization. Patient developed varicose veins a month after he received both his shots for Pfizer's COVID vaccine. Caller stated veins are swollen. It's coming down some, but patient need to know what medications could take or not take. On the left arm right inside of his elbow, he started getting some pattern of varicose veins. It was tender to some degree and people told him to massage it and put a warm pad on it. Seems like the veins were going to pop out. Patient stated he received his vaccine at a hospital. On 03May2021, reported patient was calling in regards to a covid vaccine. He developed a thrombosis on the left arm, it was varicose veins on the arm and the whole arm is swollen, it was still swollen but it was coming down. He had been using a heat pad. Initially after 2 weeks from when he got the shot he wasn't concerned, he went to the doctor and they gave him Cat scan, Ultra sound, and blood test and did not find a blood clot. He added that he had discomfort on left side Lung, but he had discomfort several years ago, an MRI in the past that showed he has a spot in the lungs, but they didn't think it was progressing. With this thing in the past 3-5 weeks he has burning pain in left lung, and it went up to the breast and lymph nodes, it is sensitive there, he has no idea what it is. Caller was asked to clarify since he stated that he had a thrombosis but all tests show no blood clot. Caller stated that he thought Thrombosis meant when the veins are popping out and gives impression of varicose veins. He had tenderness near the wrist and elbow, clarifying the inside of the elbow. If looking at the arm from the elbow up little, it looked like worms coming out, it was veins or arteries showing. The varicose veins appearance was more pronounced close to surface initially but is now minimizing. First dose date on 13Feb2021. He was not concerned after the first dose, he was fine, he thought it was a good product and it still is. Second dose date on 10Mar2021. After the second dose he felt numbness in the left arm and hot flashes, he looked down and saw the varicose veins. He did go see the primary care doctor. He also saw his heart doctor but that was an already scheduled appointment, he did not schedule it because of this situation. the heart doctor said to put a warm pad on the arm and lift the arm up over the head every so often. Caller did not have his covid vaccine card, caller stated that he has miss placed it. He found some paper but it only has his ID number and ""LHDSER"" on it. He received the vaccine in the left arm both times. All symptoms occurred on the left side. He recieved the vaccine in a hospital. Patient asked to refer to his primary care doctor as he has all the results for the tests. Investigation included CAT scan normal; Ultrasound normal; Blood test normal and MRI abnormal which showed spots in the lungs on unspecified date. The event outcome for all events was unknown. Information for batch/ lot number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1326236-1	week later diagnosed with blood clot in right leg; 102 degree fever; This is a spontaneous report from a contactable consumer (patient). This 55-year-old female patient received BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE, dose 2 via an unspecified route of administration, administered in left arm on 24Apr2021 15:30 (Batch/Lot Number: EW0161) (age at vaccination: 55 years) as single dose for COVID-19 immunisation. The patient medical history was not reported. The patient was not pregnant. The patient had received first dose of BNT162b2 on 03Apr2021 (Batch/lot number: ER8733) for COVID-19 immunisation and experienced headache, joint pain and muscle aches. Prior to vaccination, the patient was not diagnosed with COVID-19. She had not been tested for COVID-19 since the vaccination. No other vaccines within 4 weeks prior to the COVID vaccine. Concomitant medication(s) included zolpidem tartrate (AMBIEN) taken for an unspecified indication, start and stop date were not reported. Week later after the second dose, the patient was diagnosed with blood clot in right leg on 27Apr2021 and developed 102 degree fever on 27Apr2021. The patient received treatment (unspecified) for the events. The events resulted in emergency room/department or urgent care. The events were not resolved.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1326254-1	blood clot in her lung; lung problems; bronchial asthma; may have damaged her immune system; This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer (patient, self report) A female consumer of unspecified age, reported that she received the second dose of BNT162B2 (Pfizer-BioNTech Covid-19 Vaccine, Solution for injection, Batch/Lot Number and Expiration date: UNKNOWN), via an unspecified route of administration on an unknown date as single dose for COVID-19 immunisation. Patient reported that after she received the 2nd dose of the Pfizer-BioNTech Covid-19 Vaccine, she developed lung problems. She also said she had blood clot in her lung, and had bronchial asthma. Patient wants to know if there have been many reports of this nature after vaccination. She said the vaccine may have damaged her immune system, and wants to know what Pfizer can do for her. She also said the vaccine was dangerous. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1326270-1	I had thrombosis.; I felt sick; This is a spontaneous report from a contactable consumer (patient). A 38-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 1 via an unspecified route of administration on 23Apr2021 (Batch/Lot number was not reported) age at vaccination of 38-years-old, as single dose, for covid-19 immunisation. The patient medical history and concomitant medications were not reported. On 25Apr2021 the patient felt sick and went to the hospital where she was told she had thrombosis. The patient didn't have any risks for this disease. The patient received blood thinners as therapeutic measure of the event thrombosis and the doctor told her not to combine them with other medications. Outcome of the events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1326471-1	Bladder irritation two days after first shot. Some blood tinge. Resolved in a few days. Significant bladder pain, bleeding, incl clots two days after second shot. Resolved with cranberry pills after a few days. Thought maybe was due to eating jalapena peppers in my chicken nachoes. Both weekends. Then heard that women are reporting bizarre bleeding after vaccination so re evaluated and considered the bleeding could have been an adverse reaction to vaccine and decided to report my experience.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1327477-1	Patient had vaccine on the 12th. Leg pain three days later. Diagnosed with DVT six days later. No risk factors for DVT. Unknown lot number for vaccine. Not done at my faciluty
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1327724-1	Placental abruption (with heavy bleeding) occurred two days short of week 36 in first pregnancy (which was achieved via IVF with PGS-tested embryo), resulting in stillbirth on 5/6/2021; baby was 4lb 8oz upon delivery. Due date was June 4, 2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				now was dealing with low iron count and getting infusions.; now was dealing with low iron count and getting infusions.; superficial blood clot to left lower leg found last week at dr office/blood clots pt no longer having issues with clots; This is a solicited report from a non-Pfizer sponsored program (marketing program name not available) from a contactable consumer, based on information received by Pfizer. This serious solicited report was reported to Amgen on 26/FEB/2021 by a consumer from a commercial program and involves a 72 year old female patient who had superficial blood clot to left lower leg [PT: thrombosis] while receiving Enbrel, Single Dose Prefilled Autoinjector. No historical medical condition was reported. The patient's current medical condition included rheumatoid arthritis. No concomitant medications were provided. No co-suspect medications were reported.The patient began Enbrel, Single Dose Prefilled Autoinjector on an unknown date. Last week on an unknown date in FEB/2021, the patient noticed a superficial blood clot to left lower leg at physician's office. The patient was treated with unspecified medications. The outcome of the event thrombosis was reported as unknown. Action taken with Enbrel and Single Dose Prefilled Autoinjector was reported as unknown for the event thrombosis. The causal relationship between the event thrombosis and Enbrel with Single Dose Prefilled Autoinjector was not provided by the consumer.The reporter declined to provide the lot number. The reporter declined consent for follow up. No follow up attempts are possible. No further information is expected Amgen comment: This individual case report does not change the safety profile of the product. Amgen Causality Assessment: The event thrombosis leg was assessed as related to etanercept by Amgen. The reporter's assessment of thrombosis leg with the suspect product was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality assessment. Follow-up (06May2021): This is a follow-up solicited report from a non-Pfizer sponsored program from a contactable consumer, based on information received by Pfizer from Amgen, license party for etanercept (ENBREL). ADDITIONAL INFORMATION RECEIVED ON 06/MAY/2021: In this follow up, it was reported that the patient experienced blood clots pt no longer having issues with clots [PT: thrombosis] while receiving Enbrel, Single Dose Prefilled Autoinjector. The patient's current medical condition included low iron. The patient's concomitant medications included covid-19 (corona virus disease -2019) vaccine. On an unknown date, the patient received covid-19 (corona virus disease -2019) vaccine. On an unknown date in FEB/2021, the patient had blood clot (previously reported). Then on an unknown date in 2021, the patient stated that she had no longer having issues with clots. The patient stated that they thought it might be related to covid or covid vaccine. The patient stated that she now was dealing with low iron count and getting infusions. The patient stated she had low iron as a kid. The outcome of the event thrombosis was reported as recovered/resolved. The event thrombosis was resolved on an unknown date in 2021. The reporter declined consent for follow up. No follow up attempts are possible. No further information is expected. Amgen comment:This individual case report does not change the safety profile of the product. Amgen Causality Assessment Thrombosis leg is serious and related to etanercept. The reporter's assessment of thrombosis leg with the suspect product was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality assessment.; Sender's Comments: Based on the information currently available and the safety profile of the drug there is a reasonable possibility that suspect product etanercept contributed to the occurrence of the event, thrombosis leg. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1329018-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Blood clot in neck; Shortness of breath (dyspnea); Pressure on her chest; Edema to her feet; Weight gain of 2 pounds overnight; Rash on chest; Rash on chest, possibly due to tape (dermatitis contact); Heart palpitations; Oxygen saturations dropped to 85% on room air; Diarrhea; This is a solicited report based on information received by Pfizer. A contactable consumer reported that a 41-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiration date: unknown), via an unspecified route of administration on 11Jan2021 at a single dose for COVID-19 immunization; treprostinil sodium (REMODULIN), intravenous from 03Mar2020 to an unspecified date at 0.041 ug/kg, then at 0.045 ug/kg with therapy dates unspecified, and then at 0.049 ug/kg with therapy dates unspecified, for primary pulmonary arterial hypertension; and macitentan (OPSUMIT) on 11Jan2021, orally from 11Jan2021 at 10 mg, 1x/day for primary pulmonary hypertension. Medical history included ongoing primary pulmonary arterial hypertension. Concomitant medication included sildenafil citrate. On 15Jan2021, the patient had a blood clot in neck. The patient was hospitalized from 15Jan2021 to 16Jan2021 due to the blood clot in neck. It was reported that on 15Jan2021, which is 10 months 13 days after initiating IV Remodulin, the patient was hospitalized due to blood clot in her neck. The patient went to the emergency room on 15Jan2021 and she was planned to be admitted to change her catheter, later she was hospitalized on the same day and her Remodulin central line was changed as per the prior plans of getting the catheter changed. The patient was discharged on 16Jan2021. The patient then had a rash to her chest (dermatitis contact previously reported) that started on 20Jan2021. The patient believed it was from the original dressing that was placed after her catheter was replaced. The rash to left chest around her Remodulin catheter line (dermatitis contact previously reported) was also reportedly itchy at night. On an unknown date in 2021, the patient's oxygen saturations dropped to 85% on room air (oxygen saturation decreased). The patient continued to have rash on her chest (dermatitis contact previously reported). On 29Jan2021, 10 months 27 days after initiating IV Remodulin, the patient was in the emergency room due to increased shortness of breath, pressure on her chest, edema in her feet and a weight gain of 2 pounds overnight. All the tests were negative but during walking test, the patient's oxygen saturation dropped to 85% on room air. The patient was advised to wear mask continuously. 2 weeks after receiving the first COVID-19 vaccine, the patient had heart palpitations, shortness of breath, and chest pressure 2 weeks after receiving the first COVID-19 vaccine (dyspnoea and chest discomfort previously reported). The patient went to the ER (emergency room), had worked up and was seen by a pulmonologist, nothing was found, and the patient was released home. The prescribing physician's practice ordered her to wear oxygen (O2) continuously. The patient said that she only wears it when she leaves the house and at night and was fine. She felt this episode was related to the COVID-19 vaccine and was not receiving the second dose. The patient stated that her physician had took her off the Opsumit due to shortness of breath (dyspnoea previously reported). The events 'blood clot in neck', 'shortness of breath (dyspnea)', 'pressure on her chest', 'edema to her feet', 'weight gain of two pounds overnight' and 'heart palpitations', resulted in emergency room visit. The event, 'diarrhea', was controlled with diphenoxylate and atropine (LOMOTIL). The patient had a walking distance test on an unspecified date in 2021 (results were not provided) and weight was 209 lbs on an unspecified date in 2021. The events, shortness of breath (dyspnea)', 'pressure on her chest' and 'heart palpitations', was reported as serious (medically significant). The events rash on chest, and rash on chest possibly due to tape (dermatitis contact) had not resolved while the outcome of the rest of the events was unknown. The reporter's assessment of the causal relationship of the events with the suspect product was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality assessment. Follow-up (12Mar2021): This is a follow-up solicited report based on the information received by Pfizer. Additional information included medical history, suspect drug data, reaction details and clinical course details. Information on the batch/lot number has been requested. Follow-up (04May2021): This follow-up is being submitted to notify that the batch number is not available despite the follow-up attempts made. Follow-up attempts have been completed and no further information is expected.; Sender's Comments: Based on the available information the events are most likely related to an intercurrent or underlying condition which is not related to the subject drug. There is a reasonable possibility that the drug contributed to occurrence of the event rash. The follow-up information received does not alter the previous company clinical evaluation. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1329329-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329410-1	may have blood clots; since she took the first COVID vaccine, she did have some symptoms. Her face flushed, like she had a temperature and experienced fatigue; since she took the first COVID vaccine, she did have some symptoms. Her face flushed, like she had a temperature and experienced fatigue; since she took the first COVID vaccine, she did have some symptoms. Her face flushed, like she had a temperature and experienced fatigue; This is a spontaneous report from a contactable consumer received from a non-Pfizer sponsored program,, received from a contactable consumer, based on information received by Pfizer. This case is split for a 80 year old patient taking COVID vaccine and experienced face flushed, fatigue. This 80-year-old female patient received first dose of (PFIZER BIONTECH COVID-19 VACCINE, Formulation: Solution for injection, Lot number and Expiration date: unknown), dose 1 via an unspecified route of administration on an unspecified date as 1st dose, single for COVID-19 immunization. Medical history included high frequency ablation (vein), breast cancer from 2015 to an unknown date (34 radiation treatment to left breast and no recurrence of the cancer), radiotherapy (breast cancer), atrial fibrillation from Jan2016 to an unknown date, surgery (she had 6 veins removed from her legs). Concomitant medication(s) included apixaban (ELIQUIS) taken for cerebrovascular accident prophylaxis (batch no. 1745824) (oral, 5 milligram, twice a day) in 2017 or 2018, losartan taken for blood pressure (50 mg once daily), paracetamol (TYLENOL) taken for pain (as needed (but not every day)), vitamin b nos, ascorbic acid, zinc, colecalciferol and cyanocobalamin (VITAMIN B12 [CYANOCOBALAMIN]). The patient experienced since she took the first COVID vaccine, she did have some symptoms, her face flushed, like she had a temperature and experienced fatigue, and may have blood clots. The patient reported that she may have blood clots but was unsure because she did not underwent any tests to confirm. The patient was due for her second COVID-19 vaccination. The outcome of the events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329464-1	her face was swollen; Extreme pain in her head, not like a migraine (she has had migraines) it came suddenly and the pain grew stronger and she though her head was going to explode; Swollen legs accompanied brusing rash; brusing rash; Swollen legs accompanied brusing rash; Effects were similar to blood clot symptoms; The initial case was missing the following minimum criteria: unidentifiable reporter. Upon receipt of follow-up information on 04May2021, this case now contains all required information to be considered valid. This is a spontaneous report from a contactable consumer (patient). A 57-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on 30Apr2021 (Batch/Lot number was not reported) at 2nd dose, single for COVID-19 immunization. The patient's medical history included migraines. Concomitant medications were not reported. The patient previously took the first dose of BNT162B2 on unspecified date for COVID-19 Immunization. The patient experienced side effects after getting the 2 doses of the PFIZER-BIONTECH COVID-19 VACCINE, 2nd dose was given on 30Apr2021. The patient had a bad reaction, she called to report it last night, she may have a CT scan for follow-up of the residual vaccine, the effects were similar to blood clot symptoms, she had a extreme pain in her head, not like a migraine (she has had migraines) it came suddenly and the pain grew stronger and she though her head was going to explode, she almost went to the emergency room (ER). Assuming all these side effects were attributed effects that may be similar to blood clotting symptoms. Side effects experienced were as follows: Extreme headache characterized by a painful somehow compared to by the patient as she was grabbing her head and thought it would explode. She almost went to the ER for this and the pain lasted for 9 hours starting from the same night at 21:30, when she got the vaccine (30Apr2021). She mentioned she took Ibuprofen 600 mg for this and it slowly eases the pain until it went away. Swollen legs accompanied by bruising rash. This happened alongside the extreme headache (30Apr2021). Her face was swollen as well the next day (01May2021). She thought that all the mentioned above side effects were characteristics of blood clotting side effect. The outcome of the event headache was recovered on 01May2021; and unknown outcome for the other events. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329475-1	pulmonary embolism; Blood Clot; This is a spontaneous report from a contactable consumer (patient). A 59-yearold male patient received bnt162b2 (BNT162B2), dose 1 via an unspecified route of administration, administered in Arm Left on 07Apr2021 (Batch/Lot Number: ew0153) as 1st dose, single (at the age of 59 years) for covid-19 immunization. The patient's medical history and concomitant medications were not reported. The patient has not had covid-19 prior to vaccination. The patient experienced pulmonary embolism on an unspecified date and blood clot on 22Apr2021 with outcome of not recovered. The patient was hospitalized for blood clot for 2 days. The patient underwent lab tests and procedures which included blood test: negative on 26Apr2021. Therapeutic measures were taken as a result of the events incudes blood thinner. The outcome of the event blood clot was not recovered and unknown outcome for the event pulmonary embolism. The patient was tested negative for Covid-19 on 23Apr2021 (post vaccination). No follow-up attempts are possible; information about batch number already obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329495-1	numerous blood clots below the knee; left leg swelling/ It got so swollen/left foot swelling; pain with walking left leg; This is a spontaneous report from a contactable nurse reporting for herself. A 78-years-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in Right Arm on 24Feb2021 (Batch/Lot Number: EN6202) as single dose for covid-19 immunisation (Age at vaccination 78 years). The patient medical history was not reported. On 04Feb2021 the patient received the first dose of BNT162B2 and experienced sore arm. The patient's concomitant medications were not reported. The patient experienced pain with walking left leg on 28Feb2021 with outcome of recovering , left leg swelling/ it got so swollen/left foot swelling on 03Mar2021 with outcome of not recovered , numerous blood clots below the knee on 04Apr2021 with outcome of not recovered. The patient underwent lab tests and procedures which included blood test: unknown results, ultrasound joint: numerous blood clots below the knee on 04Apr2021 , vital signs measurement: unknown results. Course of the event: The patient waited for the timeframe after the second dose and had no problems. 4 days after, she experienced pain with walking in her left leg. She took Aleve for a few days. It got so swollen, she should have gone then to her Primary Medical Doctor, but she did not go until 25Mar2021. She was sent to Hospital for an Ultrasound of her leg on 04Apr2021. After the Ultrasound, she was taken to the Emergency Department at the hospital. The doctor told her she had numerous blood clots below the knee. Was prescribed Eliquis 5mg. She was instructed to take 2 tablets or 10mg twice a day, 8hrs apart for 1 week and then 1 tablet twice a day for the remainder of the 74 tablets. She completed that. She was out of them and got a refill of Eliquis 5mg on 26Apr2021. She received 60 tablets and can refill for 5 times. She is really concerned. The swelling is still so bad and so that could that mean blood clots are still in there. The ER doctor told her that the medication is dangerous. It thins the blood and can cause bleeding and the blood clot can go to her heart. Follow up information has been requested.; Sender's Comments: The causality has been assessed as related to bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in Right Arm on 24Feb2021 (Batch/Lot Number: EN6202) as single dose for covid-19 immunization, based on temporal association and profile of the product.,Linked Report(s) : 2021499984 same patient/drug, diff dose/event
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329511-1	blood clot in right leg; This is a spontaneous report from a contactable consumer (patient). A female patient of unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 15Apr2021 (Batch/Lot number was not reported) as 1st dose, single for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient received the first COVID vaccine on 15Apr2021, a week later (in Apr2021), the patient experienced a blood clot in right leg. The patient is currently on blood thinners and was scheduled to receive the 2nd vaccine on 06Mar2021. Outcome of the event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329517-1	This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE; lot number and expiry date: unknown), via an unspecified route of administration on 09Apr2021 as 1st dose, single; via an unspecified route of administration on 30Apr2021 as 2nd dose, single for COVID-19 immunization. The patient medical history and concomitant medications were not reported. The patient experienced first signs of blood clot on 28Apr2021 and ended up in hospital from 30Apr2021 to 02May2021, getting blood thinner. The patient thought he/she should report this in case it is possibly connected to vaccine. The outcome of the event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329521-1	Developed a blood clotting; This is a spontaneous report received from Pfizer sponsored program. A contactable consumer (patient) of unknown age reported that she received second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number unknown) on an unknown date at single dose for COVID-19 immunization. The first dose of vaccine was received on 24Feb2021. Medical history and concomitant drugs were not reported. The consumer reported that two weeks after she had her second dose of Pfizer vaccine she developed a blood clotting. The treatment has been very costly. The outcome of the event was unknown. Follow-up attempts are needed. Information about lot/batch number is requested.; Sender's Comments: Linked Report(s) : PFIZER INC-2021229192 same patient, different vaccine dose/event

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329533-1	Blood clot; This is a spontaneous report from a contactable healthcare professional and a physician. A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number/expiration date not reported), via an unspecified route of administration on an unspecified date as unknown, single for COVID-19 immunization. Medical history and concomitant medications were not reported. The reporter had a patient who may have had a possible side effect. It was reported that the patient may be experiencing a blood clot on an unspecified date. The reporter wanted further direction on whether or not an anticoagulant can be given. Anatomical location of blood clot is unknown. The outcome of the event was unknown. Information on the batch/lot number has been requested.; Sender's Comments: Based on available information, a possible contributory role of BNT162B2 vaccine can not be excluded for the reported event of blood clot due to temporal relationship. Case will be re-assessed upon the additional information provided. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329544-1	Blood Clot; right leg was red & hot; right leg was red & hot; itching; Pain; This is a spontaneous report from a contactable consumer. A 63-year-old patient of an unspecified gender received second dose of BNT162B2 (BNT162B2), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunization. Medical history included covid-19 from 28Mar2021 to an unknown date. The patient's concomitant medications were not reported. The patient experienced blood clot, right leg was red & hot, itching and pain all on unspecified date. On 28Mar2021 I caught covid-19. The staff next at united states penitentiary put me in quarantine until 14Apr2021. They did not do anything for me what so ever while I had covid-19. Shorty after I became better (on my own) I took the exist shot of the vaccine Pfizer. 3 weeks later in the end of this month I took the second vaccine shot. My right leg in my blood vein started burning me up. My right leg was red & hot so I went to commissary to purchase ORAJEL to put my leg to kill the pain and itching. I then put a sick call in to the prison nurse, she came the next day and said you have a blood clot and if it moves to let her know. 3 days later or more she was back in the housing unit and I ask her what she was going to do about my blood clot. She said oh you found have a blood clot you just have the big viens. Outcome of the events was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329555-1	blood clot; Also have had neck pain since the first vaccine that will not go away; extreme pain in my left leg; unable to walk; This is a spontaneous report from a contactable consumer (patient) who reported events after first and second dose of the vaccine. This case is for events after the second dose. A 33-year-old female patient (not pregnant) received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot number was not reported), via an unspecified route of administration in Arm Left on 31Mar2021 at 13:00 (at 33 years) for covid-19 immunization. Medical history included asthma. Concomitant medication included methylphenidate taken for an unspecified indication, start and stop date were not reported. The patient previously took the first dose of the vaccine on 03Mar2021 at 13:00 in Left arm for covid-19 immunization and had neck pain. On 19Apr2021 at 12:00 (also reported as 3 weeks after the final vaccination) the patient experienced suddenly extreme pain in her left leg and was unable to walk on it. Doctors found a superficial blood clot. Also had neck pain since the first vaccine that will not go away. Events required Doctor or other healthcare professional office/clinic visit and treatment (unspecified) was given. The final outcome of all the events was reported as recovering. Information on the batch number has been requested; Sender's Comments: Linked Report(s) : -PFIZER INC-2021523861 same patient/drug, different dose/event
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1331283-1	Blood clots

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332671-1	"Severe calf muscle cramps, possible blood clot; Severe calf muscle cramps, possible blood clot; This is a spontaneous report from a contactable other hcp. A 28-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in Arm Right on 24Mar2021 12:00 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunization. Medical history was reported as ""none"". The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient hasn't been tested for COVID-19. the patient previously received the first dose of BNT162B2 on 26Feb2021 at 10:00 administered in the left arm for COVID-19 immunization. The patient's concomitant medications were not reported. The patient experienced severe calf muscle cramps, possible blood clot on 01Apr2021 12:00. The outcome of events was not recovered. Patient did not receive any treatment for the events. Follow up needed, further information has been requested; Sender's Comments: A causal association between BNT162B2 and the reported events cannot be excluded based on a compatible temporal relation. Case will be reassessed upon receipt of follow-up information. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332693-1	blood clot; Sciatica; She then got sciatica in her leg and couldn't walk; charley horse in her lower leg while walking; soreness in arm / pain in her foot / lower leg/calf still hurt; This is a spontaneous report from a contactable nurse (patient). This 86 years old female patient received the second single dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot EN6203) on 04Mar2021 at 11:00, in left arm, for COVID-19 immunisation. No other vaccines were administered on the same date. The patient received the first dose of BNT162B2 vaccine intramuscular, on 11Feb2021 at 18:30 (lot EN6201). The patient received some years ago an unspecified pneumonia vaccine and an unspecified tetanus vaccine experiencing after both headache, swollen arm and rash. She didn't react to the flu vaccine. Her body didn't want any more vaccine. Concomitant medications were none. On an unspecified date in Mar2021, maybe a week or two after the second vaccination, early to mid afternoon, the patient had what felt like a charley horse in her lower leg while walking, but didn't pay any attention to it, just put ice on the leg. She then got sciatica in her leg on 24Apr2021 and couldn't walk; that was what sent her to the hospital as it was getting worse and she went to the hospital because of the pain. She just ignored it for at least a week but she was still having pain after the second week and could hardly walk due to the sciatica in that leg. At the hospital they found she had a blood clot in the calf of her leg on 28Apr2021. She was hospitalized from 28Apr2021 to 02May2021. The doctors in the hospital recommended that she should report her experience even though they didn't know if it was due to the COVID vaccine due to the proximity. The hospital doctors took care of her but her primary care provider knew that she had been in the hospital. The doctors gave her medications in the hospital that helped the pain in her foot and the sciatica but her lower leg/calf still hurt so they decided to do an ultrasound and found a blood clot. The patient still had to give herself injections in the belly for blood thinner to treat the blood clot. The patient also experienced soreness in arm on an unspecified date in 2021. Sciatica required ER visit. Blood clot was resolving. The other events outcome was unknown. The doctor said that there was a possibility that the blood clot could be related to the COVID vaccine, but it may not be related at all.; Sender's Comments: Based on the information provided by the reporter, including a prolonged onset latency of nearly two months, it appears unlikely that BNT162B2 contributed to the reported events. These are likely intercurrent medical conditions in this elderly patient. This case will be reassessed upon receipt of additional information. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Regulatory Authorities, Ethics Committees and Investigators, as appropriate.,Linked Report(s) :PFIZER INC-2021501847 same patient/drug, different vaccine dose/AE

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332754-1	felt unwell; took his oxygen level and it was at 80%; he had swelling; difficulty breathing; tachycardia; allergic reaction; tested Covid-19 positive; blood clots; he got very sick; This is a spontaneous report from a contactable consumer(patient's daughter). A 71-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 24Apr2021 (Batch/Lot number was not reported) at the age of 71-year-old as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient was fine and nothing happened until after the vaccine. The patient took the 1st dose on 24Apr2021 and the next day he got very sick, they thought it was just a reaction from the vaccine, then he was swelling said he was not feeling well. The day after the vaccine, the patient was rushed to the emergency room and confined to the ICU. He felt unwell and it got worse and worse, we took his oxygen level and it was at 80%, he had swelling, difficulty breathing and tachycardia. The reporter thought it was an allergic reaction but this morning the patient tested positive for covid, he didn't go out the only time he went out was to get the vaccine. The reporter asked If he didn't know he had covid and got the vaccine, will he get more sick. In the report they found blood clots and the doctors thought it was from the vaccine. He was diagnosed with blood clots and difficulty breathing and he tested Covid-19 positive. Events seriousness criteria reported as hospitalization. The outcome of the events was unknown. Information on the lot/ batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332784-1	Blood clot in leg; This is a spontaneous report from a contactable consumer (patient). A 66 year old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 2 via an unspecified route of administration on 29Mar2021 at 12:45 (Batch/Lot Number: EN6198) as a single dose for COVID-19 immunisation. Medical history included COVID-19 from Nov2020 to an unknown date: came down with COVID virus in November and it felt like mild flu symptoms. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 1 via an unspecified route of administration on 08Mar2021 at 12:45 (Batch/Lot Number: EN6198) as a single dose in the right shoulder for COVID-19 immunisation. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient did not have any adverse events following prior vaccinations. The patient's medical history included: father had a blood clot in his 50s. Concomitant medication (other medications taken in two weeks) were none. On an unspecified date in Apr2021, the patient experienced blood clot in leg. Relevant test included: ultrasound on 15Apr2021 showed blood clot. The clinical course was as follows: On 04Apr2021 or 05Apr2021 he started to notice a problem on his right leg. He went to see the doctor and it was determined he has a vascular problem. This problem got worse and ended up in the emergency room and was informed he had a blood clot in his leg. The clot started down by his ankle and had moved up towards half way to knee. Then moved up higher. He went to the emergency room on 15Apr2021and was diagnosed with a blood clot with an ultrasound scan that was done. He was then prescribed a blood thinner. He was informed it is a superficial thrombosis. The patient clarified the problem he started to notice with his right leg as it was like a bump, and sore. He was not sure what it was. He thought at first he may have banged his leg. Then he noticed it was sore and getting higher. He went to see his physician who referred him to a vascular surgeon. It took a while to get in to be seen by the vascular surgeon because so busy. A few days after that he went to the hospital, the emergency room because he was not getting better. That is when he found out he had a blood clot. Treatment for the blood clot in leg included, patient was prescribed rivaroxaban (XARELTO) 10mg once a day. The patient confirmed the blood clot was getting better. The outcome of the event blood clot was recovering. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332801-1	blood clots in his nose; White Spots on the Palm of Hand; allergic reaction; This is a spontaneous report from a contactable consumer (patient). A male patient of unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as 1st single dose for covid-19 immunization. The patient medical history was not reported. The patient's concomitant medications were not reported. The patient was having allergic reaction he did have blood clots in his nose that did go after some days but now he did have white spots on the palms of his hands and he wanted to know if it was like common allergic reaction. Outcome of the events was unknown. No follow-up attempts are needed; information about lot/batch number cannot be obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1333638-1	pt says she had a hot red swollen right leg that was very painful. On 4/16/21 she went to the Hospital ER. She had blood test and was given an US of the right leg where they found 2 blood clots. She was started on blood thinner. She was discharged and told to FU w/ PCP. On 4/21/21 she saw the doctor. She increased the blood thinner because she found a abdominal hematoma this size of two grapefruits. On 4/25/21 she was having trouble breathing and her leg was red and inflamed. She went back to the Hospital ER they thought she was having a pulmonary embolism. They did more blood work and chest x-ray. She was given Lovenox shots and took her off the blood thinner. She was discharged to go home and give herself these shots. On 4/27/21 she had a reaction to the Lovenox shot so she called her PCP. She saw the doctor on 4/29/21 she was told to continue using the Lovenox. She went back to Dr. on 5/4/21 and she was taken off Lovenox and put back on blood thinner Eliquis with doubled dosage. She was monitored. On 5/8/21 both of her legs were red and swollen and she went to her Doctors office. They did a blood test and was sent home. On 5/10/21 she went back to her PCP. Dr. office where she was sent to Hospital ER for direct admit. They did another US and a CT. They found she had blood clots in both legs. She was given IV fluids, antibiotics, was given heparin and taken off Eliquis. On 5/13/21 she was given warfarin along w/ heparin. She was discharged on 5/16/21. She still has pain and redness in both feet and legs and has been diagnosed w/ Cellulitis. She is now taking Oxycodone and Coumadin. She will see her PCP on 5/21/21.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1333991-1	Blot clot followed by fatal aortic thrombosis
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1334947-1	huge blood clot in her neck and a huge one in her left arm; her arm had swollen up at the elbow and was red, painful and warm to the touch; her arm had swollen up at the elbow and was red, painful and warm to the touch; her arm had swollen up at the elbow; her arm had swollen up at the elbow and was red, painful and warm to the touch; her arm had swollen up at the elbow; This is a spontaneous report from a contactable consumer (patient). A 48-year-old female patient received the second dose bnt162b2 (BNT162B2, Solution for injection, Lot Number: ER2613), via an unspecified route of administration, administered in Arm Left on 07May2021 (received at 48-years-old) as 2ND DOSE, SINGLE for COVID-19 immunisation. Medical history included fibromyalgia, colon cancer/postchemo from 2018, APS (antiphospholipid syndrome), arthritis and allergy to sulfa drugs. The patient was not diagnosed with COVID-19 prior to vaccination and was not tested for COVID-19 post vaccination. She was not pregnant. Concomitant medication(s) included pregabalin (LYRICA) taken for an unspecified indication, start and stop date were not reported; omeprazole (OMEPRAZOLE) taken for an unspecified indication, start and stop date were not reported; oxybutynin (OXYBUTYNIN) taken for an unspecified indication, start and stop date were not reported; ergocalciferol (VIT D) taken for an unspecified indication, start and stop date were not reported; cyanocobalamin (VIT B12) taken for an unspecified indication, start and stop date were not reported. Past drug history included known allergies to naproxen and oxycodone. The patient previously received the first dose of bnt162b2 (BNT162B2, Lot Number: EW0161), administered in Arm Right on 16Apr2021 12:00 for COVID-19 immunisation. The patient got her second dose of the Pfizer vaccine on Friday, 07May2021. By Monday evening, 10May2021, her arm had swollen up at the elbow and was red, painful and warm to the touch. After appointment she had on Tuesday, 11May2021, she went to the ER (emergency room) as it had gotten worse and confirmed she has a huge blood clot in her neck and a huge one in her left arm. She was put on permanent blood thinners after following up with my PCP (primary care physician). The adverse events resulted in Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Life threatening illness (immediate risk of death from the event). As corrective treatment, the patient received Lovenox, Wafarin, Zarelto. Outcome of the events was recovered with sequelae on an unspecified date in May2021. The adverse events were assessed as life-threatening.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1335964-1	"pain in his leg that felt like a clot; pain in his leg; This is a spontaneous report from a contactable consumer, the patient. A 45-year-old male patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration in the right arm on Mar2021 at 11:15 (at the age of 45-years-old) as a single dose for COVID-19 immunization. Medical history included ongoing venous insufficiency since 2019. Concomitant medications were none. The patient did not receive any other vaccinations within 4 weeks of the vaccine. The patient stated that he received his first dose on 30Mar2021 or 31Mar2021. On an unspecified date, the next day after the first vaccination, the patient experienced pain in his leg that felt like a clot for the first week and a half to two weeks. Then, the patient got a little bit better, even though he had almost gone to the emergency room, but he got better after a couple weeks. The clinical outcome of ""pain in his leg that felt like a clot "" was resolved on Apr2021. Information on the lot/batch number has been requested.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021507016 Same patient/drug, different dose/event"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1337718-1	patient developed a blood clot

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340343-1	This is a spontaneous report from a contactable consumer (patient). A female patient of an unspecified age received the first dose of BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, lot number and expiry date unknown), via an unspecified route of administration on 05May2021 as a single dose for COVID-19 immunization. Medical history included lupus, high blood pressure and severe pancreatitis wherein she had surgery all on unknown dates and unknown if ongoing. The patient's concomitant medications were not reported. The patient reported some adverse effects after the first dose of the Pfizer COVID-19 vaccine. The patient reported stomach pain and nausea from an unknown date and fears it may be a blood clot and wants to know if these are common side effects. The patient reported having stomach pain on an unspecified date and she doesn't know if it is normal. At first she thought it was because she was so nauseous, but the nausea went away. Then she thought maybe it was like a bowel thing and she was going to have issues like diarrhea, but that was not the case. The pain is pretty bad. She is wondering if that can be a side effect. She is just wondering if she should follow up with a doctor or go to the hospital, because it hurts pretty bad in 3 different sections of her stomach. Her stomach feels like her arm felt when she got the shot. It feels like she had bruising, not exactly stabbing but it is pretty painful and she is kind of concerned. She doesn't know if her doctor would have information on the vaccine for what to do, and they would probably send her to the hospital. She hopes it's not something serious. She is worried something went wrong. Her entire family got the same shot and were fine. The pain is just like where her pancreas and gallbladder are. Since she had severe pancreatitis last summer, and it hurts where she had surgery. She wants to know if it is related to the shot or something else. The outcome of the event nausea/so nauseous was recovered while the outcome of the other events was unknown. Information on the lot/batch number has been requested. Sender's Comments: A possible contributory effect of suspect BNT162B2 on reported blood clot cannot be excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340367-1	"This is a spontaneous report from a contactable consumer (patient) and a physician. A 64-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 2 via an unspecified route of administration, administered in Arm Left on 09Apr2021 in the afternoon around 2 or 3 o clock (Batch/Lot Number: EW0153) as 2ND DOSE, SINGLE for covid-19 vaccination. The vaccination was done in her building, a medical team came to them. She lives in subsidized housing. The patient's medical history included tendonitis; juvenile idiopathic arthritis from Dec 1969 and ongoing; lupus nephritis from 1989 (diagnosed ten years after SLE); systemic lupus erythematosus (SLE) from Dec 1979; discoid lupus; proliferative kidney disease stage 3; sjogren's; raynaud's; arthritis (since she was 13); bursitis; oral necrosis (this was from being on a calcium drug that they give menopausal women, it was out of her head right now when reporting. She could not think of the name. The drug was supposed to promote bone growth and keep her from getting osteoporosis); menopausal (because of chemo she started menopause at 42); vulvar lichen sclerosus; filamentary keratosis from 2015 and ongoing (this was since either Feb 2015 or Mar 2015); kidney cysts(one is benign and one is being watched); liver hepatic hemangioma; allergies (Allergies to aspirin, penicillin, dextrose IV, Biaxin, Levaquin, Cipro, band-aid adhesive, zofran, sulfa topical. The sulfa is a topical as opposed to being the sulpha with a ph.); adeno conjunctivitis from Jan2015 and ongoing (she had a really bad case of this in JAN 2015 and had to quarantine for three weeks. She lost her vision, she could not see for three days. She got a fever. It left her with a filament thing. It is just delightful. It is viral and really bad. She got it at her eye doctors office when she was there for an annual visit. She sometimes had to go there 3 times to have filament growing on her eye scrubbed off. It is wicked painful. It is quite an adventure.); she used to be three quarters but she shrunk a little; has been a couch potato for a year, it comes with the territory with lockdown and being depressed; she is immunocompromised and catches many things and is high risk She feels like a sitting duck; had issues on her palm of hands; had issues on her breasts, and chest; and various problems. Family Medical History Relevant to adverse events was none. No other vaccinations within four weeks prior to the first administration date of the suspect vaccine. No other vaccines administered on same date with the Pfizer vaccine considered as suspect. Concomitant medications included vitamin b complex (B-COMPLEX 100) 100 as dose did not clarify units, has been on for years; hydroxyzine embonate (HYDROXYZINE PAM) Dosage 25 mg, she did not remember indication, may be allergies, if she is around a cat, she takes it prophylactically; acetylsalicylic acid (ASPIRIN) low strength, 81 mg twice a week, has been on for decades, nephrologist put her on it in the 90's and She did not remember why, he said it preserves kidney function or it is good for overall health; pentoxifylline ongoing at 400 mg, 3 a day, she has been on for years, her vascular doctor prescribed; prednisone dosage:1 mg, she takes and alternates between 6 and 7 mg, If she takes lower than that, she is a ""narcoleptic"" (as reported, did not clarify); rosuvastatin Dosage:10 mg, frequency not specified, it is a nightly thing; fish oil (SUPER OMEGA 3) overall supplement that was suggested by former nephrologist at some point some; estradiol (YUVAFEM) Dosage:10 micrograms for menopause, vaginal dryness; there was no variable in her medication routine, she is on a long list of medications (she could not remember, one was blood pressure, with the last administration she was worked up and her blood pressure would not come down, she also takes Tylenol as needed), she provided the details as she knows them, but did not know full details for all of her medications.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	There was nothing new to her routine except like when Prilosec was taken off **Adverse Event Description**
				the market, she got a different medication for it. She has been on all of these for long term maintenance. Historical vaccine included bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 19MAR2021 (at the age of 64-year-old)(Batch/Lot Number: ER2613) for covid-19 vaccination, adverse events following prior vaccinations: none; and flu vaccine for immunization in last fall, 2020. She has a history of obscure side effects to medication, things that people have never heard of and she is the 1 in a million that has ever had these problems. The patient previously took ondansetron (ZOFRAN), (""a chemotherapy treatment"", given before chemotherapy), which gave her a homicidal migraine and they had to give her ice packs for her head; 9 rounds of ""Cytox"" (Cytoxin or Cytoxan) she was not sure how this was spelled, and chemotherapy (9 rounds of chemo); azathioprine (IMURAN), she does not know if it was spelled Immuran or Imuran; prednisone a number of years and she said to her rheumatologist, ""it did not make sense on an intellectual basis that the prednisone being effective in her body as it was once"" and he said he would give her IMURAN which is given to kidney patients to cause less of a reaction with the organ, and patient mentioned she had barely an immune system after all these years on prednisone, she went through 3 rounds of steroids; Heparin IV, which made her arm feel on fire on the inside, which they would not give her a flush when she had IV treatment; Dextrose IV made her skin feel hot and itchy, her mouth felt like she put something in that had been soaking in rubbing alcohol; ciprofloxacin (reported as ""Cipro""), she had a bad reaction to Cipro, she was given after she was bit by a dog on her calves, she knew it was attached to and is ""hurting her calves the rest of her life/the podiatrist said if it has not gone away, she is kind of stuck"" (as reported, not specified). The patient just wanted to give Pfizer a full picture of what she was dealing with and she is an unusual case. She added that chemo was in the 90s and that is when all of the medication reactions she mentioned were. She absolutely does not have any of the information about the products she mentioned she had previous reactions to. Every time they did ""this"", it ""hurt"" and she asked them not to do it anymore. She asked them to not do dextrose anymore with the chemo because it feels like it is on fire, and she asked them to mix it with saline. She did not have any additional information, did not have the NDC, LOT, expiration date, or manufacturer for any of these medications she reported historical adverse events on. The patient reported that she got her 2nd shot of the Pfizer covid vaccine on 09Apr2021. She has soreness at the site, more-so several hours after getting the vaccine (09Apr2021). Just her arm felt sore within moments after the second vaccine, but she did not get this after the first. She rested and took it easy that weekend. There was Sore, Tired, general lethargy, and fatigue (Apr2021): when she looked online, it really only told her what everyone knows about that you might be sore and tired and she had all that, she thought she was through with all of that. She had no chills, just general lethargy and fatigue. She was sluggish and tired and not herself that weekend (Apr 2021). She stayed home so it is difficulty to say. Her arm was definitely sore. For sluggishness and fatigue, she said that she has been a couch potato for a year, it comes with the territory with lockdown and being depressed, so it is difficult to decide if it is health or physically induced or a emotional base. She did not knowingly spike a fever, no chills or sweats. And on Monday (did not clarify dates but states it was Monday after 09APR2021, throughout the weekend she took it easy), she noticed with her foot, she had difficulty getting out of bed could not put weight on her right foot. She was alarmed. She stayed home and rested and did a lot of napping and taking it slow. It is still difficulty and moving certain ways hurts more if she flexes. She has a known history of tendonitis so she wore her brace in the meantime. She thought maybe she had tendonitis in both ankles and this was similar so maybe it was a different set of tendons. She would wear a brace, and eventually it will get better in time. She also was wearing braces like compression socks that toes stick out of. She thought she had tendonitis and that is what she has done in the past. It has been a month and she is not better. This was her left foot, most of the pain was focused on the sole of her foot towards the outer edge. That is where it started. Then it migrated over time. That was weird and had her worried. It was all in her arch in the initial spot. She notes it is not like the tendonitis she had the first time. She does not want to be an alarmist or have her doctor think she is a hypochondriac she waited. She was then rubbing her foot, the right one, and noticed it was tender on the top of her foot just like the left one. There was sore. There was pain in right foot, pain in feet. Pain in feet, her left foot had most of the pain on the top and bottom and a lot of it is along the outer part of her sole of the foot. In her right foot, the pain is mostly at the fleshy part below her toes and on top of her foot. The pain moved around into the other foot for about a month so she went to the podiatrist yesterday (06May2021) and doctor did x-rays. The night before (05May2021), the opposite leg had a line of black and blue dots on her right calf that went up her leg, she did not notice this until Wednesday (05May2021) evening because she never examines it. She soaped up a washcloth and she knew where her parts are in the bath. Her mind wanders and she was not looking. She happened to be wearing something where her legs were not covered and that is how she knew. It did not hurt. It was strange, looking like bruising in a line. The doctor said it was all like one vein and he cold feel it. She can not feel it, but he could. The doctor told her these could be clots. She still has them. Doctor said the other leg where she had the pain was only soft tissue pain. Doctor found that every place that hurt when he touched it and it still hurts where he had touched it. She was told she does not have arthritis and she has had arthritis since she was 13. The doctor did not know what it is. It was her left foot constantly, and the right foot hurt off and on during that period. The left for sure was kind of a dull ache and throbbing. The right was only a little bit. The left was worse. It appeared to be soft tissue that was hurting and he

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				said there are so many side effects of this new vaccine and to maybe try heat or ice. Ever since the patient left doctor's office until now (the time of reporting) she did not known if she need to be concerned, that if she does not know what it is, can ice or heat make it worse? She did not know the next steps. Treatment, she tried to keep it elevated because that can not hurt but that is about it. She has not done heat and ice because if the doctor does not know what it is, how does he know it will not make it worse. He asked if she could take aspirin and she said no she is allergic to aspirin. Tylenol she does not take often, only when she feels like she needs it and she has not been doing it often. When she stays off it, it does not hurt as much as when she is walking. She wanted to know if she should be worried and she wanted to add (report this) to the database if this is a side effect and if Pfizer has ever heard of this before. There was also tired, general lethargy, and fatigue. At this point when reporting, she is single and lives alone, and said that ""God forbid something happens, there is nobody to say let's get you to a hospital"". She heard about blood clots on the news. When they said blood clot yesterday, it was a red warning light going on and a siren. At the time of reporting it was Friday and she did not know that if she need to rush to the hospital. Her primary care physician (PCP) said she had never heard of this and to call Pfizer. The patient asked if she is one of a small group of people this has happened to. She asked if she should not worry and it will go away or should she get to a hospital. She thought she had a problem with her feet (foot problem) but her podiatrist does not know what was going on. She communicated with her physician office today and she said she has never heard of this. She wanted to know if her symptoms have been heard of and if she should go to the hospital. The events did not require a visit to emergency room. The patient underwent lab tests and procedures which included: height as ""5 and three quarters of an inch feet (5'0.75"")"" on an unspecified date; rapid COVID-19 test on 06May2021: negative; regular COVID-19 test on 06May2021: negative; x-ray on 06May2021 to make sure she did not break anything: unknown results. She also went because she had not done it in a year and she had something from the health department saying to get tested weekly for the virus (clarified that she was talking about the COVID tests as what she went and got done). The outcome of the events was unknown."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1340371-1	a blood clot in his left arm; possible sepsis; malnourishment; dehydration.; became so weak he slumped down; health has steadily declined; This is a spontaneous report from a contactable consumer (patient's son). A 75-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in right arm on an unspecified date (batch/lot number was not reported) as 2nd dose, single for covid-19 immunisation. Medical history included Parkinson's disease, essential thrombocythaemia, and chronic kidney disease which were all under control; and allergies to latex and iodine. The patient's concomitant medications were not reported. On 01Mar2021, shortly after second dose, the patient became so weak, he slumped down and had to be lifted by reporter's uncle, after this incident, his health has steadily declined and now he was hospitalized again on an unspecified date with a blood clot in his left arm, possible sepsis, malnourishment and dehydration. He has Parkinson disease, essential thrombocythemia, chronic kidney disease; however, all were under control until he was vaccinated with the second shot. The events resulted in doctor or other healthcare professional office/clinic visit, emergency room/department or urgent care, hospitalization, disability or permanent damage. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19 and since the vaccination, the patient had been tested for covid-19. The patient underwent lab tests and procedures which included covid test: negative on an unspecified date. Outcome of events was not recovered. Information about lot/batch number has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1340399-1	This is a spontaneous report from a contactable consumer (patient). A 35-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), via an unspecified route of administration administered in left arm on 31Mar2021 09:00 (at the age of 35-year-old) (Batch/Lot Number: ER8730) as 2ND DOSE, SINGLE for COVID-19 immunization. The patient did not receive any other vaccines within four weeks prior to the COVID-19 vaccine. The patient was not diagnosed with COVID-19 prior to vaccination. Medical history included hypertension. Concomitant medication included ascorbic acid, beta carotene, biotin, boron, calcium, calcium pantothenate, chloride, chromium, colecalciferol, copper, cyanocobalamin, folic acid, iodine, iron, lycopene, magnesium, manganese, molybdenum, nickel, nicotinamide, phosphorus, potassium, pyridoxine hydrochloride, retinol, riboflavin, selenium, silicon dioxide, thiamine, tin, tocopheryl acetate, vanadium, vitamin k nos, zinc (CENTRUM MEN). The patient previously took bactrim and experienced allergies. Historical vaccine includes BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot number: EN 6206), administered in left arm on 10Mar2021 at 08:45 AM (at the age of 35-year-old) as 1ST DOSE, SINGLE for COVID-19 immunization. On 27Apr2021 at 08:00 AM, the patient experienced bilateral embolisms (blood clots in lungs) and also had blood clot in left leg. Symptoms before diagnosis included dry cough, coughing up blood, chills, fever, body aches, and body pains. The events caused hospitalization. The events resulted in doctor or other healthcare professional office/clinic visit and emergency room/department or urgent care. The patient underwent lab tests and procedures which included COVID-19 test via nasal swab post vaccination which was negative on 29Apr2021 and on 07May2021. Therapeutic measures were taken as a result of bilateral pulmonary embolisms and had blood clot in left leg, chills, and body aches/body pains which included Lovenox injection (blood thinner) and Tylenol for pain. The patient was recovering from all events.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340433-1	"This is a spontaneous report from a contactable consumer (patient's wife). A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot Number: Unknown) as single dose for COVID-19 immunization. Medical history included dementia and he shrunk he is 5'9"" or 5'8"". Concomitant medication(s) included warfarin sodium (COUMADIN), at 4 mg once daily in the evening 6 days per week, On Fridays he takes 6 mg once in the evening. Patient's wife says her husband had one vaccination, the Pfizer vaccine and a week later her ended up in the hospital. They had to postpone his second dose because he ended up in the hospital but they have the second one coming up on the 14May2021 and she is scared for him to get it, he is adamant he is going to take it. He was taking her to the doctor for a problem she had and he told her he could not go to the bathroom and she was like what, because he has a stoma bag and she looked and it was plugged with a blood clot, he couldn't open it and drain it so she told him to get in the shower and take it off and put a new one on and she was pulling it off, it was full of blood and there was a big huge, like real liver sized clot came out of his stoma bag. She went to get the stoma bag after the thing came off she grabbed it and got it in the bag and took it out. He was taken to the hospital by the paramedics and he spent 3 days in the hospital. He is 83 now but this happened the week before his birthday, he was 82 when they noticed the blood in his stoma bad, he was a week away from 83. She provides his height and says he shrunk he is 5'9"" or 5'8"". She does not remember the date of the first dose. She says his birthday was 18Apr2021. She thinks he was admitted to the hospital on 12Apr2021 and he got the first dose the week before that. She thinks it was 12Apr2021 that he was admitted because it was just before birthday first dose week prior to that. They never made the connection. She thinks he ended up in the hospital on a Tuesday and she is trying to count from his birthday 18Apr2021, she thinks he got to come home on Thursday, 15Apr2021. She believes he went in on 12Apr2021 and was discharged on 15Apr2021, he was there 3 days. No further information provided. Her husband is not there with her. She does not have his vaccine card to provide NDC lot expiration or dose amount. he may not even have his vaccine card because he has dementia. No further information provided. He hardly takes any mediation He takes Coumadin he takes 1 per day except Fridays he takes 2, she thinks it's 4 mg and he takes 2 at night everyday except Friday he takes 3. she was guessing. She has one there called Atorvastatin that says it is for Lipitor, so that is not it. She sees Glipizide. No further information provided. She says she found the Coumadin, they are 2mg tablets and he takes 2 at night. She says the dose is not written on there because they call him every 3 weeks because he has to have a blood draw. He takes 2 tablets at night 6 days per week. She clarifies he takes 4 mg once daily in the evening 6 days per week and on Friday he takes 6 mg once in the evening. They just got report for that and it was 1.8 which is a little bit low but he lost blood and had to have a transfusion while in the hospital. It is in a white plastic bottle they got it from (Name). NDC 0093-1713-01 she does not see anything else. She thinks the pharmacy label is wrapped almost all the way around the bottle but she does not see lot number expiration. She sees they ordered it 11Feb2021 and she has a quantity that is it. Treatment received and outcome was unknown. Information on the lot/batch number has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340451-1	a blood clot came out on his stoma; This is a spontaneous report from a contactable consumer (patient's wife) from Pfizer-sponsored program . A male patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), dose 1 via an unspecified route of administration on 14Apr2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. Caller (Name) is calling on behalf of her Husband (Name). Her husband got the 1st dose on 14Apr2021 (caller not sure with the date) and after the vaccine he experienced a blood clot came out on his stoma in 2021. Caller asking if this is a reaction of the vaccine. Caller also asked if her husband can get the second dose even if he is a blood thinner. Caller also has a question for herself if she can take the 1st dose because she is allergic to Prednisone and Metformin and had a bad reaction with this medication. Therapeutic measures were taken as a result of event (blood thinner). The outcome of event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340462-1	nurse stated it could possibly be blood clots; Patient is afraid he is going to have a stroke or scared he had one after the COVID-19 Vaccines; blurred vision in right eye that comes and goes; ringing in his left ear; Very fatigued throughout the day, sleeping a lot/tired and sleeps a lot; Very fatigued throughout the day, sleeping a lot/tired and sleeps a lot; Date received First dose: Third Friday in Feb2021/Received second dose on 12Apr2021, six weeks after first dose; Date received First dose: Third Friday in Feb2021/Received second dose on 12Apr2021, six weeks after first dose; Fullness in face; Body aches; Brain fog: At times he can't do the simplest things. He will look at the clock on the wall and says he knows what it is, but can't find the words to say. Cannot carry on a conversation intelligently; Brain fog: At times he can't do the simplest things. He will look at the clock on the wall and says he knows what it is, but can't find the words to say. Cannot carry on a conversation intelligently; Brain fog: At times he can't do the simplest things. He will look at the clock on the wall and says he knows what it is, but can't find the words to say. Cannot carry on a conversation intelligently; headache all over the back of his head/Affecting his balance. In back of head, moves around, sometimes the front; headache all over the back of his head/Affecting his balance. In back of head, moves around, sometimes the front; feeling dizziness/had to stop driving; This is a spontaneous report from a contactable consumer (patient's wife). A 50-

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				spontaneous report from a contactable consumer (patient's wife). A 50- years-old male patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on 12Apr2021 (may have been in morning) (Batch/Lot Number: Unknown; Pfizer COVID-19 vaccine) as 2ND DOSE, SINGLE for covid-19 immunization, at the age at vaccination of 50 years old. The patient's medical history and concomitant medications were not reported. History and Investigations: Doesn't take any medications. Has no medical conditions, unless age was considered a medical condition. Historical vaccine included first dose of COVID-19 Vaccine reported as on Third Friday in Feb2021 (First dose: between 1430-1530), and experienced AEs: Brain fog; Headache: Thinks in the back of the head, but moved around too, sometimes in the front. Has affected his balance. Dizziness: Had to stop a couple times while driving home from vaccine facility. Aches and pains. All symptoms from the first dose went away within 3-4 days. Caller would say patient recovered completely from symptoms associated with first dose. The patient experienced nurse stated it could possibly be blood clots on an unspecified date with outcome of unknown, patient is afraid he is going to have a stroke or scared he had one after the covid-19 vaccines on an unspecified date with outcome of unknown, brain fog: at times he can't do the simplest things. he will look at the clock on the wall and says he knows what it is, but can't find the words to say. cannot carry on a conversation intelligently on 12Apr2021 with outcome of not recovered, headache all over the back of his head/affecting his balance. in back of head, moves around, sometimes the front on 12Apr2021 with outcome of not recovered, feeling dizziness/had to stop driving on 12Apr2021 with outcome of not recovered, blurred vision in right eye that comes and goes on an unspecified date with outcome of not recovered, ringing in his left ear on an unspecified date with outcome of not recovered, fullness in face on 12Apr2021 with outcome of not recovered, body aches on 12Apr2021 with outcome of not recovered, very fatigued throughout the day, sleeping a lot/tired and sleeps a lot on an unspecified date with outcome of not recovered, date received first dose: third friday in feb2021/received second dose on 12apr2021, six weeks after first dose on an unspecified date with outcome of unknown. Patient was caller's husband. Had both doses and had the same problem. His experience is very bad with the second dose and has had it for a month. Within a day after the first dose, had brain fog, headache, and dizziness where he had to stop driving a couple times. This went away within a few days. Patient and caller were concerned about the second dose. They spoke with patient's doctor and the local Health Department, both said it couldn't be from the vaccine. Patient was advised to get the second dose. Had the same symptoms with second dose, but occurred within an hour of received second dose. Symptoms have continued. Also experiencing blurred vision in right eye that comes and goes. Patient had an appointment with his doctor this week, but the doctor basically didn't look at him. Doctor advised patient to see a neurologist. Another nurse stated it could possibly be blood clots. Caller is trying to find help. Caller wanted to add with the second dose, patient experienced ringing in his left ear, fullness in his face, and headache all over the back of his head. Was feeling dizziness after second dose and had to stop driving. Patient is afraid he is going to have a stroke or scared he had one after the COVID-19 Vaccines. If the COVID-19 Vaccines woke up some other health issues, patient wants to get it taken care of. Received second dose on 12Apr2021, six weeks after first dose. Got second dose at a different location. Second dose COVID-19 Vaccine AEs: Started experiencing symptoms within an hour. Headache: Affecting his balance. In back of head, moves around, sometimes the front. Comes and goes. Brain fog: At times he can't do the simplest things. He will look at the clock on the wall and says he knows what it is, but can't find the words to say. Cannot carry on a conversation intelligently. Dizziness. Blurred vision in right eye: Doesn't know if this started on 12Apr2021 or a couple days after. States as the day goes on and he gets tired, it will worsen. It comes and goes. Ringing in left ear: Began on 12Apr2021 or a few days after. Fullness in face: Stated this comes and goes, but has stayed the same. Documented as provided. Body aches: Comes and goes. Seems to worsen as the day goes on. Very fatigued throughout the day, sleeping a lot: He's tired and sleeps a lot. It was hard for him to get out of bed. Caller stated patient's symptoms worse as the days goes on. Patient never experienced pain at the injection site. Guesses patient received COVID-19 vaccines in his left arm because the county was doing a dive-thru clinic. Since he didn't get out of his car, she believed patient would have received the vaccines in the left arm. The events REQUIRE A VISIT TO Physician office: Doctor doesn't think anything is wrong. ER: Not yet. Is considering going to the ER today. PRIOR VACCINCATIONS WTIHIN 4 WEEKS was None. RELEVANT TESTS: None. Events Clot blood and stroke considered medically significant. Follow up attempts are needed. Information of Lot/Batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340479-1	she feels pain in the lungs, in the body,in her head there is blood clot; she feels pain in the lungs, in the body,in her head there is blood clot; she feels pain in the lungs, in the body,in her head there is blood clot; every day she feels different; This is a spontaneous report from a contactable consumer (patient) from a Pfizer-sponsored program. This female patient of an unspecified age received received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in Apr2021 (Lot number was not reported) as single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. Previously the patient received the first dose of BNT162B2 on an unspecified date for COVID-19 immunisation. In Apr2021, the patient experienced that she feels pain in the lungs, in the body, in her head there is blood clot, every day she feels different with outcome of unknown. The clinical course was reported as follows: The patient had the 2nd dose of the vaccine 3 or 4 weeks ago, since then she had the symptoms, she feels something's changed, and she feels pain in the lungs, in the body, in her head there is blood clot, every day she feels different. In the hospital they do tests and X-ray but results turn out normal and nothing is wrong. She had never felt this strange in her life, she didn't feel well. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1341035-1	Fever 101.7, exhaustion, chills for about 34-48 hours after. On day 13, was hospitalized (4/28/2021) excessive rectal bleeding and fever of 101.1.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1341496-1	Leg pain initially, diagnosed with blood clots may 17. receiving treatment currently 15mg xarelto twice daily.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1342992-1	Period clots that resemble miscarriage every 20 minutes for 4+ days
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1346751-1	Tues, 2/9, entire upper arm became very red and very swollen, and some itching. Wed. 2/10, Major GI problems. Constipated, PAIN, for several hours, then Difficult multiple BMs with blood, blood clots, & mucus, Thurs, 2/11 BMs with blood clots, mucus, Thurs 2/12 BMs with fewer blood clots
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1347702-1	"On 05/05/2021 Pt received the Pfizer vaccine. The next day she got chills, headache, muscle aches. These symptoms lasted until 05/08/2021. ""I was feeling much better and started working in my garden, suddenly I felt on my right calf, and I noticed a ""big knot"". Daughter of the patient took her to the ER on 05/08/2020, where she was scheduled to a diagnostic test on 05/12/2021. On May 12, pt also received the phone call of her PCP, who diagnosed her over the phone: ""blood clot"" and prescribed her with Xarelto (Rivaroxaban), a blood thinner. A doppler test was performed to the patient that day and a blood clot was found. On May 21 Pt had an appt with her PCP, and PCP advised to report this to the CHD. Pt currently walks with a cane. her next appt is with an Hematologist to find out for how long she will be taking blood thinners."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1349560-1	Pinched middle finger, got a 1/4 diameter blood clot under skin.; This is a spontaneous report from a contactable consumer (patient). A 66-years-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in arm left on 20Apr2021 12:00 (Batch/Lot Number: er8735) as single dose for COVID-19 immunization. The patient medical history and concomitant medications were not reported. Patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. No other medications the patient received within 2 weeks of vaccination. Prior to vaccination, patient was not diagnosed with COVID-19. Since the vaccination, patient has been tested for COVID-19. The most recent COVID-19 vaccine was administered in facility. The patient experienced pinched middle finger, got a 1/4 diameter blood clot under skin on 09May2021 13:00. Patient was scheduled to go there tomorrow for the second dose. Had an accident yesterday and pinched the right middle finger. Usually when that happens, he gets a boil and it fills with a clear liquid pus. This time, it turned into a giant blood clot, well not giant but about a quarter inch in diameter in the tip of his finger. He has never had that happen before. He may have never damaged his finger in that way before. That is why it turned into a blood clot and not clear fluid. Patient was worried a little with the blood clot because he knew it has been an issue. When he first got a little worried about the blood clot, he thought he would take 2 aspirin and it might thin the blood a little bit. Patient ate some pineapple because he heard that was a blood thinner too. No treatment received for event. The patient underwent lab tests and procedures which included nasal swab: negative on 10Jun2020. Outcome of the event was unknown. Information on lot/batch number was available. Additional information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1349580-1	Patient has a blood clot; Pain sent patient to the emergency room; Patient was tired, and was having a hard time breathing. Patient was thinking it was something else; Patient was tired, and was having a hard time breathing. Patient was thinking it was something else; This is a spontaneous report from a contactable consumer. This consumer (patient's daughter) reported for a 79-year-old female patient (reporter's mother) that: Submitted by Call Centre Selected Report Type: Initial Is report related to a study or programme? No Patient Ethnicity: Unknown Is the patient also the reporter? No Reporter type: Consumer or other non-health professional Specify Consumer or other non-health professional: Reporter is patient's daughter Primary / Prescribing Healthcare Professional Info Adverse events: Dates for Patient has a blood clot: (From: Apr2021 To: Ongoing) Reporter seriousness for Patient has a blood clot: Hospitalization Dates when patient was in hospital for Patient has a blood clot: 29Apr2021 to 02May2021 Dates for Pain sent patient to the emergency room: (From: 26Apr2021 To: Ongoing) Reporter seriousness for Pain sent patient to the emergency room: Unspecified Dates for Patient was tired and was having a hard time breathing. Patient was thinking it was something else: (From: Apr2021 To: Ongoing) Reporter seriousness for Patient was tired and was having a hard time breathing. Patient was thinking it was something else: Unspecified Is Pfizer covid vaccine a Pfizer product? Yes Pfizer covid vaccine manufacturer: Unspecified Dates for Pfizer covid vaccine: (Start: 20Feb2021 Stop: Unspecified) NDC number of Pfizer covid vaccine: Unknown Expiry Date of Pfizer covid vaccine: Unknown Why was the patient taking Pfizer covid vaccine (Verbatim): Because of patient's age Other Products: No Patient History: No Investigation Assessment: No Additional Context: Reporter is calling on behalf of someone else. Reporter is calling about the Pfizer covid vaccine. Reporter is hoping for some guidance. It is reporters understanding, that if there is complications after the covid vaccine, there is compensation available. She is asking does she need to go somewhere for that. Reporter is calling about her mother. Reporter's mother is the patient. Reporter's address: Reporter was asked for her mailing address. Reporter stated that she will provide patient's mailing address, that it would be easier to provide patient's address. Primary / Prescribing Healthcare Professional Info: Declined. Patient has a blood clot: Patient is now out of the hospital, but she is on a blood thinner for the next six months. She was diagnosed with a blood clot and hospitalized on 29Apr2021. Patient began having adverse reactions two weeks prior, but patient did not realize they were related to her blood clot. Reporter does not know an exact start date. Patient is getting better with the blood thinner, but the blood clot is still there. Pain sent patient to the emergency room: Pain began on 26Apr2021. Patient is still in pain, but it is not as severe. Patient was tired and was having a hard time breathing. Patient was thinking it was something else: Began about two weeks prior to being diagnosed with a blood clot. The problem is, it is a large blood clots, and has done damage to patient's lungs. Reporter states because of the blood thinners that patient is having to take, patient is having complications. The blood thinners are very expensive. The blood thinners are interfering with other medications that patient is on. No further information provided. First dose administered on 20Feb2021 about 12:00PM in left arm. Lot number: EL9266 NDC: Unknown Expiry date: Unknown Second dose administered on 13Mar2021 at 1:30PM in left arm. Lot number: Unknown NDC: Unknown Expiry date: Unknown Patient Age at Time of Vaccination in Years: 79 Vaccination Facility Type: Other, Pop up clinic. Vaccine Administered at Military Facility? Not provided. Additional Vaccines Administered on Same Date of the Pfizer Suspect: None. Did any AE(s) require a visit to: Emergency Room? Yes. If the patient was hospitalized, how many days was the hospital stay? Four days. Physician Office? Yes. Patient has saw an oncologist, primary care physician, neurologist and pneumologist. Prior Vaccinations (within 4 weeks): None. AE(s) following prior vaccinations: Not provided. Patient's Medical History: None. Family Medical History Relevant to AE(s): None. Relevant Tests: None. Reporter is asking what all she needs to submit to Pfizer Legal. Reporter is asking if she needs a lawyer to handle this. PSCC Communication: Caller notified that all requests for compensation will need to be made in writing to Pfizer Legal. Caller was provided with Pfizer Legal Department's address of Pfizer, Inc. Legal Department Follow-up attempts are needed. Additional information is requested

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1349617-1	"Chills; sick; Fever; hard for me to walk; Atrial fibrillation; blood clot in her finger; thumb right now is real cold; the tip of my thumb is numb; tip was numb and real cold- there was no circulation going on; Irregular heartbeat; Blood pressure high; Headache; Muscle ache; This is a spontaneous report from a contactable consumer, the patient, via the Pfizer-sponsored program. A female patient of an unspecified age received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot: Unknown), via an unspecified route of administration on 19Apr2021 as a single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient previously received the first dose BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) on 19Mar2021 for COVID-19 immunization and felt fine (no adverse event). On an unspecified date in Apr2021, four days after the second vaccination, the patient was sick for seven days with chills. On an unspecified date in 2021, the patient had terrible side effects such as muscle ache and fever (""99 something"") for 7 days. On an unspecified date in 2021, the patient experienced headache, blood pressure high, irregular heartbeat, atrial fibrillation and the calf was ""real bad""- it was hard for the patient to walk. The patient called her doctor and got tested for COVID-19 two weeks after the vaccination, in May2021, with negative results. The doctor stated that the patient got so sick from the shot. On an unspecified date in 2021, the patient's thumb was like a blood clot to the thumb (blood clot in her finger), the tip was numb and real cold- there was no circulation going on. The patient went to the emergency room where an ultrasound was performed on the arm to see if there were any blood clots in there, with unknown results. The patient was treated with an unspecified ""pill"" for irregular heartbeat. The patient was scheduled for an echocardiogram and heart monitor. The clinical outcome of atrial fibrillation, blood clot in her finger, thumb right now is real cold, the tip of my thumb is numb, ""tip was numb and real cold- there was no circulation going on"", irregular heartbeat, blood pressure high, headache, muscle ache, chills, sick, fever, ""hard for me to walk"" was unknown. Information on the lot/batch number has been requested"
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1352767-1	Prolonged menstrual bleeding, unstoppable with blood clods.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1353491-1	feels like a blood clot; My right calf has felt strange ever since the second dose and it feels like a blood clot/toes also felt strange/My toes also felt strange for a few weeks but that subsided; a tightness in the leg that doesn't want to go away; This is a spontaneous report received from a contactable consumer. This consumer reported for himself that the 35-year-old male patient received second dose of bnt162b2 (BNT162B2, Pfizer COVID 19), via an unspecified route of administration, administered in Arm Left on 22Feb2021 13:00 (Batch/Lot Number: 6203) as 2nd dose, single for covid-19 immunisation. Medical history was none. The patient's concomitant medications were not reported. The patient previously took first dose of bnt162b2 (BNT162B2, Pfizer COVID 19, Lot number=9265) on 01Feb2021,01:00 PM,in Left arm for covid-19 immunisation. The patient experienced feels like a blood clot on 23Feb2021 with outcome of not recovered , my right calf has felt strange ever since the second dose and it feels like a blood clot/toes also felt strange/my toes also felt strange for a few weeks but that subsided on 23Feb2021 with outcome of not recovered , a tightness in the leg that doesn't want to go away on 23Feb2021 with outcome of not recovered. No treatment received for the events. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. No List of any other medications the patient received within 2 weeks of vaccination. The clinical course was reported as follows: My right calf has felt strange ever since the second dose and it feels like a blood clot. There is a tightness in the leg that doesn't want to go away. My toes also felt strange for a few weeks but that subsided. The adverse event resulted in Doctor or other healthcare professional office/clinic visit. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. No Known allergies. Follow up letter has been generated for further information.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1353501-1	Patient was diagnosed with blood clots confirmed by Ultrasound; This is a spontaneous report received from a contactable physician. A 48-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number EW0164), on 21Apr2021 as 2nd dose, single for COVID-19 immunization. The patient's medical history included colon cancer (patient had a history of colon cancer 2 years removed) from unspecified date. The patient's concomitant medications were not reported. The patient previously received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number not reported), on 30Mar2021 as 1st dose, single for COVID-19 immunization. It was reported that patient was diagnosed with blood clots confirmed by ultrasound during the emergency room visit on 03May2021. Patient was placed on apixaban (ELIQUIS) 5 mg, twice a day. Patient had history of colon cancer 2 years removed, and recent colonoscopy on 13Apr2021 was clear. The event was reported as non-serious. The outcome of the event was unknown.; Sender's Comments: The event of blood clot is assessed as possibly related to the suspect BNT162B2 based on temporal association, but medical history of colon cancer was confounder. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Committees, and Investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1353513-1	she did not feel good; Migraine after second dose; her heart was pounding; felt palpitations in her chest; potassium was low; dimer test that showed 0.30 clot; slight mitral regurgitation; breathing problems; shingles; This is a spontaneous report received from Pfizer sponsored program. A 74-year old contactable female consumer (patient) reported for herself that she received the second dose of BNT162B2 (Lot No. unknown) at single dose for Covid-19 immunization on 05Feb2021 at 17:30. Relevant history included Shingles when she was 42, Chicken pox. Relevant concomitant drug was unknown. The patient received the first dose of BNT162B2 (Lot No. EL3247) at 0.3 ML single dose for Covid-19 immunization on 15Jan2021, experienced Left arm Pain at injection site. Prior Vaccinations (within 4 weeks): none. The patient went to Urgent care and was diagnosed with Shingles on 15Apr2021, which was started with pain in her side, and on 30Apr2021, she was having breathing problems/ breathing was not well at 15:30-16:00. She said shingles were from the shot. The patient started shrinking before she got the vaccine. It was years before. There was no prescriber. She did not have the pain from the Shingles any longer. The pain was gone for a couple of days now. She had it for 2 1/2 weeks. She was cooking and the next day, she said she did not feel good, her heart was pounding and breathing was difficult. She went to ER and they did dimer test. They took 12 vials of blood. They told her she had .30 clot. She later stated the Dimer result was 0.30. They did chest x-ray and nothing showed. They looked at her legs too. That is why they did the sonogram. They wanted her to stay, but he gave her a sonogram of her heart. It is not enlarged and there is no fluid around it. She thought he gave referral for outpatient. She has been calling ER for an appointment. She then found that the doctor was not able to refer to an Echo cardiogram. She needs to go to the one in her home state and she is still in a different state visiting her daughter. She goes home next week and will get it then. She went for stress test Apr 2019 and emailed results said she had slight mitral regurgitation, so maybe that is where the blood is coming from. Her sister called and she told her she did not feel well. Breathing problem only lasted a day. She was cooking again and was not doing anything strenuous and felt palpitations in her chest and felt breathing issues again. It is better, but every now and then she gets palpitations. She had a migraine after second dose at 16:00 (unknown date). She went to bed at 4pm and woke up at 1am. She thought she took an Ibuprofen and did not know if it would affect vaccine or not. When she woke up, she was fine. She had shingles prior. She assumes she must have had COVID and did not know. he had the antibodies present in May 2020, when she was tested. When she googled, the people in another country were getting Shingles. She had the chicken pox when she was young. If you get the Chicken pox and the COVID, you are susceptible to shingles. She does not have any information on those people in other country to report. Her brother and sister both received the Moderna COVID vaccine. Her brother did not have any issues an her sister already reported her issues to them. No further details provided. Separate report completed for her sister. When looking up an address, she stated she needs a magnifying glass. She needed this before the vaccine. She always has her glasses to read with. ER visited on 15Apr2001 and 01May2021. On 15 They gave her ketorolac and ondansetron, potassium chloride. because her potassium was low. The outcome of event shingles, breathing problems was resolving, the outcome of other events was unknown. Follow-up attempts are needed. Information about batch/lot number is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1353535-1	Vaginal bleeding in a menopausal woman with blood clots.; Vaginal bleeding in a menopausal woman with blood clots.; Also slept for 13 hours the day following the shot and had a continual headache with nausea; Also slept for 13 hours the day following the shot and had a continual headache with nausea; Also slept for 13 hours the day following the shot and had a continual headache with nausea; This is a spontaneous report from a contactable consumer (patient). A 56-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 06May2021 at 09:45 at single dose in left arm for COVID-19 immunisation at the age of 56-year-old. Lot number EW0183. Medical history included menopausal woman. Concomitant medications were none. Historical vaccine included BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 15Apr2021 at 10:30, Lot number EW0164, dose number: 1, in left arm. On 06May2021, the patient also slept for 13 hours the day following the shot and had a continual headache with nausea. On 10May2021, the patient experienced vaginal bleeding in a menopausal woman with blood clots. It was very concerning; the patient was going to Ob/Gyn soon. The patient underwent ultrasound and biopsy. Prior to vaccination, the patient was not diagnosed with COVID-19; since the vaccination, the patient has not been tested for COVID-19. The outcome of the events was unknown. Follow up needed, further information has been requested

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				my doctor immediately said you know we want to rule out a blood clot so blood was drawn to the lab; Right ankle was swollen; There is some problem down with the leg not as bad I say but having the same pain I feel a sensation in my feet when I stand; This is a spontaneous report from a contactable consumer (patient). An 81-year-old patient of an unspecified gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection) (at 81 years of age), via an unspecified route of administration on 22Mar2021 (Batch/Lot Number: EN6206) as 2ND DOSE, SINGLE for COVID-19 immunisation; amlodipine besilate (manufacturer unknown), via an unspecified route of administration from an unspecified date (Batch/Lot number was not reported) to an unspecified date, at 10 mg, 1x/day for blood pressure management and valsartan via an unspecified route of administration from an unspecified date (Batch/Lot number was not reported) to an unspecified date, at 80 mg, 1x/day at bed time (anywhere between 9 and 10:15 PM) as blood pressure medication. Medical history included glaucoma and hypertension (Blood pressure) from an unknown date and unknown if ongoing; additional medical condition reported as felt at times like he/she want to itch ('felt itching but never scratch myself or anything like that'). Concomitant medication included bimatoprost (LUMIGAN) taken for glaucoma, start and stop date were not reported. Historical vaccine included BNT162B2 dose 1 on 01Mar2021 (at 81 years of age) for COVID-19 immunisation. The patient took the two treatments of the Pfizer vaccine. The patient had a problem from the second vaccination administered on 22Mar2021 with the LOT# EN6206. The patient took the last treatment on 22Mar2021 at the health department, which was the one that had created problems for him/her. The patient mentioned that there was no problem at all with the first treatment. The patient stated that first of all, he/she was on the blood pressure medications and one of the medications was valsartan 80 mg and the other one was amlodipine 10 mg, once a day; the patient take valsartan at bed time (anywhere between 9 and 10:15 [something like] that PM). The patient was taking amlodipine 10 mg for blood pressure, and then from reading what he/she got from the pharmacist on the valsartan, it was supposed to work by relaxing blood vessel so the blood can flow more easily lowering high blood pressure to help prevent stroke, heart attack and kidney problems; so the patient had been taking those two. It was reported that after the patient took the second Pfizer vaccine on 22Mar2021, in the evening he/she noticed that his/her right ankle was swollen not in the morning but in the evening say by anywhere from 6 to 7 PM to 8:00 PM, it was swollen occasionally and then it looked like there was some problem down with the leg not as bad he/she said but having the same pain he/she felt a sensation in the feet when he/she stands. So the patient saw his/her doctor, had an appointment with him on 20Apr2021 and at the time of the appointment which was at 2:45 in the afternoon, there was no swelling; the patient did tell him that he/she had this reaction and it started the same day he/she got the treatment but it was in the evening that she/he noticed it, not in the morning, not the afternoon but the evening, late evening. The patient did not take any treatment for these adverse events which the doctor immediately said that they want to rule out a blood clot, so blood was drawn to the lab. The patient saw the doctor on the 20th about a week later and called his office and was told report had not come in and the patient had not heard from his office since that time. The patient was making assumption right or wrong that had there been a blood clot, the office would have called him/her okay. But the doctor said that he wanted to rule that out then he said they might have need to lower the dosage of amlodipine, those were the things he remembered him saying so. But the patient's concern was not it was because the ankle does swell and then when he/she went to bed and lie down of course it stopped it did not swell, when he/she got up in the morning it was normal ankle so but the patient did not know what was going on and did not know if this was the reaction that no one else had; he/she was not sure what to do about it at this point. The patient wondered if the symptoms were direct result of his/her mixed medications that he/she had taken with the Pfizer vaccine and commented that he/she certainly had no problem with the first. The patient added concern whether there was a causal relation between the medication he/she was taking and that second vaccine, he/she does not know what. The patient's main concern was to get his/her ankle un-swollen which what happened since the last dose, no problem with the first dose but the last dose that was where it was. The patient did not know what the diagnosis that was causing the problem as he/she said the doctor had taken the had blood drawn to send to the lab to rule out the blood clot. The patient underwent lab tests and procedures which included blood test: unknown results on an unspecified date. The action taken in response to the events for amlodipine besilate and valsartan was unknown. The outcome of the events was unknown. Pfizer is a marketing authorization holder of Amlodipine Besilate in the country of incidence or the country where the product was purchased (if different). This may be a duplicate report if another marketing authorization holder of Amlodipine Besilate has submitted the same report to the regulatory authorities. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1353559-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1353752-1	He received both doses March 2021. second dose on 3/26. on 4/26 presented to ER with Epistaxis, came to about a month after Pfizer series pt went to Hospital ER 3 times that week for care of what was thought nose bleeds from picking, and later ENT for care that required travel. Pt stopped Plavix use and 6 days later on 5/1 returned to Hospital ER with serious symptoms, developed multiple thrombotic events that lead to clotting of leg/bowel/coronary. Transferred to Hospital on 5/1/2021. Multiple MI events and expired at Hospital on 5/3/2021

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1354283-1	Blood clot in right foot. Swelling. I didn't immediately consider the possible connection to the Pfizer vac. Sorry for the delay.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1354461-1	Had episode of ovarian cyst pain 1/20/21 in am. Went home, rested, pain eased a little. Able to finish work week but still bloated & cra.mpyNo nausea, no diarrhea. 1/23/21 felt like coming down with UTI. Rested, hydrated. Passed 2 small blood clots. Had a telehealth visit. Prescribed Levequin & Flagyl. Told to call PCP, needed a CT to rule out kidney stone. Saw PCP 1/27/21. Drew lab & ordered CT of pelvis. CT done 2/3/21. Consult with Oncology poss. Ovarian Ca. 3/15/21 had Robotic Total Hysterectomy- no cancer.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1355263-1	Menstrual cycle started again a week after my normal cycle ended. Abnormal menstrual cycle is still on going (started 18May2021) today is 27May2021, no sign of stopping any time soon. This abnormal menstrual cycle is extremely heavy where it is causing concern. There is a lot of blood clots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1355317-1	Sustained blood clots in left calf area approximately 6 days after receiving 1st dose.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1355766-1	patient claims she developed a blood clot approximately one week after receiving her first dose of the Pfizer Covid-19 vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1357202-1	BARD Power Port clogged with blood clot; cellulitis; Severe headache; abdominal pain; arm pain; chest pain; dose 1 administered in Leg Left; dose 1 and dose 2 on the same day; dose 1 and dose 2 on the same day; This is a spontaneous report receive from a contactable consumer (patient). A 38 years old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration as single dose, dose 1 administered in Leg Left on 19Mar2021 14:45 (Batch/Lot Number: EP7534), and dose 2 administered in Arm Right on 19Mar2021 (Batch/Lot Number: ER8729) for covid-19 immunisation. Medical history included myasthenia gravis, natural killer cell deficiency, gastroparesis, connective tissue disease, postural orthostatic tachycardia syndrome (POTS), known allergies: Gluten, dairy, xanthan gum, antibiotics, beta blockers, pain medication, latex. No other vaccine received in four weeks. It was unknown if the patient diagnosed with COVID-19 prior to vaccination. Since the vaccination, the patient had not been tested for COVID-19. Concomitant medication(s) included escitalopram oxalate (LEXAPRO); montelukast sodium (SINGULAIR); diphenhydramine hydrochloride (BENADRYL); sumatriptan succinate (IMITREX); promethazine (PHENERGAN), all taken for an unspecified indication, start and stop date were not reported. The patient experienced Severe headache 1.5 days, abdominal pain 3 days, arm pain and chest pain 7 days, BARD Power Port clogged with blood clot and subsequent cellulitis from many attempt to flush the clot (22Apr2021 04:00 PM). The adverse event result in Doctor or other healthcare professional office/clinic visit and Emergency room/department or urgent care. Treatment received for the adverse event: Home nursing visits and admitted to Emergency Room (ER). Outcome of the events was recovering/resolving.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1357215-1	Blood clot in leg and one in each lung. Admitted to hospital on 02May2021; Blood clot in leg and one in each lung. Admitted to hospital on 02May2021; This is a spontaneous report received from a contactable consumer. A 55-year-old female patient (not pregnant) received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), dose 2 via an unspecified route of administration on 29Apr2021 08:30 (at 55 years old, not pregnant) (Batch/Lot number was not reported) as 2ND DOSE, SINGLE for covid-19 immunisation. The patient's medical history was not reported. No concomitant medications. On 02May2021 08:30 patient experienced Blood clot in leg and one in each lung. Admitted to hospital on 02May2021.No other vaccine in four weeks. No other medications in two weeks. Treatment for event was Blood thinners. No covid prior vaccination. Not covid tested post vaccination. The outcome of the events was not recovered. Events resulted in visit to emergency room and physician office. Device Date: 12May2021. Information on the Lot/Batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357242-1	swollen and red leg; swollen and red leg; blood clot in right leg; Pain in right leg; her husband cannot walk, he is limping; her husband cannot walk, he is limping; This is a spontaneous report from contactable consumers (patient and wife). A 61-year-old male patient received bnt162b2 (BNT162B2 reported as PFIZER COVID-19 VACCINE), dose 2 at vaccination age of 61-year-old via an unspecified route of administration on 21Apr2021 13:00 (Lot Number: ER3732; Expiration Date: 30Nov2022) as 2nd dose, single for covid-19 immunisation. Medical history included ongoing high cholesterol and ongoing prostate [having to go to the bathroom frequently] (prostatic disorder). The patient received the first dose of bnt162b2 (BNT162B2 reported as PFIZER COVID-19 VACCINE) at vaccination age of 61-year-old on 26Mar2021 3:45 pm for covid-19 immunisation. Concomitant medications included (LIPITOR) ongoing since unspecified date (reported as 3 or 4 years ago) for blood cholesterol increased, and tamsulosin on an unspecified therapy dates (reported as he has been taking it for years now) for prostate/having to go to the bathroom frequently (prostatic disorder). The patient did not receive other vaccine in the last four weeks prior to vaccination. The patient received the vaccine from a pharmacy. The patient reported that one week after getting the second vaccine, he had pain in his right leg (28Apr2021). He stated that he did know what was going on, but he continued to work. He added that last Monday (10May2021) it got worse, so he contacted his primary care doctor and he went today (12May2021) because his leg was swollen, red and he had pain. His doctor sent him to the ER (emergency room). The patient mentioned that the ER did a CAT scan on 12May2021 and found that he had a blood clot in his right leg (reported as 10May2021). The patient's wife comes on the line and stated the reason her husband called to report was because it could be from the vaccine, but they are not sure. The wife added that her husband went to the pharmacy to pick up the blood thinner and they told him to report this. She also stated that all the side effects could happen, but he did know the side effects and he just wanted to report because the pharmacy told him to report. She mentioned that her husband started complaining his leg hurt because he has to do yard work. She states that it got worse and worse and she saw his leg get swollen and she was like you need to go to the doctor. The wife mentioned that the leg is more swollen and painful, but is the same, her husband cannot walk, he is limping on an unspecified date in 2021. She stated that her husband has just started taking medication, yesterday and today. The outcome of the event thrombosis leg was not recovered while the outcome of the other events was unknown.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357246-1	large blood clots during menstruation; irregular menstrual cycle; This is a spontaneous report from a contactable consumer (patient). A 34-year-old female patient received BNT162B2 (Lot Number: er8731), via an unspecified route of administration in Apr2021 as single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. The patient experienced irregular menstrual cycle and large blood clots during menstruation, both on 04May2021. The events resulted in Doctor or other healthcare professional office/clinic visit. The outcome of the events was recovering.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1357258-1	"embolic stroke; blood clot; His major issue was speech and wasn't able to formulate any words; left sided weakness on his leg; This is a spontaneous report from a contactable nurse (patient's daughter). A 79-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 30Mar2021 11:00 (Batch/Lot number was not reported) as 2nd dose, single dose (at the age of 79-years-old) for COVID-19 immunisation. Medical history included transient ischaemic attack (TIA) from 2019 to an unknown date, mild hypertension, idiopathic neuropathy: in the left leg, started in the beginning of the year from 2019, melanoma removed in the 1970s, enlarged prostate, neuropathy, cholesterol and fluid retention. Ongoing concomitant medications included tamsulosin taken for enlarged prostate; gabapentin taken for neuropathy; candesartan taken for hypertension; pravastatin taken for cholesterol; acetylsalicylic acid (BABY ASPIRIN); and furosemide (LASIX [FUROSEMIDE]) taken for fluid retention. The patient has been taking all of his meds for quite some time, over 5 years. The patient did not have prior vaccinations within 4 weeks. The patient received the first dose of BNT162B2 on 09Mar2021 10:30 for COVID-19 immunization. The patient completed the Pfizer COVID vaccine series, the second shot was on 30Mar2021. On 09Apr2021, the patient had an embolic stroke. The patient's daughter stated that the patient has a history of TIA in 2019 and was concerned. She does not know if it was related. She added it might be that if people have previous risk, there may be some precautions, like to take a full dose of aspirin before, a week before. She sent an e-mail last 05May2021 and hasn't gotten a response back. The patient went to the ER due to embolic stroke and was in the hospital Friday (09Apr2021) through Wednesday. The patient went to the stroke rehab center for 3 weeks. The daughter clarified that the patient was in the hospital until 14 or 15Apr2021 before being transferred (also reported as 5 or 6 days and in rehab for three weeks). Then was discharged last 06May2021. The patient's condition seemed to be getting better. His major issue was speech and wasn't able to formulate any words. He was able to speak now. His rehab would be a lot of neurological rehab and speech therapy. The patient previously had left sided weakness on his leg from some random thing that happened. It didn't seem to get worse. It really affected his speech and cognition. The patient's daughter thought that the embolic stroke was related to the Pfizer Covid vaccine and added that thought was in some way but cannot say in what way. She thought that it exacerbated some type of blood clotting. She stated that he never had blood clotting issues before and was the type of person that would take a baby aspirin and get a nose bleed, he was the opposite of a clotter. An MRI in the hospital showed blood clot and stroke. He had blood work but does not know what it showed and presumed he was fine (unknown results). He has never really had any kind of blood work issues and does not know why he was on cholesterol medication. The patient required emergency room visit for embolic stroke. The outcome of the events was unknown. Information on the lot/batch number has been requested.; Sender's Comments: Considering a plausible temporal relationship and known product safety profile, a possible contributory role of suspect product BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) to the reported ""embolic stroke"" cannot be excluded. Previous medical history and Old age is a risk factor for the onset of stroke. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1357265-1	"I have a blood clot in hurry I have blood clots regularly; Blood is way too thick so he give triple dose; This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received bnt162b2 (BNT162B2), via an unspecified route of administration on 01May2021 (batch/lot number and expiration date not reported) as UNKNOWN, SINGLE for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient stated, ""I received COVID 19 vaccination on 01May2021 at my Doctor's Office and yesterday (12May2021), I went to see my Oncologist. I am on blood thinners, its Coumadin. I get Warfarin which is you know (incomplete sentence, thus, not clarified and was not captured). Anyway, I take Coumadin, my blood is way too thick, so he gave triple dose and I went back today, and it is still bad, and I am taking another triple dose tonight and going back. What are the safety issues in getting Coumadin and getting that vaccination?"" When probed for follow up, consumer stated, ""Yes, it be great if they did. I need to find something we need to correct because I have a blood clot in hurry. I have blood clots regularly if I do not keep my blood thin."" Patient stated, ""Have you heard other people reporting this problem or was it the concern in the beginning or what can be done in other words?"" Outcome of the events was unknown. Information on the lot/batch number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357269-1	I am feeling a clot in my left armpit after 1 day of having 2nd dose of COVID19 Pfizer; This is a spontaneous report from a contactable consumer (patient). A 35-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration and administered in left arm at age of 35-years, on 11May2021 18:00 (Lot Number: EW0179) as single dose for covid-19 immunization. The patient medical history was not reported. The patient was not pregnant at time of vaccination. Concomitant medications were none (the patient did not receive any other medications within 2 weeks of vaccination). The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Historical vaccine included the first dose of BNT162B2 on an unknown date for covid-19 immunization. The patient reported that she was feeling a clot in her left armpit at 08:00 on 13May2021 (reported as after 1 day of having 2nd dose of COVID19 Pfizer). The patient was not diagnosed with COVID-19 prior to vaccination and had not been tested for COVID-19 since the vaccination. Event outcome was not recovered. No follow-up attempts needed. No further information expected.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357288-1	had a superficial blood clot in my left calf between the first and second vaccines; This is a spontaneous report from a contactable consumer (Patient). A 51-years-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 01Apr2021 16:00 (Batch/Lot Number: ER8733) at the age of 51-years-old as SINGLE DOSE for covid-19 immunization. The patient medical history was not reported. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient's concomitant medication was none, patient did not received other medications within 2 weeks of vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient had a superficial blood clot in his left calf between the first and second vaccines on 16Apr2021. Event resulted in Doctor or other healthcare professional office/clinic visit. Treatment received for the event included had a shot of enoxaparin sodium (LOVENOX) and am taking rivaroxaban (XARELTO). The patient underwent lab tests and procedures which included blood test: negative on 06May2021, Nasal swab covid-19 test: negative on 27Apr2021. The outcome of the event was not recovered.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357347-1	Pulmonary embolism; Two massive bilateral clots; This is a spontaneous report received from a contactable physician (patient). A 75 years old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration at the age of 74 years old, administered in arm left on 23Feb2021 (Lot Number: EL9204) as single dose for covid-19 immunization. Medical history included, chronic obstructive pulmonary disease (COPD), history of cancers. No known allergies. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient had received other medications within 2 weeks of vaccination. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient previously took the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration at the age of 74 years old, administered in arm left on 02Feb2021 (Lot Number: EN6200) as single dose for covid-19 immunisation. The patient experienced pulmonary embolism on 19Apr2021, two massive bilateral clots on 19Apr2021. Events resulted in Emergency room/department or urgent care, Hospitalization, Life threatening illness (immediate risk of death from the event), Disability or permanent damage. The patient was hospitalized for all events for 6 days. The patient underwent lab tests and procedures which included Nasal swab: negative on 03Feb2021. Therapeutic measures were taken as a result of pulmonary embolism, two massive bilateral clots and included Heparin and Xarelta. The outcome of the events was recovered with Seroquel in 2021.; Sender's Comments: Based on the temporal relation, the association between pulmonary embolism, thrombosis and vaccination with BNT162B2 cannot be completely ruled out. The contribution of patient's age cannot be completely ruled out in the occurrence of the events. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1357350-1	Blood clots in lung and leg; Blood clots in lung and leg; Dizziness; Weakness; Headache; shortness of breath; This is a spontaneous report from a contactable consumer (patient). A 36-year-old non-pregnant female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in the left arm on 29Apr2021 16:00 (Batch/Lot Number: EW0162) as 2nd dose, single dose for COVID-19 immunization. The patient received the first dose of COVID-19 vaccine on 08Apr2021 17:45 left arm for COVID-19 immunization. Medical history included fibromyalgia and had covid-19 prior to vaccination. Concomitant medications included gabapentin; ethinylestradiol, norgestimate (ORTHO-CYCLEN); and diclofenac sodium (DICLOFENAC SODIUM); all taken for an unspecified indication, start and stop date were not reported. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. On 30Apr2021 05:00, the patient had blood clots in lung and leg, dizziness, weakness, headache (still after a month) and shortness of breath. As a result of the events, the patient visited doctor or other healthcare professional and has been to the emergency room. The events required hospitalization and was considered a life-threatening illness (immediate risk of death from the event). The patient was hospitalized for two days. The patient was put on blood thinners as treatment for the events. The patient

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1360548-1	"3 clots were discovered on the groin area of the left leg; patient has two stunts in the right leg (1 in thigh and 1 in calf) and reported that it had been uncomfortable but not painful; This is a spontaneous report from a contactable pharmacist. A 66-year-old male patient received the second dose of BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, Lot Number: ER8732), via an unspecified route of administration, administered in the right deltoid (reported as right arm) on 26Mar2021 at 03:55 as a single dose for COVID-19 immunization. Medical history included two stunts in the right leg (1 in thigh and 1 in calf) on an unknown date. The patient had no allergies. The patient's concomitant medications were not reported. The patient previously received the first dose of BNT162B2 on 24Feb2021 (lot number:EN6201), intramuscular in the right arm at 65 years of age. The patient went to his vascular doctor on an unspecified date (reported as 05Oct2021) and 3 clots were discovered on the groin area of the left leg. The MD said it did not warrant going to the ER. The patient has two stunts in the right leg (1 in thigh and 1 in calf) and reported that it had been uncomfortable but not painful. The MD placed him on Eliquis 10 mg, BID for 7 days then reduced the dose at 5 mg BID thereafter for about 5 months. The outcome of the events was unknown.; Sender's Comments: ""The causal association of the events of æclots on the groin area of the left leg' and æleg uncomfortable' with the suspect drug BNT162B2 cannot be excluded due to limited information in the case. The case will be reassessed once new information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate."""
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1360642-1	I didn't know if this could be a blood clot forming as result of the vaccination or this just regard could still be a blood clot.; I have too swelling in my lower left leg just below the knee; This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot Number: EW0162; NDC / UPC number of COVID Vaccine: Unknown; Expiration Date: 31Jul2021), via an unspecified route of administration on 20Apr2021 as 2nd dose, single for covid-19 immunisation. Medical history included blood clot and stated as I do have blood clot in my past case history. My mother died at blood clot to the heart. Concomitant medications were not reported. Historical vaccine included first dose of BNT162B2 (Lot Number: ER8734; Expiration date: 31Jul2021) for COVID-19 immunisation. On 24May2021, the patient experienced i didn't know if this could be a blood clot forming as result of the vaccination or this just regard could still be a blood clot and i have too swelling in my lower left leg just below the knee. Additional Context was received as, I was calling in regard to my vaccination did I had. The first shot I had on 30Mar2021 and then the second one was on 20Apr2021. Further patient stated, that's what I am really not sure about I am not sure of this, if this is a COVID related vaccine issue or not. I have swelling too in my lower left leg just below the knee. I noticed this first time yesterday and notice this again today few minutes ago looking better and I didn't know if this could be a blood clot forming as result of the vaccination or this just regard could still be a blood clot and I wasn't sure if there was any relation to the vaccine. Patient stated, I am going to try to see my doctor tomorrow, helps to get into see him tomorrow. Like I said I am trying to see my doctor tomorrow. I am trying to see him hopefully he will able to help me. I have concerned about the swelling because I do have blood clot in my past case history. My mother died at blood clot to the heart I had in never and stroke and that is why I concerned. So that's why I go to the doctor tomorrow. Outcome of the events was recovering. Further probing could not be done as consumer was not willing to complete the report. Hence limited information available over the call.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1361781-1	Major pain and headache in left arm resulted in major stroke within 24 hours
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1362357-1	Within 1 week of receiving COVID vaccine,I started having left leg pain that lasted until hospitalization. I?ve had sciatica before and thought it was this. After second COVID vaccine(03/02/2021),started getting shortness of breath that continued to get worse. Finally, in mid-April, along with the shortness of breath and left leg pain, I started getting right,mid back pain that radiated to right chest and shoulder. On April 21,2021, I could no longer tolerate the chest and back pain and drove myself to ER and Urgent Care. The Dr. ordered a CT Angio of chest and determined I had bilateral pulmonary embolism and sent me by ambulance to Hospital.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1362428-1	On 5/28/2021 I started to develop shortness of breath. It continued to worsen and I went to the hospital on 5/30/2021 and it was discovered that I had a pulmonary embolism and after additional ultrasounds 3 DVT blood clots in my right leg.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"two blood clots inside his brain, and one on the outside/ three blood clots in and near his brain; two blood clots inside his brain, and one on the outside/ three blood clots in and near his brain; pulled neck muscle; neck pain; fevers/ fever; severe headaches/incessant headaches; swollen lymph nodes; extreme migraines/ migraine headaches; could not move his neck without the assistance of his hands/ unable to freely move his neck; eyes are still swollen; Neck swelling; The initial case was missing the following minimum criteria: no identifiable reporter. Upon receipt of follow-up information on 24May2021, this case now contains all required information to be considered valid. This is a spontaneous report from a Pfizer-sponsored program. A contactable consumer, physician and non-contactable consumers reported that a 17-year-old male patient (also reported as 18 years old) received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 21Apr2021 (Batch/Lot number was not reported) as 1st dose, single dose for COVID-19 immunisation. Medical history included prior COVID-19 infection. Before the shot, the patient was 100 percent healthy, playing and practicing basketball. There was nothing wrong with him. He didn't have a sore throat (or) any injury. The patient's concomitant medications were not reported. On an unspecified date, the patient was hospitalized with three blood clots in and near his brain that developed after he received the first dose of Pfizer-BioNTech COVID vaccine. The symptoms started after getting his first COVID-19 vaccination dose, and symptoms were initially dismissed as a pulled neck muscle. He received the vaccine on 21Apr2021 and began experiencing neck pain, fever and severe headaches on an unspecified date (also reported as ""one day later""). The patient's pediatrician initially dismissed the symptoms as a pulled neck muscle. However, the patient's mother was convinced it was something else. After more than a week of symptoms and being unable to freely move his neck, the family got this diagnosis: two blood clots inside his brain, and one on the outside. The patient was healthy and well before. It was unclear how long the patient will be in the hospital. It was also stated that the patient experienced swollen lymph nodes and migraine headaches. Further reported that the day after his COVID-19 vaccine shot (on 22Apr2021), the patient felt his neck swelling. In the coming days, he suffered from severe headaches. He also could not move his neck without the assistance of his hands few days after the shot (unknown date). Patient suffered from fevers and incessant headaches. He started to feel swelling and pain in his neck (""that night""). Days later, he got extreme migraines, so the patient's mother took him to the emergency room. It was reported that the patient plays competitive basketball seven days a week, which means high contact physical activity that could have possibly aggravated the swelling caused by the vaccine. On Thursday (unknown date), the patient left the ICU, but his eyes are still swollen and the road ahead is uncertain. On an unspecified date, the patient was out of intensive care unit (ICU) but the clots aren't gone. He has an magnetic resonance imaging (MRI, unknown date, unknown results) to see if they shrunk. The patient is currently at home recovering (as reported). Doctor recommended that patient have a COVID Nucleocapsid antibody level drawn to determine if he had been exposed previously to the Corona virus. On an unspecified date, the patient underwent test for nucleocapsid antibodies, patient is POSITIVE for SARS-CoV-2 nucleocapsid antibodies. This indicates that he had been previously infected with Corona virus prior to his receiving the BNT162 vaccine. There is a very serious concern that previously infected persons may be at higher risk of developing post-vaccine complications that un-immune persons. The doctor believed that the prior infection in combination with the vaccine led to the blood clots in the brain. Outcome of event ""two blood clots inside his brain, and one on the outside/ three blood clots in and near his brain"" was not recovered, while other events were unknown. Information on the lot/batch number had been requested."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1364281-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1365501-1	on Tuesday 05/18/2021 i take my 2th dose of Vaccine and on 05/19/2021 at around 2100 pm i start to develop high fever at 103-104 with abdominal cramps, same way i was on the 05/20/2021 and on Friday morning 05/21/2021 i start bleeding with large clots and wasn't able to get from the bed, and around noon my waters broke following with a lot of blood and clots,i had to deliver my baby girl at home at 15 weeks, 911 was called, because the placenta didn't come out, so i was rushed to the ER Baby was Delivered at 13:30pm on 05/21/2021
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1365955-1	Significant vaginal bleeding and vaginal blood clots. Extremely heavy bleeding during mid-cycle (between periods). No prior history of heavy bleeding nor any vaginal blood clots. Started approx 24 hours after first shot and continued for several days, with bleeding and clots. Scheduled virtual doctors visit immediately and followed up with in-person pelvic exam and ultrasound. Normal ultrasound results. Stopped birth control pills for several days, then restarted, then soon after changed to new birth control.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1367162-1	Bruises on my thighs, looks like blood clots; Bruises on my thighs, looks like blood clots; This is a spontaneous report from a contactable consumer (patient). A 42-years-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration, administered in Left Arm on 17Apr2021 11:00 at the age of 42-years-old as single dose for covid-19 immunisation. Medical history included anaemia, iron deficiency. No known allergies. Patient was not pregnant, not pregnant at time of vaccination. Prior to vaccination, the patient was not diagnosed with COVID-19. Concomitant medications included iron, other vitamins such as biotin and minerals nos, vitamins nos (PRENATAL VITAMINS) within 2 weeks of vaccination. The patient previously received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date as single dose for covid-19 immunisation. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced bruises on thighs, looked like blood clots on 17Apr2021. No treatment received for the events. Since the vaccination, the patient had not been tested for COVID-19. Outcome of the events was not recovered. The events assessed as non-serious. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1367204-1	felt like a little fuzzy; Lot of discomfort; warm spots on her skin; Headache; Blood clots; itchy red rash; This is a spontaneous report from a contactable consumer (parent of the patient). A 13-year-old female patient received the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Lot Number: EW0185), via an unspecified route of administration on 14May2021 as 1st dose, single for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. On an unspecified date the patient experienced blood clots, really warm spots on her skin in one of her legs, front of her thigh and her knees, her arms the tip part between her elbow and wrist on both of them, like where you can feel it's warm to the touch, lot of discomfort, headache and she kind of feel like a little fuzzy like in her thinking, she said she felt like she was forgetting things like what you guys were saying. On 16May2021, she developed an itchy red rash on her legs and over her knees and on the tops of her arms from her wrist to her elbows. She has taken Benadryl to treat but it did not help. She is calling to see if this has been reported before. The outcome of events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1367764-1	He got a superficial blood clot in left arm ?He's got the shot May 21 and I noticed he had two hard knots on left arm and there was probably about a half inch between them and as hard as a rock on that following Monday may 24,2021 , he said it was sore some more when you touched it we was told to keep an ion it make sure it didn't get any worse, turn red, swell, feel hot to the touch, and then to put a warm compresses on it ??
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1368535-1	Pfizer-BioNTech COVID-19 Vaccine EUA States exactly 2 weeks after received 2nd dose of Pfizer vaccine, lost feeling in his right leg. Went to the hospital. Continued to have neurological symptoms. States was diagnosed with Bell's Palsy and blood clot in right leg. States cannot ambulate without walker now. States doctors are unsure of cause of neurological issues.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1368820-1	I started a very heavy period that lasted 25 days. I usually have a light 2 day period like clockwork. No birth control.. very normal my whole life until my shot. I had heavy bleeding along with clots of blood.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1370626-1	"stroke; blood clots on her legs, lungs and head; blood clots on her legs, lungs and head; blood clots on her legs, lungs and head; This is a spontaneous report from a contactable consumer (patient son). A 77-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on 31Mar2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunization. Medical history was none. Historical vaccine included BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) as 1st dose on 10Mar2021 for Covid-19 immunization. After having the first dose she told him that she could not make a whole 20 minute walk as she used to, she could for 10 minutes as she felt tired. She presented a consistent cough, that was not ""bad"". The patient's concomitant medications were not reported. Caller stated that he would like to consult about side effects of the Pfizer Covid-19 vaccine that may have caused his mother to be passing away right now. Stated that they both received the first dose on 10Mar2021 and the second dose on 31Mar2021. A month and a week later she went to her sister's house and this Tuesday he received a phone call where they stated that she had a stroke due to blood clots on her legs, lungs and head. They just left the hospital and the nurse stated that she could have hours or a couple of days left. He would like to consult if there are any reports of these side effects. The outcome of events was unknown. Information on the lot/batch number has been requested.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021568883 same patient, different dose/events"

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1370660-1	Blood clot disorder; This is a spontaneous report from Pfizer sponsored program. A contactable female consumer (patient) reported for herself that she received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 on an unspecified date and dose 2 on 19May2021, both via an unspecified route of administration (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. Caller went on her 2nd dose today 19May2021, she was asked a question if she had a blood clot disorder, she was not asked about this on the 1st dose, and apparently, she was having a blood clot disorder on an unspecified date. She asked the people in the facility if she needed to be concerned, and they said that she didn't need to worry about it, however she wanted to make sure that everything was fine or should she watch out for something. Outcome of event was unknown. No follow-up attempts are possible. Information about lot/batch number cannot be obtained.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1370663-1	The first Pfizer shot landed me in the hospital with blood clots and pneumonia; The first Pfizer shot landed me in the hospital with blood clots and pneumonia; This is a spontaneous report from a contactable consumer (patient). A 55-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as 1ST DOSE, SINGLE for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The first Pfizer shot landed patient in the hospital with blood clots and pneumonia. The outcome of events was unknown. Information about lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1371264-1	-tightness in inner thigh- day or so after first dose -tightness in calf worsened over the course of a couple weeks -swelling and soreness in calf gradually increased until doctor visit -doctor visit and ultrasound found multiple blood clots in mid thigh and calf -doctor prescribed blood thinner -continued swelling and tightness in thigh and calf
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1371386-1	30yof, 36w pregnant, came to clinic on 6/1 with 2 week history of lower leg tenderness and edema. Doppler confirmed on 6/1 that patient had an acute proximal DVT in the left lower extremity, with occlusive thrombus within the distal femoral vein. She denies personal history of clot; this is her 2nd pregnancy; she notes no family history of clot; this pregnancy is complicated by excessive weight gain (current BMI 32.28 kg/m2). Pt received Pfizer COVID vaccine on 4/24/21.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1371513-1	After vaccination I still had output out of my Ilestomy bag (diarrhea). By Friday evening I began omitting with extreme nausea, incredible chills mixed with incredible sweats (where I saturated the bed and wanted to die). I was too weak to get out of bed and go to the ER in the City. So I emailed my GI clinic to tell her what happened on Mon March 5th and was told to go to the ER for evaluation. I went to the ER on the 6th. I asked her if she could schedule a CTE to see if the Chron's was flaring / if I had a blockage. I continued to drink clear broths. Then I had sips of a smoothie and it triggered the nausea vomiting's, extreme pain of #25 but without chill/night sweats. I went to the ER on the 6th, they did a CT which showed I had an acute Chron's flair going up a foot of my Ilestomy bag and inflamed causing the bowel obstruction and was discharged to follow-up with my GI. my leg continued to hurt and when they did the contrast test in the ER my leg turned dark and all veins were really dark. On the 9th I saw Dr and told him about the leg pain and what was going on and he told me to follow up with my GI. I tried to get a port in my chest because of the continued diarrhea. On the 13th in the morning I was called to go to the Hospital and went the same day, the nurse was preparing me for all the tests to be done the 14th. I asked the Nurse to look at my leg and she did and then left and got other clinicians and they confirmed I had multiple blood clots. They treated me with high dose Heparin for the blood clots. I couldn?t because I was in so much pain. I was in the hospital for 8 days and left the 20th of April. I had falls in the hospital. I continue to have difficulty eating and losing a lot of weight. And now getting the Entivio infusions every 4 weeks.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1371903-1	Irregular menstruation and menorrhagia following COVID-19 vaccine on 4/28/2021. Had irregular breakthrough bleeding on 5/7 that lasted 3 days and then again on 5/20 - 5/27 that lasted greater than 7 days with very heavy clots which is irregular for patient. Patient convinced that it is due to COVID-19 vaccine. She reports compliance with daily OCP.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1371978-1	The patient made two visits to primary Dr. with swollen feet and ankles. Visits were a few weeks apart. Dr had Lab done the second visit and didn't see anything The next month Right foot was continuing to be swollen and had pain on bottom of foot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1372065-1	Blurry vision and(flashing light in peripheral vision)right eye the next day after receiving the second dose on April 27,2021. Flashing light lasted for a week. Afterwards just blurry vision until May 30th vision became distorted and dark. Visited eye specialist 06/04/2021 and was advised to report to the CDC that blood clot was found in right eye. Treatment is now needed that involves injections to the right eye . (3 treatments in span of 10 weeks) with possible laser treatment if injections are not as effective. Eye Specialist -

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1373734-1	States her cholesterol is always 75 and is now 200; facial droop or pain in her face; noticed her arms started getting numb and swelling up; noticed her arms started getting numb and swelling up; facial droop or pain in her face; talk with the slur; tingling in her fingers; ended up in the emergency room and had a stroke; she had blood clots on the brain from the stroke; she went into the ER her blood pressure was 200/180; This is a spontaneous report from a contactable consumer (patient) and a contactable physician. A 51-year-old female patient received bnt162b2 (Lot Number: ER8730; Expiration Date: 31Jul2021), second dose via an unspecified route of administration, administered in left arm on 11Apr2021 (early in the morning) at the age of 51-year-old as 2nd dose, single for covid-19 immunization. Medical history included ventricular septal defect from an unknown date and unknown if ongoing (she was born with a hole in her heart but that had closed up and when she had the stroke, the hole opened back up and they told her she had blood clots on the brain from the stroke). There were no concomitant medications. The patient previous received first dose of bnt162b2 (Batch/Lot Number: ER2613; Expiration Date: 31Jul2021), via an unspecified route of administration, administered in left arm on 21Mar2021 single dose for covid-19 immunization. The patient experienced stroke on 20Apr2021, she had blood clots on the brain from the stroke on 20Apr2021, blood pressure was 200/180 on 20Apr2021, her cholesterol is always 75 and is now 200 on an unspecified date, facial droop on an unspecified date, her arms started getting numb and swelling up on an unspecified date, pain in her face on an unspecified date, talk with the slur/ slurred speech on an unspecified date. The events her arms started getting numb and swelling up, pain in her face and talk with the slur/ slurred speech were serious per hospitalization; stroke, she had blood clots on the brain from the stroke, blood pressure was 200/180, her cholesterol is always 75 and is now 200 and facial droop were serious per hospitalization, medically significant. The patient was hospitalized for from 20Apr2021 to an unknown date. The events require a visit to emergency room and physician office. Caller states she is calling about the Pfizer shot and had the 2nd dose on 11Apr2021 and noticed her arms started getting numb and swelling up and she did not do anything about it but 2 days after that she called the facility where she had the Pfizer shot and they said to call her HCP and the HCP was out of town so she spoke with a nurse and was told to go to urgent care at that time. States she went to urgent care and they said her arm was swollen and to rub alcohol on it and she did that and a couple of days after that ended up in the emergency room and had a stroke. States the numbness and swelling was just the whole left side of her body and she went to the emergency room at the hospital on 20Apr2021 but on 16Apr2021 she went to the urgent care; states the numbness and swelling is better but comes and goes with the tingling in her fingers and the numbness, but her face is doing much better and she does not have the facial droop or pain in her face anymore; states her speech is back and she does not talk with the slur anymore. States the facial droop, face pain and slurred speech began when she went to the ER and that was on 20Apr2021 and is now completely resolved. Caller states the 1st dose of her Pfizer Vaccine was administered on 21Mar2021 with lot ER2613, the 2nd dose was administered on 11Apr2021 with lot ER8730; states both doses of the Pfizer Vaccine were administered in her left arm and her patient card does not have the NDC written on it but the expiry date for both doses of the vaccine is 31Jul2021. States the whole year of 2020 she was doing fine till she took the Pfizer Vaccine and that shot everything up regarding her cholesterol levels and Blood pressure which are all high; states her cholesterol is always 75 and is now 200; states she has high blood pressure and when she went into the ER her blood pressure was 200/180; states she was born with a hole in her heart but that had closed up and when she had the stroke, the hole opened back up and they told her she had blood clots on the brain from the stroke. The patient underwent lab tests and procedures which included cholesterol: 200 on 10May2021 (her cholesterol is always 75 and is now 200), blood pressure: 200/180 on 20Apr2021. The outcome of events for stroke, she had blood clots on the brain from the stroke, her blood pressure was 200/180, her cholesterol is always 75 and is now 200, tingling in her fingers were unknown, for other events was resolved on an unspecified date. Vaccination Facility Type: school. No Additional Vaccines Administered on Same Date of the Pfizer Suspect. No Prior Vaccinations (within 4 weeks).

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1373779-1	a blook clot; blockage in the Left Anterior Descending artery; symptoms of shortness of breath; This is a spontaneous report from a non-contactable consumer or other non hcp. A 58-year-old male patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on 25Feb2021 17:30 (at the age of 57-years-old) as 2nd dose, single for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient previously took first dose of bnt162b2 on 02Feb2021and experienced cardiac blockage. The patient did not take any other vaccines within 4 weeks prior to the COVID vaccine. On 15Apr2021 the patient experienced a blook clot, blockage in the left anterior descending artery and symptoms of shortness of breath. The events were reported as serious (life threatening). It was reported that 15Apr2021, emergency trip to ER, blockage in the Left Anterior Descending artery. This blockage was only 40% blocked in Feb2021, and suddenly in April became 99% blocked. Dr suspected a blook clot when symptoms of shortness of breath appeared suddenly. The events resulted in Doctor or other healthcare professional office/clinic visit and Emergency room/department or urgent care. Blockage in the Left Anterior Descending artery was treated with cardiac stents. The patient underwent lab tests and procedures which included COVID-19 test tested performed on an unspecified date (result not reported). The outcome of events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021600543 same patient, different vaccine dose/events
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1374118-1	I have an extremely heavy period now, I woke up and there was a huge blood clot in my underwear and I literally gushed blood I?ve never seen that in all my 27 years of living and I?m afraid of what the next few days of my period will be like.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1374299-1	PT STARTED EXPERIENCING INTENSE HEADACHES ON THE MORNING OF 5/21/2021. THE HEADACHES CONTINUED ON AND OFF FOR SEVERAL DAYS UNTIL ON 5/24/2021 AT 3 PM, SHE HAD A DEBILITATING HEADACHE WITH DIZZINESS THAT COULD NOT BE RELIEVED. THEN ON THE AFTERNOON OF 5/25/2021, SHE STARTED EXPERIENCING VISUAL CHANGES WITH BLACK SPOTS IN HER VISION. SHE WAS ABLE TO SEE HER DOCTOR ON 5/27/2021 AT 10 AM. HE SENT HER FOR A CT SCAN OF HER HEAD WHICH WAS FOLLOWED BY A MRI OF HER BRAIN. THE MRI AND CT SCAN REVEALED THAT SHE HAD A LEFT FRONTAL ISCHEMIC STROKE. SHE WAS ADMITTED TO THE HOSPITAL ON 5/27 AND DISCHARGED HOME ON THE AFTERNOON OF 5/28. SHE HAD A CT ANGIOGRAM OF THE BRAIN, AN ECHO OF THE HEART, ULTRASOUND OF THE CAROTIDS AND A HYPERCOAGULATION BLOOD PANEL. ALL OF WHICH WERE NORMAL AND SHOWED NO EVIDENCE OF WHERE THE BLOOD CLOTS CAME FROM THAT CAUSED HER STROKE. SHE CONTINUES TO HAVE HEADACHES WITH DIZZINESS, FATIGUE, AND NOW BAD ANXIETY.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1376394-1	Developed a R ICA acute thrombus (seen on CTA) and arrived with significant left sided weakness as a code stroke on 6/5. Received tPA and went for thrombectomy to evaluate the acute R ICA occlusion further which was found to be patent (thought to be due to tPA effect). After tPA, pt has complete resolution of weakness. MRI brain found a R striatum acute infarct. Pt was on an estrogen oral contraceptive medication, nonsmoker. Healthy with no hematologic conditions (no prior bleeding or clotting issues). No chronic medical conditions and healthy (would run 6 miles a day, etc), on no other medications or over the counter supplements.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1376861-1	Elevating fever, extreme fatigue, body aches- 9th day bleeding while urinating: 10th day- BLOOD CLOT while urinating 2nd vaccine - milder fever, less fatigue, fewer body aches. 5th day, blood in urine. End of side effects.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1377891-1	I went to chemo therapy in 2018 for colon cancer after colon resection surgery. During chemo my first port failed and caused a huge blood clot in my neck and left arm. I have had three other events in my life of having blood clots, I did not know getting the vaccine would make me so susceptible for more. I got a huge blood clot in my neck and another one in my left arm after receiving the vaccination, my left arm swelled up huge at the elbow, with pain. Blood clots are not always painful for me, depends on where it?s at. I also have swollen lymph nodes in my armpits from the vaccine, confirmed by my oncologist. I went to the ER for the blood clots and again returned two days later to make sure I did not have PE, however the nodules in my lungs had enlarged and my lymph nodes are swollen. This is well after two weeks after I got the second dose of the vaccine, however, the blood clots take time to get as big as they are, so this probably started after the first injection, perhaps just got worse after the second one. I am now stuck on blood thinners forever because of this vaccine, I will not be getting any more of these shots.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378000-1	After 2nd dose, about 24 hours later my period started (I?m about 4 days early). I usually have extremely heavy periods, but right now it?s thick blood clots. Having bad endo stabbing pain. My period lasted 12 days after taking my 2nd dose. Which was 3 days longer than usual. And now, 17 days later, I started menstrual cycle again (too early)! I have the biggest & most blood clots that I have ever seen & heavy period.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378092-1	Had 5 (periods instead of 3) in 3 months, one lasting 13 days. All were heavier and more blood clots then normal. My cycle was 28 days and now seems to be 21 days.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378241-1	Multiple DVTs in both legs 2 PEs in lungs Difficulty breathing poop
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378382-1	2/23 vaccination Within 12 hours, I started shaking, spiked fever, HR increase, SOB, out of body experience, scary experience. Called 911 transported to ER. Monitored cardiac unit for 24 hours, HA. Two months of HAs. I went home and I don't recall exact how everything transpired. I do remember they wanted follow up with PCP. I developed blood clots in R arm, platelets were high since getting vaccinated. *originally diagnosed with colitis but now diagnosed with Chrohns disease. 1/16 I did have a personal injury in a parking lot that exacerbated the issues, I was vaccinated after. *In transition with new doctors now and have not really discussed the potential of this being caused by vaccine, focused more on pain management. **before this vaccine, I was relatively healthy before, no issues.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378384-1	in the pt's words received my 1st (5/5/21) and 2nd (5/26/21) doses of Pfizer at County Fairgrounds. I want to document my experience as I had some pretty severe reactions to my second dose. Evening of the 26th I got chills and a fever of 102.9F. I took one ibuprofen and the fever went away. The next morning I had a slight headache. Around noon I got intense chest pains/pressure that lasted 30 minutes. I could not take a full breath as it was very painful. On the 28th, I woke up with a terrible headache for no apparent reason. Some ibuprofen helped. After a few hours it was gone. At 2am on the 29th, I woke up with very intense stomach cramps. I went to the bathroom and was able to pass out some stool over the course of about 30 minutes in the bathroom, and the cramps went away. On May 31st, I got what seemed to be a blood clot in my left index finger. It went away after about 20 minutes but still felt sore the rest of the day.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1378524-1	Ongoing migraine headaches that are not relieved. Loss of peripheral vision in left eye. Constant ear ringing. Blood clot in right arm. Ongoing dizziness and vertigo. Possible intracranial hypertension. I am being tested next week. These symptoms have been ongoing and have not been relieved. I was given Eliquis for the blood clot, which I no longer take. I take Trokendi 100 mg daily which is being increased for the migraine. I am also suffering with chronic fatigue.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1379019-1	Severe and excruciating cramping in my left leg that I have thrombosis in. The last time I had this is when a clot developed again and hematologist then put me in warfarin(was in Xeralto previously)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1379717-1	4/30 - Five minutes after the vaccine, I passed out. I felt uncontrollably shaky and my vision went white. I was not anxious prior to getting the vaccine and felt terror afterwards. 5/1 - I developed severe depression and suicidal ideation. I reported to those around me that I did not feel safe. I am not diagnosed with depression and did not experience depression prior to the vaccine. I felt suicidal, completely hopeless, anxious, and was uncontrollably bursting into tears without an identifiable trigger. 5/2 - I developed shooting pain in my legs and arms. It felt like a pulsating, stabbing pain that throbbed. It was unlike any pain I had experienced ever before. 5/3 - I developed a metallic taste in my mouth that could not be washed away with water or food. It was extremely noticeable. 5/4 - My period came a week early. I have been on birth control for seven years and have never experienced any changes in my menstrual cycle. I made no other lifestyle or health changes other than the vaccine. The period was extremely heavy, full of clots, and was painful. It lasted longer than usual, until about 5/9. 5/16 - I developed another period only a week after the first period. This period was equally as heavy and long-lasting. It remained until about 5/21. Following the vaccine, I began suffering from heart palpitations, leg cramps, and random pains. I did not experience these prior to the vaccine. I cannot identify the date these symptoms began, but it was within the first week of taking the vaccine.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381046-1	and multiple golf ball-sized clots.; Extreme menstrual period; High pain, extremely heavy flow; High pain, extremely heavy flow; This is spontaneous report from a contactable consumer (patient). A 35-year-old female patient received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE) via an unspecified route of administration, administered in Arm Right on 07Apr2021 11:15 (Lot Number: ER8734) at the age of 35-year-old, as 2nd dose, single for covid-19 immunisation. Medical history included known allergies Latex and Penicillin. Prior to vaccination, the patient was not diagnosed with COVID-19. Historical vaccine included first dose of bnt162b2 (lot number: EN6206) on 17Mar2021 11:15, administered in right arm, for covid-19 immunisation. The patient's concomitant medications were not reported. The patient had extreme menstrual period (unlike her typical cycle) experienced about 18 days after the second shot. High pain, extremely heavy flow (more than an oz and hour sustained for almost two full days), and multiple golf ball-sized clots. Events occurred on 25Apr2021 07:30. No treatment was taken for adverse event (AE). Since the vaccination, the patient has not been tested for COVID-19. The outcome of events was recovered. No follow-up attempts are possible; information about lot/batch number obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381062-1	blood clots; she did get really sick / she was displaying a lot of symptoms; fever; whole body was aching; fatigue; nauseated; bruises on the legs / now she's getting more / she would say they have worsened; This is a spontaneous report received from a contactable consumer (patient's mother). A 19-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EW0182), via an unspecified route of administration in arm left on 14May2021 at single dose for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. Patient previously received flu shot and experienced nausea. The patient experienced blood clots on an unspecified date, bruises on the legs, she was getting more, she would say they have worsened in May2021, fever on 15May2021, whole body was aching on 15May2021, fatigue on 15May2021, nauseated on 15May2021, and she did get really sick, she was displaying a lot of symptoms on an unspecified date. Patient started having bruises on the legs and thought that it might be related to blood clots. She mentioned that she also experienced the common side effects but she was not concerned of that as people she knew also had those symptoms but was more concerned about the bruising. On patient's legs, she had a bunch of bruises. Reporter was just really nervous about her daughter's bruises. When patient was working, she was not noticing, but today patient noticed her legs and said reporter it was a lot of them. Patient did get really sick with her first shot, she had a fever, she couldn't go to work the next day. Reporter was not sure if the patient should get the second dose on 04Jun2021. Patient's whole body was aching, she was nauseated, she was displaying a lot of symptoms. Reporter checked online and they said it was a normal thing. Patient came down and showed reporter the bruises. Patient wasn't really paying attention to them. Patient had one or two bruises when she noticed and at the time of the report she was getting more. They had worsened. Patient took ibuprofen (ADVIL) for fever and she felt better. Outcome of events blood clots and she did get really sick, she was displaying a lot of symptoms was unknown, of event bruises on the legs, she was getting more, she would say they have worsened was not resolved, and of other events was resolved.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381070-1	I am leaning towards calcified blood clots, however that has not been confirmed yet; felt a bump under my skin in my left arm/more painful bumps had appeared; This is a spontaneous report from a contactable healthcare professional. A 37-year-old female patient received the second dose of BNT162B2 (lot number: EN6205), via an unspecified route of administration in arm left on 18Mar2021 14:00 at single dose for COVID-19 immunisation. Medical history included known allergies: Strawberries. Concomitant medications included muscle relaxers, and anti-inflammatory. Patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Patient previously received the first dose of BNT162B2 (lot number: EN6200) on 24Feb2021 14:00 in left arm for COVID-19 immunization. Shortly after, maybe a week, after getting the second dose patient experienced a bump under her skin in the left arm on 01Apr2021 16:00. Over the last few weeks more painful bumps had appeared. Most of these hard lumps were in her left hand. She had one that she knew of in her right hand. Patient met with her PC doctor and the doctor sent her for X rays and ultrasound. Every tech and doctor that had looked at her hands had never seen anything like this. These masses had been confirmed with the ultrasound and more were found during that visit. Patient was moving forward with getting a magnetic resonance imaging (MRI) of her hands. Patient was leaning towards calcified blood clots (01Apr2021 16:00), however that had not been confirmed yet. Adverse events result in doctor or other healthcare professional office/clinic visit. Nasal swab on 24Apr2021 was negative. Treatment was received for the adverse events. Outcome of the events was not resolved. Information about lot/batch number was available. Further information was expected.; Sender's Comments: Based on temporal association, a causal association between the reported event and BNT162B2 cannot be fully excluded. Case will be reassessed when additional information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and investigators, as appropriate.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1381101-1	feel a constant and migrating thrombosis/palpitations on the right side of my head; palpations; mildly painful -not quite a headache; painful sensations in the lower left leg; a slight discomfort of the arm where the shot was administered; This is a spontaneous report from a contactable consumer (patient). A 56-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration, administered in right arm on 11May2021 10:30 (at the age of 56 years old) (Lot Number: EW0182) as 1st dose, single for COVID-19 immunization. Medical history included high blood pressure, allergies: contrast dye. There were no concomitant medications. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. There were not any other medications the patient received within 2 weeks of vaccination. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient has not been tested for COVID-19. Patient reported that there was a slight discomfort of the arm where the shot was administered (11May2021) - not enough to worry about & expected. The next morning (12May2021) he felt painful sensations in the lower left leg - sensations similar to stress fractures and palpations. It subsided and has not returned. About three/four days later (reported as 14May2021 18:00) continuing two weeks leading to the second dose - he felt a constant and migrating thrombosis/palpitations on the right side of his head. It was mildly painful - not quite a headache yet persistent and localized to the RT temporal lobe above the temple & slightly above that region. It became exacerbated by critical thinking when it occurred & persisted for long periods of time. The cognitive effect has become slightly debilitating & stressful. No treatment received. The outcome of the event 'a slight discomfort of the arm where the shot was administered' was unknown, of the event 'painful sensations in the lower left leg' was recovered, of the other events was not recovered.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1381126-1	12 days after first dose vomited clotted blood; 12 days after first dose vomited clotted blood; chest tightness; extreme fatigue to the point of barely being able to walk continued for several weeks.; extreme fatigue to the point of barely being able to walk continued for several weeks.; bleeding; elevated HR; BP; elevated HR; BP; difficulty breathing; This is a spontaneous report from a contactable consumer (patient). A 53-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE lot number: EN6202) via an unspecified route of administration, administered on the left arm on 01Mar2021 as 1st dose, single for COVID-19 immunization. Medical history included Echo from Nov2020, exposure to A. flavus in 2014 with development of fungal ball in sphenoid cavity causing brain injury and all body damage, Autonomic disorder developed tachycardia. Concomitant medications were not reported. It was reported that 12 days after first dose on 13Mar2021, patient vomited clotted blood; hospitalized. The patient was hospitalized from 13Mar2021 to 15Mar2021. Cleared and discharged after a day unknown cause of bleeding on 16Mar2021. Elevated HR; BP; difficulty breathing, chest tightness and extreme fatigue to the point of barely being able to walk continued for several weeks. Events were considered serious, disability. The events resulted to a doctor or other healthcare professional office/clinic visit and emergency room/department or urgent care visit. No treatment was received. Covid test post vaccination: Nasal Swab on 22Mar2021 and 10Apr2021 Negative. Prior to vaccination, the patient was not diagnosed with COVID-19. Outcome of the event vomited clotted blood was recovered on 15Mar2021 while not recovered for the other events.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1381200-1	she is bleeding into the eye and that to her is just, it keeps getting worse; thrombo; elevated potassium; had bruising in the other leg; doctor did PTT, confirmed as PTT, and it was elevated; This is a spontaneous report from a contactable consumer (patient). An 81-year-old non pregnant female patient received the second dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; formulation: Solution for injection, Lot Number, Expiry date UNKNOWN), via an unspecified route of administration at left arm on an unknown date in Feb2021 (at the age of 81 years old) as a single dose for COVID 19 immunisation. Medical history was not reported. The patient received the medications within 2 weeks of vaccination was Asperin. Prior to the vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. Patient received first dose of BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; formulation: Solution for injection, Lot Number, Expiry date UNKNOWN), via an unspecified route of administration on an unknown date in Feb2021 experienced adverse events such as large bruising on one leg, after the next dose, bruising on the other leg. She has bruising on both lower limbs, a bleeding disorder. She bled from her eye at the ophthalmologists office. On an unspecified date patient second one, then had bruising in the other leg. She went to the hematologist, and to the doctor. The doctor did PTT, confirmed as PTT, and it was elevated. Reported added that patient also had an elevated potassium. She states this was like what you would get if patient were taking warfarin. it was very concerning now, not only does she have bruising in the leg, but she was bleeding into the eye and that to her was just, it keeps getting worse. She has nothing else to add to that report except that her mom has an elevated PTT, TPTT and states she does not know, it was thrombo. At 81, her mom walks around an hour a day, does aerobics. She was not seeing a hematologist and she was concerned. If she cut herself, she would bleed out. The fact she was bleeding into the eyeball, she has to google the acronym but it was basically clotting issues it was resulting in. No additional details provided; Reporter did not know the name of the elevated test. Information about lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381206-1	She is worried it is a kind of blood clot.; She is worried it is a kind of blood clot.; Swollen lymph nodes in right leg; This is a spontaneous report from a contactable consumer(Patient). A 61-years-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Solution for injection, Lot number was not reported), dose 2 via an unspecified route of administration, administered in Arm Left on 23May2021 as 2nd DOSE, Single for covid-19 immunization. Medical history included diagnosed allergies, compromised immune status, respiratory illness, genetic/chromosomal abnormalities, endocrine abnormalities, obesity. No family medical history reported. The patient's concomitant medications were not reported. Patient previously took historical vaccine of first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE; Solution for injection, Lot number was not reported), via an unspecified route of administration, administered in Arm left on 02May2021 as single dose for covid-19 immunization. The patient did not receive any other vaccine in four weeks. Prior to vaccination patient was not diagnosed with COVID. It was reported that, on 26May2021, after getting the second dose of the Covid vaccine, she had swollen lymph nodes in her right leg. She was worried and asks what to do and She asked if she needs to go to Urgent Care. she is worried it is a kind of blood clot. She asks if Pfizer will call her doctor and then call her. All she had were swollen lymph nodes. They are getting big and big. She confirms they are getting bigger. No relevant tests reported. The outcome of the events was not recovered. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381304-1	stomach pain last about for 2 or 3 days, stomach pain came back after second dose; The pain was so bad he couldn't stand up.; The x-ray showed he had a big black spot in his stomach and the doctor told him it was a blood clot.; This is a spontaneous report from a contactable consumer (patient) and a physician. A 68-year-old male patient received the first dose of BNT162B2 (PFIZER COVID-19 VACCINE, lot number: EN6198) on 26Feb2021, and the second dose of BNT162B2 (lot number: EN6208) in arm on 19Mar2021 , both at the age of 68 years old at single dose for covid-19 immunisation. Medical history was none. The patient's concomitant medications were not reported. The patient previously received flu vaccine in Oct2020 or Nov2020. The patient had his first dose of BNT162B2 on 26Feb2021 and about 4 days later he had stomach pain on 02Mar2021. It lasted about 2 or 3 days then the stomach pain let up. The patient thought the stomach pain was over. Then the patient received his second dose of BNT162B2 and in about the same amount of time after receiving BNT162B2, the stomach pain came back. On Mar2021, the pain was so bad he couldn't stand up. The patient went to the doctor and he was given 4 ground up Aspirin to take. The patient screamed when the doctor pressed on one spot of his stomach. The patient was taken to the x-ray room and his stomach was x-rayed. He was asked where the pain was and when the doctor pushed on that spot, his back arched up in the air because it hurt. The x-ray showed he had a big black spot in his stomach and the doctor told him it was a blood clot. The doctor on call said it was a blood clot and his regular doctor told him it was a blood clot. The doctor's determined the blood clot was from BNT162B2 because the blood clot happened right after he received the vaccine. Besides his regular doctor who is listed in the formal field, he also saw another doctor when he went in that day with the stomach pain. He was told his regular doctor was busy and he had to see another doctor. The patient didn't have any blood drawn at the doctor just the x-ray of his stomach. The doctor said he had a blood clot, a black spot, about the size of a silver dollar in his stomach. The doctor clamped her index finger and thumb together in a circle to show him how big the black spot was. The x-ray technician and two doctors told him it was a blood clot. He had the 4 aspirin ground up and the doctor told him the aspirin got rid of the blood clot. He also took some pills for 2 to 3 days but he can't recall the name of the pills. He says the x-ray tech zeroed in on the spot that was causing him pain in his stomach. The outcome of events was unknown. No follow-up attempts are needed. No further information is expected.; Sender's Comments: Considering the temporal association, a causal association between administration of BNT162B2 and the onset of blood clot cannot be fully excluded. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to Committees and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1381828-1	Within 24 hours, my right foot swelled, especially the middle of the sole, but also the toes and top of the foot. After a two-week delay due to a misdiagnosis by my PCP, I was seen at an urgent care center. An ultrasound of the right leg discovered a blood clot. I was placed on Eliquis 10 mg twice a day for the next 7 days followed by 5 mg twice a day. After 7 days, the swelling was reduced by 90%. The swelling has remained at that level since then. I changed my PCP on May 21 and was cleared for most activities. She advised me to continues with the Eliquis for three months from May 3, with no need for further testing.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1382265-1	PATIENT DEVELOPED SEVERE ABDOMINAL PAIN, N/V, FEELING FAINT LESS THAN 24 HOURS AFTER ADMINISTRATION (THE NIGHT OF 6/3/21). ON 6/4/21 PATIENT WENT TO ER AND WAS GIVEN IV FLUIDS. ON 6/5 PATIENT PASSED BLOOD CLOTS IN STOOL THAT BECAME MORE SEVERE ON 6/6/21. 6/6/21 PATIENT RETURNED TO ER AND WAS DX WITH COLITIS AND A GI BLEED. PATIENT STATES A KNOWN ALLERGY TO POLYETHYLENE GLYCOL - TOOK BENADRYL BEFORE RECEIVING EACH DOSE OF PFIZER VACCINE PER THE GUIDANCE OF HER PRIMARY CARE PHYSICIAN. PATIENT HAD NO REACTION TO FIRST DOSE.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1382490-1	Patient presented for 2nd dose and reported having a blood clot the day after receiving her first dose. She is unsure if it was due to the vaccine or her vascular malformation. She did not seek medical help or receive a diagnosis from a medical professional. States the clot dissolved on its own. Pfizer Covid vaccine
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1382933-1	About three days after vaccine, I started bleeding and getting blood clots. I went to ER to make sure I wasn't miscarrying. I wasn't that day. Everything looked okay. Baby was measuring much smaller than it should be. I kept bleeding lightly for a couple of days. Tuesday I was bleeding heavily - I went to ER again. Hormone levels - were going up but not as they should be. From Tuesday - Friday, I didn't bleed. I started bleeding on Saturday again. And I'm still bleeding. I called my OB today and she is going to call me back in the morning to let me know what to do. I was last seen on Friday at ER again. Pregnancy History - One other pregnancy - carried twins fine. Estimated date of delivery for this pregnancy - January 17, 2022.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1384248-1	Patient stated that she has blood clots in her legs due to COVID vaccine; This is a solicited report from a non-Pfizer sponsored program, received from a contactable consumer, based on information received by Pfizer. This 38-year-old female patient was involved in a patient support program. The patient (patient ID: redacted) received apixaban. The report describes a case of thrombosis (Patient stated that she has blood clots in her legs due to COVID vaccine). Co-suspect products included COVID-19 vaccine for an unknown indication. On an unknown date, the patient started apixaban (unknown route), (unspecified dose and frequency). On an unknown date, the patient experienced thrombosis (seriousness criterion medically significant). The action taken with apixaban (unknown) was unknown. At the time of the report, thrombosis outcome was unknown. The patient received apixaban for unknown indication. The patient reported that she had blood clot in her legs due to COVID vaccine. For apixaban (unknown), the reporter considered thrombosis to be not related. Further follow up was not possible, as no further information was available. The reporter's assessment of the causal relationship of the event with the suspect product COVID-19 vaccine was not provided at the time of this report. Since no determination has been received, the case is managed based on the company causality assessment.; Sender's Comments: BMS Medical Evaluation Comment: This patient had thrombosis after receiving apixaban study therapy. Patient also got vaccinated for COVID-19. Based on the limited information available regarding the therapy details, event onset date, action taken with the suspect and outcome of the event, it cannot be ascertained with the reasonable possibility that the apixaban could have contributed to the event thrombosis.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1384585-1	blood clot in his leg; swelling in his leg; sore arm; his muscle aches; didn't dilute it and he's received 6 times over the amount; didn't dilute it and he's received 6 times over the amount; Diarrhea; real tired; his blood pressure was real high; breathing problems; This is a spontaneous report received via a Pfizer-sponsored program. A contactable consumer reported that a 46-year-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Batch/Lot number was not reported) via an unspecified route of administration on 20Apr2021 (at the age of 46-year-old) as 1st dose, single for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. On 20Apr2021, patient experienced diarrhea, was real tired and his blood pressure was real high. It was also reported that the vaccine wasn't diluted and the patient received 6 times over the amount on 20Apr2021. On an unspecified date, patient had sore arm and his muscles ached. On Apr2021, patient had breathing problems and was treated with breathing treatments three times a day and was put on steriods and inhalers. On May2021, patient had a lab work that confirmed that he had a blood clot in his leg, which has worsened because he has some swelling in his leg, the same leg that they found the blood clot in. The patient was hospitalized for the event blood clot in his leg and was given blood thinners. The patient underwent lab tests and procedures which included CAT scan of head and lungs both with unknown results on May2021. Outcome of the events blood clot in his leg, swelling in his leg and breathing problems was not recovered while outcome of all other events was unknown. Information on the lot/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1384595-1	received the first dose of the Pfizer COVID-19 vaccine and also recently started taking Depo-Provera. After receiving the first dose of the Pfizer COVID-19 vaccine her daughter developed a blood clot; tired and frail; feels lost and light headed; cramping pain; This is a spontaneous report from a contactable consumer (Patient's mother). A 17-year-old female patient received first dose of BNT162B2 (BNT162B2) via an unspecified route of administration, administered in Arm Right on 03May2021 15:00 (Batch/Lot number was not reported) as 1ST DOSE SINGLE for covid-19 immunization and medroxyprogesterone acetate (DEPO-PROVERA), route of administration, start and stop date, batch/lot number and dose were not reported for contraception. The patient's medical history and concomitant medications were not reported. The patient experienced received the first dose of the pfizer covid-19 vaccine and also recently started taking depo-provera. after receiving the first dose of the pfizer covid-19 vaccine her daughter developed a blood clot, tired and frail, feels lost and light headed and had cramping pain all on unspecified dates. It was reported that after receiving the first dose of the Pfizer COVID-19 vaccine her daughter developed a blood clot. Her daughter, who is currently on Depo Provera got her first dose of the Covid 19 vaccine. The daughter has been experiencing blood clotting and would like to further document this experience. Is also seeking medical advice to inform her daughter. Her daughter Has been on Depo Provera for 4 to 5 months. Since she had the Covid vaccine, she has been having bad blood clotting and has been having a continuous period. When she got the shot, she has been getting blood clots straight for three weeks. Patient is Tired and frail: She is a waitress, so all of this has made it hard. It started right when she got the shot and has pretty much stayed the same. She will feel like she is lost when she wakes up and is light headed from the loss of blood. It occurs when she is doing activities, like working. Patient took some Midol one day for cramping pain but it barely did anything. The action taken for medroxyprogesterone acetate was unknown. Outcome of tired and frail was not recovered. Outcome of all the other events was unknown. Information on Lot/Batch number was not available. Additional information has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				"Anytime I exerted myself it get worse so I went to the hospital and they admitted me because they couldn't control my angina; Bypass had developed a huge blood clot; Hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed.; Chest Pain; Nose bleed; Light headedness/light headed and dizzy; Hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed.; Hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed.; My arm hurt; This is a spontaneous report from a contactable consumer (reported for herself) via the Pfizer sponsored program. A 44-years-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in Arm Left upper on 07Apr2021 (Batch/Lot Number: ER8735) as 2ND DOSE, SINGLE for covid-19 immunization, at the age at vaccination of 44 years old. Medical history included coronary artery disease, Microvascular disease, migraine, Bypass of LAD artery with my LIMA artery, angioplasty, had open heart surgery before, all from an unknown date and unknown if ongoing. Historical vaccine included first dose of Pfizer COVID-19 Vaccine on 10Mar2021 at the age at vaccination of 43 years old (got the first shot in right upper arm) for covid-19 immunization. Concomitant medications included clopidogrel bisulfate (PLAVIX); lisinopril; amlodipine; rosuvastatin calcium (CRESTOR); fluoxetine hydrochloride (PROZAC); clonazepam (KLONOPIN), Nexletol (further clarification unknown). The patient experienced anytime i exerted myself it get worse so i went to the hospital and they admitted me because they couldn't control my angina on Apr2021 with outcome of unknown, bypass had developed a huge blood clot on Apr2021 with outcome of unknown, chest pain on 19Apr2021 with outcome of not recovered, nose bleed on 08Apr2021 with outcome of recovered on unknown date, light headedness/light headed and dizzy on Apr2021 with outcome of unknown, hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed (hematoma) on Apr2021 with outcome of recovered on 19Apr2021, hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed (Bruising of arm) on an unspecified date with outcome of unknown, hematoma on her arm a couple of days later after that she got it was swollen, bruises she got nose bleed (Swelling arm) on Apr2021 with outcome of unknown, my arm hurt on Apr2021 with outcome of unknown. The patient underwent lab tests which included lipids: unknown (had lipid channel), something checking for heart stress: unknown, both on unknown date. Therapeutic measures were taken as a result of angina attack, chest pain, nose bleed, hematoma, Bruising of arm, Swelling arm and Pain in arm. Seriousness for events ""anytime i exerted myself it get worse so i went to the hospital and they admitted me because they couldn't control my angina"" and ""bypass had developed a huge blood clot"" was hospitalization, medically significant. The patient was hospitalized for both events from 20Apr2021 to 23Apr2021. Events chest pain, Light headedness/light headed and dizzy resulted in Emergency Room Visit. Additional information: Transferring agent stated, ""I have patient on the line today and the reason for my transfer today is that she is having adverse effects from the COVID-19 Vaccine, she did clarify it is the Pfizer COVID Vaccine and she is experiencing hematoma on her arm a couple of days later after that it was swollen, bruises she got nose bleed as well and on 19th April that she was experiencing chest pain she went to ER and they were thinking it was cause of the COVID Vaccine as well."" Consumer stated, ""On 07Apr2021, I have the LOT number and everything of the vaccine I received for both of them on 07Apr2021, I got the Pfizer the second Pfizer shot and I developed a hematoma on my arm which was as big as a baseball. It was really swollen and I take blood thinners and on the next day on 04-08, I started getting a lot of nose bleed. I never get a nose bleed and so I called my doctors and asked what should I do and she said stop taking your blood thinners so I stopped taking the blood thinners and then may be after 7 days, I started taking my blood thinners again. So, it took about 3 weeks for the hematoma to go away on 04-19th, I had a huge chest pain, I have had open heart surgery before and I had to go to the ER because I felt light headed and dizzy. My arm hurt, my chest, I took nitroglycerine (treatment) and it made still little better but anytime I exerted myself it gets worse so I went to the hospital and they admitted me because they couldn't control my angina and my cardiologist did an angioplasty (Further clarification unknown) and he found that my 2019, I had a bypass of LAD artery with my LIMA artery (Medical History) and he found that my bypass had developed a huge blood clot and failed. So, basically I am looking at go in and having another bypass using my LIMA artery but the surgeon, the coronary surgeon that I talking he wanted me to call Pfizer to let you guys know because he said that that is so weird that it failed and the shot and everything and he just wanted you guys to be aware and you know that's what we wanted safety because I know that when I went in I had a lot of a questions and none of my doctor could answer. It says I shouldn't take it if I am on blood thinners but they said it is still better than you getting COVID so I went ahead and took the shot but actually now I have to get a bypass again. It might be from the vaccine, it might not be from the vaccine but the timings is just you close together there."" Patient took Nitroglycerin, she was on now but that's because of the chest pain so that was on before she had the chest pain when she had the shot (Further not clarified). When probed of the event is still ongoing: Consumer stated, ""Yes, because my By pass graph is now failed."" Treatment: Patient took Tranexa (name not clarified). It was a medicine for her chest pain. Outcome: Consumer stated, ""As in nose bleed and bleeding are gone, now it is just chest pain"". Follow-up attempts are completed. No further information is expected."
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1384638-1	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1385234-1	Blood Clot from heart to brain. Caused a TIA
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1385337-1	Was living happily and independently in a retirement community. After the second dose because losing his memory, focus, depression, followed by 2 strokes, brain bleed, blood clots, seizures, and a Dx of Parkinson's and Parkinson's dementia. He saw doctors monthly and never had any signs of any of these things.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1385500-1	Very heavy period began, 2 weeks ahead of schedule. Much heavier bleeding began after about 12 hours and has continued for 3 days. This level of bleeding is not typical. Many blood clots have also been passed.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1386375-1	AFTER THE FIRST VACCINE: HERPES BASED BLISTERS IN MOUTH TO THE POINT THAT TAKING OR EATING WAS VERY PAINFUL. SAW MY PA ON 03/25/21. VALTREX PRESCRIBED. BLISTERS WERE MUCH BETTER WITHIN 48HOURS. 03/27/21 UNEXPECTEDLY HAD MY PERIOD. I HAD JUST FINISHED UP MY PERIOD WHEN I HAD THE VACCINE ON 03/18/21. THIS PERIOD WAS UNLIKE ANY OTHER. THE BLEEDING WAS INCREDIBLY HEAVY, FULL OF CLOTS AND LASTED 9 DAYS. MY USUAL CYCLE IS 5. AFTER SECOND DOSE: ANOTHER UNEXPECTED PERIOD. THIS TIME NOT AS HEAVY, NOT AS MANY CLOTS AND DID NOT LAST AS LONG. SORE UNDERNEATH ARM AND LYMPH NODE UNDER ARM SWELLED.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1386595-1	Wednesday, May 19, 2021 1st COVID-19 vaccination shot Summary: Wednesday, May 19, 2021 received my 1st COVID-19 vaccination shot. My body tried to reject it. I threw up 5 times. Ensued many unwanted side effects. I don?t think it would be good to get the 2nd vaccine dose. Given I am not fully vaccinated, I need an alternative solution to prevent being infected with the virus. Help!!!! Please Help!!! Bad Side Effects: Stuffy Nose Congested Nose Headaches Vomiting Nauseous Upset Stomach Unsettled Stomach Unusual Perspiration Pain in Left Arm Abnormal Period Menstrual Blood Clots Shoulder Pain/Tension Upper Back Pain/Tension Chest Pain/Tension Shooting Pain in Chest Heart Palpitations? Numbness Runny Nose Sneezing Profusely Allergies? Hoarse Voice Winded Dry Mouth Existing Worsened: Cold Sores Stomach Noises Snoring? Foot Numbness Ear Ringing Eye Floaters Joints Popping Sound Other Observations: Stomach Bloated Increased Flatulence Pseudo Conclusion: Given my body?s adverse reaction I don?t think it is wise to get the 2nd shot. It is obvious my body does not like it. I am disappointed because I really did not plan to get vaccinated; however, I felt I had no other option when the government made the announcement. When the government announced No masks for those vaccinated, I know there would be some NOT vaccinated & NOT wearing a mask thus putting me at risk. I am okay with wearing a mask for another year or even for the next 5 years. The doctors have said me wearing a mask is protecting others not necessarily myself. I concluded even if I continue to wear a mask there would be others (NO mask & NOT vaccinated) not following the rules potentially exposing me to the virus. Hence the only way to protect myself is to get vaccinated. The vaccine appears to have affected so many areas of my body. I regret getting the 1st shot, I want to remove every trace of it from my anatomy. Please let me know if it is possible to rid the vaccine along with all the bad side effects from my body. I do not want to subject my body to anymore of this vaccine; however, I still need a reliable solution to avoid being infected with the virus. I don?t know what to do. Details: Wednesday, May 19, 2021, I received my 1st COVID-19 vaccination shot around 3:30pm. The medical staff had all those being vaccinated to sit for at least 15 minutes for observation to be monitored. I stayed for monitoring for close to an hour (wearing my masks). I felt fine, I noticed my right nostril felt stuffy, congested with a slight burn sensation. I noticed a slight headache when I left. I had about 3 errands to do. 2 hours later I wasn?t feeling well. In addition to a stronger headache my stomach felt queasy. Around 6:30pm I had to pullover to vomit; I threw off my mask and threw up once and then a 2nd time. I sat in my car taking deep breaths until I felt I could make it home. I make it home around 7pm in time to throw up 3 more times, for a total of 5 times. The vomit tasted acidic, sour, citrusy, and metallic at times, and salty. I was only throwing up the vaccine it was mostly clear with white alka seltzer looking deposits. I last ate some crackers around 12 noon. Vomiting is NOT normal for me. I can?t remember the last time I threw up, many many many years ago. l had a headache, queazy stomach, and was exhausted; about 7:30pm tried to sleep. I got up around 10:30pm ate a few saltines and drank a little water but I still didn?t feel good so I didn?t eat anything else. Rough night trying to sleep, tossing and turning. Next day (Thursday day 2) I felt slightly better, not comfortable enough for real food so I drank plenty of water in small amounts and ate saltines; headache subsided. I also had more than usual underarm perspiration with a strong unfamiliar odor. Wednesday, May 19, 2021 1st COVID-19 vaccination shot It appears the vaccine affected my menstrual cycle. My period usually lasts for 3-5 days max, usually light blood flow on days 4 and 5. It was a normal light day 5 when I received the vaccine. After the vaccine, I began having heavy blood flow. The heavy blood flow is accompanied by blood clots. In the past I?ve had a blood clot here and there but not like this. Menstrual blood clots for 9 days after being vaccinated. This is NOT normal for my body. My period was 14 days. Prior to being vaccinated, I NEVER had a period last this long. To my surprise the vaccine affected my female organs. I am right handed, so the nurse recommended using my left arm for vaccination. Left arm where vaccine injected was very sore. Arm pain subsided after 3 days. Pain soreness tension primarily on my

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	left side, shoulder, upper back, chest area. There is a shooting pain on left **Adverse Event Description**
				shoulder, down back to breast, chest heart area; pain comes and goes. Heart palpitations? Experienced this for over a week about 10 days. Numbness in left hand when I sleep on my left side, wake with my left hand numb. Numbness in both hands when I try to sleep on my back. Numbness in my feet after standing for about 15 minutes. Numbness in my leg and buttocks, primarily left side after sitting for an extended period of time. Before being vaccinated I only experienced numbness in my right foot after being on my feet for an extended period of over 4 hours. Last year my right foot/heel would began to hurt after being on my feet for an extended period of time (> 4 hours). Then there would be pins and needles sensation. The solution would be to rest and elevate my feet. I also replaced my shoes (proper support). Post vaccination even before the time my feet would hurt now my right foot goes numb within 30 minutes of standing even while wearing supportive shoes. My left butt cheek goes numb when I am sitting with my legs elevated; this was my most comfortable position pre-vaccination. Since the vaccine my stomach feels upset or unsettled especially in the morning. I am also noticing bloating and more gas than usual. My stomach is also excessively more noisy. Before the vaccine my stomach would make noises when I was hungry or hadn?t eaten, but now there are so many stomach noises almost nonstop all day and night long all the time My nose is feeling congested with a slight burn sensation since getting the vaccination. Immediately after getting the shot my right nostril felt stuffy, congested with a slight burn sensation. This congested feeling comes and goes, especially noticeable when wearing a mask. Prior to being vaccinated I had not experience this issue even as I have worn multiple masks for over a year with NO stuffy nose. Saturday, May 29, 2021 (day 10 post vaccination) I get cold symptoms; runny nose and sneezing. I haven?t had a cold in years; I don?t really get colds; if so the symptoms are mild and quickly dissipate. Now my nose has been running pretty much non-stop causing me to blow my nose just as often. I am also sneezing a lot. This is NOT normal for me. I drank plenty of liquids; however, running nose and sneezing continued leading to hoarse voice. These symptoms prevented me from sleeping. I don?t know if these are cold or allergy symptoms. I had NO allergies before the vaccine. These symptoms finally subsided after 3 or 4 days. My neighbors are waking up in the middle of the night (2am, 3am, 4am, 5am). I began to wonder if I am waking my neighbors possibly with snoring. I don?t really know, no evidence. Throat feels winded and dry when talking especially when I try to whisper. I have to drink a lot of water. Ringing in my ear started In 2019. I thought this phantom sound went away; however, post vaccination I?m hearing the ringing sound again. It seems worse than before. 2018 diagnosed with eye floaters. I am noticing floaters more since getting the vaccine. I usually see the eye floaters when going to light from dark. Post vaccination I am noticing the floaters even in dim light; not normal. I started getting Cold sores in 2006. I thought I had them somewhat under control. Since the vaccine l keep getting cold sores, this may be due to my menstruation. Recently within the past year, my shoulder and elbow joints sometimes make this popping sound when I exercise/move my arm. Post vaccination my shoulders and elbows are making the popping sound every time I move my arm. Now sometimes my knees and ankles are also making a popping .
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1386605-1	Blood Clots and Bleeding

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1387713-1	"blood clot already eliminated; pain; This is really miserable because it persisting and it is always there; I feel like frustrated; light pink rashes in the back from knee above in both legs; both of my legs below the knees had really bad itches; mild pain in the back of the upper left leg; sharp pain at a spot in my left hip joint; Sore in the injection area; This is a spontaneous report from a contactable consumer (patient). A 62-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration, administered in Arm Left on 23Apr2021 15:30 (Batch/Lot Number: EW0169; Expiration Date: 31Aug2021) as 1ST DOSE, SINGLE DOSE for covid-19 immunisation, age at vaccination 62 years old. Patient is not pregnant at vaccination time. Medical history included diabetes mellitus from Sep2020 to and blood pressure (abnormal) from an unknown date. Concomitant medications included metformin taken for diabetes mellitus, start and stop date were not reported; dapagliflozin propanediol monohydrate (FARXIGA) taken for diabetes mellitus, start and stop date were not reported; levothyroxine (LEVOTHYROXINE) taken for an unspecified indication, start and stop date were not reported; lisinopril (LISINOPRIL) taken for blood pressure (abnormal), start and stop date were not reported. Clinical course was reported as follows: On 24Apr2021 noon (12:00), patient suddenly felt a shar at a spot in her left hip joint. It was then moved to the mild pain in the back of the upper left leg, which never disappeared. The pain there got a lot worse, so severe, also reported as now just a mild pain. Patient went to bed around 12 am on 25Apr2021, both legs below the knees had really bad itches; by next day 6 AM it was gone and never returned. On 26Apr2021, in late afternoon, in the back of both of the legs started bad it itches, and patient saw light pink rashes in the back from knee above in both legs, which disappeared by midnight, and never again noted. Sore in the injection area (Apr2021) similar to flu vaccine, which was gone after a few days. Patient stated adverse reaction getting worse and asked how it can be resolved because he can't function normally because of adverse effect. Patient stated that this is really miserable because it is persisting, and it is always there. He stated he is a normal person his breath, he still sees, still read, and is sound. He feels he is in constant pain, mild pain is annoying, but this week is so worse. Patient stated his ""blood clot already eliminated"" (onset date unspecified, event not clarified). Patient felt frustrated. Second dose was scheduled on 14May2021, but he didn't take it, he cancelled the second dose because of pain. Treatment was not given for the events (does not take medications). Outcome of the events ""blood clot already eliminated"" was recovered on an unspecified date, ""both of my legs below the knees had really bad itches"" was recovered on 26Apr2021, ""light pink rashes in the back from knee above in both legs"" and ""Sore in the injection area"" was recovered on Apr2021. Outcome of ""sharp pain at a spot in left hip joint"" and ""mild pain in the back of the upper left leg"" was not recovered (persisting for three weeks). Outcome of the other events was unknown. Seriousness reported as non-serious. Follow up needed, further information has been requested."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1387722-1	massive blood clot from her thigh to her ankle; incapable of walking; serious pain in my left thigh/muscle strain; serious pain in my left thigh/muscle strain; leg fell asleep; This is a spontaneous report from a contactable consumer (patient). A 24-year-old female patient received the second dose of BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, lot number and expiry date unknown), via an unspecified route of administration, administered in the left arm on 11May2021 at 14:45 as a single dose for COVID-19 immunization. The patient's medical history was not reported. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient was taking unspecified birth control medication. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient was on her second week of completing the vaccine. The patient reported that on Thursday night, 20May2021, she was just laying down on her couch watching Netflix with her laptop on her lap, her leg fell asleep but she didn't think anything of it because once she started walking around it went back fine. However, on 21May2021, Friday morning, she was having serious pain in her left thigh. She believed it was a muscle strain. The patient did a cold compress, warm bath and took Tylenol from Friday to Saturday. She thought it got better but by 23May2021, Sunday night, she was incapable of walking on her own and needed assistance just to go to the bathroom. On 24May2021, she went to Urgent Care and they stated that she may have a blood clot so she went to the ER (Emergency Room). While at the ER they said she had a massive blood clot from her thigh to her ankle. She would need a procedure done to remove it and was later admitted into the hospital and scheduled for the procedure. Blood thinners and suctioning out the blood clot was done as treatment. The patient been tested for COVID-19 (nasal swab) on 24May2021 with a result of negative. The outcome of the events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1387756-1	Sharp chest pain; Shortness of breath; Multiple occlusive thrombus in right leg(distal femoral, and popliteal vein with nonocclusive in right mid femoral vein); Right leg swelling; This is a spontaneous report from a contactable Other Health Professional. A 53-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on Apr2021 (Batch/Lot number was not reported) as unknown, single dose for COVID-19 immunisation. Medical history included Acute cholecystitis without obstruction from an unknown date and unknown if ongoing. The patient's concomitant medications were not reported. The patient had not been diagnosed with covid-19 prior vaccination. It was unknown if patient had been tested for covid-19 post vaccination. He had no known allergies. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient experienced multiple occlusive thrombus in right leg(distal femoral, and popliteal vein with nonocclusive in right mid femoral vein) and right leg swelling on 13Apr2021; shortness of breath on 17Apr2021 and sharp chest pain on 18Apr2021, all with outcome of Recovered/Resolved with Sequel. Further events description stated that on 13Apr2021 the patient developed right leg swelling; on 17Apr2021 developed shortness of breath; on 18Apr2021 he woke with sharp chest pain. He went to ER and found to have multiple occlusive thrombus in right leg (distal femoral, and popliteal vein with nonocclusive in right mid femoral vein). CT Chest in Apr2021 revealed emboli in bilateral lower lobes, right middle and right upper lobes. The patient was hospitalized for the events for 3 days. Therapeutic measures were taken as a result of the events. The patient had heparin drip and ELIQUIS. The events resulted to Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, Hospitalization, Life threatening illness (immediate risk of death from the event). The events recovered with lasting effects on an unspecified date. Information about Lot/Batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1387768-1	Blood clot caused stroke on right side of the body 4 weeks after the 2nd dose of vaccination; Blood clot caused stroke on right side of the body 4 weeks after the 2nd dose of vaccination; The patient also received the first dose of other vaccine on 18Apr2021.; This is a spontaneous report from a contactable consumer (patient). A 76-year-old female patient received the second dose of BNT162B2 (PFIZER COVID-19 VACCINE, lot number: unknown), at the age of 76 years old, in left arm on 18Apr2021 14:30 at single dose for covid-19 immunisation. Medical history included high blood pressure. The patient was not pregnant at the time of vaccination. The patient was not diagnosed with covid-19 prior to vaccination. The patient received high blood pressure medicine within 2 weeks of vaccination. The patient did not receive any other vaccines within 4 weeks prior to vaccination. The patient previously received the first dose of BNT162B2 (lot number: unknown), at the age of 76 years old, in left arm on 28Mar2021 14:30 at single dose for covid-19 immunisation. The patient also received the first dose of other vaccine on 18Apr2021. The patient experienced blood clot caused stroke on right side of the body 4 weeks after the second dose of vaccination on 15May2021 20:00. The patient was hospitalized for the events blood clot caused stroke on right side of the body for 14 days. The events blood clot caused stroke on right side of the body caused disability. The patient had not been tested for covid-19 since the vaccination. Therapeutic measures were taken as a result of blood clot caused stroke on right side of the body 4 weeks and included treatment with tissue plasminogen activator (TPA) shot, magnetic resonance imaging (MRI) and therapy. The outcome of events blood clot caused stroke was recovering. The outcome of event received other vaccine on 18Apr2021 was unknown. Information on the batch/lot number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388159-1	On March 23, 2021, I began having an extremely heavy menstrual flow, passing clots and requiring protection to be changed at least hourly which lasted approximately three weeks. There has been no period since that time. Previous periods were irregular presumably related to peri-menopause, but were never heavy enough to change protection more often than every 4-5 hours and never lasted longer that 4-5 days.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388169-1	Blood clots in left leg from groin area to knee area. Swollen and painful left leg. Prescribed Eliquis blood thinner. Currently in treatment phase with symptoms basically the same.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388271-1	Blood clot in the leg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388491-1	Pt called pharmacy on 06/10/2021 to report he suffered a blood clot of unknown etiology on 05/16/2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388554-1	The same day after after receiving the vaccine I noticed while I was doing yard work my leg was swollen, and so I shook it off as if it was nothing. Later on I realized that it didn't get smaller my leg started to get bigger. So I called the doctor and they advised me to come in immediately to do an ultrasound. They later on told me that the swelling was due to a blood clot that formed and they prescribed Xarelto 15 mg 2x day and with that prescription I didn't see any improvement so they prescribed me to do 20mg 1x day and I'm just now seeing results now.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1388607-1	Chest pains while working out since receiving the vaccine. Ultimately, two blood clots were formed and caused a cardiac arrest May 25th, 2021.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1389143-1	I am a woman who has PCOS. My cycle is extremely inconsistent and never last more than a day maybe once every year or 2. Two weeks after my 2nd dose of the Covid vaccine I started my menstral cycle , experiencing cramps, discomfort, pressure in my uterus, blood clots daily, and so far it?s been for 8 days. I am not alarmed yet because I understand this might be natural but I am documenting this because it is out of the ordinary in my unique case. I also experienced what seemed like a stomach flu for about 2 days On the 6th and 7th of June. Experiencing diarrhea, nausea, no fever, but heavy fatigue. I no longer feel sick but my period is still active.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1390933-1	She had 2 blood clots in her right leg, and she was now on Eliquis; she had right leg swelling; she had right leg pain; flu; This is a spontaneous report from a Pfizer-sponsored program from a contactable consumer. A female patient of an unspecified age received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration, administered in right arm on 06May2021 (Batch/Lot Number: Not Provided) as a single dose, for covid-19 immunisation, with first dose given 15Apr2021. Medical history and concomitant medications were not reported. The patient said she had the flu 3 days after getting her second COVID-19 Vaccine, had right leg swelling, right leg pain, and had 2 blood clots in her right leg. The patient said she was now on apixaban (ELIQUIS), and financial assistance, due to being on unemployment. The outcome of the events was not reported.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1390945-1	Blood clotting; This is a spontaneous report from a contactable consumer (patient). A 44-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Batch/Lot Number: EW0164), via an unspecified route of administration, administered in arm left on 21Apr2021 at 09:00 (at the age of 44-years-old) as 2nd dose, single dose for COVID-19 immunisation. Medical history included high blood pressure and COVID-19 from an unknown date and unknown if ongoing. Concomitant medication included desvenlafaxine succinate (PRISTIQ) taken for an unspecified indication, start and stop date were not reported. The patient previously received the first dose of BNT162B2 (Batch/Lot Number: FR2613), administered in arm left on 25Mar2021 at 09:30 AM, for COVID-19 immunisation. The patient experienced blood clotting on 16May2021 at 22:00. The patient underwent lab tests and procedures which included sars-cov-1 test: positive on an unspecified date (prior to vaccination). There was no treatment received for the event. The outcome of the event was recovering. No follow-up attempts are possible. No further information is expected. Information on lot number already obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				a possible clot behind her right calf muscle in her right leg; Right eye is now twitching every so often but not constantly; She was afraid that she was going to have a stroke; Even gets headaches; Extreme pressure and pain in her sinus, lymph, right side of her face and her ear; Extreme pressure and pain in her sinus, lymph, right side of her face and her ear; The right side of her face and her head don't feel right; Pain and pressure in her lymph gland on the right side of her throat; Has extreme tiredness and fatigue; Almost felt like she had a possible clot behind her right calf muscle in her right leg. It was a pain near that calf muscle between her knee and the bottom of her heel; She felt slightly dizzy; Slept more; She didn't go back for the 2nd dose because she considers her reaction to be severe; some neurologic side effects; extreme pressure and pain in her right lymph gland, sinus cavities, her eye, her ear, her temple; This is a spontaneous report from a non-contactable consumer (patient herself). A female patient of an unspecified age received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Formulation: Solution for injection, Lot number was: EW0162; Expiry date: unknown) via an unspecified route of administration, administered in Arm Right on 16Apr2021 as 1ST DOSE, SINGLE DOSE for COVID-19 immunization. The patient medical history include anaphylactic shock in 2010 died and was brought back and concomitant medications were not reported. The patient had not taken any other products. She worked in the drug industry for so many years and she did not take medications other than occasionally she would take vitamins or supplements like fish oil or krill oil and maybe vitamin D but nothing on a regular basis and she had not been on any prescription drugs for a very long time. Caller said that she was calling because she got the vaccination on 16APR2021 and she had very severe adverse reactions within 24 hours of getting it. She got her COVID vaccine late Friday afternoon and by Saturday night she had experienced extreme pressure and pain in her right lymph gland, sinus cavities, her eye, her ear, her temple, the right side of her head and her right gland still was not normal. She was trying to avoid going to the doctor because she had a high deductible but she probably would have to go because her right eye was now twitching every so often but not constantly. She really regrated getting the vaccine. She didn't go into a computer system to sign up for her vaccine, someone else did it for her and whatever she agreed to she did not agree to it anymore. She didn't go back for the 2nd dose because she considers her reaction to be severe because she had been extremely healthy for the past few years and not needed to go to the doctor for hardly anything so she was afraid that she was going to have a stroke. Now, over a month and a half later her right gland did not feel right. She even got headaches and the right side of her face and her head did not feel right. She did experience an anaphylactic shock in 2010 died and was brought back, no further details provided. When she got her shot, they kept her for 1/2 an hour because of this. She felt slightly dizzy but for the most part she seemed okay but 24 hours later she literally was curled up and she thought that she was going to die. It was extreme pressure and pain in her sinus, lymph, right side of her face and her ear. Everything began approximately 24 hours after the COVID vaccination except for the eye twitching, that only started about a week ago and it was not constant but she had never experienced anything like that in her life before and it had happened a dozen times in the past two weeks.She was concerned that she might have some neurologic side effects. She said all of her other side effects have subsided somewhat but the real persistent side effect was the pain and pressure in her lymph gland on the right side of her throat. That had been pretty persistent since she got her vaccine, as far as the others, they came and went. She also had extreme tiredness and fatigue. She had slept more since she got the COVID vaccine than normal and she woke up and was awoke for a few hours and then she was tired again and wanted to sleep. The other day she probably slept for 16 out of the 24 hours that day which was quite excessive, it was not normal. This was not constant though. The tiredness was constant. She was sleeping more than she was used to sleeping or that she ever had. In the recent past she did not normally sleep as much. She said that the eye twitching didn't occur today, it occurred yesterday and the day before but she had never experienced anything like that before. Her 2nd dose was supposed to be 07MAY2021 . Another thing that she had that was highly unusual was that she almost felt like she had a possible clot behind her right calf muscle in her right leg. It was a pain near that calf muscle between her knee and the bottom of her heel and it was pretty persistent for few days as well. She didn't bump it on anything. She used a massage cream to massage it out which helped a little bit. It was a little bit strange. The outcome of all events was unknown. followup attempts are possible, additional information has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1391000-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1391010-1	Stroke; she had blood clots on her brain; headache; her left leg was numb like it was asleep; blurry vision; This is a spontaneous report from a contactable consumer (patient). A 27-year-old female patient received second dose of bnt162b2 (BNT162B2, solution for injection, Lot Number: ER8736) in left arm via an unknown route of administration on 24Apr2021 at 10:10 (at the age 27-year-old) as single dose for covid-19 immunization. The patient medical history and concomitant medications were not reported. The patient previously received the first dose of bnt162b2 (BNT162B2, solution for injection, Lot Number: ER8727) right arm, via an unknown route of administration on 31Mar2021 as single dose for covid-19 immunization and experienced no adverse event. On 14May2021 at around 07:00 or 08:00, the patient experienced stroke. The patient was hospitalized for stroke from 15May2021 to May2021. On an unknown date, the patient had blood clots on her brain, headache, her left leg was numb like it was asleep, blurry vision. It was reported that patient was recently got her vision back to where she can stand the light long enough to do the research and call around. The stroke happened on 14May2021 and patient went to the hospital on 15May2021 for 5 days because she thought it was just a headache. The patient cannot find the exact date she was discharged. The vision was the first thing that went, then headache started. When patient woke up early in the morning her left leg was numb like it was asleep. Patient used the restroom and did her morning routine, and when she sat back down had the worst headache of her life. They said she had blood clots on her brain. When patient got there, they couldn't stop it because it had already happened and the damage was done. It was reported that patient knew she was going to be travelling and would start working around children. Patient was exposed to a lot of people and thought at the time it was the best thing to go ahead and get the vaccine to go back to a new normal. Patient had bloodwork test and drew so much blood trying to figure out how but results showed unknown. The most common stroke was a blood clot on the brain. Everything matches from the blurry vision to the headaches. Patient declines to report on any of the people she read about, she is just calling to report on herself. The stroke was early because her aunt had a stroke literally the week before and her wake was in the morning. She was getting ready for that. The patient had no other vaccinations within four weeks prior to the first administration date of the suspect vaccines. Patient's lab test included blood work: unknown on an unspecified date. The outcome of the events was unknown. PSCC Communication was as provided contact information for Pfizer Legal Department and report reference number to the caller. Caller declined to complete reports on the people she read about online. Information on Lot/Batch number was available. Additional Information has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1391022-1	Incarcerated and ischemic bowel due to massive thrombosis of portal vein and SMV; Incarcerated and ischemic bowel due to massive thrombosis of portal vein and SMV; This is a spontaneous report from a contactable consumer. A 59-year-old female patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on 23Apr2021 (Batch/Lot Number: ER8736) as single dose at the age of 59-year-old for COVID-19 immunisation. Medical history included pre-myelofibrosis aspirin on daily baby. No known allergies. Concomitant medication(s) included acetylsalicylic acid (BABY ASPIRIN) taken for an unspecified indication, start and stop date were not reported. The patient received bnt162b2 (BNT162B2), dose 1 on 02Apr2021 (Batch/Lot Number: ER2613) as single dose at the age of 59-year-old for COVID-19 immunisation. The patient had not received any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient experienced incarcerated and ischemic bowel due to massive thrombosis of portal vein and SMV (hospitalization, disability, life threatening) on 09May2021 with outcome of recovering. The adverse events resulted in any Doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care. The patient was hospitalized for the events for 26 days. The patient underwent lab tests and procedures which included Sars-cov-2 test: negative on 24May2021 Nasal Swab. The treatment received for the adverse event was emergency bowel resection. Since the vaccination, has the patient been tested for COVID-19. No follow-up attempts are possible. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1391968-1	Started getting pain in my right leg on May 1st, seen my Family Dr. on 4/29/21 clean bill of health. Pain wouldn?t go away was going to my chiropractor the whole month of May 2 times a week, for my right leg, I favored that leg never missing work, I?m a childcare provider. On 31May2021 the pain went down my leg and behind my knee just throbbing, I was walking up steps and a loud snap happens in my leg dropped me right there couldn?t move. Was taken to the ER, where they found a blood clot in my leg which caused pressure to build up on my knee which ruptured my meniscus, damaged joints, and cartilage in my knee. They put me on blood thinners, have to have surgery and can?t for 3 months I have to be off blood thinners. Can?t work and was forced to close my daycare business of 18 yrs and sell everything due to this injury. My Dr has linked it to the vaccine my blood wok showed high levels of hormones. Never ever had an issue until I got second shot. So now I?m trying to get disability, not looking good due to me being self employed. So who is gonna step up and help me financially, I was open threw the whole pandemic and this is how I go out of Business! Devastating to say the least?. My Community is shook up over this, so I took one for the team my family refuses to get the vaccine and others have decided NO after hearing my story.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1392423-1	Blood clot in left leg. Currently on blood thinners to address it. I have no history of blood clots before this,
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1392762-1	Extreme fatigue, heart pains, dizziness,foggy head, heart attack, blood clots and the list goes on
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1394086-1	Seriousness: Does not really know what it is, it could be thrombosis; Caller has experienced redness/redness in thumb and finger tips; tingling/tingling in left hand; decrease sensation in his finger tips/numbness and decreased sensation in left hand; He is still experiencing all of these and has gotten worse; This is a spontaneous report from a contactable physician (patient). A 53-year-old male patient received bnt162b2 (BNT162B2), dose 1 via an unspecified route of administration, administered in Arm Left on 13May2021 (Batch/Lot Number: EW0183) in the morning between 10:00 and 11:00 am as single dose for covid-19 immunisation at the age of 53-year-old. Medical history included an open reduction internal fixation, surgery on his right clavicle 5 years ago, which is what made him want to get the vaccine. Concomitant drugs included Multivitamin, Vitamin A and Vitamin D. The patient had an exam in December and all of the test results were normal. On 16May or 17May2021, the patient has experienced redness in thumb and fingertips, numbness, and decreased sensation in left hand, tingling in left hand. Symptoms have been ongoing for 10 days; it has progressively gotten worse. It was reported that the symptoms started 2 days after getting the vaccine on 13May2021, was 16May2021 or 17May2021. He takes multi vitamins and minerals and thought it may be related to that. He takes high doses of Vitamin A and Vitamin D. He actually thought it was due to the Vitamin A and Vitamin D. He stopped the vitamins for a few days. Treatment included: Thought maybe, apart from the vitamins, it may be because he was washing his hands so much that it may be from that. He used Aveno lotion to try and lubricate them. The outcome of event Thrombosis was unknown, outcome of other events was not recovered. The reported considered the events were related to the vaccine as it was the left finger tips, especially the thumb, compared to the right. That's why he thinks it is related to the vaccine. The clinical course was reported as: The redness was first, then the tingling. He doesn't remember the exact day. The numbness and decreased sensation were afterward. He is still experiencing all of these and has gotten worse. The thumb on the left hand looks pink and wrinkled. When comparing the right see the difference tell they are different. Seriousness: Does not really know what it is, it could be thrombosis. The redness in the thumb and finger tips is in the right as well but is not as bad as in the left. Follow attempts needed. Further information is expected.; Sender's Comments: Based on plausible dose -event relationship post vaccination and no other alternate explanation the causal role of BNT162B2 cannot be excluded for the reported event of thrombosis. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to regulatory authorities, Ethics Committees, and Investigators, as appropriate
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1394144-1	after I got my 2nd vaccine I was diagnosed with blood clot; This is a spontaneous report from a contactable consumer (patient). A 74-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Batch/Lot Number: EW0173; Expiration Date: 31Aug2021), via intramuscular route of administration, administered in Arm Left on 20May2021 13:00 (at the age of 74-year-old) as 2nd dose, single dose for COVID-19 immunisation. Medical history included Blood pressure from an unknown date and unknown if ongoing. Concomitant medication included nebivolol hydrochloride (BYSTOLIC) taken for blood pressure, start and stop date were not reported. The patient previously received first dose of BNT162B2 (Lot number: EW0172; 74-year age at time vaccination) intramuscular, administered in Arm Left on 29Apr2021 (approximately 1pm afternoon) for COVID-19 immunisation. Patient stated, I do not know if it was related to the vaccine, but I thought it was interesting that 2 weeks to the day after I got my 2nd vaccine I was diagnosed with blood clot (28May2021), and I never had a blood clot I am on blood thinners now, I just said that was interesting I don't know where it came from I had no injury, I am very active. there was no reason for me to have the blood clot other than I do get the vaccine 2 weeks prior to the diagnoses. Again stated, I got the vaccine on 20May, I started noticing it on 28May. Patient stated, I did not go to the hospital to get diagnosed I had ultrasound. Patient stated diagnosis date of blood clot was 03Jun2021 and Eliquis (each pill is 5 milligrams and I have to take 4pills a day) used for blood thinner. The patient underwent lab tests and procedures which included ultrasound: blood clot on 03Jun2021.Therapeutic measures were taken as a result of after patient got my 2nd vaccine i was diagnosed with blood clot. The outcome of event was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1395109-1	Sharp pain in left leg toe for 30 minutes then a bumpy feeling. Next day morning showed up a bruise kind of spot (blood clot). No numbness, itchiness or pain in that area.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1396010-1	"Hemoptysis. On 06/05, a one-time event (old, dark clots) occurred. On 06/06, there were several instances, some older clots, some lighter/fresher blood. Then 4 days of no blood, but ""normal"" (usual) infected sputum colors (preexisting lung infection; see continuation page). Then on 06/11 beginning @ 7:57am, fresh (light) blood streaks/tinged sputum, which occurred a number of times over the span of approximately 5 hours, until ""normal"" pale yellow infected sputum was expectorated. Then on 06/12 beginning @ 9:30am, fresh blood again, which again occurred a number of times over the span of about 5 hours until the usual infected colors of sputum were expectorated. Additional information for Item 18: I have a history of MAI/MAC infection and resultant hemoptysis since 2006. However, until the first Pfizer dose, it had been 2 years since the last incident of hemoptysis, which was due, at least in large part, to malnutrition spanning a couple of months, for reasons unrelated to the lungs. I believe my most recent hemoptysis episodes are a result of my immune system being distracted by the Covid-19 vaccine, thus allowing the Mycobacterium avium (and/or other) to more effectively attack my lung tissue. There was no hemoptysis yesterday (06/13), but this has me concerned about the second dose causing additional, possibly worse, problems for my immune system, as it needs to be concentrating on my lung infection full-time; it has been keeping the MAC (and/or other) generally under control, assuming hemoptysis is a good indicator, which my X-rays over the years of monotherapy with Ethambutol suggest is the case. I discontinued Ethambutol on 06/15/2012, with no further hemoptysis until 04/26/2019, and then no hemoptysis until 11 days after the first Pfizer Covid-19 dose. It seems reasonable to conclude that the vaccine led to these latest hemoptysis events and perhaps allowed the preexisting bacterial infection to cause additional lung damage.The physician noted above is my current pulmonologist. I was to have follow-up lung CT scan and lung function tests, but did not do so in 2020 because of Covid-19 concerns. I had planned to see Dr. and my other doctors after being fully-vaccinated, but since lung function tests cannot be performed on patients with active/recent hemoptysis, this may cause an even further delay in my obtaining appropriate diagnostics and treatment."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1396536-1	53 yo woman with no PMH was diagnosed with COVID-19 by PCR on 5/7/21. She had a mild illness with cough and fevers but did not require hospitalization. Her symptoms lasted about 1 week. She got her first dose of Pfizer COVID vaccine on 5/29 around 11 am. Within 4 hours (around 3 pm) she started to develop extreme fatigue, which she attributed to vaccine. This was followed by nausea, vomiting and diarrhea later that same day. The symptoms got worse on 5/30. Her diarrhea/GI symptoms resolved on 5/31. That day, she had a fever to 101.3. She continued to have fatigue, and started to have dyspnea, and presented to hospital on 6/1/21. On arrival to ED her vitals were 88/41, 36.1, 125, 18, 95% on RA (13:22). She become more hypotensive with nadir BP 63/48, HR 109 at 17:30, at which time she was started on norepinephrine. Physical exam at admission was notable for clear lungs, tenderness to palpation in the midepigastric and LUQ areas, and the absence of rash. Admission labs: WBC 31.3, Hb 14.2, Hct 43.0, Plt 144, MCV 96, 85% neutrophils, BNP 3117, Na 132, K 3.8, CO2 19, Cl 95, BUN 35, Create 4.62, gluc 133, protein 6.1, alb 2.2, TB 5.5, DB 2.9, Alk phos 159, ALT 47, AST 33, lactate 4.2, troponin I 0.047, random cortisol 93. Strep pneumo and legionella antigens negative. Hepatitis serologies, HIV test, all negative. COVID test by PCR was positive on 6/1/21. She was admitted to the ICU. The day following admission (6/2) CRP 32.4, ESR 106, LDH 399, ferritin 1626, procalcitonin 54.85. CT A/P unremarkable except for bilateral patchy airspace opacities. RUQ US unremarkable, including Dopplers. Echocardiogram (TTE) with LV systolic function mod-severely globally reduced, EF 35%, no valvular disease. Dopplers of LE showed small non-occlusive thrombus in R common femoral. The patient was seen by Infectious Diseases on HD#2 (6/2/21) and IVIG was recommended and administered due to suspicion for MIS-A. She had also been started on cefepime, vancomycin and metronidazole on admission. Blood, urine and resp cultures were negative. She continued to have high norepinephrine requirements, reaching 0.34 mcg/kg/min on 6/3. Due to ongoing shock (cardiogenic vs septic), a Swan-Ganz was placed on 6/4/21. RAP 11 mmHg, RV 38/10 mmHg, PA 40/20 (27) mmHg, PCWP 19 mm Hg, CO 5.75, CI 2.8, SVR 1100, MvO2: 70%. These were consistent with distributive/septic shock rather than cardiogenic. The patient was able to be weaned from pressors on 6/5/21 at 11:55. Throughout her entire hospital stay, she was comfortable with no respiratory distress, and no abnormalities in mentation. She was transferred to the general medical floor and completed 7 days of antibiotics for empiric sepsis treatment. Labs at discharge on 6/8 showed procalcitonin 5.43, CRP 8.8, LDH 362, Ferritin 1511, TP 5.9, ALT 17, AST 29, Alk phs 158, TB 1.3, Alb 1.6, Create 0.98, WBC 32.5, Hb 7.8, Hct 22.9, Plt 100, neutrophils 70%. She was feeling well without complaints, and will follow up as an outpatient. She was prescribed Xarelto for DVT.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1396655-1	large blood clot in right leg/groin, xarelto po, consult hematologist and vascular surgeon, still waiting for treatment to resolve blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1396688-1	Had some crazy nosebleeds beginning 4 days after my first dose. I ended up pulling a gigantic blood clot out of my nose during the first one. It was terrifying. I?ve had 6 nosebleeds since my first dose and I?ve never had issues with nosebleeds before. They last a long time too and each one comes with a clot. I also have bruises all over my legs and that have appeared randomly and if I get cut I take forever to stop bleeding.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1398544-1	"Had thrombosis, Blood clots in leg; This is a spontaneous report from a contactable consumer (patient). A 67-year-old female patient received bnt162b2 (Lot Number: EN6198; Expiration Date: Jun2021), dose 1 via an unspecified route of administration, administered in arm left on 18Mar2021 at the age of 67-year-old as single dose for covid-19 immunisation. Medical history included diabetes. The patient's concomitant medications were not reported. Three days after vaccination, the patient had thrombosis, blood clots in her leg and had gone to the Emergency room on 21Mar2021. It started at night but she didn't know what it was, may be the same day she got vaccine (18Mar2021). The patient was taking ""blood thinners"" (medication name not clarified) for the Blood clots. The patient underwent lab tests and procedures which included ultrasound: unknown results on an unknown date. The outcome of event was unknown."
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1398569-1	Blood Clots in left leg; This is a spontaneous report from a contactable 74-year-old male consumer (patient) reported for himself that he received his second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Batch/Lot number: ER8733) via an unspecified route of administration into the left arm on 07Apr2021 (at the age of 73-year-old) as single dose for COVID-19 immunisation. The patient medical history included blood pressure and blood disorder. Concomitant medications included lisinopril for blood pressure and baby aspirin to keep his blood thin. He started taking lisinopril 10-15 years ago. His blood pressure was under control with it. He stated he has been taking baby aspirin for years. Doctor advised him to quit taking it while he was taking Eliquis. He stopped taking it on 20Apr2021 when he was diagnosed with blood clots. The patient previously received his first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Batch/Lot number: EN6205, Expiry date: unknown) via an unspecified route of administration into the left arm on 17Mar2021 as single dose for COVID-19 immunisation. He stated that on 07Apr2021 he received his second shot. On 20Apr2021, the patient reported of blood clots in left leg that showed up after the Pfizer Covid vaccine which was continuing. Never had a blood clot in his life until now. On 20Apr2021, he went to the doctor and was told he had blood clots in left leg. His doctor sent him to the emergency room for testing. The doctor told him he was on blood thinners until August and put him on Eliquis. On 20Apr2021, test results showed sonogram positive for blood clots in left leg. He was not admitted to the hospital as there were no beds available. They were full and kept him there for about 6-8 hours and sent him home. Report was not related to a study or programme. The patient took Pfizer Covid-19 Vaccine because of COVID-19. Investigation assessment done. Patient provided age at time of blood clot diagnosis as 74. No further information provided. No additional vaccines were administered on same date of the Pfizer suspect nor any prior vaccinations within four weeks. Outcome of the event was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1398577-1	COVID-19 positive after vaccination.; COVID-19 positive after vaccination.; Had a clot and thrombectomy; This is a spontaneous report from contactable other-HCP (patient). A patient of an unspecified age and gender received bnt162b2 (PFIZER-BIONTEC COVID-19 VACCINE, Solution for injection, Lot Number: Unknown) via an unspecified route of administration on unspecified as a single dose for COVID-19 immunization. Patient's medical history included thrombectomy and concomitant medication was not reported. On an unspecified date, patient had COVID-19 positive after vaccination and had a clot and thrombectomy. Treated with Xaralto and had another clot. Event took place after use of product. The outcome of events was unknown. Information about lot/batch number has been requested.; Sender's Comments: Based on the limited information currently provided the lack of efficacy of the administered vaccine BNT162B2 and also the causal association between the reported event clot with the usage of the vaccine cannot be excluded. The contributory role of the patient's medical history needs to be taken into consideration. The case will be reassessed once new information is available. The impact of this report on the benefit/risk profile of the Pfizer product is evaluated as part of Pfizer procedures for safety evaluation, including the review and analysis of aggregate data for adverse events. Any safety concern identified as part of this review, as well as any appropriate action in response, will be promptly notified to RAs, Ethics Committees, and Investigators, as appropriate.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1399489-1	Chest pain and head ache was taken to the ER on 4/16/21
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1399986-1	Shortness of breath , chest pain , afib placed on blood thinner administered oxygen . Admitted to intensive care. Blood clots in both legs and both lungs
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1400337-1	On May 18, 2021, Patient said he was experiencing severe back pain and that both of his arms had gone numb and he had vomited. Patient called his mother to come help him with the baby because he wasn't feeling well. When she got there 30 minutes later he was not breathing. They were not able to revive him. He died of a blood clot in the front of his heart.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1402683-1	large blood clots, minor chest pain

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1403576-1	I developed pain up and down my right arm quite suddenly, basically from the shoulder all the way in to my wrist and fingers off and on, it got progressively worse it was very painful, I went to the doctor and they ordered an ultrasound to ruled out a blood clot. They had to do the ultrasound on the ER and they found a clot in my basilic vein. They gave me a pain med so I could sleep and add heat and Ibuprofen. I am being referred to another doctor because the pain is coming back and they are going to do more tests. I have donated blood about a month before I had the vaccine and the ER doctor thought this may be because of that.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1403597-1	After receiving the vaccine I had the soreness in my arm and a small fever of 99. Over the course of the next few weeks I was experiencing severe eye pain which caused me to start losing site in my right eye and also microclots in my eyes, my eye doctor stated that the vaccine was not the cause of my conditions with my eye. But other than the soreness of my arm where the vaccine was administered and a slight fever I really didn't have any other problems.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1404274-1	Death, sudden heart attack or blood clot
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1405801-1	Blood clot in the right lower leg; This is a spontaneous report from a contactable consumer (patient). This 58-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) second dose at the age of 58-year-old (lot number=EN 6201) on 12Feb2021 at 12:30 PM as single dose on Left arm for COVID-19 immunization. Prior to vaccination, the patient was not diagnosed with COVID-19. Medical history included Diabetes, CAD, arthritis and known allergies Yes. The patient was not Pregnant at Time of Vaccination. Concomitant medications were not reported. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient received other medications within 2 weeks of vaccination. The patient previously took bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) first dose at the age of 58-year-old (lot number=EL1283) on 22Jan2021 at 12:30 PM as single dose on Right arm for COVID-19 immunization. The patient experienced adverse event of Blood clot in the right lower leg on 10Mar2021 at 04:00 PM. The adverse event result in Doctor or other healthcare professional office/clinic visit. The treatment of Blood thinner received for the adverse event. Since the vaccination, the patient has not been tested for COVID-19. The outcome of the event was Unknown. No follow-up attempts are needed. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1405811-1	blood clot; high levels of d dimer in blood; very dizzy; can barely walk; migraine; fatigue; chills; temperature changes; pain in body switching from left to right and vice verse; chest pain; shallow breathing; This is a spontaneous report from a non-contactable consumer. This consumer (patient) reported that a 29-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) at the age of 29-years, via an unspecified route of administration on 20May2021 at single dose for COVID-19 immunization, in hospital. Medical history was not reported. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. The patient was not pregnant at time of vaccination. Concomitant medications included gabapentin, lorazepam (ATIVAN), norethisterone (NORETHINDRONE), omeprazole. The patient previously received all antibiotics except for doxycycline. The patient went to hospital to check for blood clot; high levels of d dimer in blood; very dizzy, can barely walk; migraine; fatigue; chills and temperature changes; pain in body switching from left to right and vice verse; chest pain and shallow breathing; serious side effects 1 week later. The events occurred on 27May2021. The adverse events resulted in doctor or other healthcare professional office/clinic visit, resulted in emergency room/department or urgent care. Outcome of the events was not recovered/not resolved. Since the vaccination, the patient has not been tested for COVID-19. No follow-up attempts are possible; information about lot/batch number cannot be obtained

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1405814-1	hospital with AFib; Has clots in both legs/Developed one blood clot in arm; fell; Has clots in heart, in multiple areas of heart; This is a spontaneous report from a contactable consumer (patient's daughter). A 75-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on 07Apr2021 (Batch/Lot Number: ER8733) as 2nd dose, single for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration in arm on 10Mar2021 (Lot Number: EN6204) for covid-19 immunisation. The patient in hospital with AFib on 21May2021. The patient fell on 17May2021, it was unsure if the patient fell on 17May2021 from the AFib. The patient had been hospitalized since 21May2021. It was unknow if the reason the patient fell was because the patient went into a different rhythm. The patient got sick and got AFib. The AFib was still ongoing and the patient was still hospitalized. The patient had clots in both legs, it began when the patient was hospitalized on 21May2021. The patient had no history of blood clots. The patient had clots in heart, in multiple areas of heart. All the blood clots were acquired while the patient was in the hospital. The patient developed one blood clot in arm on 31May2021 after being on apixaban (ELIQUIS), the patient acquired a clot in one arm. The other clots started on 28May2021 and 29May2021. In past weekend the patient went in to get a TEE and they could not perform the TEE because the callers mother had blood clots in her heart. Reporter seriousness for events hospital with AFib and clots in both legs/clots in heart, in multiple areas of heart/developed one blood clot in arm was unspecified. The outcome of events was unknown. No follow up attempts are possible. No further information is expected.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1405827-1	Did Xray found a big black spot on top of left side of stomach. dr said it was blood clot from the vaccine.; Severe stomach pain; This is a spontaneous report received from a contactable consumer (patient). This 68-year-old male patient received bnt162b2, dose 2 via an unspecified route of administration on 19Mar2021 (Batch/Lot number was not reported) as 2nd dose, single at the age of 68-year-old for covid-19 immunisation. The patient received the 1st dose of bnt162b2 on unknown date for covid-19 immunisation. The patient medical history and concomitant medications were not reported. The patient experienced severe stomach pain. Did Xray found a big black spot on top of left side of stomach. Dr said it was blood clot from the vaccine. Happened week after taking shot. The product was used by the patient. The outcome of event was unknown. Information on the lot/batch number has been requested.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1405835-1	Blood clot left leg development; This is a spontaneous report from a contactable consumer (patient). A 69-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot Number: EN6200), intramuscular at the shoulder/upper arm on 19Feb2021 11:30 (at the age of 69-years-old) as second dose, single for COVID-19 immunization. Medical history included diabetes and hypertension from 2012 and ongoing, and hypothyroidism from 1980 and ongoing. Concomitant medications included metformin taken for diabetes from 2012 and ongoing; thyroid (ARMOUR THYROID) taken for hypothyroidism from 1980 and ongoing; clonidine taken for hypertension from 2019 and ongoing; and liothyronine sodium taken for hypothyroidism from 2015 and ongoing. The patient had no prior vaccinations. The patient previously received BNT16B2 (lot number: EL9265), intramuscular at the shoulder/upper arm on 29Jan2021 11:30 (at the age of 69-years-old) at first dose, single for COVID-19 immunization and experienced severe pain. The patient experienced blood clot left leg development on 08May2021. The event was reported as serious due to persistent/significant disability/incapacity and required physician office and emergency room visit. Therapeutic measure taken as a result of blood clot left leg included Eliquis. Outcome of the event was not recovered.; Sender's Comments: Linked Report(s) : US-PFIZER INC-2021218434 same patient/drug, different dose/events

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				The heart rate was 245; A blood clot in the caller's neck; It was thought he might have lymphoma; Felt like he had a golf ball in his left arm/that little golf ball in his arm started traveling to his whole body- his back, his chest, legs, calves; Sweating; Didn't feel too great; Breathing was off/trouble breathing; Throat was hurting/sore throat; Neck was swollen; This is a spontaneous report from a contactable consumer. This consumer (patient) reported for himself that: A 38-year-old male patient received first dose of BNT162B2 (Solution for injection; Lot number: ER8131 or ER8231 and Expiration date were not reported) via an unspecified route of administration, administered in left arm on 18Apr2021 (at the age of 38-year-old) as first dose, single dose for COVID-19 immunization. The patient medical history and concomitant medications were not reported. On 18Apr2021, he went to get the Covid shot through name withheld which Pfizer sponsors and it was done at name withheld. He went there and took the shot. He was told to wait 15 minutes and then left. When he got to car, immediately he felt like he had a golf ball in his left arm. The next couple of days, that little golf ball in his arm started traveling to his whole body. He felt it in his back, his chest, he felt it in his legs. Literally, in the next 3 days, he felt it literally travel through his whole body. He felt it in his calves, it was crazy also mentioned they felt sore or like ball in the arm. He started sweating and didn't feel too great. He thought it was standard side effects and didn't think much of it. A couple of days past by and then Thursday morning, he woke up and his breathing was very off. He clarifies his throat was hurting. He thought it was a sore throat and took some cough medicine and cough drops. Then the next day, it wasn't going away and was still hurting, wasn't getting any better. His breathing was very rough. If he was just standing still like laying down, then felt okay. But if went to move or do something it was very difficult. Again, expecting these symptoms, some people have said these symptoms and he didn't' think much of it. The next morning patient woke up and breathing felt fine. His neck was swollen. He thought maybe he had a sore throat or something. He was taking cough drops to soothe his throat the following day, the throat was still hurting, and it was the same. The patient called the doctor and did a telehealth. The patient told the doctor when he received the first dose Pfizer Covid Vaccine and all his symptoms. The doctor said it is a little unpredictable, nothing the doctor hasn't heard. He went to the doctor office the next day and was hooking to a nebulizer for breathing and prescribed an asthma pump. The doctor thought patient had Lymphoma in the neck. The doctor wanted to go the following day for an ultrasound. He went to the ultrasound and they checked the neck and immediately there were red flags. His heart rate was checked, and the heart rate was 245. The patient was hospitalized from 27Apr2021 to 08May2021. The minute they saw heart rate and started putting intravenous and received treatment to low down heart rate. The heart rate went down to 170 or 165. He had x-rays and a CAT scan done. He had fluids in his heart and fluid in the lung and a blood clot in neck. He had a hole cut open on left side to lungs and the lungs were drained. He went back to the ICU, they ran another CAT scan and X-ray and gave the medication. He does not know if this medication was antibiotics or what the medication was. The CDC received this information because the patient case was so rare. When the CAT scan and X-rays were run again, the fluid is right back up in the lungs. Then he was taken to surgery for the heart. An incision was cut from where itches is to back on the side, a very big incision. The heart was gone into to cut a little piece around the heart. Something was cut out so the heart could drain, and fluid could get out of the heart. While all of this is happening, he wasn't getting any better, was getting worse. The patient health deteriorated, breathing was about 10 percent and was hooked up to a breathing machine. He was sedated at that time and not awake anymore. The machine was keeping him alive. Now, every day for basically five days he was asleep and not knowing what was going on and was tested for Leukemia, cancer, and A to Z every disease you can think of to try to find common ground. Everything came back negative; nothing was wrong with the caller. He was given this or that medication every day and was sedated for about 5 days. Every day of the sedation was getting worse. He was dying and time was ticking. On the 5th day, everything was getting worse. The woman kicked everyone out of the room and went into his mouthpiece on the ventilator that was keeping the alive doing the breathing for the patient. The woman pulled something out of his lung. The woman started pumping him with steroids. What the woman took out of his lung stopped the fluid from going in. The fluid wouldn't fill back up in the lung. Once the woman started giving the steroids, the steroids killed the Pfizer Covid Vaccine. The steroids reversed the Pfizer Covid Vaccine. He was started breathing on his own again. After a couple of hours, the patient finally woke up. The patient is now recovered and has been home the past few weeks on a whole bunch of medication. He is on blood thinners, steroids, and stomach stuff. The outcome of the events was unknown. Information on Lot/Batch number was available. Additional information has been requested.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1405838-1	
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1406787-1	8 weeks after second shot, I developed 2 superficial blood clots in my left leg. At the time, I was taking a daily low-dose of aspirin as recommended by my doctor for people my age. I experienced a little over a week of pain before the clots dissolved. I have no history of blood clots.
THROMBOSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1406917-1	Had a large boil at the shot site, developed blood clots in arm which caused alot of pain.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1407072-1	Stroke from blood clot 4 weeks after the second shot.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1407223-1	Heart stent 4/26/2021 Blood Clot left leg diagnose 5/13/2021 Xarelto 20 mg
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1407228-1	Beginning the month after the first injection, my menstrual cycle became extremely heavy, with a large volume of very big clots. I have never experienced this in 20 years of having a period. My cycle went from 4 days to 7, and I have to wear an adult diaper during these days now. I have had bloodwork and an ultrasound, which showed increased thickness in uterine lining. Now having to have a biopsy done. This has continued with every cycle following my first injection.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1408197-1	I started experiencing pain in my calf on 4/21 that same night of the vaccination. There was a tight feeling in my knee and every time I took a step with my right leg there was pain that radiated down my calf. It almost felt like a muscle pain so I ignored it for a few days. By 4/24 the pain had gotten so bad that I didn't want to walk anywhere and cried when I walked. My mom came and picked me up and took me to the urgent care center. They asked me a few questions and told me that I needed to go to the ER to get checked for a blood clot. I went to hospital on 4/24, and an ultrasound of my knee was done. The doctor diagnosed me with a blood clot and told me to stop taking the Xulane which was my birth control at the time. They told me to follow up with my OBGYN and a hematologist. (I had been on my birth control for about 6 months and I have never had any issues with blood clots and they do not run in my family. The NP is who told me to report my blood clot to the CDC because she is a little suspicious that the shot is where it come from.)
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1408210-1	Received second dose above on 1/8/21. Developed a blood clot in left leg early February, Went to doctor after a week or so on February 8. Bad Swelling in left calf

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1409717-1	she had pain in her knee; it was hard for her to walk; blood clot found in right leg; This is a solicited report from a non-Pfizer sponsored program, PSP-ELIQUIS-CONDUENT-USA: PSP-ELIQUIS-CONDUENT-USA-CW3050920, from a contactable consumer, based on information received by Pfizer from Bristol-Myers Squibb (manufacturer control number: US-BRISTOL-MYERS SQUIBB COMPANY-BMS-2021-051474). This case was split from master case 2021623975 for Covid-19 vaccine with she had pain in her knee and it was hard for her to walk. An 88-year-old elderly female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Solution for Injection, Batch/Lot number was not reported) via an unspecified route of administration on an unspecified date at an unknown, single dose for covid-19 immunisation. The patient medical history was not reported. The patient had no allergies. Concomitant medication included apixaban (ELIQUIS) (batch no. 1798944, expiration date: SEP 2023) via oral route at 5 milligram, twice a day taken for venous thromboembolism from an unspecified date in Jan 2021. On an unspecified date she had pain in her knee, it was hard for her to walk and on an unspecified date in Jan2021 the patient experienced blood clot found in right leg. The patient underwent lab tests and procedures which included blood pressure measurement: normal. This 88-year-old female patient was involved in a patient support program described. The patient received APIXABAN. The report describes a case of HOSPITALISATION. The occurrence of additional non-serious events is detailed below. In January 2021, the patient started APIXABAN (batch no. 1798944) (Oral), (5 milligram, twice a day). On an unknown date, the patient underwent hospitalisation (seriousness criterion hospitalization), OEDEMA PERIPHERAL (Swelling in her feet/fluid in her feet) and arthralgia (Bad knee pain). The patient was hospitalized for 2 days due to Hospitalisation. The action taken with apixaban(oral) was unknown. At the time of the report, hospitalisation and oedema peripheral outcome was unknown and arthralgia did not resolve. The patient reported that she was in hospital for two days. The consumer stated her feet swells up. After receiving the coronavirus vaccine she had pain in her knee and it was hard for her to walk. Patient was inquiring if she can take green leafy vegetables with Eliquis. Patient started on Eliquis for a blood clot found in her right leg in Jan-2021. Patient reported that she believes the blood clot was there longer because her feet would become swollen. Patient re-ported that her feet were swollen today however, it will go away tomorrow. The swelling will go away for three or four days, sometimes up to a week. Patient reported that there was nothing all winter, the swelling was down. There was increased swelling with the warmer weather. Doctor told her the blood clot should go away. She went back to the doctor because she still had swelling. The doctor told her she has fluid in her feet. Her doctor told her to follow up with her primary physician because she never had a blood clot before and she should have her medications checked. Patient reported that after receiving the vaccine she can run but she still feels pain in her knee. The strength of apixaban was 5 mg For APIXABAN(Oral), the reporter did not provide any causality assessments. The primary reporter was the consumer Most recent follow up information received from the consumer incorporated above includes: 26-May-2021: Event Hospitalisation was added, Event verbatim was amended from Swelling in her feet to Swelling in her feet/fluid in her feet, Covid-19 vaccine was changed from suspect to concomitant drug, product attributes, event attributes, patient demographic details and narrative updated. BMS Medical evaluation comment: This patient had hospitalized for unknown reason after apixaban therapy. Based on the limited information regarding hospitalization details, laboratory reports, treatment details, it cannot be ascertained that the suspect drug contributed to the reported event. Outcome of the events was unknown.
THROMBOSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1410234-1	Blood clot?> loss of consciousness-> artificial coma ?> death 10 days after vaccination
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1152079-1	Patient developed calf pain in the morning of 3/18/21, pain for 2 days and then the pain went away. Right lower leg started swelling on or around the 19th and patient went to the urgent care on the morning of 03/22/21. Patient was sent to imagining center to have venous doppler where they in turn sent her to the ER after having a positive venous doppler study. Blood clot noted behind right knee.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1167457-1	clot; swelling and pain in foot / foot was swollen/swelling and pain in calf; swelling and pain in Ankle/ Ankle was swollen; pain in foot, calf and anckle / foot was sore; pain in foot, calf and anckle / foot was sore; This case was reported by a consumer via other and described the occurrence of clot blood in a female patient who received Herpes zoster (Shingles vaccine) for prophylaxis. Co-suspect products included Flu Seasonal QIV Dresden (Influenza vaccine Quadrivalent unspecified season) for prophylaxis and COVID 19 VACCINE for prophylaxis. Concomitant products included COVID 19 VACCINE. In November 2020, the patient received Shingles vaccine. On an unknown date, the patient received Influenza vaccine Quadrivalent unspecified season. On 2nd February 2021, the patient received the 2nd dose of COVID 19 VACCINE. On 15th February 2021, between 2 and 4 months after receiving Shingles vaccine and less than 2 years after receiving Influenza vaccine Quadrivalent unspecified season, the patient experienced clot blood (serious criteria GSK medically significant). In February 2021, the patient experienced swelling of feet, ankle swelling, foot pain and pain ankle. The patient was treated with rivaroxaban (Xarelto). On an unknown date, the outcome of the clot blood was unknown and the outcome of the swelling of feet, ankle swelling, foot pain and pain ankle were recovering/resolving. It was unknown if the reporter considered the clot blood, swelling of feet, ankle swelling, foot pain and pain ankle to be related to Shingles vaccine and Influenza vaccine Quadrivalent unspecified season. Additional details were provided as follows: The reporter was the patient's son. The age at vaccination was not reported. The patient did not have medical history and family history. The patient got the Flu vaccine every year. The patient received 2nd dose of Covid vaccine in left arm. The reporter did not have NDC, lot and expiry date for the patient's 1st and 2nd Covid vaccine. The patient was told by the nurse administering the Covid vaccine that the 2nd dose would be a little bit stronger. Either on 04th February 2021 or 05th February 2021, less than a week after receiving Covid vaccine, less than 4 months after receiving Shingles vaccine and less than 2 years after receiving Flu vaccine, the patient experienced severe swelling and pain in foot, ankle and calf. The patient first noticed left foot was swollen. The patient stated her foot was sore. The reporter did not know if it went to her ankle and then from there. Doctor advised the patient the patient had a clot and was put on Xarelto. An angiography and catheter were suggested to see if she had something closing in her. The patient would be going back in few weeks to see about the clot. The patient had never had this clotting concern before. Reporter clarified that, the patient was going to have another procedure such as angiography or catheter, but the patient had an MRI on 15th February 2021, 13 days after receiving Covid vaccine, and was diagnosed with the clot. The patient spoke with doctor who decided to put the patient on Xarelto. No information of NDC, lot and expiry date or dosing of Xarelto was provided. The reporter thought the patient's clot was weird and odd. At the time of reporting, the pain and swelling were gotten better. No further details were provided. The reporter consented to follow up. It was unknown if the reporter considered the clot blood, swelling of feet, ankle swelling, foot pain and pain ankle to be related to Covid vaccine.
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1201471-1	Blood clot on 4/2 in left calf
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1209444-1	Blood clots in my lungs
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1214759-1	Pain, swelling left knee found to be a blood clot when he went to the Emergency Room at hospital. Visit was on 3/31/2021 Treatment with rivaroxaban and bed rest. Pain and muscle spasms persist as of 4/15/2021
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1245000-1	Blood clots in the lungs, pulmonary embolism
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1276503-1	LEG PAIN DIAGNOSED AS BLOOD CLOT; HEADACHE; MILD FEVER; This spontaneous report received from a patient concerned a patient of unspecified age and sex. The patient's weight, height, and medical history were not reported. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, and batch number: 201A21A expiry: unknown) dose was not reported, 1 in total administered on 09-APR-2021 to left arm for prophylactic vaccination. No concomitant medications were reported. On an unspecified date, in APR-2021, the patient experienced mild fever (99 degrees fahrenheit), headache and leg pain diagnosed as blood clot (leg pain) on 21-APR-2021. On an unspecified date in APR-2021, laboratory data included: Body temperature (NR: not provided) 99 F, and Diagnostic ultrasound (NR: not provided) unknown. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the leg pain diagnosed as blood clot, headache and mild fever was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0:2021045225-covid-19 vaccine ad26.cov2.s-Leg pain diagnosed as blood clot. This event(s) is considered unassessable. The event(s) has a compatible/suggestive temporal relationship, is unlabeled, and has unknown scientific plausibility. There is no information on any other factors potentially associated with the event(s).
THROMBOSIS	COVID19 (COVID19 (UNKNOWN)) (1202)	UNKNOWN MANUFACTURER	1362296-1	Patient went to Emergency room at local hospital with leg pain. Evaluated and discovered clot in left leg and is being treated with Eloquis

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
THROMBOTIC STROKE	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0919546-1	thrombotic stroke -necessitating hospitalization; and craniotomy; required mechanical ventilator for 2 days. Patient now extubated, breathing on her own. Patient remains hospitalized with marked deficits (aphasic)
THROMBOTIC STROKE	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1207086-1	Thrombotic CVA of L hemisphere and spleen. CVA occurred overnight and was complete when patient found. Therefore, no treatment. Result R side paresis arm and leg and aphasia/apraxia.
THROMBOTIC STROKE	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1285786-1	Thrombotic Stroke
THROMBOTIC STROKE	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1044569-1	THROMBOTIC STROKE IN THE DISTRIBUTION OF THE LEFT MCA DISTRIBUTION
THROMBOTIC STROKE	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1256998-1	Within a couple days of the vaccine my husband had a small TIA. This included numbness in his left arm. The numbness subsided. On the 12th, the numbness returned which extended from his left ear down his left arm and down his left side to about the waist. We took him to the ER. A brain MRI showed he had a thrombotic stroke.
THROMBOTIC STROKE	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1278550-1	thrombotic strokes; right cerebellar stroke; vertebral artery dissection/ tore a vertebral artery in her neck; two sub acute strokes; vertigo; nausea; weakness; fatigue; sore arm; mild chills; body aches; This is a spontaneous report from a contactable consumer (patient). A 37-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; Lot Number: EN5318) via an unspecified route of administration, administered in Arm Left, at the age of 37-year-old, on 04Feb2021 16:30, as SINGLE DOSE for covid-19 immunisation. Medical history was not reported. The patient did not receive any other vaccines 4 weeks prior. Concomitant medication included ongoing drospirenone, ethinylestradiol (DROSPIRENONE/ETHINYLESTRADIOL) taken for contraception. The patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection; Lot Number: EL3302) single dose, at the age of 37-year-old, on 14Jan2021 1500PM in the left arm, for covid-19 immunisation and experienced fatigue and a sore arm. On 05Feb2021, the patient experienced fatigue, sore arm, mild chills, and body aches. On 08Mar2021, the patient was hospitalized because of vertebral artery dissection after an intense workout that morning/ she tore a vertebral artery in her neck which she immediately clotted and led to her having a stroke. The patient had vertigo, nausea and weakness on 08Mar2021 and went to the ER and they found she had a thrombolytic stroke in the cerebellum as a result of the vertebral artery dissection. The patient was admitted to the hospital on 08Mar2021 then she was discharged 11Mar2021. While she was in the hospital, they did an MRI and found two sub-acute thrombolytic strokes or blood clots in her brain. The patient tore the artery in her neck which led to the right cerebellar stroke; her blood was really clotty but they could tell by the imaging that the two subacute blood clots that led to having the other two strokes were there approximately three weeks before but not much older than that (without exhibiting symptoms, so she had three thrombotic strokes in total). Since she was discharged she has been out of work and will be on FML for two to three months and going to physical therapy from the effects of the strokes. Adds while she was in the hospital they were trying to determine the cause of the two sub-acute strokes that were unrelated to the dissection. They did several tests including: EKG which was normal; TEE scope down her throat to look at her heart which was normal; and blood work to check for a clotting disorder which was also all normal. So far nothing has been found. Outcome of mild chills, body aches, and sore arm was recovered on 06Mar2021; outcome of vertebral artery dissection and right cerebellar stroke was recovering; while it was unknown for the other events.
THROMBOTIC STROKE	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1340517-1	"Had a stroke from blood clot; she lost weight; This is a spontaneous report from a contactable consumer (mother) that: A 77-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: ER8272) via an unspecified route of administration on 31Mar2021, as a single for COVID-19 immunization. Previously patient received first dose of BNT162B2 (covid-19 vaccine, Batch/Lot Number: not clear) via an unspecified route of administration on 10Mar2021 as single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. Reporter stated, ""I don't know so I just was try to find out or who I report or ask about. I convinced my mother to get her Covid vaccine, so she got her second shot on 31Mar we got it together. Yesterday (17May2021) my mom had a stroke from blood clot, and I was just asking myself if it could have had anything to do with from Pfizer shot as she never had any problem like that and now, she was in the hospital from the stroke, and I was just wondering if me talking her and getting her vaccine could possibly had to do with this. Reporter further stated, Absolutely, I have question as far as if that was the cause, was there any special treatment that she should be getting maybe I can at least inform the doctor about that anything. I am out of my mind right now because I felt that I talk to her getting the vaccine because I want to her to be safe, she was 77 yrs. old (Not clarified) she never had any medical problem I feel guilty. Height and weight: Reporter stated, She was approximately 5'6. I don't know the weight as she lost weight, she was not obese or anything like that, but I honestly don't know her weight. The outcome for the events was unknown."

Note: Submitting a report to VAERS does not mean that healthcare personnel or the vaccine caused or contributed to the adverse event (possible side effect).

Notes:

Caveats: VAERS accepts reports of adverse events and reactions that occur following vaccination. Healthcare providers, vaccine manufacturers, and the public can submit reports to VAERS. While very important in monitoring vaccine safety, VAERS reports alone cannot be used to determine if a vaccine caused or contributed to an adverse event or illness. The reports may contain information that is incomplete, inaccurate, coincidental, or unverifiable. Most reports to VAERS are voluntary, which means they are subject to biases. This creates specific limitations on how the data can be used scientifically. Data from VAERS reports should always be interpreted with these limitations in mind.

The strengths of VAERS are that it is national in scope and can quickly provide an early warning of a safety problem with a vaccine. As part of CDC and FDA's multi-system approach to post-licensure vaccine safety monitoring, VAERS is designed to rapidly detect unusual or unexpected patterns of adverse events, also known as "safety signals." If a safety signal is found in VAERS, further studies can be done in safety systems such as the CDC's Vaccine Safety Datalink (VSD) or the Clinical Immunization Safety Assessment (CISA) project. These systems do not have the same limitations as VAERS, and can better assess health risks and possible connections between adverse events and a vaccine.

Key considerations and limitations of VAERS data:

- Vaccine providers are encouraged to report any clinically significant health problem following vaccination to VAERS, whether or not they believe the vaccine was the cause.
- Reports may include incomplete, inaccurate, coincidental and unverified information.
- The number of reports alone cannot be interpreted or used to reach conclusions about the existence, severity, frequency, or rates of problems associated with vaccines.
- VAERS data are limited to vaccine adverse event reports received between 1990 and the most recent date for which data are available.
- VAERS data do not represent all known safety information for a vaccine and should be interpreted in the context of other scientific information.

Some items may have more than 1 occurrence in any single event report, such as Symptoms, Vaccine Products, Manufacturers, and Event Categories. If data are grouped by any of these items, then the number in the Events Reported column may exceed the total number of unique events. If percentages are shown, then the associated percentage of total unique event reports will exceed 100% in such cases. For example, the number of Symptoms mentioned is likely to exceed the number of events reported, because many reports include more than 1 Symptom. When more then 1 Symptom occurs in a single report, then the percentage of Symptoms to unique events is more than 100%. More information. (/wonder/help/vaers.html#Suppress)

Data contains VAERS reports processed as of 06/18/2021. The VAERS data in WONDER are updated weekly, yet the VAERS system receives continuous updates including revisions and new reports for preceding time periods. Duplicate event reports and/or reports determined to be false are removed from VAERS. More information. (/wonder/help/vaers.html#Reporting)

For more information on how many persons have been vaccinated in the US for COVID19 to date, see https://covid.cdc.gov/covid-data-tracker/#vaccinations/ (https://covid.cdc.gov/covid-data-tracker/#vaccinations/)

Help: See The Vaccine Adverse Event Reporting System (VAERS) Documentation (/wonder/help/vaers.html) for more information.

Query Date: Jun 27, 2021 5:06:28 PM

Suggested Citation:

United States Department of Health and Human Services (DHHS), Public Health Service (PHS), Centers for Disease Control (CDC) / Food and Drug Administration (FDA), Vaccine Adverse Event Reporting System (VAERS) 1990 - 06/18/2021, CDC WONDER On-line Database. Accessed at http://wonder.cdc.gov/vaers.html on Jun 27, 2021 5:06:28 PM

Query Criteria:

State / Territory: The United States/Territories/Unknown
Symptoms: THROMBOSIS; THROMBOTIC STROKE
Vaccine Products: COVID19 VACCINE (COVID19)
VAERS ID: All
Group By: Symptoms; Vaccine; Vaccine Manufacturer; VAERS ID
Show Totals: False
Show Zero Values: Disabled