The Vaccine Adverse Event Reporting System (VAERS) Results

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1146222-1	started getting numbness in hands-fingers and lips and then numbness to feet and fell to ground with inability to walk - no strength in legs knees or arms. left facial drooping and slurring of speech and severe weakness - paralyzed
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1157503-1	PARALYZED FROM WAIST TO ALL THE WAY DOWN LEGS; 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: Unk) dose was not reported, administered on MAR-2021 for prophylactic vaccination. The batch number was not reported and has been requested. No concomitant medications were reported. On an unspecified date, the subject experienced paralyzed from waist to all the way down legs. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of paralyzed from waist to all the way down legs was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0: 20210347177-covid-19 vaccine ad26.cov2.s-paralyzed from waist to all the way down 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).
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1163209-1	Numbness in extremities, paralysis full body. Pneumonia left lung. Now diagnosed with Guillain-Barre syndrome, due to reaction to covid-19 vaccine
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1182506-1	4/6/21 I awoke with symptoms of numbness/paralysis on left side of face limited to upper lip and cheek area and both eyes would not stop watering. Movements to drink or make facial expressions were limited due to the numbness. Because I have sensitive skin I thought I may have had some type of reaction to something. So I waited to see if it would clear up through the day. I was not experiencing any other symptoms. By that evening it was apparent that I was not getting better and possibly getting worse because I began to experience some noticeable drooping in my cheek area. I decided to call my doctor in the morning. Because I am currently out of town, my home doctor suggested I see a local ER to rule out a stroke and determine if I was experiencing Bell's Palsy. I went to an ER clinic on 4/7/21
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1185249-1	Taste of medicine in mouth/throat - 1:30pm Swelling and numbing of right side of face begins - 2:30pm Benadryl taken - 2:53pm Numbing spread to both sides of throat, difficulty swallowing - 3:15pm 24/7 Nurse Hotline at Medical Center called - 3:16pm Nurse hotline advised to call 911 immediately Paramedics arrive - 3:23pm Bell?s palsy symptoms begin - 3:25pm Transported via ambulance to Emergency Room - 3:26pm Bell?s palsy worsens, facial contractions continue Prednisone given in emergency room Prescription of Methylprednisolone Tablets, USP is prescribed for 6 days, 21 pills, 4mg each Continue use of 25mg of Benadryl as needed Follow up with neurologist recommended and scheduled for 04/09/2021 Day 2, 04/08/2021 - slept off and on, felt terrible, facial twitching, forehead pain from twitching, right eye uncomfortable. Day 3, current day, waiting to see neurologist, right eye pain increasing and blurriness of vision in that eye.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1186151-1	Severe COVID-19 infection with ARDS. Requiring mechanical ventilation, paralysis and flolan nebulization.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1193804-1	8 hours later , I had 101 fever , shivers , nausea and full on body aches . Since feeling so bad , next day I had a thc gummy to help with pain . That night , I could not move my legs ... a paralysis feeling went from feet to face . Half of my fave was crooked and I could barely talk .
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1201895-1	Patient states that on the night of 4/9/2021 around 10PM she was laying down and felt paralyzed and could not move for 6 hours. Patient states that she was unable to call 911 because she could not move.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1202379-1	Significant flaccid paralysis in the right arm and right leg with some difficulty swallowing
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1210110-1	Fever of 102 degrees starting same night as the one dose COVID Janssen vaccine along with excrutiating headache. Took Tylenol each day of fever. Rested as headache made me very nauseas. Took Zofran for nausea. Fever lasted four days. Sever headache lasted 5 days. Lower leg cramps plus chills with sweating began on day 5.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1211052-1	My father received his shot on March 13, 2021. He drove to move home and on March 28th started having his right arm go numb but did not tell us. He asked to go the ER March 29th at 6:45am saying his chest hurt and was having a hard time breathing, this has happened before since having Covid in September. I called 911 after he became dizzy and could not walk. While at the ER suffered a massive stroke that paralyzed him from his nose down over a course of 3 days. His brain stem was affected and he lost the ability to swallow. After being on a ventilator for 72 hours he was removed from it and died less then 30 mins later from drowning in his own saliva on April 1, 2021.

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PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1220300-1	I will begin by noting that I am known for high pain tolerance and have been a strong, healthy, professional athlete for thirty years. The evening of the vaccination, I went to bed at 9pm feeling odd. By midnight, I became paralyzed with excruciating pain throughout my torso and arms with such an extreme fever that it felt as if my heart was going to stop. I had crushing chest pain which inhibiting my breathing ability in a way that could only be compared to being crushed under a burning truck. The paralyzation was so intense, with pre-existing pain multiplied by a thousand, that I couldn't even speak to alert my husband, who was right beside me. This continued for several hours until it subsided enough for me to pass out from exhaustion. The following day I battled residual aches throughout my torso and arms, accompanied by a mild fever. The trauma of fearing for my life that night and over the past week, exacerbated by the news of this vaccine I received being paused for issues related to my demographic, had made me too afraid to report my extreme side effects until now. With the encouragement and support of my family, I have worked up the courage to do so. It was the scariest, most intense pain and fever I have ever experienced in my life all at once and no one I have described this to has ever heard of the severity of what happened to me.
PARALYSIS	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.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1261220-1	PARALYSIS; UNABLE TO WALK/ WALKING DIFFICULTY; CHILLS; FEVER; HEADACHE; 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 not reported, batch number: Unknown) dose, start therapy date were not reported, 1total, 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 paralyzed after receiving vaccine, the patient was hospitalized for 7 days. The patient did not had blood clot, but had difficulty walking, fever, chills, headache and was unable to walk. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from paralysis, fever, chills, headache, and unable to walk/ walking difficulty. This report was serious (Hospitalization Caused / Prolonged).; Sender's Comments: 20210430056-Covid-19 vaccine ad26.cov2.s-Paralysis. This events is 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.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1295644-1	on 4/10/21 headache, feet felt numb, continued to get worse until almost total paralysis of body. He is still currently in hospital getting treatments. ,
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1303947-1	BACK ACHE; MUSCLE SPASM; HEAVY DARK CLOTTY PERIOD; PARALYSIS; PINS AND NEEDLES SENSATION; NUMBING; SWEATS; SORENESS; NAUSEA; This spontaneous report received from a consumer concerned a 26 year old female. The patient's height, and weight were not reported. The patient's concurrent conditions included birth control implanted in arm that is old and needs to be taken out, alcohol user, and non smoker, and other pre-existing medical conditions included the patient had no drug allergies nor illicit drug use. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, route of admin not reported, batch number: 1808982, and batch number: 1808982 expiry: UNKNOWN) dose was not reported, 1 total, administered on 03-MAY-2021 at 09:00 on left arm for prophylactic vaccination. No concomitant medications were reported. On 3-May-2021 at 09:00, the patient was vaccinated. At 17:00, the patient had pain, nausea, sweats, and soreness. At 18:00, she had pins and needles sensation, numbing, and paralysis. Later, she had sweats. Paramedics had checked the patient blood pressure and heart rate and they were within normal limits. The patient was still experiencing backache, muscle spasm, pain and heavy dark clotty period which was unusual. The patient had birth control implanted in arm that was old and needed to be taken out. Laboratory data included: Blood pressure and Heart rate was found normal. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from soreness, back ache, muscle spasm, and heavy dark clotty period, and the outcome of nausea, sweats, pins and needles sensation, numbing and paralysis was not reported. This report was serious (Other Medically Important Condition).; Sender's Comments: V0 20210511206-COVID-19 VACCINE AD26.COV2.S- Paralysis. 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).
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1322162-1	"Patient claims 3 minutes after vaccination, his body felt like ""it was on fire"", he started sweating and he felt like passing out. He also claims he could not move or speak for 30 minutes. The patient then claims he was completely fine after 1 hour and has no reaction today."

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	(1203)			completely fine after 1 hour and has no reaction today.
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1325858-1	PARALYSIS ON BOTH SIDES OF HIS BODY; This spontaneous report received from a consumer via social media platform 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,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. The patient was paralyzed on both sides of body for a month and had no improvement. The patient had vaccine and on day 5 after vaccination issue had started. On day 9 after vaccination, the patient was paralyzed and still had at the time of this report. The patient's physician stated the patient was perfectly healthy no medical issues, it was the vaccine that caused it. The patient was not better at the time of this report. The action taken with Covid-19 vaccine ad26.cov2.s was not applicable. The patient had not recovered from paralysis on both sides. This report was serious (Other Medically Important Condition).; Sender's Comments: V0. 20210524291- Covid-19 vaccine ad26.cov2.s-Paralysis. 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).
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1340597-1	I had a brain tia-stroke. Blood clots. Paralyzation.loss of vision. Loss of voice
PARALYSIS	COVID19 (COVID19 (JANSSEN)) (1203)	JANSSEN	1400856-1	Pt received shot, went home, within 1 hour of having shot he felt extremely high. The feeling escalated quickly, lasted about a 1/2 hour, and then subsided. He described his symptoms as feeling paralyzed, as he couldn't move, even to reach his phone to call for help, but his mind was alert. He felt tired and achy and slept for 3 hours.

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				"FELT PARALYSED; DIAGNOSED WITH GUILLEN BARRE SYNDROME; INJURED RIGHT FOOT; NUMBNESS IN FACE; LEFT LEG WEAKNESS; SWELLING IN LEGS; TIREDNESS; FEELINGS OF WEAKNESS; NUMBNESS IN LEFT HAND; This spontaneous report received from a consumer concerned a 46 year old female. The patient's weight was 113 kilograms, and height was 66 inches. The patient's pre-existing medical conditions included the patient had no known allergies. The patient received covid-19 vaccine ad26.cov2.s (suspension for injection, intramuscular, batch number: 201A21A, and expiry: 07-AUG-2021) dose was not reported, administered on 01-MAY-2021 to left arm for prophylactic vaccination. Concomitant medications included immunoglobulin human normal for guillen barre syndrome. On 01-MAY-2021, the subject experienced feelings of weakness, numbness in left hand. On 28-MAY-2021, the subject experienced tiredness. On 29-MAY-2021, the subject experienced left leg weakness and swelling in legs. On 02-JUN-2021, the subject experienced numbness in face. On 09-JUN-2021, the subject experienced diagnosed with guillen barre syndrome, and was hospitalized. On an unspecified date, the subject experienced injured right foot, and felt paralysed, and was hospitalized on 08-Jun-2021. On 01-MAY-2021, the patient developed numbness in her hand on the left side and felt weak. The patient travelled overseas and had a connection to a country and her husband stated that she started to have swelling in her legs. When she reached her destination, she was admitted to the hospital and was diagnosed with Guillen Barre Syndrome by the neurologist per her husband's recollection. In the hospital she was given Immunoglobulin G for 5-7 days and she did regain some feeling in her legs. Based on patient's husband recollection this may have occurred ""maybe tired due to travelling,"" but when she arrived to the country that was when her left leg weak and requested a wheelchair. The patient then injured her ""right foot"" in another country, but ""she didn't feel it and she requested help because she couldn't walk."" per her husband. After she reached her final destination, she felt tired and then she completely fell. It was reported that she could not ""control herself and felt paralyzed."" On 08-JUN-2021, she entered the hospital and was still in hospital at the time of this report. Her husband states that he ""does not know any side effect from any vaccine"" that caused this in the past. Lastly, he mentioned ""she may have numbness in her face when she reached original country"". He also stated that the on 04-JUN-2021, doctors conducted a computerized tomography scan and did not find a blood clot. The action taken with covid-19 vaccine ad26.cov2.s was not applicable. The outcome of the feelings of weakness, swelling in legs, diagnosed with guillen barre syndrome, numbness in left hand, tiredness, left leg weakness, numbness in face, injured right foot and felt paralysed was not reported. This report was serious (Hospitalization Caused / Prolonged, and Other Medically Important Condition).; Sender's Comments: V0: This is a spontaneous report received from consumer concerning a 46 year old female, unknown ethnicity, who experienced weakness and numbness of the left hand on the same day after receiving the Janssen Covid-19 vaccine and was diagnosed with Guillain-Barre Syndrome 39 days after. Patient's height was 66 inches and weight was 113 kilograms. Past medical, previous surgery, current illnesses and medications taken as well as history of allergy, smoking, drug abuse and alcohol abuse were not reported. On the same day as the vaccine was given, patient developed weakness and numbness on the left hand. Patient still travelled. Twenty-eight (28) days post vaccine, patient experienced left leg weakness and swelling on the legs. Patient was noted to feel tired the day before. Thirty-two (32) days post vaccine, patient developed numbness on the face. Computerized Tomography (CT) scan was done and there was no blood clot found. Thirty-eight (38) days post vaccine, patient felt 'paralyzed' and injured her right foot. Patient was brought to the hospital the following day and was subsequently admitted. Patient was diagnosed with Guillain-Barre Syndrome and was given IVIG. Information regarding other potential etiologies was insufficient. Considering the temporal relationship, the event is assessed to have an indeterminate relationship with vaccination."
PARALYSIS	**COVID19 (COVID19 (JANSSEN)) (1203)**	**JANSSEN**	1405260-1	
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0930167-1	Treated in ED the same day as vaccination for left facial tingling and numbness. Diagnosis from ED of Left 5th nerve Transient Palsy. Resolved in 72 hours.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0968886-1	Two hours after injection I got very cold and got uncontrollable trembles with total loss of muscle use .
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0978081-1	Axillary nerve palsy x 2 weeks. Supraspinatus able to abduct arm 15 degrees and then no further abduction. Positive drop arm test. Treatment - observation, improved day 11 from vaccination, resolved after day 14 from vaccination.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0979064-1	No hx of seizures. Woke up with severe abdominal pain and need to have bowel movement. Diarrhea 1x wtih nausea. Severe ear ringing, headache, confusion, and out of body feeling. Loss of control over large muscles, shuffling gait, unable to open eyes or speak, impaired LOC, anxiety and emotions, diaphoresis and hyperventilation.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0981431-1	Bells Palsy left face. left facial numbness and partial paralysis
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	0987242-1	Paralysis, weakness, numbness in bilateral lower extremities. Symptoms started 1/26/2021 resulting in hospitalization. Discharged on 1/28/2021 with Out-patient physical therapy

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PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	0989665-1	about 6 hours post vaccine I started having chills and a headache. 11 hours post vaccine my hands became paralyzed in severe contracture. My face and feet and legs were number. I could barely breathe. I had to be transferred by Ambulance to Hospital
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1005403-1	Full body paralysis from 6pm until 5:45 am following day could talk and mind active
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1005853-1	Major side effect: Paralysis in left foot. Weak left leg. Second vaccine - Jan 28, 2021 at 4:00 PM. Typical side effects were noticed on Jan 29, that being chills, fever, dizziness, weakness in both legs, itchy arm at vaccine site. Jan 30 side effects were dizziness, stumbly, chills and trouble walking with left leg. Legs were still weak, left was worse. I had paralysis in my left foot. My left foot would not move back, only forward so when I walked my toes drug the ground. I called Doctor. He told me to see a neurologist and go to ER if symptoms progressed. I have made the appointment for Feb 16, 2021. Foot is slowly improving. Left leg is still weak and there is some numbness in left leg and left foot.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1012955-1	My upper right side was completely in pain and stiff. The pain was shooting up my neck and reaching underneath my right breast. Soon afterward, I started catching the chills, a came down with a fever. My head was booming and I felt overly weak and nauseous. The pain in my arm was completely paralyzing to the point where even when I moved my head and was completely in pain. I had covid before in April and this experience was worse.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1019474-1	event started: 1-21-2021 at approx 6-7pm; left arm tingling and intermittent numbness, then at 1-22-2021 at 2-3am muscles cramps and unable to move toes bilateral feet and a HA and nausea. 1-23-2021 started with confusion, wondered off at a grocery store and was able to think straight while paying at a register. other than that no other episode.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1020784-1	Paralysis swelling. Seizure, lost eye sight, partial speech ! And is bed riding and loss of balance
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1038396-1	paralysis, hallucinations, lack of speech, no bowel control, slept for 2 days straight, no mental reasoning or control of bodily functions
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1039884-1	Lost control of bladder; Could not walk; Paralyzed from the waist down for about 24 hours; Received the 2nd dose on 22Jan; Diarrhea; Vomiting; A spontaneous report was received from a consumer, concerning a 67-years-old female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced paralyzed from the waist down for about 24 hours/PT: Paralysis, received the 2nd dose on 22 Jan 2021/PT: inappropriate schedule of product administration, lost control of bladder/PT: urinary incontinence, could not walk/PT: gait inability, diarrhea/PT: diarrhoea, and vomiting/PT: vomiting. The patient's medical history was not provided. Concomitant medications included omeprazole, fluoxetine hydrochloride, propranolol, colestipol, acetylsalicylic acid, and calcium. The patient received their first of two planned doses of mRNA-1273 (Lot number: 037K20A) on 28 Dec 2020. On 22 Jan 2021, the same day as the onset of the events, the patient received their second of two planned doses of mRNA-1273 (Lot number: 032L20A) intramuscularly for prophylaxis of COVID-19 infection. On 22 Jan 2021, following the second vaccination, the patient experienced vomiting and diarrhea. On 23 Jan 2021, the patient could not walk, was paralyzed from the waist down for about 24 hours, and lost control of her bladder. Treatment information for the events was not provided. The patient received both scheduled doses of mRNA-1273; therefore, action taken with the drug in response to the events was not applicable. The outcome of the event, paralyzed from the waist down, was considered recovered/resolved on 24 Jan 2021. The outcome of the event, received second dose on 22 Jan 2021, was considered resolved on 22 Jan 2021. The outcome of the events, vomiting, diarrhea, could not walk, and lost control of bladder, was unknown.; Reporter's Comments: This case concerns a 67 year old female subject, who experienced a serious unexpected event of paralysis after second dose of mRNA1273 (Lot# 032L20A). Very limited information regarding this event has been provided at this time. Further information has been requested.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1040736-1	On the day following receiving the first vaccine, January 20, 2021 at 2:30 in the afternoon, Tom's legs and feet became paralyzed and he was unable to walk requiring a wheel chair. His doctor was called and after determining he did not have another stroke, it was suggested to rest and wait for 24 hours. By 8 AM the following morning, January 21, 2021. The very same sequence of events occurred after his second dose on February 16, 2021.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1042447-1	Intense pain at the injection site and radiating down right arm for 24 hours after the injection. 72 hours after injection paralysis of right wrist possible radial nerve damage.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1043901-1	I was paralised from waist down for 1.5 Days with the loss to control over holding my urine
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1046200-1	headache, fever, chills, nausea, body aches, sore arm, muscle spasms- hands and arms contracting (hands in fists and arms to chest like you see in individuals with CP) unable to move for 10+ minutes

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PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1048812-1	8 days after vaccination - stroke/full right side of body paralyzed - clot buster drug at ER
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1050142-1	Paralysis for 2 days. Unable to use either arm, barely able to bear weight or walk. Had to be fed, given water, and helped to go to the bathroom.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1057247-1	Patient was sitting for 10minutes when husband reported she started to slurr speech and inability to communicate Upon entering the room patient has head tilted against wall with arms and legs spread out Patient was responsive, but weak, reporting she felt paralyzed and couldn't open eyes Vitals taken, vitals we're stable Pulse ox-98%, BP-144/90, Pulse 90, glucose-110 After 5 minutes, patient was still weak, unable to open eyes, pupils had no response to light EMS called 1 hour post vaccine, symptoms have improved as per husband ER doctor told patient symptoms should resolve over course of day
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1057331-1	Lot # 011L20A- patient reported L arm reddness, swelling and warmth. Lot # 031L20A- patient reported L arm reddness, swelling and warmth, along with palsy of the L side of her face.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1057377-1	On the evening of the first vaccination, my sister experienced chills and fever. The next morning she experienced a kind of paralysis, could not sit up or stand up, was falling out of bed and out of her chair, had to be carried to the bathroom, although she usually can walk by herself. This lasted about a day. I took her to the ER where they excluded other possible causes of her condition and simiply attributed it to the vaccine. No treatment was prescribed.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1059306-1	This morning upon awaking was unable to rise from bed, semi-paralysis, peed in the bes as was unable to move any extremities to allow me to get up! Fever, chills, badly shaking, unable to walk unaided. some confused thinking. Unable to call out for help or recall how long I had remained motionless.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1059619-1	Within several hours my arm began getting really sore and I became extremely tired. I laid down for a nap in the recliner. I awoke 2 hours later (around 9:30 p.m.) with the whole left side of my body temporarily paralyzed. The whole side felt heavy/sore like my arm - almost as if the vaccine had spread down my side. It really scared me because I initially couldn't move. I finally worked my way into a sitting position, then stood up and walked it off, then was fine. No more problems with it. Just today (2/27) I am starting to feel better as far as fatigue and soreness and nausea go. No more paralysis.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1060792-1	Severe joint and muscle paralysis. Fever for 24 hours. Remained in bed for 2 days.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1069121-1	Pt received first COVID vaccine on 1/23 and on 2/12 developed persistent periorbital headache and on 2/14 developed OS ptosis and diagnosed w/CN3 palsy
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1069744-1	Cranial nerve 6 (Abducens) palsy, left side
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1072900-1	He received the vaccine and after 2 hours he went home and was paralyzed from the waist down. He tried to stand up and then he passed out on the floor for about 10-15 minutes. He went to sleep and woke up around 5 hours later. He then tried it again and urinated on himself and he fell down on the floor, passed out without getting up until the morning. He did call his doctor who was not available, spoke to the nurse and advised to go to the hospital. He is weak and cannot drive so he did not go to the hospital. He slept the rest of that day and the following day. Then when he saw his doctor about 2 days ago, he described his reaction and told to report the reaction and that he would report it himself. He was advised not to take the 2nd vaccination. He is no longer having the heavy reaction, his muscles are aching and his vision is blurred, especially on his back and the shoulder. His doctor did do blood work at the office visit.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1074236-1	I have been experiencing some tingling on the left side of my face since receiving the first COVID-19 vaccination dose. It flare after receiving the second COVID-19 vaccination dose. The level of tingling varies throughout the day. It has not resulted in noticeable paralysis.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1079103-1	Patient reports getting her 2nd vaccine near 3PM and that evening had symptoms including high fever and paralysis. Paralysis lasted until 9am and patient reports symptoms including rigors, fever, pain and lethargy lasting about 3 days with the symptom of lethargy continuing. VEARS reporting information given to client who is a nurse and she states she will self report.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1081556-1	States took the CoVid vaccine a week ago in his rt arm (Not currently in LINKS). Began with facial droop and numbness - did have a hx of Bells palsy 12 years ago. Obvious facial asymmetry noted with nerve paralysis on the right and is unable to fully close eyelid and the side of his moth is drooping and unable to close completely.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1081740-1	Paralyzed from the waist down for a few minutes and fell down; Feels disoriented; Sometimes can't even stand up; A spontaneous report () was received from a physician concerning a 77-year-old, male patient who received Moderna's COVID-19 vaccine (mRNA-1273) and was paralyzed from the waist down and fell down/paralysis, feels disoriented/disorientation and sometimes can't even stand up/dysstasia. The patient's medical history was not provided. Products known to have been used by the patient, within two weeks prior to the event, included poly vitamins. On 10 Feb 2021, the patient received their first of two planned doses of mRNA-1273 (Lot number: 015M20A) intramuscularly for prophylaxis of COVID-19 infection. On 10 Feb 2021, approximately a few hours post vaccination, the patient experienced paralysis from the waist down for approximately 15 minutes which made her fall down. This event occurred two times. The patient stated she felt disoriented and could not stand up. Treatment information was not provided. The outcome of the events, was paralyzed from the waist down and fell down/paralysis, feels disoriented/disorientation and sometimes can't even stand up/dysstasia, were considered as 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1082220-1	Moderna COVID-19 Vaccine EUA Woke up at 4AM with a high fever (103F), arm/leg numbness and paralysis, nausea, vomiting, headache, and dizziness. Numbness, paralysis, and vomiting only lasted about 15-20 minutes. Fever and headache lasted 18 hours from onset.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1083315-1	"By 3:00 he had a whole body chill. He was then incoherent, lost all control of all body muscles and was incontinent both bladder and bowel. He continued as such when Dr. said to give him Prednisone along with the Benadryl he was receiving every six hours starting at 3:30 p.m. 3/8/2021. He was given a Prednisone at 4:00 p.m. 3/9/2021 and again at 8:00 p.m. on 3/9/2021 along with his Benadryl. He was unable to ambulate or even turn his body over in bed. He was able sip water and sports drink but ate nothing since noon on 3/8/2021. It wasn't until early Sunday morning that he was able to move his legs and get to a walker and use the toilet. He slow, very slowly has improved since then. Late Sunday he was able to get up, and sit in a chair. Later he was able to take a sit down shower and shampoo and be an to feel ""nearly human"" again. He has slowly progressed since then and Monday morning was able to navigate by himself and to descend and ascend a 13 stair step to the basement. In discussing the 24 hours he spent in bed he was unaware of any of the things going on such as not being able to move, but knew he was in pain if he moved his legs. He also had a temperature of 102.3 which rose to 102.9 and then with cold compress on his forehead and time his fever dropped to 95.4 and the back up to a normal of 97.5. These are similar symptoms he had after the first injection of the Covid 19 Moderna vaccine and he had had a similar reaction in 2017 from the first flu shot he ever had in his life. He felt he needed to take this vaccine in order to be able to ""have a life"" even though we assumed he would have such a bad reaction. He is slowly improving and if it is like the first injection it will take him nearly two weeks before he can ambulate at the level he was before the vaccination. Still, he is grateful to Center for being so proactive as to have available the vaccine and it being more convenient for cancer patients than waiting in line at a mass inoculation center. He would appreciate an acknowledgement that this information has been received and could possibly passed along to help others in his situation. Thanks."
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1084103-1	Palsy of the face and left side extremities. Stroke like symptoms.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1086874-1	Paralysis of entire body; A spontaneous report was received from a consumer concerning a 79-year-old, male patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced paralysis of his entire body. The patient's medical history was not provided. Concomitant products included Evolocumab, blood pressure medication and metformin. The patient received the first of two planned doses of mRNA-1273 (Batch number: unknown). The date of administration of first dose of vaccine was not mentioned. On 19 Feb 2021, approximately one day prior to the onset of the event, the patient received the second dose of two planned doses of mRNA-1273 (Batch number:006M20A) intramuscularly for prophylaxis of COVID-19 infection. On 20 Feb 2021, Patient experienced paralysis of his entire body at night and the ambulance had to come to put him back in bed. Treatment information was not provided. 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 event paralysis of his entire body, was considered resolved on 21 Feb 2021.; 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.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1086931-1	Shingles in the ear; Passed out; Couldn't mover her arm, head, legs, chest, felt paralyzed; Felt weak; A spontaneous report was received from a consumer concerning a female patient of 75 year old, who was received Moderna's COVID-19 vaccine(mRNA-1273) and experienced loss of consciousness, paralysis, weakness and shingles in the ear. The patients medical history, as provided by the reporter, included myasthenia gravis. Concomitant medications reported included lidocaine patch and prednisone. On 10 feb 2021,prior to the onset of events, the patient received their first of two planned dose of mRNA-1273(Lot number:010M20A) intramuscularly on an unknown arm for prophylaxis of COVID-19 infection. On 11 feb 2021, one day after vaccination,patient experienced loss of consciousness,paralysis and weakness. The event loss of consciousness and paralysis was also considered to be medically significant. On 16 feb 2021, patient experienced shingles in the ear. Treatment for the events included acyclovir and ice. Action taken with mRNA-1273 in response to the event was not reported. The outcome for the events loss of consciousness, paralysis, weakness and shingles in the ear were considered as unknown.; Reporter's Comments: Based on the current available information, a strong temporal association between the use of the product and the start date of the events, a causal relationship with the events cannot be excluded.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1087234-1	Joint pain in elbow and wrist paralysis in fingers
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1087769-1	Here is my story. Part 1: I had my second Moderna vaccine on 2/16. 36 hours later I was laying In bed when I all of a sudden felt very strange- sort of faint - like I was going to pass out. Then I started feeling like my Body was going numb. We called 911. I was taken to the ER. What followed was about 15-20 ?neurological? episodes (every ten minutes or so) where my entire body from head to toe became paralyzed- I couldn?t move any of my limbs. I also had no sight. I was not blacked out Bc I could hear everything that was happening. It was also accompanied by extreme chest pain, numbness, contratcing of muscles in chest cavity that felt almost like a heart attack. Each individual episode lasted about 3-5 minutes and then slowly the paralysis would undo and I?d regain feeling. And then the full episodes would start over a few minutes later. This went on for a few hours repeatedly while in the ER. I was admitted for testing and monitoring for two days (EEG, EKG, MRI, Catscan) all clear. Once discharged I had lingering neurological symptoms but not the full paralysis episodes. Lingering symptoms included severe chest pain, numbness/weaknesses predominant on the left side as well as other random areas. I was very weak and tired. I started to improve very slowly - by day 10 post hospitalization I was starting to feel like things were turning the corner and maybe I was on my way to recovery. Part 2: I started to finally feel more improved from the lingering neurological issues and then all of a sudden around 5:30PM I was in the kitchen making dinner for my kids on 3/2 and the episodes started again. Called 911. Again, every ten minutes or so they came on (maybe 15-18 episodes) of complete paralysis- this time accompanied by seizure like rapid eye movement from side to side and fluttering of the eyes. I was readmitted to the hospital where the episodes repeated to happen again, maybe 15-20 times. I was admitted. They did a 24 hour video EEG which ?caught? the episodes on video and sound but did not appear negatively on the EEG in terms of abnormal brain activity. I?ve been released as they say there is nothing to do. I have lingering muscle weakness in my eyes- I can?t focus my eyeballs or ?track? on an eye exam and continued muscular chest pain. I also have a lot of PTSD from the trauma of the events.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1098995-1	Within 3 hours of getting g the vaccine my blood pressure went up, my breathing became difficult & the whole left side of my body went into paralysis. I was taken to the ER, later that day was admitted to the hospital. I spent 5 days there getting tested. I was released with 0 use of my left arm or hand, and my left leg was weak. It?s been 3 months and I still can?t use my left arm. I am seeing multiple different doctors to try & help me.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1100271-1	Lost control of muscles; ended up blacking out; could not move; could not talk; A spontaneous report was received from a Consumer concerning about a 82 years old female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced Paralysis, Syncope, Mobility decreased and Aphasia. The patient's medical history was provided as diabetes, dementia as current illness. Relevant concomitant medication was reported as blood thinner for blood clots. On 08 FEB 2021, prior to the onset of event, the patient received their first dose of two planned doses of mRNA-1273 (lot/batch: unknown) for prophylaxis of COVID-19 infection. On 10 FEB 2021, the patient experienced Paralysis, Syncope, Mobility decreased and Aphasia. Due to these events patient's required hospitalization and on oxygen. Laboratory details was not provided. Treatment information was not provided. Action taken with mRNA-1273 in response to the events was unknown. The outcome of the event's Paralysis, Syncope, Mobility decreased, and Aphasia was as unknown.; Reporter's Comments: The events were consistent with increased risk of cerebrovascular complications associated with elderly age of patient. Company assessed the events to be unlikely related to company product.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1101161-1	couldn't move her body at all; lay there unable to move or crawl or even pull herself up for 6-7 hours before she was discovered; no recollection of this episode of what happened to her or what she was doing at the time it occurred; she fell to her garage floor; speech was gargled; brain fog and difficulty; residual confusion; local reaction limited arm movement; local reactions such as pain; A spontaneous report was received from a consumer concerning a 77-year-old, female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced local reactions such as pain and limited arm movement (vaccination site movement impairment/vaccination site pain), fell to garage floor (fall), couldn't move body at all (paralysis), lay there unable to move or crawl or even pull self up for 6-7 hours before discovered and taken to hospital (paralysis), brain fog and difficulty (feeling abnormal), no recollection of this episode of what happened or what she was doing at the time it occurred (amnesia), speech was gargled (dysarthria), and brain fog is improved now but still has some residual confusion (delirium). The patient's medical history was not provided. Relevant concomitant medications were not reported. On 27 FEB 2021, prior to the onset of events, the patient received their second dose of mRNA-1273 (lot/batch: unknown) intramuscularly in the right arm for prophylaxis of COVID-19 infection. On 27 FEB 2021, the patient experienced vaccination site movement impairment and vaccination site pain. On 28 FEB 2021, the patient experienced paralysis, amnesia, fall, dysarthria, and feeling abnormal and due to these events hospitalization was required. On 04 MAR 2021, the patient experienced Delirium. Laboratory details was provided as CT Scan on 28 FEB 2021 and the result was normal. Treatment information was not provided. 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. At the time of this report, the outcome of the event's, local reactions such as pain and limited arm movement,fell to garage floor, couldn't move body at all , lay there unable to move or crawl or even pull self up for 6-7 hours before discovered and taken to hospital, brain fog and difficulty, no recollection of this episode of what happened or what she was doing at the time it occurred, speech was gargled, and brain fog is improved now but still has some residual confusion, were considered 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1101794-1	Felt extremely warm, started feeling lightheaded, started sweating profusely, vision started blacking out, got very close to blacking out and fainting. I was seated at the time, waiting the 15 minutes for observation after the shot, symptoms happened about 10 minutes into the waiting period. I was given some cold water right away, over the next 10 minutes my vision started coming back and I stopped sweating. Was told I looked flush when they first came over but was looking much better after another 10 mins. I stayed an additional 30 minutes for observation after the event and had friends come take me home and get my car. I'm still feeling very slightly light headed and somewhat tired 2 days later, but otherwise feeling great.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1104792-1	Neurological impairment - unable to move extremities; hospitalized and symptoms continuing
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1106173-1	He was paralyzed,couldn't use his legs, hand, got collapsed; Lack of strength; Injection site was annoying, rubbing, scratchy; down deep in his muscle a slight feel pain at injection site; extreme dizziness lasted for a day and a half.; shortness of breath; A spontaneous report was received from a consumer concerning himself, 70 years old, male patient who experienced he was paralyzed, couldn't use his legs, hand, got collapsed/paralysis, injection site was annoying, rubbing, scratchy/injection site pruritus, down deep in his muscle a slight feel pain at injection site/injection site pain, shortness of breath/dyspnoea, lack of strength/weakness, extreme dizziness lasted for a day and a half/dizziness. The patient's medical history was not provided. Concomitant product use was not provided by the reporter. The patient received their first of two planned doses of mRNA-1273 (Batch number: Unknown) on 24 Feb 2021 intramuscularly for prophylaxis of COVID-19 infection. The patient was paralyzed, he couldn't use his legs, hand, got collapsed. Injection site was annoying, rubbing, scratchy, down deep in his muscle a slight feel pain. He also had shortness of breath, lack of strength and extreme dizziness lasted for a day and a half. Treatment of the events was not provided. Action taken with mRNA-1273 in response to the event was not provided. The outcome of the event he was paralyzed, couldn't use his legs, hand, got collapsed, injection site was annoying, rubbing, scratchy and down deep in his muscle a slight feel pain at injection site was unknown and of events shortness of breath, lack of strength, extreme dizziness lasted for a day and a half was recovered.; Reporter's Comments: Based on the information provided, a strong temporal association and the established safety profile of mRNA-1273 vaccine, a causal association between the reported events and the product use cannot be excluded. Injection site pain and injection site pruritus are consistent with the known safety profile of the vaccine

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1111267-1	Limp; Paralyzed (pain was so bad he couldn't move it); Low grade fever; Severe left hip pain; A spontaneous report was received from a consumer who was also a 85-year old, male patient who received Moderna's COVID-19 Vaccine (mRNA-1273) and experienced left hip became paralyzed (pain was so bad he couldn't move it) (paralysis), low grade fever (pyrexia), limp (gait disturbance), and severe left hip pain (arthralgia). The patient's medical history was not provided. No relevant concomitant medications were reported. On 09 Jan 2021, approximately 10 hours prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (lot 012L20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 09 Jan 2021, the patient woke up at 10:00 pm with severe left hip pain. His left hip became paralyzed (pain was so bad he couldn't move it), and he took 2 ibuprofens then went to bed. On 10 Jan 2021, he was still in pain with a low grade fever of 98.8 which is 1 to 2 degrees higher than normal for him. He reported using his wife's walker to ambulate and took 4 ibuprofens which helped his hip. On 11 Jan 2021, the patient was still in pain, had a limp, and ran a low fever until the afternoon. Treatment for the events included ibuprofen. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the event, low grade fever, was considered resolved on 11 Jan 2021. The outcome of the events, left hip became paralyzed (pain was so bad he couldn't move it), low grade fever and limp, was unknown.; Reporter's Comments: Based on the information provided, the events of low grade fever, gait disturbance and severe left hip pain are temporally associated with the administration of mRNA-1273 and a causal association cannot be excluded. Fever and arthralgia are consistent with the known safety profile of the product. The event of paralysis is considered unlikely pending additional information.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1116038-1	weakness, paralysis, administered stress dose IV dexamethasone, regained strength in 1 hour
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1122899-1	pt states that 11 hours after taking the 2nd dose she collapsed falling to the floor and was paralyzed and had urinary incontinence for 20 hours. EMT came to put pt back in bed. EMT came once again and put pt in wheelchair to get her to a car so she can be taken to Hospital ER. They did blood work which potassium was low, urinalysis test showed bladder infection, and CT was normal. Pt released after 4 hours and told to FU w/ her neurologist. Pt saw Neuro Dr. on 3/17/2021 said she must have had some sort of reaction to the vaccine.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1127847-1	Participant felt flushed, feverish, fatigued with general aches and dry cough over the weekend after receiving injection, took acetaminophen and cough syrup on Monday. He became short of breath on 1/20/2021 and was hypoxic on oximeter check, was sent to the ER. He was intubated in ER and went into respiratory failure with sepsis due to COVID19. He was treated with tocilzumab, became paralyzed and DVT in left lower extremity was found. HE required pressors and diuresis, he developed AKI and hyperkalemia. On 2/21 he was in multi-organ failure. His level of cognition decreased until he was no longer responsive and he died on 2/24/2021.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1129665-1	Arms, legs and body was paralyzed; Nerve pain that travel from her head to the back; Excruciating pain,Pain is still remaining; Not being able to get up from bed; Lightning pain in the back; A spontaneous report was received from a 76 year old female patient concerning herself, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced excruciating pain and the feeling of not being able to get up from bed, because her arms, legs and body was paralyzed, nerve pain that travel from her head to the back and patient had lightning pain in the back. The patient's medical history was not provided. Concomitant medications were not reported. On 26 Feb 2021, the patient received the first of two planned doses of mRNA-1273 intramuscularly on left arm for prophylaxis of COVID-19 infection. On 01 Mar 2021, after receiving the first dose of the mRNA-1273, the patient experienced excruciating pain, feeling of not being able to get up from bed, because her arms, legs and body was paralyzed (Other Important Medically Significant). This pain was still remained, kind of nerve pain that travel from her head to the back and also the patient had lightning pain in the back. Laboratory investigation details was not provided. Treatment details included Tylenol for pain. Action taken with mRNA-1273 in response to the events was not reported. The outcome of events, excruciating pain, feeling of not being able to get up from bed, because her arms, legs and body was paralyzed, nerve pain that travel from her head to the back and lightning pain in the back was not recovered/not resolved at the time of reporting.; 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1144725-1	PALSY OF THE RIGHT MOUTH FOR 2 DAYS (GONE); TREATED WITH ORAL STEROID RASH AT THE INJECTION SITE (STILL HAVE); TREATING WITH TOPICAL STEROID AND ITCH CREAM HEADACHE JOINT PAIN

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1153147-1	Felt like she was paralyzed; Could not move or lift arm; Weak; Fever of 100.4; A spontaneous report was received from a consumer concerning a 84-year old female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced felt like she was paralyzed, could not move or lift arm, fever of 100.4, and weak. The patient's medical history mentions COVID-19 in Dec 2020 and blood pressure. Concomitant medications provided were blood pressure medicine NOS and vitamins. On 09 Mar 2021, prior to the onset of the events, the patient received their first of the two planned doses of mRNA-1273 (Batch number: 030a21a) intramuscularly for prophylaxis of COVID-19 infection. On 09 Mar 2021, patient stated she had a bad reaction. Patient stated it felt like she was paralyzed, and she could not move or lift arm. Patient stated she had a fever of 100.4 and weak . The event, felt like she was paralyzed was deemed medically significant. Treatment information was not provided. Lab tests were not provided. Action taken with respect to mRNA-1273 was not reported. The outcome of the events, felt like she was paralyzed, could not move or lift arm, fever of 100.4 and weak 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1153960-1	Told her not get the second dose of the vaccine; Doctor told her she had a bad reaction to the first dose; Partially paralyzed for about 4/5 hours on her bottom half; Little bit itchy û hair, ears, and eyes; A spontaneous report was received from a consumer concerning a 78-year-old, female patient, who experienced partial paralysis on her bottom half/paralysis, bad reaction to the first dose/vaccine complication, not to take second dose/ intentional dose omission, itchy- hair, ears, and eyes/pruritus. The patient's medical history included COVID-19 in Dec 2020 and high blood pressure. Products known to have been used by the patient, within two weeks prior to the event included high blood pressure pills. On 09 Mar 2021, the patient received their first of two planned doses of mRNA-1273 (Lot number: 036A21A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 10 Mar 2021, the patient was partially paralyzed for 4-5hrs on her bottom half. The patient had same symptoms when prior exposed to COVID-19 in Dec 2020 and experienced little itchy- hair, ears and eyes. Brain scan, lungs, heart, blood tests and neurological functions were tested when patient consulted a health care provider (HCP) hence was advised not to take second dose. Treatment for the event included Benadryl. Action taken with mRNA-1273 in response to the events was not provided. The outcome of the event, partial Paralysis on her bottom half/Paralysis was recovered. The outcome of the events,bad reaction to the first dose/vaccine complication, not to take second dose/ intentional dose omission, itchy- hair, ears, and eyes/pruritus were considered 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1154398-1	The right side of my face has lost most movement above the mouth. Paralysis for short.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1154735-1	Patient has Multiple Sclerosis and had a fever of 100.4, began to exhibit pain in all extremities, pain when others touch like fibromyalgia. Patient felt paralyzed and could not move out of bed without assistance. Had problems such as balance issues, full body pain and discomfort, dizziness, spinal pain from neck to lower lumbar, burning sensation in lower extremities down to toes.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1155206-1	Developed left facial palsy approximately 48 hours after injection. Presented to local ER on 3-21-21 and diagnosed with Bells Palsy
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1157863-1	Acute-onset vertical binocular diplopia found on 2/19/21 exam by me to be a left fourth nerve palsy. By follow up on 3/19/21 the diplopia has essentially resolved and the left fourth nerve palsy was measured to be nearly resolved as well.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1161994-1	Became paralyzed; Every single muscle became tensed; Sore arm; severe reaction/ vaccination adverse reaction; Nauseous; A spontaneous report was received from a consumer concerning herself, a 22-year old female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced every single muscle became tensed, became paralyzed ,sore arm, nauseous and a severe reaction. The patient's medical history was not provided. Concomitant medication use was not provided by reporter. On 02 Mar 2021,prior to the onset of the events, the patient received their first of two planned doses of mRNA-1273 (Lot number 023A21A) intramuscularly for COVID-19 infection prophylaxis. On 02 Mar 2021, after first dose of the Moderna vaccine, the patient experienced every single muscle became tensed, became paralyzed ,sore arm, nauseous. Treatment details included acetaminophen. Action taken with mRNA-1273 in response to the events was not reported. The outcome of the events, every single muscle became tensed, became paralyzed ,sore arm, nauseous, and a severe reaction was considered 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1163510-1	Symptoms began on 3/24/21 and progressed until 3/28/21. Diagnosis is he suffered a spinal stroke, involving T6-T10. He is paralyzed from just below his nipple line, all the way down.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1165006-1	Moderna Covid 19 Vaccine EAU severe flu-like symptoms escalating to incontinence, cumulating in fainting and being unable to move any body part except one index finger. After that subsided, foggy brain, fever. Eventually a day of diarrea.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1168998-1	"TIA; Right side of her face was drooping; Left arm was paralyzed; Slurred speech; Dizzy; Confused; A spontaneous report was received from a consumer concerning a 46-year-old, female patient, who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced TIA (transient ischaemic attack), dizzy (dizziness), her left arm was paralyzed (paralysis), right side of her face was drooping (facial paralysis), slurred speech (dysarthria), and she was confused (confusional state). The patient denied having any allergies, past medical history or concomitant medications. Therefore, no medical history or concomitant medications were reported. On 14-Jan-2021, the patient received the first of two planned doses of mRNA-1273 (Lot #: 251L20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 10-Feb-2021, approximately twenty-four days prior to the onset of events, the patient received the second of two planned doses of mRNA-1273 (Lot #: 16M20A) intramuscularly in the left arm for prophylaxis of COVID-19 infection. On 06-Mar-2021 the patient had a ""TIA"" (transient ischemic attack). She was sitting down and when she got up, she got dizzy and her left arm was paralyzed. The right side of her face was drooping, she had slurred speech and she was confused. The patient went to the hospital where she was hospitalized for 2 days. The MD (medical doctor) said she had a ""7 mm focal diffusion."" Treatment information was not provided. 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 outcomes of the events, TIA, dizzy, her left arm was paralyzed, right side of her face was drooping, slurred speech, and she was confused, 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."
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1169787-1	The patient received her FIRST dose of Moderna COVID vaccine on 1/26/21 and SECOND dose on 2/23/21. On 3/8, she first started to experience shortness of breath, which was attributable to cardiac etiology and diastolic heart failure at the time, but which in retrospect may have been her earliest symptom of new onset myasthenia gravis, which is the AE I am wanting to report. She was hospitalized from 3/8-3/13, and underwent extensive cardiac workup that was negative. She again developed shortness of breath on 3/17/21 and was re-admitted to the hospital on 3/18/21 for diuresis for presumed diastolic heart failure exacerbation, received diuresis from 3/18-3/22/21 but then developed new onset diplopia, at which pint I evaluated her in neurological consultation on 3/22. Her brain MRI disclosed an UNrelated incidental small left cerebellar infract, but her neurological exam showed a new R lateral rectus palsy and new moderate to severe left eyelid ptosis, suspicious for myasthenia gravis. Anti-acetylcholineseterase antibodies---binding, blocking, and modulating, were all sent off and came back all strongly positive on 3/26-27/21. The patient then called stating she had developed recurrent shortness of breath, new onset dysphagia, and worsening double vision. I urgently re-evaluated her in outpatient follow-up in my office on 4/2/21 and sent her immediately to the hospital to be readmitted, where she remains as of this reporting date and she has been receiving IVIG with improvement in her symptoms. She did NOT have any previous known history of eyelid ptosis, dysphagia, diplopia, or muscle weakness. In retrospect, the patient states that her current symptomatology began around 3/7/21 with the benefit of hindsight. I am reporting this event because it is very unusual for an 82 year old WOMAN to develop new onset of myasthenia gravis, and it is a known autoimmune disease, and a vaccine would be a plausible trigger for its development, or trigger for its first symptomatic manifestation. I was also especially concerned because my fellow neurologist, ALSO had a patient develop myasthenia gravis about 2 weeks or so following their second dose of the Pfizer COVID vaccine. It would be an EXTRAORDINARY coincidence for two instances of new onset myasthenia gravis to occur 2 weeks post- vaccination at essentially the same time in an institution of our size, so we both felt obliged to report both cases ASAP to the VAERS. I was not able to reach the patient in her hospital room today to get additional specifics about where she got the vaccine, its lot number etc., however she is a retired and has indicated her willingness to provide you with any and all additional information you need, and I am also personally very willing to provide any additional information about her eventual outcome.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1170061-1	Pt had 1st episode of Bells - right side symptoms on 12-27-20. She had a very dense paralysis which not completely resolve at this time. She had her 1st Moderna shot on 03-10-21 and on 03-19-21 c/o onset of similar symptoms, but on the left side of her face. The current episode was not as bad as the initial episode in December. Currently mild residual symptoms, mostly noted when smiling.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1175024-1	Paralysis of her whole body for 30 seconds; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (Paralysis of her whole body for 30 seconds) in a 57-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 025A21A) for COVID-19 immunisation. The patient's past medical history included No adverse effect (no medical history reported.). On 23-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 24-Mar-2021, the patient experienced PARALYSIS (Paralysis of her whole body for 30 seconds) (seriousness criterion medically significant). On 24-Mar-2021, PARALYSIS (Paralysis of her whole body for 30 seconds) had resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown Route) was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown Route), 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; 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
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1178244-1	Guillain-BarrT syndrome; Tingling and numb toes and fingers, which spread to elbows and knees; Tingling and numb toes and fingers , which spread to elbows and knees; Progressed to paralysis; Problems breathing on her own; A spontaneous report was received from a consumer concerning his relative a 39-years-old female patient who received Moderna's COVID-19 vaccine (mRNA-1273) and experienced tingling, numb toes and fingers which spread to elbows and knees, difficulty breathing progressed to paralysis. The patient's medical history was not provided. No relevant concomitant medications were reported. On 23 Dec 2020, approximately three weeks prior to the onset of the events, the patient received her first of two planned doses of mRNA-1273 (batch: 011J20A) intramuscularly for prophylaxis of COVID-19 infection. On 14 Jan 2021, the patient experienced tingling, numb toes and fingers, which spread to elbows and knees and progressed to paralysis which required hospitalization and patient was diagnosed with guillain-barrT syndrome. On 25 Jan 2021, the patient was intubated with mechanical breathing because she had problems breathing on her own. patient was hospitalized for a long-term acute care hospital. Treatment for the event included patient was intubated with mechanical breathing. The second dose of mRNA-1273 was discontinued in response to the events. The outcome of the events, tingling, numb toes and fingers which spread to elbows and knees, progressed to paralysis, guillain-barrT syndrome, was intubated with mechanical breathing and had problems breathing her own were 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 events, a causal relationship cannot be excluded. However, Further information is requested.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1179191-1	Upper extremity dysfunction; felt like it was paralyzed; Left arm hurt; pain in shoulder down to armpit; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (felt like it was paralyzed) in a 36-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 027L20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Asthma, Migraine and Sinus disorder. Concomitant products included MONTELUKAST SODIUM (SINGULAIR) and SALBUTAMOL (ALBUTEROL HFA) for Asthma, IBUPROFEN for Migraine. On 06-Jan-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) .5 ml. On 06-Jan-2021, the patient experienced PARALYSIS (felt like it was paralyzed) (seriousness criterion medically significant), PAIN IN EXTREMITY (Left arm hurt) and PAIN (pain in shoulder down to armpit). On an unknown date, the patient experienced MUSCULOSKELETAL DISORDER (Upper extremity dysfunction). At the time of the report, PARALYSIS (felt like it was paralyzed) and PAIN IN EXTREMITY (Left arm hurt) had resolved and PAIN (pain in shoulder down to armpit) and MUSCULOSKELETAL DISORDER (Upper extremity dysfunction) outcome 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 events, a causal relationship cannot be excluded.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1181372-1	Paralyzed, sharp cramping and then makes it difficult to breathe or move as it will worsen in the chest area. First episode happened 30 minutes after injection and then again on the 4/7/21 . Lasts for 20-25 minutes. Woke up in 4/8/21 to a red patch in the same area where injection was administered.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1184935-1	sagging, flacid left eye and cheek. slight paralysis in left leg about an hour then symptoms went away

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1189615-1	water and blood coming out the legs; paralyzed feeling in my legs completely, cant even walk around the house; uncontrollable diarrhea; severely decreased appetite; severe bloating; high body temperature of 103; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of HAEMORRHAGE (water and blood coming out the legs) and PARALYSIS (paralyzed feeling in my legs completely, cant even walk around the house) in an 82-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 001B21A) 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 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 HAEMORRHAGE (water and blood coming out the legs) (seriousness criterion medically significant), PARALYSIS (paralyzed feeling in my legs completely, cant even walk around the house) (seriousness criterion medically significant), DIARRHOEA (uncontrollable diarrhea), DECREASED APPETITE (severely decreased appetite), ABDOMINAL DISTENSION (severe bloating) and PYREXIA (high body temperature of 103). At the time of the report, HAEMORRHAGE (water and blood coming out the legs), PARALYSIS (paralyzed feeling in my legs completely, cant even walk around the house), DIARRHOEA (uncontrollable diarrhea), DECREASED APPETITE (severely decreased appetite), ABDOMINAL DISTENSION (severe bloating) and PYREXIA (high body temperature of 103) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Treatment information was included TORSEMIDE, CEFUROXIME and MUPIROCIN cream. Based on the current available information which includes a temporal association between the use of the product and onset of the reported events, and excluding other etiologies, a causal relationship cannot be excluded.; Sender's Comments: Based on the current available information which includes a temporal association between the use of the product and onset of the reported events, and excluding other etiologies, a causal relationship cannot be excluded.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1191505-1	After 20 minutes patient felt blurriness and could not move her body for a few seconds , after sitting and drinking some water, patient felt much better and was able to walk
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1193548-1	I first noticed an increase in mucus in my lungs, causing me to cough up phlegm. The vision issues have increased the MS Optic Nuritis problems, loosing vision in right eye! But its the headaches I can't stand. They come and go, and sometimes its sooo bad that I can't move my head and body. Sitting in the dark helps. Thus having MS and lowered immune system has the headaches causing me to be immobile during episodes, sometimes lasting only an hour, sometimes all day! And chills, but not as bad as the headaches. I don't want to lose my vision again due to having to do these vaccines every year with these side affects.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1204093-1	Severe pain and palsy on left arm and neck. Pain not resolved with a variety of treatments according to patient.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1205803-1	Difficulty breathing, Total body muscle paralysis Ongoing
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1207891-1	Took the moderna 1st test and within two hours of the vaccine, between the forefinger of my left hand and the thumb began to swell, tighten. I observed it, it appeared to be a large clot, it stiffened almost paralyzing the area between these two fingers, it hurt, and then as suddenly as it came it left. No further on the first vaccine. On the second vaccine in the same hand, a day after the shot, my left hand swelled from the wrist to the fingers but especially in the area of the back of the hand. it nearly paralyzed moment in the hand, for at least two days. it hurt extemely when I would attempt to move the hand back at the wrist, to move the hand in any way. I took two motrins on the second day and it went away as suddenly as it began. I have not had any other problem to date
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1210876-1	First I developed a skin rash on both my arms, neck and face - when to urgent care and received treatment - shot, medcines to take home [resolved itself in a few days. At the same time I started experiencing pins, needles, burning, pain in leg and arm muscles that attacted my up and lower extremities. This all started about a week after I took the first shot. I went to my a general practioner - explained my crisis and she sent me home on steroids' and tramadol. It helped some but then got worst - I then went to emg room and they put me on Gabapentin 100 mg. It helped some. Ended up going to the emg room because the paralis came up in my neck , throat, mouth, lip and chest. Gave me Gabapentin and Tramadol.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1218043-1	Temporary paralysis in my right hand. Around 9am today, my hand began making these involuntary movements and suddenly got very tense, almost like a claw. I had no control over my hand and it remained in this distorted position for about 1 minute before my hand relaxed back to its normal state.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1219269-1	This was the Johnson and Johnson one shot. My mother received it and today is in the hospital with acculsive blockage in left brain, paralyzed from stroke and is being operated on. I believe it is a blockage from the vaccine. She was old but extremely cognizant, independent, healthy, strong of sound mind. This is why I am reporting it. If she dies I will report again.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1230343-1	Patient reported a total paralysis from arm to toes with unbearable pain for a week. Cannot walk when in pain.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1233580-1	Full body convulsions and Paralysis. Left-arm went numb first and traveled throughout the body. Convulsions started in left hand and also traveled throughout the body. It took six minutes from the shot to the first convulsion. Convulsions didn't stop until lorazepam was administered. Benadryl was administered immediately on-site in pill form. The hospital also gave additional Benadryl as well. Blood sugar dropped to 50 within the first half-hour. Extreme eye sensitivity to light and extreme exhaustion afterward. Pain felt extreme as well. Many side effects such as very low pain tolerance, head ache, light sensitivity, confusion, loss of memory, severe muscle pains, muscle numbness throughout body, blood sugar dropping, etc.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1235609-1	Stroke 6 days after receiving the second dose of the vaccine; Paralysed left side; Fell on the floor unresponsive; Fell on the floor Unresponsive; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of CEREBROVASCULAR ACCIDENT (Stroke 6 days after receiving the second dose of the vaccine), PARALYSIS (Paralysed left side) and UNRESPONSIVE TO STIMULI (Fell on the floor unresponsive) in an 86-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 reported). Concomitant products included ACETYLSALICYLIC ACID (ASPIRIN 81) for an unknown indication. On 04-Mar-2021, the patient received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 10-Mar-2021, the patient experienced CEREBROVASCULAR ACCIDENT (Stroke 6 days after receiving the second dose of the vaccine) (seriousness criteria hospitalization and medically significant), PARALYSIS (Paralysed left side) (seriousness criteria hospitalization and disability), UNRESPONSIVE TO STIMULI (Fell on the floor unresponsive) (seriousness criteria hospitalization and medically significant) and FALL (Fell on the floor Unresponsive). The patient was hospitalized on 10-Mar-2021 due to CEREBROVASCULAR ACCIDENT, PARALYSIS and UNRESPONSIVE TO STIMULI. The patient was treated with Surgery for Cerebrovascular accident and Surgery for Paralysis. At the time of the report, CEREBROVASCULAR ACCIDENT (Stroke 6 days after receiving the second dose of the vaccine), PARALYSIS (Paralysed left side), UNRESPONSIVE TO STIMULI (Fell on the floor unresponsive) and FALL (Fell on the floor Unresponsive) outcome was unknown. Not Provided Medical history was not reported. The patient was taking unspecified Antihypertensive medication. The patient was placed on a feeding tube and was transferred to a Nursing home and would require medical service for the rest of her life. 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245365-1	CVA isolated; Diplopia (left eye); Left with a palsy; Inability to walk confidently; Left cranial nerves VI, and III show deficit's; Cannot see out of left eye; This spontaneous case was reported by a health care professional (subsequently medically confirmed) and describes the occurrence of CEREBROVASCULAR ACCIDENT (CVA isolated), DIPLOPIA (Diplopia (left eye)), PARALYSIS (Left with a palsy), GAIT DISTURBANCE (Inability to walk confidently), CRANIAL NERVE DISORDER (Left cranial nerves VI, and III show deficit's) and VISUAL IMPAIRMENT (Cannot see out of left eye) in a 66-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 immunisation. Concurrent medical conditions included Obesity (BMI 45-45.9). On 13-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 01-Apr-2021, the patient experienced CEREBROVASCULAR ACCIDENT (CVA isolated) (seriousness criteria hospitalization, disability, medically significant and life threatening) and DIPLOPIA (Diplopia (left eye)) (seriousness criteria hospitalization, disability and medically significant). On 12-Apr-2021, the patient experienced PARALYSIS (Left with a palsy) (seriousness criteria disability and medically significant), GAIT DISTURBANCE (Inability to walk confidently) (seriousness criteria disability and medically significant), CRANIAL NERVE DISORDER (Left cranial nerves VI, and III show deficit's) (seriousness criteria disability and medically significant) and VISUAL IMPAIRMENT (Cannot see out of left eye) (seriousness criteria disability and medically significant). At the time of the report, CEREBROVASCULAR ACCIDENT (CVA isolated), DIPLOPIA (Diplopia (left eye)), PARALYSIS (Left with a palsy), GAIT DISTURBANCE (Inability to walk confidently), CRANIAL NERVE DISORDER (Left cranial nerves VI, and III show deficit's) and VISUAL IMPAIRMENT (Cannot see out of left eye) 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. The Patient was Treated with Lovenox, Coumadin, Brimonidine eye drops.The patient has been taking High Blood Pressure Medication,Severe Obesity medications and Hyperlipidemic Medications.The patient has no known Allergies. 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
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1245464-1	Trouble standing; Able to walk into the kitchen awkwardly; It felt like I was almost paralyzed/It was very difficult to control my legs and arms as if I had relapsed into losing the feeling in my whole body; Fell into a chair; Felt warm and had a fever of 100.3; Soreness in left arm; Hard to sleep; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (It felt like I was almost paralyzed/It was very difficult to control my legs and arms as if I had relapsed into losing the feeling in my whole body) in a 55-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 022M20A and 042L20A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Multiple sclerosis (began new DMT, Tysabri, in June 2020, monthly infusion) and Anemia. Concomitant products included IRON for Anemia, NATALIZUMAB (TYSABRI) for Multiple sclerosis, VITAMIN D NOS and CALCIUM, ERGOCALCIFEROL (CALCIUM +D [CALCIUM;ERGOCALCIFEROL]) for an unknown indication. On 26-Jan-2021 at 1:30 PM, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 23-Feb-2021 at 2:30 PM, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 23-Feb-2021, the patient experienced PAIN IN EXTREMITY (Soreness in left arm). 23-Feb-2021, the patient experienced INSOMNIA (Hard to sleep). On 24-Feb-2021, the patient experienced FALL (Fell into a chair) and PYREXIA (Felt warm and had a fever of 100.3). On 24-Feb-2021 at 7:00 AM, the patient experienced PARALYSIS (It felt like I was almost paralyzed/It was very difficult to control my legs and arms as if I had relapsed into losing the feeling in my whole body) (seriousness criterion medically significant), DYSSTASIA (Trouble standing) and GAIT DISTURBANCE (Able to walk into the kitchen awkwardly). At the time of the report, PARALYSIS (It felt like I was almost paralyzed/It was very difficult to control my legs and arms as if I had relapsed into losing the feeling in my whole body), PAIN IN EXTREMITY (Soreness in left arm), INSOMNIA (Hard to sleep), DYSSTASIA (Trouble standing), GAIT DISTURBANCE (Able to walk into the kitchen awkwardly), FALL (Fell into a chair) and PYREXIA (Felt warm and had a fever of 100.3) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 24-Feb-2021, Body temperature: 100.3 (High) Increased. Treatment information was not provided. Company Comment: This case concerns a 55-year-old female with a serious unexpected event of paralysis, and nonserious unexpected pain in extremity, dysstasia, gait disturbance, fall, insomnia and expected pyrexia. Event onset 2 days after first dose mRNA-1273. Events resolved. Based on 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-2021-025381 (Patient Link).; Sender's Comments: This case concerns a 55-year-old female with a serious unexpected event of paralysis, and nonserious unexpected pain in extremity, dysstasia, gait disturbance, fall, insomnia and expected pyrexia. Event onset 2 days after first dose mRNA-1273. Events resolved. Based on 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1246262-1	I had a massive stroke at work a few weeks after the 1st vaccine shot. Rushed to hospital for emergency surgery to remove blood clot in right side of brain. My paralysis was reversed after surgery but lingering headaches exist.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1249626-1	Paralysis from left arm down to feet; Unbearable pain; Could not walk; This spontaneous case was reported by a health care professional (subsequently medically confirmed) and describes the occurrence of PARALYSIS (Paralysis from left arm down to feet) in a 37-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 006B21A) 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). On 03-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On an unknown date, the patient experienced PARALYSIS (Paralysis from left arm down to feet) (seriousness criterion medically significant), PAIN (Unbearable pain) and GAIT INABILITY (Could not walk). At the time of the report, PARALYSIS (Paralysis from left arm down to feet), PAIN (Unbearable pain) and GAIT INABILITY (Could not walk) 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 product use was not provided. Treatment information was unknown. 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1256463-1	Shot hurt immediately 4:00 pm by 11:00pm my neck, arm, and hand hurting very badly by 12:00 midnight have severe pain and my hand is paralyzed this continues until 2:00 am pain began to lessen around 3:00 am was able to go to sleep . My neck and hand continued to hurt for 2 days . My arm hurt for 5 days .
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1256967-1	Transverse myelitis, on ventilator paralyzed from chest down

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1267473-1	She got her vaccine, she was feeling fine, the person noticed that she was in hiking gear and her dog as she had just had a 5 mile hike. He had a drawing feeling in her left arm, which she ignored and otherwise felt fine. Got the dog in the car, started to drive home, and got on her exit and within 3-5 minutes to get on the road going to her street felt a little weird and had blurry vision, and then it was really blurry and then she realized she was having a reaction. There was dirt on the right side of the road and turned her car there and remember hitting the hazard lights and not having any vision at all. Her left side was completely numb, and her right arm was extremely heavy and she was able to get her phone, but could move her fingers still and kept clicking on her phone until she got the Town Center where she got the vaccine. She got a hold of them and got a hold of someone, and then she was completely paralyzed with a swollen tongue as well, and not able to be understood. She stayed with her on the phone and was hardly able to make out the signs to let them know where possibly was so that they could find her. She could barely feel her fingers at that point, and was just able to move her eyes. She also felt fuzzy and was able to communicate. The lady got hold of 9-1-1 and able to look up her phone # from the vaccine center who had her address and information. It was a 3 way phone call, and it took them 35-45 minutes to find her, and two police officers passed by her right next to her with hazzards on. Finally she heard the fire truck and was able to tell the lady that the ambulance went by her and they finally found her. Her pulse was irratic and was able to tell them that she was not able to move and that her chest was heavy and took her to the ambulance. She was able to move her arm a little bit and then it fell back down again. Her vital signs were not normal so they were going to take her to the hospital. They texted her exboyfriend to come and get her dog and he came and got her car and dog. They took her to the hospital and they were not able to get vein access on the ambulance, and they put her in a wheelchair and not able to hold her head up. She was in the wheelchair for about 4 1/2 hours without any vein access, no blood drawn and nobody attending to her. Finally a nurse came in and put her in another wheelchair and pushed her to the other side and was able to get a little more comfortable. The nurse came in and said something about her Ex-boyfriend who is a heart surgeon at that particular hospital. Once she was on this new wheelchair and was finally able to move her toes, hands, and fingers and little by little sensation came back as she was a gymnast for 30 years and was flexing to try to gain motion. She was able to finally get up and weakly walk and they released her after giving her a Benadryl. Since going home she feels heavy (arms feels like cement), weaker, and does not feel completely herself, and went for a walk yesterday (which she is normally able to hike 5-8 miles) and has been dropping things due to either weakness or nerve issues. It has taken a toll on her and it?s hard for her to take her dogs up and down the stairs without extreme fatigue. Loss of appetite and loss of strength. She is a writer and is going to publish a paper regarding this as she feels she was not given the right information and proper treatment at the hospital.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1267960-1	Received the first shot of covid vaccine at 8:45 am. later that day I was having all of the known/common side effects, but I was also having heart palpitations. (5pm -9pm? fell asleep) I was alone and could not move for several hours. I woke the next day still feeling most of the common side effects but the heart palpitations had subsided. I was very scared and this was only my first does. I am afraid that getting the second dose will put me in the hospital or much worse.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1269836-1	I'm paralyzed from my chest down!
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1273283-1	4-25-21 9:45 PM - full body paralysis for 48 hours. As of 4-30-21 9:40 am can walk, can move hands....still issue with balance and dexterity. Approx at 90% capacity.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1275864-1	9 days after the Moderna covid 19 vaccine: Severe headache, steadily increased over 24 hours Unable to move head ER visit, transferred to hospital Treatments: IV anti bacterial, IV anti viral, dexamethasone, morphine, toradol DIAGNOSIS Meningitis Hospitalized overnight Patient recovered within 5 days
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1278201-1	12/31 vaccination Later that afternoon, I started with body aches, hard time breathing. My husband took me to urgent care and put me on O2 for a while. A couple days later, same episode. Off and on, depending how active I was, it would happen. I would use my inhaler when needed. I went to talk to dr. Went back to urgent care . They got me a O2 machine and nebulizer. Stayed in at hotel and admitted into the hospital. I had to be incubated and inpatient till 1/29/ 2021. I was told I was COVID +. I really don't remember. I couldn't walk, move, I was too weak. I am still going thru physical therapy, appt to see lung doctor, on O2 machine. I go to PT 1x a week because I don't have enough oxygen bottles. *PCM ordering CT and to see lung specialist ; putting in referral *Referral OBGYN, stomach issues/pain *feet numbing and tingling; I still lose balance

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1281067-1	body wide tingling; both hands became clenched and paralyzed; tachycardic; hurting pretty badly/shooting pain down both legs; not feeling so good; shortness of breath; Arm soreness; severe chills; extreme muscle pain; body wide muscle aches; intense headache; been in bed all day; fever of 102; This spontaneous case was reported by an other health care professional (subsequently medically confirmed) and describes the occurrence of PARALYSIS (both hands became clenched and paralyzed) in a 35-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 031B21A) 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 first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 06-Apr-2021, the patient experienced MALAISE (not feeling so good), PAIN (hurting pretty badly/shooting pain down both legs), DYSPNOEA (shortness of breath), PAIN IN EXTREMITY (Arm soreness), CHILLS (severe chills), MYALGIA (extreme muscle pain; body wide muscle aches), HEADACHE (intense headache), FATIGUE (been in bed all day) and PYREXIA (fever of 102). On 07-Apr-2021, the patient experienced PARALYSIS (both hands became clenched and paralyzed) (seriousness criterion medically significant), TACHYCARDIA (tachycardic) and PARAESTHESIA (body wide tingling). At the time of the report, PARALYSIS (both hands became clenched and paralyzed), MALAISE (not feeling so good), TACHYCARDIA (tachycardic), PAIN (hurting pretty badly/shooting pain down both legs), PARAESTHESIA (body wide tingling), DYSPNOEA (shortness of breath), PAIN IN EXTREMITY (Arm soreness), CHILLS (severe chills), MYALGIA (extreme muscle pain; body wide muscle aches), HEADACHE (intense headache), FATIGUE (been in bed all day) and PYREXIA (fever of 102) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 06-Apr-2021, Body temperature: 102 f High. On 07-Apr-2021, Electrocardiogram: 108 diffuse NS ST/T abnormal. 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 reporter states that the patient was en route to the hospital because both hands became clenched and paralyzed during her ER visit. No treatments were mentioned by the reporter. No concomitant medications were reported. Most recent FOLLOW-UP information incorporated above includes: On 24-Apr-2021: Significant follow up- new events 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1283710-1	AFIB Palsey to the optic nerve, double vision
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1286246-1	paralyzed and could not move except to stay flat on my back for two days
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1288044-1	Acute ecephalopathy, mental confusion, trouble with motor skills (balance, can't walk, shaking, some paralysis, tremors), mood changes, delusions, possible hallucinations, etc.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1292508-1	Pt reports that on 2/28/2021 pt woke up at 7 am and was unable to move. Pt reports that she was paralyzed for about 3 hours. Pt was vaccinated on 2/27/2021 at a vaccine site. Location was High school and was administered by hospital. This was given in the left arm.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1293608-1	My son his paralyzed t12 and his left hip swelled up within 24 hours of injection, then he used a hand massager to loosen it up and the leg and knee became bright red, swollen even more, and the pain in the leg became unbearable he couldn't move without intense pain, he was taken to ER, where he was treated for possible infection, and they noticed his hemoglobin was dropping and his leg was getting larger, he received 3 units of blood during his stay, A hematoma developed and was growing in the left thigh. He was given Vancomycin, then Doxy. The blood stabilized day 6 and he was discharged day 7 to home to continued recovery.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1294833-1	Thought was going to die with first shot; Feeling paralyzed and laid in bed for 3 days; About 7-8 hours her arm was killing; Lot of her previous pain from the accident had come back; Unusual reaction; Every bone and muscle in her body was hurting; Headache; Every bone and muscle in her body was hurting; This spontaneous case was reported by a consumer and describes the occurrence of NEAR DEATH EXPERIENCE (Thought was going to die with first shot) and PARALYSIS (Feeling paralyzed and laid in bed for 3 days) in a 72-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 Accident automobile. Concomitant products included ALPRAZOLAM (XANAX), HYDROCODONE BITARTRATE, PARACETAMOL (VICODIN) and MELATONIN for an unknown indication. On 20-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 17-Apr-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 NEAR DEATH EXPERIENCE (Thought was going to die with first shot) (seriousness criterion medically significant), PARALYSIS (Feeling paralyzed and laid in bed for 3 days) (seriousness criterion medically significant), PAIN IN EXTREMITY (About 7-8 hours her arm was killing), PAIN (Lot of her previous pain from the accident had come back), ADVERSE REACTION (Unusual reaction), BONE PAIN (Every bone and muscle in her body was hurting), HEADACHE (Headache) and MYALGIA (Every bone and muscle in her body was hurting). At the time of the report, NEAR DEATH EXPERIENCE (Thought was going to die with first shot), PARALYSIS (Feeling paralyzed and laid in bed for 3 days), PAIN IN EXTREMITY (About 7-8 hours her arm was killing), PAIN (Lot of her previous pain from the accident had come back), ADVERSE REACTION (Unusual reaction), BONE PAIN (Every bone and muscle in her body was hurting), HEADACHE (Headache) and MYALGIA (Every bone and muscle in her body was hurting) outcome was unknown. mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) dosing remained unchanged. 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-2021-099253 (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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1295836-1	States three weeks later on 4/23/2021 at 7pm radiating pain began from right shoulder to just above right elbow, severe pain, weakness, periods of paralysis. Self treated with plant based steroids and began to improve 5/1/2021. States on 5/4/2021 pain back up to shoulder, milder pain, still sore today, 5/7/2021. Has not and does not plan to see doctor for pain or discomfort. Receiving #2 Moderna COVID-19 vaccine today.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1299652-1	I have experienced a dramatic FLARE of Osteoarthritis in both knees and hips to the point that I have had to start using a cane to walk. Previous to this, the Osteoarthritis was only obvious in my left knee, but was in remission and I had not had to take any pain medication for months. After the second Moderna vaccination, the Osteoarthritis has flared and become extremely painful, even while in the bed lying down. It now is very painful in both the right and left knee and both, the right and left hip sockets. I have to take 2 Excedrin Extra Strength pills every six hours to lessen the pain. I also now have to use a cane for walking because it is too painful to bend my knees and support my weight. I am morbid obese - at 297 pounds (down from 391 pounds on September 30, 2019).
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1302692-1	He reported a GBS-type reaction where his tingling in his legs got worse and he was paralyzed from the waist down to the point that he couldn't walk. These symptoms lasted about a week. He brought this up later after the side effects had resolved. He also reported a transient amnesia episode where he forgot where he was driving locally and had to have some one come pick him up 2 weeks after the 2nd vaccine. Both of these side effects occurred after the second vaccine.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1303246-1	5 mins after initial shot, rapid heart rate that subdued in approximately 15 min. About 12 hours after the shot, experienced swelling in face and closure of throat, excessive sweating, fatigue and disorientation/dizziness and muscle paralysis/weakness in the back of neck, needing to use my hand to lift my head up.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1306323-1	Paralyzed from knees down; Diarrhea; Fever of 101.9¦F; Vomiting; Chills; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PARALYSIS (Paralyzed from knees down) in a 38-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 COVID-19 on 21-Dec-2020. Concomitant products included GABAPENTIN, DOMPERIDONE, LEVETIRACETAM and OMEPRAZOLE for an unknown indication. On 14-Apr-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-Apr-2021, the patient experienced PARALYSIS (Paralyzed from knees down) (seriousness criteria hospitalization and medically significant), DIARRHOEA (Diarrhea), PYREXIA (Fever of 101.9¦F), VOMITING (Vomiting) and CHILLS (Chills). The patient was hospitalized from 15-Apr-2021 to 16-Apr-2021 due to PARALYSIS. On 16-Apr-2021, DIARRHOEA (Diarrhea), PYREXIA (Fever of 101.9¦F), VOMITING (Vomiting) and CHILLS (Chills) had resolved. On 17-Apr-2021, PARALYSIS (Paralyzed from knees down) had resolved. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 21-Dec-2020, SARS-CoV-2 test: positive (Positive) Positive. On 15-Apr-2021, Blood test: normal (normal) Normal. On 15-Apr-2021, Body temperature: 101.9 (High) 101.9 Fahrenheit. On 15-Apr-2021, Influenza virus test: negative (normal) Normal. On 15-Apr-2021, SARS-CoV-2 test: negative (Negative) Negative. 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 provided was paracetamol The lab tests were reported as MBR, infection of the spine, tumor of the spine, inflammation of the spine. The results were provided as Normal. The event paralysis reoccurred on 16 APR 2021 at 08.00 pm and recovered on 17 APR 2-21. Again the event occurred on 17 APR 2021 at 08.00 pm and got recovered on 18 APR 2021. 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.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1306355-1	experience paralysis all left side; Fever; This spontaneous case was reported by a nurse and describes the occurrence of PARALYSIS (experience paralysis all left side) in a 54-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. Concurrent medical conditions included Systemic lupus erythematosus (lupus immunosuppressant disorder). Concomitant products included PREDNISONE for an unknown indication. On 30-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) (unknown route) 1 dosage form. On 01-May-2021, the patient experienced PARALYSIS (experience paralysis all left side) (seriousness criterion medically significant) and PYREXIA (Fever). At the time of the report, PARALYSIS (experience paralysis all left side) and PYREXIA (Fever) outcome was unknown. For mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown), the reporter did not provide any causality assessments. Treatment information was not provided. 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.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1309392-1	Approximately 2 1/2 weeks after receiving my first dose of the Moderna vaccine, I woke up on January 16 in severe pain in my right shoulder and arm. Approximately two hours later I developed paralysis in my right fourth and fifth finger. I eventually saw a neurologist who diagnosed me as having Turner parsonage syndrome. The pain has resolved but I continued to have paralysis in my right fourth and fifth finger and the lateral portion of my right hand.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1310981-1	Event at time of dose: vasovagal syncope, left/right hands/arms paralysis, mid and lower stomach/intestines start of paralysis; treatment: 50 m Benadryl shot, visit to ER; event approximately 2 weeks after dose: both calves tingling, left leg tingling/pain, left forearm tingling, left pinky and ring finger numbness; treatment: doctor visit, MRI, heart EKG, stress test, echocardiogram.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1326622-1	Myelitis transverse. Paralyzed for at least 2 days
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1327374-1	The patient began experiencing a flushing feeling and headache approximately 30 minutes following the administration of the Moderna COVID-19 vaccine on 5.7.21, later on that day white blisters began to appear on her hands and feet along with severe joint pain. The patient took 3 ibuprofen and 1 percocet to combat the joint pain. When the patient went to bed she had a fever of 102 degrees. Upon waking the patient experienced full body paralysis and was then taken to the ER. Once administered to the ER, the patient stated that she was given 3 different antibiotics. Once the patient was moved to the floor where she had remained in the hospital for 7 days where she was given corticosteroids to combat the severe joint pain and pressure experienced between head and neck. The patient stated that she is scheduled to be on antibiotics for 3 months but her symptoms have resolved

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				scheduled to be on antibiotics for 3 months but her symptoms have resolved.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1333688-1	Roughly 48 hours after second shot of Modern COVID19 vaccine (lot 013L20A) on 2/14/2021 started to notice motor weakness and loss sensation in legs adding to generalized weakness post-vaccine, on 2/17/2021 went to Hospital (DC) emergency room because patient identified symptoms as ascending paralysis of lower extremities, 2/17/2021 physical exam revealed Guillan-Barre syndrome w/ flaccid paralysis of lower extremities, sensory deficit, and absent deep tendon reflexes, MRI of brain through lumbar spine were within normal limits other than minor degenerative changes, lumbar puncture w/in normal limits, received 5 doses of IVIG starting on 2/18/2021, primary evaluation and treatment at hospital by NP-C and Dr. of Stroke Unit, on 2/25/2021 was transferred to Rehabilitation Hospital (DC), released on 3/5/2021 w/ plan for in-home health physical and occupation therapy (continued through 5/6/2021), seen by NP-C as well as Dr. at rehabilitation hospital outpatient during April 2021, started outpatient physical and occupational therapy at different Hospital on 5/7/2021, saw new Dr. on 5/19/2021
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1349113-1	Loosing vision in right eye; Can't move my head and body; Condition aggravated; To be immobile during episodes; Increase in mucus in my lungs; Cough up phlegm; Cough; Pain; Headache; Chills; This regulatory authority case was reported by an other health care professional and describes the occurrence of BLINDNESS UNILATERAL (Loosing vision in right eye) and PARALYSIS (Can't move my head and body) in a 64-year-old female patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 045A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. The patient's past medical history included Chronic lung disease (lungs are really sensitive) since an unknown date, Non-smoker since an unknown date, Cold (head) since an unknown date, Bronchitis (due to head colds) since an unknown date and Pneumonia. Previously administered products included for Immunisation: Shingles (extremely ill and 4 hours from being admitted to hospital with bad pneumonia after immune system dropped so low). Past adverse reactions to the above products included Pneumonia with Shingles. Concurrent medical conditions included Sulfonamide allergy, Allergic fungal rhinosinusitis (AFS), Hives and Multiple sclerosis (headaches had been recorded. The 8 lesions in frontal lobe causing more issues, thus the pain.). Concomitant products included METOPROLOL SUCCINATE, BUSPIRONE HCL, CYCLOBENZAPRINE, DULOXETINE and TRAZODONE for an unknown indication. On 20-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 BLINDNESS UNILATERAL (Loosing vision in right eye) (seriousness criteria disability and medically significant), PARALYSIS (Can't move my head and body) (seriousness criterion medically significant), CONDITION AGGRAVATED (Condition aggravated), MOBILITY DECREASED (To be immobile during episodes), BRONCHIAL SECRETION RETENTION (Increase in mucus in my lungs), PRODUCTIVE COUGH (Cough up phlegm), COUGH (Cough), PAIN (Pain), HEADACHE (Headache) and CHILLS (Chills). At the time of the report, BLINDNESS UNILATERAL (Loosing vision in right eye), PARALYSIS (Can't move my head and body), CONDITION AGGRAVATED (Condition aggravated), MOBILITY DECREASED (To be immobile during episodes), BRONCHIAL SECRETION RETENTION (Increase in mucus in my lungs), PRODUCTIVE COUGH (Cough up phlegm), COUGH (Cough), PAIN (Pain), HEADACHE (Headache) and CHILLS (Chills) had not 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. 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 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.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1351266-1	Transverse Myelitis relapsed. I was able to use a walker, now I am back in a wheelchair. I lost about 80% of my mobility overnight.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1353114-1	Losing breathe; Paralysis of face; Bells Palsy; Guillain-Baire syndrome,; Numbness in hands; Could not swallow; Tingly in hands; Impaired vision; Speech impairment; Could not walk; Paralysis in his head from neck up.; Pneumonia; Totally impaired; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (Paralysis in his head from neck up.), PNEUMONIA (Pneumonia), DYSPNOEA (Losing breathe), FACIAL PARALYSIS (Paralysis of face), BELL'S PALSY (Bells Palsy), GUILLAIN-BARRE SYNDROME (Guillain-Baire syndrome,), HYPOAESTHESIA (Numbness in hands), DYSPHAGIA (Could not swallow), LOSS OF PERSONAL INDEPENDENCE IN DAILY ACTIVITIES (Totally impaired), PARAESTHESIA (Tingly in hands), VISUAL IMPAIRMENT (Impaired vision), SPEECH DISORDER (Speech impairment) and GAIT DISTURBANCE (Could not walk) in a 61-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) for COVID-19 vaccination. The patient's past medical history included Plasmapheresis in April 2021, Feeding tube insertion in April 2021, Tracheostomy tube insertion in April 2021 and Mechanical ventilation in April 2021. On 16-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 13-Apr-2021, received second dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) dosage was changed to 1 dosage form. On 15-Apr-2021, the patient experienced PARALYSIS (Paralysis in his head from neck up.) (seriousness criteria hospitalization, disability, medically significant and intervention required), PNEUMONIA (Pneumonia) (seriousness criterion medically significant), DYSPNOEA (Losing breathe) (seriousness criteria hospitalization, medically significant, life threatening and intervention required), FACIAL PARALYSIS (Paralysis of face) (seriousness criterion hospitalization), BELL'S PALSY (Bells Palsy) (seriousness criterion hospitalization), GUILLAIN-BARRE SYNDROME (Guillain-Baire syndrome,) (seriousness criteria hospitalization, disability, medically significant, life threatening and intervention required), HYPOAESTHESIA (Numbness in hands) (seriousness criteria hospitalization and medically significant), DYSPHAGIA (Could not swallow) (seriousness criteria hospitalization and medically significant), LOSS OF PERSONAL INDEPENDENCE IN DAILY ACTIVITIES (Totally impaired) (seriousness criterion disability), PARAESTHESIA (Tingly in hands) (seriousness criterion hospitalization), VISUAL IMPAIRMENT (Impaired vision) (seriousness criterion hospitalization), SPEECH DISORDER (Speech impairment) (seriousness criterion hospitalization) and GAIT DISTURBANCE (Could not walk) (seriousness criterion hospitalization). At the time of the report, PARALYSIS (Paralysis in his head from neck up.) had not resolved and PNEUMONIA (Pneumonia), DYSPNOEA (Losing breathe), FACIAL PARALYSIS (Paralysis of face), BELL'S PALSY (Bells Palsy), GUILLAIN-BARRE SYNDROME (Guillain-Baire syndrome,), HYPOAESTHESIA (Numbness in hands), DYSPHAGIA (Could not swallow), LOSS OF PERSONAL INDEPENDENCE IN DAILY ACTIVITIES (Totally impaired), PARAESTHESIA (Tingly in hands), VISUAL IMPAIRMENT (Impaired vision), SPEECH DISORDER (Speech impairment) and GAIT DISTURBANCE (Could not walk) outcome was unknown. Non-drug therapy included: The patient had 7 plasmapheresis, trachea to breathe, feeding tube and put on a ventilator. 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-2021-141559 (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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1363986-1	Strength less than half in his right hand; Paralyses started 72 hours after vaccine; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (Paralyses started 72 hours after vaccine) in a 42-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 023C21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. No Medical History information was reported. On 15-May-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) 1 dosage form. On 15-May-2021, the patient experienced ASTHENIA (Strength less than half in his right hand). In May 2021, the patient experienced PARALYSIS (Paralyses started 72 hours after vaccine) (seriousness criterion medically significant). At the time of the report, PARALYSIS (Paralyses started 72 hours after vaccine) and ASTHENIA (Strength less than half in his right hand) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. No concomitant medication reported. Patient received covid vaccine on 15 May 2021. Per father son paralyses started 72 hours after strength less than half in his right hand. No treatment information was provided Company Comment : 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.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1368220-1	Core and lower extremity paralysis, speech impairment, presumed multiple sclerosis exacerbation, loss of ability to sit, loss of ability to stand, loss of ability to walk. Treatment: hospitalization, IV steroids, physical occupational and speech therapy. Other treatments as well. Outcomes as of 2 June 2021: No return to prior function, slight improvement in speech.
PARALYSIS	COVID19 (COVID19 (MODERNA)) (1201)	MODERNA	1372124-1	I had fever of 102, chills, headache, etc. I expected that. But I also had about five or ten minutes of total paralysis. I could not move. I could not lift my hand. Thankfully, it went away but I have been ?foggy? for the two and a half months since the second dose.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1373291-1	transverse myelitis; paralyzed waist down; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of MYELITIS TRANSVERSE (transverse myelitis) and PARALYSIS (paralyzed waist down) in a 69-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch no. 040A21A) for COVID-19 vaccination. No medical history was provided by the reporter. Concomitant products included DOCUSATE SODIUM (STOOL SOFTENER) for Constipation. On 09-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 MYELITIS TRANSVERSE (transverse myelitis) (seriousness criteria hospitalization prolonged, disability and medically significant) and PARALYSIS (paralyzed waist down) (seriousness criteria hospitalization prolonged, disability and medically significant). The patient was hospitalized from sometime in March 2021 to sometime in 2021 due to MYELITIS TRANSVERSE, and then on sometime in March 2021 due to PARALYSIS. At the time of the report, MYELITIS TRANSVERSE (transverse myelitis) outcome was unknown and PARALYSIS (paralyzed waist down) had not resolved. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Intramuscular) was unknown. Additional concomitant medications included a non-specific antidepressant and rosacea 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. Further information has been requested. Most recent FOLLOW-UP information incorporated above includes: On 25-May-2021: Non significant follow up; 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.
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1396638-1	Palsy. Drooping of lift side of face and twitching of eye and left eye doesn?t close. About a week and a half
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1398514-1	palsy after 1 week of taking moderna covid -19 vaccine; This spontaneous case was reported by a consumer and describes the occurrence of PARALYSIS (palsy after 1 week of taking moderna covid -19 vaccine) 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 PARALYSIS (palsy after 1 week of taking moderna covid -19 vaccine) (seriousness criterion medically significant). At the time of the report, PARALYSIS (palsy after 1 week of taking moderna covid -19 vaccine) outcome was unknown. The action taken with mRNA-1273 (Moderna COVID-19 Vaccine) (Unknown) was unknown. The patient experienced palsy a week after taking Moderna Covid-19 vaccine. No treatment or concomitant medications were reported by the reporter. 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
				Gets cold very easy; Out of breath; Strep through body; Swelled all over body/left foot still swollen; Weak; Paralyzed for over two weeks; Possible A-fib; Arthritis aggravated; Face was swollen; Hands were swollen; Not feeling well; This spontaneous case was reported by a consumer (subsequently medically confirmed) and describes the occurrence of PARALYSIS (Paralyzed for over two weeks), FEELING COLD (Gets cold very easy), DYSPNOEA (Out of breath), STREPTOCOCCAL INFECTION (Strep through body), SWELLING (Swelled all over body/left foot still swollen), ASTHENIA (Weak) and ATRIAL FIBRILLATION (Possible A-fib) in an 83-year-old male patient who received mRNA-1273 (Moderna COVID-19 Vaccine) (batch nos. 025A21A and 025A21A) for COVID-19 vaccination. The occurrence of additional non-serious events is detailed below. Concurrent medical conditions included Arthritis and Cane user. Concomitant products included HYDROCHLOROTHIAZIDE, FINASTERIDE, OLMESARTAN, SIMVASTATIN, OMEPRAZOLE MAGNESIUM (PRILOSEC [OMEPRAZOLE MAGNESIUM]), COLECALCIFEROL (VITAMIN D [COLECALCIFEROL]) and PARACETAMOL (TYLENOL) for an unknown indication. On 03-Mar-2021, the patient received first dose of mRNA-1273 (Moderna COVID-19 Vaccine) (unknown route) 1 dosage form. On 31-Mar-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 MALAISE (Not feeling well). On 04-Apr-2021, after starting mRNA-1273 (Moderna COVID-19 Vaccine), the patient experienced ARTHRITIS (Arthritis aggravated), SWELLING FACE (Face was swollen) and PERIPHERAL SWELLING (Hands were swollen). On an unknown date, the patient experienced PARALYSIS (Paralyzed for over two weeks) (seriousness criteria hospitalization and medically significant), FEELING COLD (Gets cold very easy) (seriousness criterion hospitalization), DYSPNOEA (Out of breath) (seriousness criterion hospitalization), STREPTOCOCCAL INFECTION (Strep through body) (seriousness criterion hospitalization), SWELLING (Swelled all over body/left foot still swollen) (seriousness criterion hospitalization), ASTHENIA (Weak) (seriousness criterion hospitalization) and ATRIAL FIBRILLATION (Possible A-fib) (seriousness criterion medically significant). The patient was hospitalized for 34 days due to PARALYSIS, then for 36 days due to ASTHENIA, DYSPNOEA, FEELING COLD, STREPTOCOCCAL INFECTION and SWELLING. The patient was treated with MORPHINE in April 2021 at an unspecified dose and frequency and Physical therapy for Paralysis. At the time of the report, PARALYSIS (Paralyzed for over two weeks) had resolved and FEELING COLD (Gets cold very easy), DYSPNOEA (Out of breath), STREPTOCOCCAL INFECTION (Strep through body), SWELLING (Swelled all over body/left foot still swollen), ASTHENIA (Weak), ATRIAL FIBRILLATION (Possible A-fib), ARTHRITIS (Arthritis aggravated), MALAISE (Not feeling well), SWELLING FACE (Face was swollen) and PERIPHERAL SWELLING (Hands were swollen) outcome was unknown. DIAGNOSTIC RESULTS (normal ranges are provided in parenthesis if available): On 04-Apr-2021, Hepatic enzyme: elevated (High) Patient experienced elevated liver enzymes.. In 2021, Heart rate: high (High) Heart rate- increased possible a Fib. Concomitant medication also includes an unspecified blood thinner. Patient was prescribed with Tramadol which was provided at discharge. 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, patient's hx of arthritis and being a cane user are confounding factors that may play a possible contributory role providing an alternative explanation. FU1 received additional serious AEs after second dose does not change company comment. Further information has been requested. This case was linked to US-MODERNATX, INC.-MOD-2021-067304, US-MODERNATX, INC.-MOD-2021-067204 (E2B Linked Report). Most recent FOLLOW-UP information incorporated above includes: On 07-Jun-2021: Follow up received on 07-JUN-2021, upgraded case to serious with the addition of event paralysis and hospitalization.; 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, patient's hx of arthritis and being a cane user are confounding factors that may play a possible contributory role providing an alternative explanation. FU1 received additional serious AEs after second dose does not change company comment. Further information has been requested. US-MODERNATX, INC.-MOD-2021-067304:First dose case US-MODERNATX, INC.-MOD-2021-067204:Crosslinked (family member)
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1401699-1	
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1403479-1	Beginning the first week in April developed headache, nausea, generalized abdominal discomfort, dizziness, paresthesias, dyspnea. Eventually developed severe epipastric pain for 5 days which was associated with left 6th intermittent palsy (normal eye exam.). Have continued with a dysequilibrium with a tendency to to the right. Sensation in legs that they are not feeling correctly and hence I walk in a wide-based gate. All laboratory has been normal or near normal except for triglycerides (500, non-fasting)
PARALYSIS	**COVID19 (COVID19 (MODERNA)) (1201)**	**MODERNA**	1404020-1	Difficulty breathing. Paralysis body aches and cramping. Fever. I was unable to breathe. Airway closed up and was unable to take deep breaths. I went to urgent care and the refused to see me as I the triage nurse said had a fever of 102 degrees Fahrenheit. I had severe bodily cramps and paralysis. I still have difficulty breathing and it's the 16th. The first dose also presented me w/ the inability to breathe however went away after about 7 days. This second dose seems to have given me ongoing lung issues.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0918637-1	Right sided jaw pain started 12/22/20 in the am progressed through the evening. When I woke up I had numbness and weakness in the right of my tongue which I quickly realized included the right portion of my face. Included tongue was numb and I couldn't move it. An Er trip and CT scan confirmed it was Bells Palsy. Today, My symptoms continue. Top of my head all the way to the way to bottom of neck is experiencing paralysis. Can move tongue now and it's partly numb but it's cooperating a bit more now. Nerve pain has worsened all over all the right side of face. Feels like someone has kicked face and I have a deep bruise. It's a deep ache. Nothing will take away the ache away. My eye doesn't lubricate like it should. So I have to either manually close it and wet it manually or tape it closed.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0926808-1	The morning after I received the shot in my right arm, I woke up and jumped out of bed to turn off my alarm. My right leg buckled and completely gave out. It felt like there was not a muscle or bone in the leg. I melted down on top of my leg, twisting my ankle and my knee. I ended up with my knee twisted behind me as I hit the floor and my nightstand. I don't know if you would call it muscle fatigue or weakness, it felt like muscle paralysis, but it only lasted less than a minute before it was gone, however, the damage was already done. I hurt my knee very badly because I had no control over my leg muscles. I've never had anything like that happen before.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0929154-1	"I was pretty sick, I felt palsy; I was pretty sick, I felt palsy; I developed the fever for all about 30 hours after the vaccination; Headache; This is a spontaneous report from a contactable physician. This physician reported for himself that a 67-years-old male patient received BNT162B2 (BioNTech Covid 19 vaccine; Batch/lot number: EL1284), via an unspecified route of administration on 30Dec2020 at SINGLE DOSE for ""Because I am a physician, a healthcare provider"" (covid-19 immunization). Medical history included had Covid back on the 4th of December. There were no concomitant medications. Physician stated he was wondering he just took your Pfizer Covid vaccination, the first one. He took it couple of days ago. He took it on the 30th of this month, two days ago. Physician further stated he developed the fever for all about 30 hours after the vaccination. Just because he developed the fever with the first one, it was like a 101.7 that was about 36 hours. So the question was just because he developed the fever with the first one, he was pretty sick, he meant he felt palsy, you know headache and all that it was like probably developed the same with the second one? Patient weight was maybe 125 pounds. When probed if vaccine was prescribed by any Physician, Physician stated he went himself. Causality by physician stated as ""he think so"". Lab work reported as He did not have the results yet. He just look a Covid screening test because he had Covid back on the 04Dec. So, he just took a Covid screening test. Treatment received included aspirin. The outcome of all events was unknown.; Sender's Comments: The causal relationship between BNT162B2 and the event palsy cannot be excluded as the information available in this report is limited and does not allow a medically meaningful assessment. This case will be reassessed once 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."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0932371-1	The patient received the vaccine in the left deltoid at 07:45 on 1/7/21. The patient reports that on 1/8/21 at around 0200, she was awoken with a sharp pain in her left arm followed by complete paralysis of her left hand for about 15 seconds. Did not re-occur. Did not seek treatment. Reported to Dr. and no further issues seen.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0940838-1	10 minutes after the shot was given, I suddenly felt my heart was racing and a tingling sensation started from my left arm and radiated to my whole body. The sensation was very intense it felt like my whole body was being paralyzed. I was fading out because of high heart rate. My heart rate was 160 and I heard someone said my blood pressure was high. I was transferred to the ER and given epinephrine, anti-histamine, steroids.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0946749-1	7:00PM fatigued, burning up fever 100., ibuprofen/tylenol dose; Sunday afternoon nausea, loss control of body, anxious, feeling of fainting, unable to move-paralyzed, pressed button for medical help, ambulance arrived, pt transported to ER -- 102. temp ambulance, RN at hosp temp 98., pt was shaky, 8:30PM erratic heartbeat per admitting doctor - pt admitted. Pt PCP/Cardiologist contacted, kept on heart monitor, pt discharged Monday afternoon. 1/14/21 chest pain, nausea, 102. fever. symptoms
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0960439-1	Cardiac event; Paralysis; Fever; Numbness; Chest Pains; Dizziness; Weakness; This is a spontaneous report from a contactable consumer (patient). A 30-years-old female patient started to receive first dose bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration in Left arm on 15Jan2021 14:15 at SINGLE DOSE for covid-19 immunisation. The patient was not pregnant. Medical history included tachycardia, Pre-ventricular contractions, allergies to Latex, covid-19 (reported as covid prior vaccination: Yes). Concomitant medication included metoprolol and multivitamin. No other vaccine received in four weeks. On 15Jan2021 14:30, the patient experienced chest pains, dizziness, weakness. On 15Jan2021 18:20, the patient experienced cardiac event, paralysis, fever, numbness, chest pains. The 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 the events. No covid tested post vaccination. The outcome of the events was recovering. Information on batch/lot number was requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0965215-1	Systemic: Patient experienced temporary paralysis from the shoulders down lasting approximately 20 minutes-Severe
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0976778-1	heart started to race; tingling sensation from her left arm that radiated to all over her body to the point where she felt paralyzed; tingling sensation from her left arm that radiated to all over her body to the point where she felt paralyzed; heart rate jumped to 160; heart rate jumped to 160 and blood pressure went up; choking sensation on her neck; This is a spontaneous report from a contactable healthcare professional. A 42-year-old non-pregnant female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, lot number EL1284/expiration date), dose number 2 via an unspecified route of administration on 12Jan2021 at a single dose on the left arm for COVID-19 immunization. Medical history was not reported. The patient did not have COVID prior vaccination. The patient previously took nitroglycerin and experienced drug allergy. Historical vaccine included BNT162B2 (lot number EL0140, dose number 1) for immunization on 22Dec2020 after which the patient experienced mild headache. Concomitant medications were not reported. There were no other vaccines received within 4 weeks. The patient had no adverse reaction other than mild headache after the first dose, so she was very relaxed and she felt comfortable getting the 2nd dose. However, after 10 minutes after the second dose, her heart started to race and she started to feel a tingling sensation from her left arm that radiated to all over her body to the point where she felt paralyzed. Her heart rate jumped to 160 and her blood pressure went up. She couldn't even open her eyes nor speak a word. She was transported to the ER and was given steroids and anti-histamine. She refused to get an epinephrine shot at first, but she started to feel choking sensation on her neck so she asked for an epi shot. The patient had not been tested for COVID post-vaccination. The outcome of the events was recovering.; Sender's Comments: Based on temporal association, a possible contributory role of BNT162B2 cannot be excluded for reported events heart racing, tingling sensation, paralyzed, heart rate high, blood pressure increased and chocking sensation. 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.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	0994675-1	"numbness has moved up her neck; she touches her neck, it hurts a little; maintains numbness and ""no control of her arm""/can barely move her right arm; paralyzed at the shoulder from the elbow up;it's numb; can barely move her right arm; paralyzed at the shoulder from the elbow up; flu like symptoms; Threw up a couple times; mild pain at the injection site; This is a spontaneous report from a contactable nurse reporting for herself. A 31-years-old female patient received the first dose of BNT162B2 Pfizer-BioNTech COVID-19 Vaccine, Batch/lot number: EL3247, Expiry date 31May2021, via an unspecified route of administration in right arm on 16Jan2021 at 07:30 at single dose for COVID-19 immunization as frontline worker. The patient had no relevant medical history. There were no concomitant medications. The patient received BNT162B2 Pfizer-BioNTech COVID-19 Vaccine in Hospital. The patient reported that she was fine all day Saturday (16Jan2021), she just had some very mild pain at the injection site started a couple hours after getting injection. She developed flu like symptoms on the following day (17Jan2021) at 18:00PM and threw up a couple times. On 18Jan2021 at noon she noticed that she could barely move her arm: anything above her elbow on her right arm was not moving; it was paralyzed at the shoulder from the elbow up. If someone were to pick it up and let go, she couldn't control it at all. She also stated that she had no pain at all in that arm; she couldn't feel anything at all; it was numb. When she woke up on 19Jan2021 the numbness had moved up her neck, instead of being localized to her arm. She stated that when she touched her neck, it hurt a little, but it was not super significant. She stated that it was debilitating at that point as she was maintaining numbness and ""no control of her arm"". The nurse reported that this event had not required yet a visit to physician, but she has a feeling it will. She will probably have to go somewhere due to her paralyzed arm, but she had not yet. At the time of the report the event ""No control of her arm""/can barely move her right arm; paralyzed at the shoulder from the elbow up; it's numb"" had not resolved yet; the event ""mild pain at the injection site"" resolved on 17Jan2021, the events ""Flu like symptoms"" and ""Threw up a couple times"" were recovering and outcome of the events ""Numbness has moved up her neck"" and ""She touches her neck, it hurts a little"" was unknown. The reporter assessed the events ""Pain injection site"" and ""Flu like symptoms"" as non-serious and assessed the event ""Can barely move her right arm; paralyzed at the shoulder from the elbow up; it's numb"" as disabling. She assessed all the reported events as related to BNT162B2 Pfizer-BioNTech COVID-19 Vaccine (method of assessment: Global Introspection).; Sender's Comments: Based on the temporal relationship, the association between the events numbness of upper extremity and arm paralysis with BNT162b2 use 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
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1011604-1	Temporary paralysis right side (face, arm & leg). Lasted approximately 4 hours.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1013160-1	Note:Tested positive to CV19 27th October 2020 Pharmacy. CV19 Symptoms: High fever and fatigue. No loss sense of taste and/or smell. No breathing problems. 5th February 2021 Pfizer 1st dose CV19 vaccine. Scary symptoms started in grocery store parking lot around 6PM (after vaccine at 8:35A) my legs felt like they were partially paralyzed and I would not be able to walk to the car. I struggled to walk. Got home and laid down. Thought it was side effects to vaccine and it would pass if I went to sleep. Woke up and felt totally paralyzed and it required a lot of effort to turn on my side to try to reach my phone. All my muscles and joints ached. I am never sick and not a complainer.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1016727-1	"Pt. describes severe pain in her left arm within 12 hours of administration. She also reports a feeling of ""paralysis"" and severe ""sluggishness."" She has mental foginess. Her symptoms lasted 4-5hrs. This occurred again about 24hrs after administration."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1022718-1	Systemic: Other- paralysis. Stroke ruled out at Emergency room. symptoms lasted 2 days and began morning after immunization
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1028519-1	Fever, chills, muscular pain, chest pain, shaking, paralyzed, vomiting
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1035517-1	"psychedelic feeling; horrible palsy, as if her muscle was being pulled like a rubber band; horrible palsy, as if her muscle was being pulled like a rubber band; the worse bowel movement. She had to go the bathroom a lot; Her right arm was the worse with the pain and then her hands would kind of move in a kind of crazy way where it was tightening; muscle cramping and pulling; Her toes would contort and go backwards.; she feels like the left arm is going to do something, its like the last to be pulled; shingles; muscles became tight/contort; minor nausea; Stomach cramps; Gas; Indigestion; became weird in her head, almost like a lightheadedness; This is a spontaneous report from a contactable consumer. A 73-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EL9261), intramuscular on 30Jan2021 at 10:00 at a single dose on right arm for COVID-19 immunisation. Medical history included fever, being ill (deathly ill and it lasted for three weeks) and bug bites from an unknown date and unknown if ongoing. There were no concomitant medications. The patient is the type of person that takes a lot for her to take medications. She is allergic to a lot of things; she always has crazy reactions. She can't take morphine. It took her 7 pills to get the right one for high blood pressure. She has the craziest one-day reactions. She received the COVID-19 vaccine on Saturday much against her will, knowing she had COVID but not diagnosed in March. She had a fever, she was deathly ill and it lasted for three weeks. She had a nasal swab done and it came back negative. She doesn't think that was right, she believes she had COVID-19 in Mar2020. Her daughter is younger, and she had the same thing, she couldn't breathe good for 3 weeks. She had pneumonia. She knows she had COVID-19. She was trying to lose weight. She was given her first COVID-19 vaccine by a doctor. Everything was going okay at first. About an hour and a half later, she became weird in her head, almost like a lightheadedness. She can't describe it. She has never taken drugs. She doesn't drink alcohol at all. She has like a virgin body. She states plus she is red headed, which it is known red heads have a harder time with medication. As the day goes on, she just felt lightheaded all the rest of the day. The next day (Sunday, 31Jan2021), she wakes up and it hits her like a ton of bricks. She woke in the morning and seemed to be okay and then it was like she was pregnant and blew up. She started cramping really bad and had gas. Everybody has gas. She was a little nauseated as well. She experienced lots of indigestion, cramping in her stomach like she was going to have a baby or the worse bowel movement. She had to go the bathroom a lot. She had a little bit of nausea. She had nausea pills because she had a gallbladder taken out and she took nausea pills for 2-3 days. She didn't take any medication for any of these symptoms she started to experience on Sunday. However, she did start popping Gas-X. She was taking two Gas-X every 3 or 4 hours. The caller states it helped. She had lots of gas; gas was just coming out of her bottom. She had the gas, cramps, her stomach hurt. She was able to get through it, which that is good. Then Monday, 01Feb2021 was the day she had weird things happen to her that she was kind of concerned about. Her muscles started to draw, that is the best way she can describe it. Her feet would just draw up weird. Her arms and hands would do crazy things and contort or draw up. This hurts because the muscles are like a rubber band pulling. It was weird, and she didn't like it. It wasn't like when she has a horse and has to get up and stand, it wasn't like that at all. Her muscles were tightening up like a rubber band. Her right arm was the worse with the pain and then her hands would kind of move in a kind of crazy way where it was tightening. She thought to herself, ""what is this Parkinson's?"". She doesn't even know. She still has some stomach issues, but it is easing up. The gas is still coming out. It wasn't debilitating or anything. She has some minor nausea. What scared her more was the muscle cramping and pulling. Her toes would contort and go backwards. The muscle tightness occurred yesterday, 01Feb2021. It happened all day and was sporadic. Her toes moved in a different way like a

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				happened all day and was sporadic. Her toes moved in a different way like a rubber band pulling on muscle. Nothing has happened today. Everything was yesterday. Her whole body basically was affected, just different sections at different times. Today, she feels like the left arm is going to do something, it's like the last to be pulled. Her hands and fingers stayed the same. She explains like the upper arm area from her elbow to her shoulder part. She can't even think of what that area is, but it's the part between her elbow and shoulder. She has been doing lots of exercises and moving, hoping the part of her elbow to her should don't atrophy or won't pull. She hasn't told anybody about it. She doesn't like it. It was very frightening. She is scheduled for her second shot and she is scared to death to go back. She mentions her nerves are shot. From stress, she had previously broken out instantly. She thought it was a bug biting her. She never had gone to the doctor but after about 4 weeks, she thought she had these horrible bug bites. She was shaking her clothes and she almost wanted to strip it was so bad. She thought it was severe bug bites. She couldn't figure it out. She finally went to the doctor after a week. It was a Saturday. The doctor told her it was shingles. Explained that it itches a lot. She went on medication. She clarifies she went on acyclovir 800mg, three pills a day. She called her doctor knowing that she had shingles and how scared she is with medicine, the doctor assured her she could get the COVID-19 shot as scheduled. The medicine for shingles (acyclovir) was finished up the Friday before she received the COVID-19 vaccine. She got the shingles shot at 60 but everybody is now saying to get the new shot for shingles which she has never done obviously since she got the shingles. She plans to get the shingles shot in the summer. She is not worried about it. She took 3 pills a day and that was fine. She is fine, meaning every day she did normal things going through the whole process. She states she plans to go to the bank. She asked if anybody has reported the reaction she experienced after receiving the first dose of the Covid vaccine. She received the first dose of the vaccine last Saturday, in which within an hour and a half her ""brain got weird, like a psychedelic feeling and it felt like I was getting an injection in the operating room to go to sleep."" On Monday she experienced, ""horrible palsy, as if her muscle was being pulled like a rubber band."" Stated that it wasn't a horse feeling. Her hands and fingers would pull and contort in different positions, in which they would remain tight and stay in that position, and then she would be able to make a closed fist. On Monday night, her left arm contorted (which is not administration arm) and her muscle became really tight from her elbow to her wrist leading to her hand contorting as well. She describes the muscle tension as if a rubber band was being pulled, it would have burst. This muscle pulling hurt, in which she took 3 ibuprofen (ADVIL). She has crazy reactions with almost anything that she takes. She has never smoked or drank. The patient underwent lab tests and procedures which included COVID-19: unknown on an unspecified date and unknown on Mar2020, SARS-COV-2 test: negative on an unspecified date. Therapeutic measures were taken as a result of gas (flatulence), muscles became tight/contort, minor nausea, horrible palsy, as if her muscle was being pulled like a rubber band and shingles. The outcome of the event 'the worse bowel movement. She had to go the bathroom a lot' was not recovered while of the rest of the events was unknown."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1035537-1	developed paralysis in the opposite part of the body; Her hand, fingers and knee all became swollen; Her hand, fingers and knee all became swollen; This is a spontaneous report from a contactable consumer via Pfizer Sponsored program Pfizer First Connect. A female patient of an unspecified age received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 02Feb2021 at single dose for COVID-19 immunization. The patient medical history and concomitant medications were not reported. The patient took the first dose of the vaccine on an unknown date for COVID-19 immunization. In Feb2021, about 12 hours after getting the vaccine, the patient developed paralysis in the opposite part of the body where she didn't get the vaccine. Her hand, fingers and knee all became swollen. She had the symptoms on her right and was given the vaccine in her left. The final outcome of the events was unknown.Information on lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1036675-1	61 yo F with history of bilateral lung transplant 6/23/17 presented to ED on 2/4/21 with chief complaint of worsening shortness of breath, nausea and diarrhea for past week since receiving since receiving COVID-19 vaccine (Pfizer) on 1/28/21. Upon arrival to triage she was obviously dyspneic with significantly low oxygen saturations. O2 sats on arrival were 65%, improved to mid 90's with O2 6 liters per NC. Admitting diagnosis: hypoxic respiratory failure post COVID vaccine. Lab work shows an elevation of the BUN and creatinine at 31 and 1.71 which is slightly higher than her usual baseline levels. BNP is elevated at 2 448 with a mildly elevated troponin. Procalcitonin is also elevated. Patient's white blood cell count is 11.07. Full viral panel including COVID-19 is not detected. All blood cultures and respiratory cultures were negative. Patient chest x-ray shows numerous bilateral patchy opacities which is significantly different from her previous chest x-ray here. Empiric rejection treatment initiated including high dose methylprednisolone, plasmapheresis, IVIG, Thymoglobulin. She continued to decline and ultimately required intubation, proning and paralyzing on 2/8/2021 and then VV ECMO cannulation on 2/13/2021. EGD done 2/14/2021 as unable to pass the TEE probe during cannulation prior day (unable to complete due to abnormal anatomy). Acute pupil exam change in the early am hours of 2/15/2021 prompted urgent head CT which revealed catastrophic brain bleed. Brainstem reflexes were lost soon after. Despite placing an EVD emergently at bedside, brain stem reflexes were not recovered. GOL engaged and patient not an organ donation candidate. Therefore discussion with sister at bedside resulted in decision for cessation of life support. Patient expired shortly after support withdrawn and pronounced dead on 2/15/2021 at 11:11 AM.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1042343-1	could not lift arm/move body; her tongue started to be tingly; fuzzy feeling; tongue swollen a bit; hives; hard to swallow; weird taste in mouth; a high grade fever over 103, went down on 03Feb2021 to 102, 101, 99, then 98.6; chills uncontrollably; heart racing a bit; joint-muscle pain; headache; dizziness; swollen lymphnodes; red rash on chest/arms/stomach with some hives; This is a spontaneous report from a contactable consumer reporting for herself. This 42-year-old non-pregnant female patient received the second single dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot EN9581) on 02Feb2021 at 10:30, in left arm, for COVID-19 immunisation. The first BNT162B2 vaccine dose was administered on 12Jan2021 at 13:30, lot number EL3246, in left arm, for COVID-19 immunisation. Medical history included heart murmur, acute sinusitis, allergies to acetaminophen/oxycodone (PERCOCET), cephalexin (KEFLEX), codeine, trees, pollen, grass, mold, cats, some dogs. The patient had no COVID prior to vaccination and COVID was not tested after vaccination. Concomitant medications were not reported. The patient received the second vaccine dose on 02Feb2021 at the hospital and was monitored for 30 minutes. Within that time her tongue started to be tingly, she experienced fuzzy feeling, tongue swollen a bit with hives, hard to swallow, weird taste in mouth. The nurse on staff walked her to the ED, she was admitted and treated for allergic reaction to the vaccine and received prednisone for 5 days and diphenhydramine (BENADRYL). Epipen if needed. Later that night, on 02Feb2021, the patient had a high grade fever over 103, chills uncontrollably, heart racing a bit, could not lift arm/move body, joint-muscle pain, headache, dizziness, swollen lymphnodes, hard to swallow, every side effect on sheet just really bad. Her fever eventually went down on 03Feb2021 to 102, 101, 99, then 98.6. She still had all other symptoms not as severe, later that night she had a bit of a red rash on chest/arms/stomach with some hives, swallowing was still a bit hard. She continued to take BENADRYL and steroid. 06Feb2021 was her last day of steroid. As of 06Feb2021 she was feeling a bit better with no fever, swallowing a bit hard but other symptoms were less. The events resulted in Emergency room/department or urgent care. Fever resolved on an unspecified date in Feb2021. The other events were resolving.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1054524-1	Headache, worsening of his neurologic status, dizziness, change in gait, shuffling feet, then unable to walk, more sleepy, unable to move extremities, unable to speak, tremors
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1055759-1	Loss of use right leg. Paralysis, numbness, tingling, severe pain, and muscle spasms. Acute onset without other trauma
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1056639-1	"Left eye 6 nerve palsy and could not move my eye laterally; This is a spontaneous report from a contactable physician (patient) A 77-year-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection) on 27Jan2021 at single dose via an unspecified route of administration for COVID-19 immunization. Relevant medical history and concomitant medications were not reported. On 01Feb2021, patient started to notice, left eye 6 oculomotor nerve (""abduscens"") palsy and could not move his eye laterally. Patient also asked information about 6 oculomotor nerve and possible interaction with vaccine. At the time of the reporting event outcome was unknown. Information about 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 event cannot be excluded, due to 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 to Regulatory Authorities, Ethics Committees and Investigators, as appropriate."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1056640-1	Lt parietal occlusion; DVT; Right paralysis; This is a spontaneous report from a contactable Nurse reporting for her husband. A 71-years-old male patient received the first dose of bnt162b2 (BNT162B2; Lot # EL 1284) vaccine , intramuscular in the left deltoid on 22Jan2021 17:00 at single dose for Covid-19 immunisation . The patient medical history was not reported. Concomitant medication included apixaban (APIXABAN), acetylsalicylic acid (ASPIRIN) atorvastatin (ATORVASTATIN), cyanocobalamin (CYANOCOBALAMIN), metoprolol tartrate (METOPROLOL TARTRATE) , pantoprazole (PANTOPRAZOLE), sumatriptan (IMITREX [SUMATRIPTAN]), triazolam (TRIAZOLAM). The patient experienced DVT (deep vein thrombosis) on 26Jan2021 with outcome of not recovered , left parietal occlusion (ischaemic stroke) on 26Jan2021 05:30 with outcome of unknown , right paralysis on an unspecified date with outcome of unknown. The patient was hospitalized for DVT (deep vein thrombosis) and stroke from 26Jan2021 to 30Jan2021. The patient underwent lab tests and procedures including blood pressure diastolic: 84 mmhg on 30Jan2021 , blood pressure systolic: 141 mmhg on 30Jan2021 , body mass index: 26.4684 kg/m2 on 26Jan2021 , body temperature: 98.2 ¦F on 30Jan2021, heart rate: 55 bpm on 30Jan2021 , magnetic resonance imaging: acute left parietal lacunar infarct, Lower extremity ultrasound: left popliteal vein DVT, oxygen saturation: 95 % on 30Jan2021 , respiratory rate: 18 br/min on 30Jan2021. The reporter considered the reported events to be possibly related to BNT162B2 vaccine. Follow up information has been requested.; Sender's Comments: Based on the limited information currently available, a possible contributory role of the suspect drug in the reported events cannot be completely excluded given the known suspect drug profile and/or implied 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 Regul atory Authorities, Ethics Committees and Investigators, as appropriate.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1062962-1	[COVID-19 mRNA vaccine (Pfizer-BioNtech] treatment under Emergency Use Authorization (EUA)
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1067034-1	"Pt reported he got the second COVID vaccine at 8:33 am at the Center at an event run by Health Clinic. At 10 am he began to feel unwell describing ""disorientation"" which upon further explanation seems he is describing difficulty with his proprioception, ie not able to tell where his body is in space, and disequilibrium, ie having to stop and hold on to something while walking. This lasted most of the day and subsided by evening. He has continued to have ongoing shorter episodes 3-4 times per day lasting 5-10 minutes. In addition he has noted increasing weakness in his right thigh, particularly when going from sitting to standing and the through swing while walking. He describes these as ""paralytic attacks"" that occur once or twice per day and last hours. Both symptoms are similar to those he had post CVA in 2019 and greatly improved with PT and home exercises. In addition he has noted fecal urgency and fecal incontinence in the last couple of weeks, he is unsure if this is directly related to the vaccine."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1070730-1	hands, feet, toes and entire body, felt almost paralyzed; not being able to turn over in bed; could not bend her fingers or toes; muscles throughout her body remained stiff and tight; could not bend her fingers or toes; muscles throughout her body remained stiff and tight; tense muscle aches; Chills; headache; This is a spontaneous report from a contactable consumer (patient). This consumer reported that a 72-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EN6200) at the age of 72-years, via an unspecified route of administration in left arm at 11:00 AM on 16Feb2021 at single dose for COVID-19 immunisation, in hospital. Medical history included pituitary microadenoma from 1969 (52 years) and in good health other than the fibromyalgia; allergies to latex. Patient was not pregnant, was not pregnant at the time of vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID-19 vaccine. Prior to vaccination, the patient was not diagnosed with COVID-19. Concomitant medications included desipramine; duloxetine hydrochloride (CYMBALTA); ergocalciferol (VIT D); zinc. The patient previously received first single dose of BNT162B2 (lot number: EK9231) at the age of 72-years via an unspecified route of administration in left arm at 04:00 PM on 25Jan2021 for COVID-19 immunisation. Patient reported that she had been diagnosed with fibromyalgia and saw a rheumatologist that specializes in treating fibromyalgia. At 01:00 AM on 17Feb2021, fourteen hours later, she awoke to not being able to turn over in bed. Her hands, feet, toes and entire body, felt almost paralyzed. She could not bend her fingers or toes. Her muscles throughout her body remained stiff and tight for the next 16 hours. Chills, headache but no fever. Patient took paracetamol (TYLENOL) every four hours, during this time. It did not help with the tense muscle aches. Later on 17Feb2021 (that same day), approximately 19:00 (7:00 PM), her muscles started to relax. Patient was writing this the day after and she felt fine. She could hardly wait for the booster shot. Outcome of the events was recovered/resolved in Feb2021. Since the vaccination, the patient has not been tested for COVID-19.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1078523-1	Extreme shaking, leg weakness, slight fever. I couldn?t move at all. It was like I was paralyzed.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1081039-1	Patient presented with acute onset of rapidly progressing ascending weakness and paralysis and associated decrease in sensation
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1084251-1	I got paralyzed .. could not stand on my legs ... I was wobbly
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1090776-1	beeps, noise in her ears; muscle and joint pain; muscle and joint pain; tongue was a little numb; face lips; felt paralyzed/Felt like I couldn't move it like paralyzed a little bit; whole left side of body started going numb/Left side was numb around 7 or 8 that night/all left side was numb; pains in all of her face; pain and soreness in arm/arm was painful; tachycard, high heart rate; very tired, sleepy; very tired, sleepy; headache, heavy in her head; pain in both ears; This is a spontaneous report from a contactable consumer (patient). This 71-years-Old female patient received the first dose of (Pfizer Biontech Covid 19 Vaccine, Lot No. EN6198) intramuscularly at single dose on 23Feb2021 for COVID-19 immunisation. Relevant history and concomitant drugs were unknown. Past dug included flu vaccine in Sep2020. The patient received her first dose of Covid 19 vaccine on 23Feb2021 at 3:30 in the afternoon. She came home and felt, not bad at any time but more or less towards 6:00 or 7:00 at night she started feeling bad. Her whole left side of body started going numb around 7 or 8 that night. Her arm was painful, face lips and all left side was numb. Felt like she couldn't move it like paralyzed a little bit. Clarified as whole left side, even her leg, her feet everything on that left side. She started feeling pains in bones of head, especially her face, clarified all of her face and her left side was numb even her lips on the left side were numb. Her tongue was a little numb. She had muscle and joint pain. The pain, of course, in the arm, but that was normal. Clarified pain in the arm as the injection site. She didn't feel any swell there. She did not take medication for anything. She used a lotion. The lotion was trividol aloe vera arnica caleudula and helped to relive the pain. The arm was still a little painful but it's not bad but that feeling made her tachycard, clarified as high heart rate, feeling like boom boom boom, but she did not have any shortness of breath. The high heart rate was gone the same day. She had no fever or chills. She felt very tired and slept for about 10 hours but during the sleepy she started feeling a headache she couldn't get up because she was so tired and she felt pain in both ears. She had not been to the doctor in a while. She was still old fashioned so she carries everything in a little telephone book. While trying to read information caller states she had to change her eyes. She still felt numbness in left side of left leg but she thought it might be because she had not been exercising. When she was waking up around 9am in the morning, it was yesterday that she felt better in regards to the paralyzed feeling. When pains in her face came she felt and she thought what is this? It only lasted for a little while. It was during the night. The pain and soreness in arm at injection site is still there, she did not feel any swelling, or hot at that site. She confirmed she still felt a little tired, sleepy. It's getting better. She may feel a little heavy in her head and would like to go to sleep, but she was not sleeping. She had many things to do and knew she had to be up. The headache went away and was gone on 24Feb2021. It was persisting for a while, not too hard or heavy, but it resolved by 2 or 3 in the afternoon on the 24th. In regarding to pain in ears she mentioned she no longer had. She had beeps, noise in her ears even at the time she went to bed it was better. The outcome of events paralyzed, pain in arm, tired, sleepy, was recovering, the outcome of events tachycard, pains in all of her face, pain in both ears was resovled, the outcome of other events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1091676-1	optical neuritis; inability to move, chew, eat, etc.; inability to move, chew, eat, etc.; neurological damage; unable to speak; changes in her eyes; Her right eye was bulging; Neither eye looked in the same direction, or able to track objects; known allergies: Apparently the Covid Vaccine by Pfizer; This is a spontaneous report from a contactable consumer. The consumer reported two reports for two separate vaccine doses for the same patient. This is the first of two reports. A 74-year-old female patient (not pregnant) received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, lot/batch number and expiration date not provided), via an unspecified route of administration on 28Jan2021 at single dose for COVID-19 immunisation. Medical history included advanced stages of Multiple Sclerosis, Diabetes, depressant, covid prior vaccination. The patient's concomitant medications included diabetic meds (pills), anti-depressants, etc. The patient previously received the first dose of BNT162B2 in Dec2020 at age of 74 years old for COVID-19 immunization and experienced neurological setback and it wasn't terribly obvious at that time. Facility type vaccine was Nursing Home/Senior Living Facility. No other vaccine in four weeks. The patient (mother of the reporter) suffered a neurological setback following the first injection, although it wasn't terribly obvious at that time, in comparison to what occurred after she received the second vaccine. The patient suffered major neurological damage after the second vaccine; unable to speak, followed by an inability to move, chew, eat, etc. The reporter also noticed optical neuritis or changes in her eyes. Her right eye was bulging. Neither eye looked in the same direction, or able to track objects. The patient was currently in a long term acute hospital (pending clarification) with a tracheostomy and a food peg. She had advanced stages of Multiple Sclerosis. Reporter had since learned she had been advised by her MS doctors NOT to receive flu shots. The reporter had known this and would not have listened to the doctors at the nursing home, who recommended she receive the Covid Vaccine. This was serious. The reporter was awaiting medical records, which would provide exact dates, times, etc of the administration of the vaccines. Adverse events resulted in Disability or permanent damage. Known allergies: Apparently the Covid Vaccine by Pfizer. If treatment adverse events: unknown. The outcome of the events was not recovered. The patient underwent lab tests and procedures which included unknown covid test: negative in Jan2021 (post vaccination). The outcome of the events was not recovered. Information on the lot/batch number has been requested.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1092879-1	(3/10-3/11) Fatigue, headache & body aches on day of vaccine and overnight (3/11) 12:00pm lightheadedness and fatigue (took nap) (3/11) 3:10pm overall weakness (took 99mg Potassium tablet) and sudden onset of paralysis that lasted for 1 1/2 hrs (3/11) 6:05pm still feeling fatigued and a little weak, but strong enough to get out of bed on own
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1098619-1	Pain and paralysis in all 4 extremities. Burning and stabbing sensation, bulging veins in hand. Was hospitalized from 2/28 until 3/5. Left hospital with no diagnosis and no real relief
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1106603-1	100 year old patient in reasonable health (reading, socializing, doing Zoom calls, etc.) took second Pfizer vaccine on February 5, 2021. On the morning of February 22, 2021 the patient suffered a major hemorrhagic stroke. He suffered severe paralysis, could not speak, and suffered from severe pain. Within 24 hours he was moved to Hospice Care. A day later on February 24, 2021 he died.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1121599-1	I'm paralyzed; Anxiety; chills; Shaking but not fever; This is a spontaneous report from a contactable consumer (patient). A 48-year-old female patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot Number: EN6205), via an unspecified route of administration on 06Mar2021 as single dose for COVID-19 immunization. Medical history included hypertension, brain injury (doesn't have a good memory), menopause and menopausal problem. The patient is on a lot of medications (unspecified medication) for her Brain injury. The patient previously received the first dose of BNT162B2 on unspecified date for COVID-19 immunization. The patient had her second dose on 06Mar2021 and had been having bad like chills and shaking too but not fever. The patient was shaking bad (not clarified) there but she doesn't know if it's part of her menopausal problem from my menopause (as reported) but she was shaking. The patient mentioned that she was paralyzed. The patient stated that she was taking some Ativan for anxiety, she has been so anxious; and she had been taking Benadryl. The outcome of the events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				mild peripheral neuropathy/sensory peripheral neuropathy around her gums and lips on her left side, as well as tingling on 4 of the 5 fingers of her left hand; paralysis in her lips and gums all the way down to her finger tips on the side of the injection which was the left side; portion of the trigeminal nerve on the left side seemed to have been impacted; blood pressure had sky rocketed; This is a spontaneous report from a contactable physician and a contactable consumer (patient). A 71-year-old female patient received first dose of BNT162B2 (Pfizer-Biontech Covid-19 Vaccine), lot no. EL9269, via an unspecified route of administration on 13Feb2021 at a single dose for COVID-19 immunisation. Medical history was not reported. There were no concomitant medications. The physician asked if there were any reports of paresthesia around the gums and/or tingling in the hands and fingers. The patient received the first dose of the Pfizer COVID-19 vaccine. About 1 week after receiving the vaccine (20Feb2021), she experienced symptom complex, consistent with a mild peripheral neuropathy/sensory peripheral neuropathy around her gums and lips on her left side, as well as tingling on 4 of the 5 fingers of her left hand, She was admitted and a full workup for central disorders and vascular disorders was negative (Feb2021). But her symptoms were continuing intermittently. She had numbness of the gums. A portion of the trigeminal nerve on the left side seemed to have been impacted. Also, peri-orally on the left side. The physician asked if this symptom complex has been reported and if there were reports of these mild sensory peripheral neuropathies coming from the vaccine. The physician stated that any info on dental paresthesia or any reports of other incidents of sensory paresthesia would be very helpful. The patient also called about the COVID 19 vaccine. On 13Feb, she got the vaccine in and then one week later on the following Saturday, the date would have been 20Feb, she started having a sensation of tingling and paralysis in her lips and gums all the way down to her finger tips on the side of the injection which was the left side and that has continued. She went to see the doctor the following Monday 22Feb and not knowing what it was, she sent her to the ER she was kept overnight, she was discharged 23Feb but her symptoms persisted even through today. These sensations were happening on average 3 times per day, sometimes many more anywhere from 3 minutes to 1 hour 45 minutes then it goes away. She did not go to her primary care. She confirmed her first dose was given 13Feb and the tingling and paralysis started the 20Feb it was still ongoing at this time. It has persisted but she can't tell whether it was getting better or not because sometimes it will last 3 minutes sometimes it will last 1 hour and 45 minutes. It started to effect other parts of her so she thinks it might be getting a little bit worse. It was starting to effect other parts of her body the progression goes, it started at the tip of her tongue and gums and then shortly within 30 seconds or less than a minute it spread to her upper and lower lip only on the left side and that was how it was on her gums as well it was only on the left side then she had the tingling in her finger tips on the left hand. She specified that she experienced these sensations on the tip of her tongue, left gums, lower left lip, and into the fingers on the left hand. She would like to know if these symptoms have been reported in association with the vaccine and if so, how long she can expect them to last. She stated that yesterday, a physician called in on her behalf for more information and said he received literature from 29Jan regarding reports that included information about dental paresthesia after the vaccine. She was planning not to get the second vaccine. She was not taking medications, she was in good health she did go and was admitted in the ER, they did a work up. Her blood pressure had sky rocketed on Feb2021 and since she mentioned sensation they ruled out a stroke. On Feb2021, they did blood work, MRI and CT scan and everything was they did not find anything linking to a stroke. She spent the night in observation. She was in the ER, they did put her in the main hospital but it was for observation from the emergency room. She went in on the 22Feb and was discharged home on 23Feb. She asked where this goes from here, these symptoms were concerning to her. She did not find any incidences at least in general of reporting this, asked if there were these kinds of things being reported, she didn't know how long it was going to last or if it will get worse. The outcome of blood pressure had sky rocketed was unknown while outcome of other events was not recovered.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1123133-1	
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1127357-1	While I was waiting during the 15 minute window, I started to feel light headed and saw black spots. I told the nurse I didn't feel well and wanted to wait a little longer. Within 10 minutes or so, I started to experience tingling sensation all over my face and then it spread to my neck and then to the rest of my body. My face started to feel stiff and I became really hot. My tongue felt swollen and my entire jaw felt very stiff. I lost my ability to speak and then eventually had full body paralysis. I could not move any limb. I could hear and see, but I could not verbalize a response or move any part of my body. I remained in this state for about 1.5 hour. Afterwards, I regained my motor skills and my speech.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1127472-1	Fever, low oxygen saturation, severe encephalopathy, kidney failure, bilateral pulmonary infiltrates, sepsis, tachycardic, acidotic, intubated on ventilator since admission, paralyzed/sedated, ABG results showed high CO2 and O2 retention, edema, electrolyte imbalance, ARDS, low hemoglobin and hematocrit levels, blood transfusion needed
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1127584-1	Paralysis to bilateral hands for 2-3minutes
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH))**	**PFIZER\BIONTECH**	1128853-1	"palsy; She was having a lot of congestion; Tingly feeling in the right side of the face/tingling/tingling feeling through her arm; Hypersensitivity in the face; Felt pain rise from the right side of her neck to her head; Blood pressure went up/blood pressure was 153/92/Her blood pressure was

Symptoms	(1200) Vaccine	Vaccine Manufacturer	VAERS ID	checked three times and the lowest was 142/90; Felt dehydrated; She felt **Adverse Event Description**
				hot/feeling extreme heat on her face/burning; Thought she had a little sweat.; Red flushness on her chest; Tingling feeling under her throat; Face swelled up a little bit; Felt numbness across her face; She felt a slight itchiness on the right side of her face; Her face looked very red, specifically her cheeks and forehead; Fatigued, she felt like she was drugged; She couldn't fall asleep; Extreme GERD; Extreme racing heartbeat; This is a spontaneous report from a contactable consumer via Medical Information Team. A 48-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection, Lot number was not reported), dose 1 via an unspecified route of administration (reported left side) on an unspecified date as single dose for COVID-19 immunization. Medical history included felt a little hot prior to the COVID vaccine from an unknown date, neck pain, extreme GERD, extreme racing heartbeat, right eye proptosis, and immune issues from an unknown date. Concomitant medication(s) included oxymetazoline hydrochloride (CLARITIN ALLERGIC); levothyroxine sodium (SYNTHROID); liothyronine sodium; hydroxychloroquine sulfate (PLAQUENIL S); and famotidine (PEPCID), all taken for an unspecified indication, start and stop date were not reported. The patient previously took diphenhydramine hydrochloride (BENADRYL) and had a side effect. On an unspecified date, the patient reported having these feelings in both arms, more so a tingly feeling in the right side of the face, and hypersensitivity in the face. She had a pain that goes from the right side of the face to the back of the brain. Her blood pressure went up. She also had a lot of liquid, and felt dehydrated almost felt like her right side was going to drool. She wanted to know if this could be palsy or a nerve condition. She stated she's concerned about getting second dose because she was still having these feelings. The patient wanted to know if it would be safe to use the EpiPen her Doctor prescribed to her, since her blood pressure went up high, instead of going down, like it usually would happen in the severe cases of allergic reactions; for that and the tingly face, the doctor recommended that she receive the second dose in the hospital. She stated she received her first dose at a convention center and wanted to know if Pfizer could provide her with a letter saying she would need to receive her second dose in a hospital. She was supposed to get the second vaccine on Thursday (unspecified date). She was concerned because of a reaction she immediately had and has had since that timeframe. She felt a little hot prior to the COVID Vaccine. After she got the COVID Vaccine, she felt hot. She was sitting there waiting and felt a little hot on her forehead. She thought she had a little sweat. She also thought she was being crazy because she was hot before she got to the COVID vaccine location. Twenty-nine minutes in she started feeling extreme heat on her face. She was burning and tingling. She felt she was having a lot of congestion. She saw her face turned very red. She had to use a lot of ice because it felt like she was on fire. She was there for two hours being monitored. Her blood pressure was 153/92 when it normally was 100/70. Her blood pressure was checked three times and the lowest was 142/90. She stated she is not allergic to Benadryl, but when she was a little kid she had a side effect so she doesn't use Benadryl. She was given Pepcid immediately after COVID Vaccine. She had red flushness on her chest. She had a tingling feeling under her throat. Her face swelled up a little bit, but enough where she could feel it. She stated she had neck pain prior to the COVID vaccine. Fifteen minutes after the COVID vaccine, she felt pain rise from the right side of her neck to her head. She also felt numbness across her face. It didn't go away. After two and a half hours at the COVID vaccine facility, she was told to go home, and someone would call to check on her. She felt a slight itchiness on the right side of her face. Her face looked very red, specifically her cheeks and forehead. She was so fatigued, she felt like she was drugged. She took Claritin. She couldn't fall asleep. She has extreme GERD and would wake up with an extreme racing heartbeat. She also had this happen prior to the COVID vaccine. She stated she takes Omeprazole, basically the prescription version of Prilosec. She stated her GERD is inflammation-based and the COVID vaccine must have triggered the GERD. Eight hours after the COVID vaccine, she fell asleep for five minutes, woke up, looked in the mirror and there was a big, red rash area on the whole right eye area on the right side of her face above her eye, but not to the top of her forehead. She hasn't been diagnosed with the cause of what is going on with her right eye. She does have right eye proptosis prior to vaccine so maybe the whole right side of her face was targeted because she has immune issues on that side prior to vaccine. On the next four days, she got zero sleep and had a hard time sleeping. She went to her primary doctor on Monday (unspecified date) and her blood pressure was 140-something over 80-90. Five days after the COVID vaccine, her blood pressure returned back to normal. This whole time she was taking Claritin. She could feel tingling feeling through her arm. She almost feels like she had been shocked or electrocuted. She had that continually feeling in her face and arm since the COVID vaccine. She did see an allergist who stated caller should have been given an EpiPen immediately. She has never used and EpiPen before. She saw the allergist seven to eight days later. The allergist took blood to see if there was still a histamine response, but her results were normal. She was still feeling those weird feelings of being electrocuted or shocked. The right side of her face is hypersensitive. It comes and goes. She did see her endocrinologist who stated maybe it's a neurological response to the COVID vaccine. She stated the doctor had seen what happened to her before, but it didn't last long. This doctor saw the face would go away after thirty minutes and reporting the ones that come back for the second time, their high blood pressures lasted for twenty-four hours. Her high blood pressure lasted for 5 days. The past couple of days her blood pressure has been very low and then high. It is all over the place, which is not normal for her. The outcome of the event ""blood pressure went up/blood pressure was 153/92/her blood pressure was checked three times and the lowest was 142/90"" was recovered, and

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				unknown outcome for the remaining events. Information on the lot/batch number has been requested."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1133561-1	Extreme exacerbation of CIDP symptoms. Weakness and paralysis of extremities, inability to walk, loss of sensation in hands and feet, marked increase in neuropathic pain
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1138897-1	35 wks pregnant, noticed these symptoms the morning after having the vaccine: right side of the face is slightly paralyzed, when blinking only the left eyes blink and right eye does not fully close, right side of the tongue slightly numb but still able to taste, when smiling or manipulating mouth only the left side is fully engaged, right side if about 70% engaged, feel tightness on the back of right side of the head
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1139570-1	"Pfizer COVID-19 Vaccine EUA Patient presents 3/23/21 for medical monitoring of syncopal episode s/p COVID - 19 vaccine # 2. Lowered to the floor by vaccine staff. Patient reports symptoms of weakness, other wise states she """"feels fine"""". Two additional syncopal episodes witnessed by myself when patient attempts to change position. Past medical history is significant for mild persistent allergic asthma, pre-diabetes, OSA, obesity and BPPV. Patient reports history of same reaction s/p previous vaccine. At a approximately 0837 patient was noted to have a worsening rash present to her face. EpiPen administered via left lateral thigh at 0838. EMS called to transport patient the ED for further evaluation. Treatment administered: epipen (lot 0FM406, exp 02/22) left lateral thigh at 0838. Upon arrival to the ED, patient's temperature is 98.8¦ F, heart rate 92-103, respiratory rate 30, saturating 99% on room air with blood pressure 164/91. The patient's laboratory studies were largely unremarkable. While in the ED, the nurse was getting ready to get the patient up to the commode when the patient started to """"not feel right again."""" She asked her to administer the medications for allergic reaction and reports wall the 2nd medication was being infused, she felt paralyzed and unable to talk, although she was able to hear everything and was aware, she was unable to move or speak for a short amount of time. CT head with completed at this time and showed no acute intracranial findings. Chest x-ray unremarkable. EKG showed normal sinus rhythm with sinus arrhythmia, heart rate 80 and prolonged QT at 470. The patient was administered Solu-Medrol 125 mg IV, Pepcid 20 mg IV, Benadryl 50 mg IV, normal saline 1 L IV. Pt was admitted to hospital for observation. The patient was seen on the unit laying in bed with her husband at the bedside. She reports that she is feeling slightly better and hungry and has never felt at paralyzed/unable to speak feeling before and is very nervous about this. She was wondering if she had a seizure, although she reports feeling aware the entire time and did not lose bowel or bladder function. We discussed that that is not typical for a seizure, but even if we did an EEG at this time, it would not be able to tell us what occurred in the past. We discussed continue the Benadryl, Pepcid, and IV steroids for today and likely decreasing them tomorrow. We discussed the workup of syncopal episodes as well. The patient and her husband were agreeable to this plan. Patient was discharged from the hospital on 03/24/2021 in a stable medical condition. Patient was instructed to follow-up with her PCP in 1 week."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1140678-1	paralysed for 10 hours, in both arms and legs; This is a spontaneous report from a contactable consumer (patient). A 75-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiry date were not reported), via an unspecified route of administration on an unspecified date at 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 VACCINE, lot number and expiry date were not reported), on an unspecified date for COVID-19 immunization. The patient took both doses of the Pfizer covid vaccine, but after the second dose he was paralysed for 10 hours, in both arms and legs on an unspecified date. Outcome of the event was recovered on an unspecified date. Information about lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1140955-1	Paralysis of arm and hand on side (L) where vaccine was administered. Extreme pain (Fire -like) shooting up and down the leg on the opposite side (R) with very bad joint pain. This went off & on through the night/early morning in the first 22 hrs after vaccination. Woke with most pain gone, L hand red and sore, minor headache. Through out the following week I have experienced difficult pain and instability of some joints.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1144572-1	Paralyzed in his arms and legs; Legs were weak; Severe shivers; This is a spontaneous report from a contactable consumer, the patient. A 75-year-old male patient received the second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot number: EN6198), via an unspecified route of administration in the right arm on 02Mar2021 at 09:30 (at the age of 75-years old) as a single dose for COVID-19 immunization. Medical history included dizziness, hyperuricemia (he produces too much uric acid and it causes kidney stones), back pain, COVID-19 in Dec2020 and was cleared by the end of Dec2020. Ongoing concomitant medications included diazepam for dizziness since 1978, allopurinol for hyperuricemia, oxycodone hydrochloride/paracetamol (PERCOCET) for back pain since 7-8 years ago. There were no other vaccines administered on the same day or within 4 weeks prior to the vaccination. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE; Lot number: EL9262) on 09Feb2021 at 09:30 (at the age of 75-years-old) in the left arm and experienced muscle aches. On 02Mar2021 at 21:00, the patient experienced severe shivers. On 03Mar2021 at 02:30, the patient woke up to go to the bathroom and noticed his legs were weak. On 03Mar2021 at 07:00 he was paralyzed in his arms and legs- he could feel his muscles but couldn't make them move. He just laid there for ten hours, and it was scary, and he thought that over a period of time the vaccine would get filtered out. After that, he was fully recovered and able to move. The events did not require a visit to the physician or the emergency room. The patient was concerned that he got the second shot too soon after having COVID-19 in Dec2020. The clinical outcome of legs were weak and paralyzed in his arms and legs was resolved on 03Mar2021, severe shivers was resolved on 04Mar2021.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1144973-1	Felt drunk within 2.5 hours of getting vaccine. By 4pm same day, I spiked 100.7 fever. By 10pm, I had severe muscle and joint pain. Woke up next day, still had fever. I was very fatigued. Two days post vaccine, I was nauseas. By 11:30am, my face went numb. Followed by numbness in my arms and legs. Dr said parenthesis and post injection inflammatory reaction.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1153319-1	blood clot formation that went to my brain; stroke; numbness; paralyzed; vision is messed up; trouble concentrating; This is a spontaneous report from a contactable consumer (patient) . This 70-year-old male patient received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, Lot number: unknown, may be EN8199 or EN6199), via an unspecified route of administration on 11Mar2021 at single dose in right arm about 16:00 for COVID-19 immunisation. Medical history included ongoing probably overweight and blood clot about 20 years ago. The patient's concomitant medications were not reported. It was reported that, the patient who was administered the first dose of Pfizer-BioNTech COVID-19 Vaccine on 11Mar2021. He reported on 14Mar2021 he got a blood clot and had a stroke. He asked if he should or should not still get the second dose of the Pfizer-BioNTech COVID-19 Vaccine related to these events. He had in the hospital from 14Mar2021 to 15Mar2021. He was supposed to have an appointment with his Family Care Physician, Doctor next Monday or Tuesday. He had to go to the emergency room because of these events where he was admitted to the hospital from 14Mar2021-15Mar2021 when he was discharged home under family care. They let him go so soon on 15Mar2021 because he insisted on it, if he was just going to be laying in the hospital he could be laying at home. There was not much they could do in the hospital except observe him, monitoring symptoms of when his numbness and the being paralyzed like in the face for improvement, then they figured the clots were clearing up. They put him on an Aspirin and other unknown things to keep events from reoccurring soon after these events. He was still does not feel 100%, his vision was messed up, he has trouble concentrating, vision was blurred sometimes; that was what happened, he knew he had a stroke when his vision went south on him on 14Mar2021. While in the hospital they did everything except no x-rays; he had CT scan, lots of bloodwork and stuff like that. He does not have further information on those tests or results. The seriousness of the events blood clot formation that went to my brain, stroke, numbness and paralysed was reported as serious (hospitalized). The outcome of events was reported as unknown. Information about lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1153404-1	Loss of hearing to last less than 5 min. followed by ringing ears through the following day. Uncontrolled palsy for head and neck intermittently through the injection day. I lost my voice and had spasms in my larynx. For several hours I had a hard time choosing my words and speaking which was the scariest. I could not focus enough to even wash my hands on injection day and had to be driven home from work. I got dehydrated because water tasted horrible. The next morning I am better on all fronts except the ringing in my ears wont stop

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1153519-1	"I couldn't move for about 4 hours, I was slight paralyzed laying on the floor; IV (intravenous) by needle; This is a spontaneous report from a contactable consumer (patient). A 78-year-old male patient received bnt162b2 (BNT162B2), dose 2 intravenous (at the age of 78-years-old), administered in Deltoid Left (reported as left shoulder) on 15Mar2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunization. Medical history included none. The patient's concomitant medications were not reported. The patient previously received bnt162b2 (BNT162B2), dose 1 via an unspecified route of administration (at the age of 78-years-old), administered on 22Feb2021 (Batch/Lot number was not reported) as single dose for COVID-19 immunization. The patient stated, ""I received my second Pfizer shot (captured as Unspecified medication) on Monday (15Mar2021) and on Monday night (15Mar2021) I woke up on the floor and I couldn't move for about 4 hours, now finally got back into the bed by 5 o'clock in the morning and I was slight paralyzed laying on the floor and I felt it. You may did know about this because it is probably related to the vaccine. They have improved drastically (events), after yesterday I tried to get some sleep because I laid on the floor for 4 hours, but today I feel alright."" The route of administration was ""IV (intravenous) by needle."" No treatment received for the events. The outcome of the events ""I couldn't move for about 4 hours, I was slight paralyzed laying on the floor"" was recovered on an unspecified date. The event Paralyzed was considered as serious (medically significant). Information about Batch/Lot number has been requested."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1153561-1	"I had a stroke on the right side of my brain; My whole left side is paralyzed; This is a spontaneous report from a contactable consumer (patient). A 68-year-old male patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in arm left on 19Mar2021 at 10:30 (Lot Number: EN7539) (at the age of 68-year-old) as single dose for COVID-19 immunisation. Medical history included diabetes from an unknown date and unknown if ongoing and pituitary tumor from 2004 to an unknown date. Concomitant medication included metformin taken for an unspecified indication, start and stop date were not reported. The patient previously received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration, administered in arm left on 26Feb2021 at 10:30 (Lot Number: EN6205) (at the age of 68-year-old) as single dose for COVID-19 immunisation. On 19Mar2021 at 21:00, the patient reported that: ""I had a stroke on the right side of my brain. My whole left side is paralyzed. I am an invalid walker. This has changed every aspect of my life"". The events were serious, after emergency room visit the patient was hospitalized for 2 days, the events were also life-threatening and disabling. Therapeutic measures were taken as a result of the events and included physical therapy. On an unspecified date, the patient underwent lab tests and procedure which included electrocardiogram (ECG), chest X-rays, magnetic resonance imaging (MRI), all results were unknown. The patient outcome of the events was not recovered."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1154077-1	paralyzed with no sensation at all in her legs till her waist; headache; fever 101.2; weakness; chills; fell; This is a spontaneous report from a contactable Physician. A 69-year-old female no pregnant patient received second dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 10Mar2021 10:30 AM at single dose for COVID-19 immunisation. No pregnant at time of vaccination. Medical history included prior spinal cord trauma with rt foot drop. No Known allergies. No COVID prior vaccination. The patient received the first dose BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) at 69-year-old, via an unspecified route of administration at single dose for COVID-19 immunisation. The patient started the next day on 11Mar2021 02:00 PM, after the 2nd shot with headache, fever 101.2 and weakness. Took Tylenol and went to have a nap. She woke up with still fever and chills, when she tried to get up she fell. She was currently in the hospital, paralyzed with no sensation at all in her legs till her waist. AEs resulted in Emergency room/department or urgent care, Hospitalization for 5 days, disability or permanent damage. Lab data included Nasal Swab on 11Mar2021: Negative. No COVID tested post vaccination. Solumedrol 1000mg IV were received. Treatments were received for the events. Outcome of the event was recovering. Information on the lot/batch number has been requested.; Sender's Comments: Based on the temporal association, a possible contributory role of BNT162B2 to the development of 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.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1155146-1	Completely unable to move my body and mouth for a couple of hours. Severe headache, severe tingling in my legs.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1160276-1	The day after the vaccination I had spasms in my left eye which continued every day for two weeks. They came and went several times a day. The diminished over the course of the two weeks. I also had some paralysis in my left eye. I was given some Vitamin B12 and the symptoms resolved. I got the second dose of the vaccine and did not have any problems with side effects.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1163722-1	Bells Palsey left side face, numbness, slight paralysis- took Benadryl when I got home.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1165960-1	felt paralyzed and nauseous; felt paralyzed and nauseous; fever; This is a spontaneous report from a contactable consumer. A 78-years-old non-pregnant female patient received BNT162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE), via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) (at the age of 78-years-old) as a single dose for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient had been tested for COVID-19 via antibodies test on an unspecified date with positive result. On an unspecified date, a day after receiving the vaccine, the patient developed a fever and said that she felt paralyzed and nauseous. The patient needed to go to the hospital in order to recover and tested positive for antibodies. The outcome of events, fever, felt paralyzed and nauseous was unknown. The lot number for the vaccine, BNT162b2, was not provided and will be requested during follow up.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1168791-1	Patient reports the following symptoms 24 hrs after receiving the Pfizer Covid vaccine: extreme fatigue, chills, sweats, fever, and paralysis on left side. Patient has Multiple Sclerosis and has issues with gait and movement on left side at times. He states paralysis was an exaggeration of what he normally experiences at base line.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1180858-1	Systemic: Dizziness / Lightheadness-Severe, Systemic: Exhaustion / Lethargy-Severe, Systemic: Numbness (specify: facial area, extremities)-Severe, Additional Details: 24 hours after vaccination patient experienced paralysis and was rushed to ER where ER physician said reaction was most probably due to the vaccine. patient recovered within 2 weeks. MD said patient can get 2nd dose but patient has refused second appointment for vaccination,
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1191851-1	paralysis; This is a spontaneous report from a contactable nurse via medical information team. A female patient (nurse) of an unspecified age received second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, formulation: solution for injection) via an unspecified route of administration on 11Jan2021 as a single dose for covid-19 immunization. The patient's medical history and concomitant medications were not reported. It was reported that her rheumatologist was treating another nurse that now has paralysis due to the vaccine. On 09Feb2021, the patient experienced paralysis. The clinical outcome of the paralysis was unknown. Information related to batch/lot number has been requested.; Sender's Comments: Based on the information currently available, a possible contributory role of the suspect drug to the reported event paralysis cannot be completely excluded based on temporal association and known drug safety profile. 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.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1201247-1	Paralysis (CMS/HCC) death
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1205956-1	started with watery diarrhea, followed by typical fatigue low grade fever migraine and muscle aches by 8pm 4/6/21. by 2am 4/7/21 felt my eyes roll back, lost 2ish hours of consciousness, checked temperate and was 103.7. i felt paralyzed by 4ish am. could not move lower half before 7am, regaining weak movement in upper body. by 8am could barely move my lower half, using a cane to get to the restroom a few steps away from bed. arm that was injected, throat, nose, eyes were all swollen. primary doctor and pharmacist both said to go to ER. went there once i was able to secure a ride and childcare. ER doctor requested blood work and covid test. covid test was negative, blood work came back normal. had a bloody nose before i came home. diagnosed there with post vaccination reaction. told to take benadryl and ibuprofen/tylenol. each day i woke after vaccination unable to move my neck (including today). it takes me several hours to be able to move my neck muscles. waking up 4/8/21 my right side was temporarily paralyzed but regained slow achy movement throughout the day. each day after vaccination joint pain somewhere in my body takes place and nerve pain is often associated. 4/10/21-4/13/21 i have experienced sciatic nerve pain down both sides all the way down to my toes. my nose and throat are still swollen. i am still having to take benadryl daily due to throat swelling. my throat and neck itch.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1206432-1	I got an Embolic stroke on 4/4/2021- 9 days after I got the vaccines from Hospital. On 4/3/2021 afternoon, I suddenly couldn?t see well, felt dizziness. At 3 AM on 4/4/4021, called 911 because I couldn?t feel my left side. I couldn?t see well on my left eyes, couldn?t feel left hand, left leg, left face, and the rest of my left side are numbed and I am not able to move after I was brought to the hospital. I am now at rehab Center.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1213332-1	"After first dose, ""normal"" side effects of extremely sore arm, headache lasting one day. After second dose, fever to 101.5 at 1:00am after receiving shot at 3:00pm previous day. Extreme arm and shoulder pain. Waking up again at 3:00am experienced FULL BODY paralysis. Estimate fever at 103+ but was unable to move to verify. Eyes were in pain and blurry vision. unable to reach phone to call 911. These symptoms lasted 1 hour and son called to check on me and rushed over. Fever was at 101 but immediate does of motrin and then later tylenol, broke fever. Arm remained immobile for full day. Now it is mobile but still painful to lift overhead. All other symptoms subsided."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1214456-1	Critical Limb Ischemia Symptoms: Progressive BL lower extremity and back pain starting on 3/20/2021. She presented to the ED twice and DVT ultrasound and lumbar spine CT showed no acute findings. Symptoms progressed to the point of severe pain on 4/1/2021 with toe numbness and paralysis. CTA of BL lower extremities was performed showing bilateral arterial thrombus. She underwent bilateral common femoral artery thromboembolectomy with vascular surgery. Of note, patient was on eliquis 5 mg BID given history of PE in 2018. She was compliant with this medication.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1220770-1	Guest started hyperventilating after shot, complained of loss of motor function and felt the pain of being paralyzed. And also had hypoxia.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1222774-1	Observed resident with left side weakness and facial drop. neurological sign checked and left side found flaccid and unable to move. 911 called. Notified of pt sent via 911 to hospital
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1223409-1	Patient experienced paralysis from the lower chest to toes 12 hours after vaccination. She was taken by ambulance to the Hospital, then transferred to another Hospital. She will now be transferred to the a Clinic.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1224331-1	"couldn't stand on his feet; Fell awake; screaming so bad; Chills; Almost paralyzed; Pain; like somebody cutting his side, on his lower abdomen on the right side/underside of the lower abdomen/slight pain every single day, sometime excruciating; couldn't walk after vaccine; This is a spontaneous report from a contactable consumer (patient). This 74-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration administered on the arm left on 01Mar2021 (Batch/Lot number was not reported, reported as ""maybe I don't know 6198, it's GN or EN"") as a single dose for COVID-19 immunization. Medical history included atrial fibrillation, broken back, breathing problem because of his heart and lung, his lungs have problem, he had some nodules in it, and kidney stone once 25 years ago, 5 little stones. The patient's concomitant medications were not reported. The patient had a reaction, it is the worst reaction he has ever had in his life, the pain he had, he was almost paralyzed and couldn't walk after vaccine and he had to get a wheelchair because he couldn't stand on his feet. He ended up in the emergency room. The vaccine put him out of work, and now he can work some. He is trying to make it up in two days, today and tomorrow to finish up. He was very upset and wanted to know why this happened to him. The patient stated that other people he talked to had no reaction, if they had a fever or anything it was gone in a coupled days, but he ended up in the emergency room. On 01Mar2021, patient had his first dose and at 02:30, he got out of the facility and came out okay with no problem. At 21:00, all of a sudden he saw that he could not stand up on his feet and he fell awake. He waited in the bed, people around him got worried. Eventually, he asked them to bring wheelchair, maybe he can drive. He drove home with his wife to the emergency from his house and he got admitted in emergency and he was screaming so bad, he did not believe in his life, he never heard somebody doing like this. He felt like somebody getting chills like somebody cutting his side, on his lower abdomen on the right side. It's underside of the lower abdomen, he had slight pain every single day, sometime excruciating, he took Ibuprofen for it. He thought he had a kidney stone because he had it once 25 years ago, 5 little stones, so they took him to emergency, took him to the CT scan. They didn't find anything, but the pain was excruciating and in the emergency room everybody thought the crazy man is here who is screaming. He couldn't control his scream. He couldn't walk, he couldn't move. Every movement was like you put a knife on the cutting, it was that bad. His second dose was due on 22Mar2021. He went to the pharmacy and asked them the number and let them know that he had a reaction like this and they called somebody and said to cancel the shot. They cancelled it. That day for 3 days, patient had the pain, it was slowly because he had to take eventually oxycodone. The doctor advised in the emergency room that nothing was wrong, just to slow down the pain but he was not allowed to take the oxycodone. He took the 10 mg oxycodone and the pain which was an hour and a half was a little bit down. After 3 hours, it was a little better, he didn't want to scream, it was gone but the pain was still there for at least 3 days, really bad. He was taking ibuprofen and a muscle relaxant. He did all those things and still I have the pain left and he did not know what to do with that. The lab test he had, he had been okay, he did not have any problem. He got the result on 22Mar2021. Outcome of the events pain and abdominal pain was not recovered and of the other events was unknown. Information about lot/batch number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1224846-1	complete body went paralyzed; lost vision; throwing up; fever; This is a spontaneous report from a contactable consumer. A 41-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: En6207), dose 1 via an unspecified route of administration, administered in left arm on 23Mar2021 08:30 as single dose for covid-19 immunisation. Medical history was reported as none. There were no concomitant medications. The patient started throwing up, fever, complete body went paralyzed and lost vision all it the ER on 05Apr2021 13:00. The events were resulted in emergency room /department or urgent care. There is some treatment (saline and ondansetron) for the events. The patient did not have covid prior vaccination, did not test covid post vaccination. The patient received Covid test post vaccination via Nasal Swab on 05Apr2021, covid test result was negative. There is no known allergies, no other medical history, no other vaccine in four week, no other medications in two weeks. The outcome of the events was resloving.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1227876-1	a friend did experience that, from his back down he could not move his legs; This is spontaneous report received from contactable consumer, reporting for friend (patient). A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE), dose 2 via an unspecified route of administration on an unknown 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. When getting the second, the patient did experience that, from his back down he could not move his legs and had to go to the hospital, in 4 or 5 days he left. The outcome of the event was not reported. The lot number for the vaccine, BNT162B2, was not provided and will be requested during follow up.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1227879-1	Paralysis; Right lip swelling; This is spontaneous report from a contactable consumer (patient). A non-pregnant 35-year-old female patient 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 in left arm, on 01Apr2021 at 05:15 PM (17:15), 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 on 04Mar2021 for COVID-19 immunization (left arm). 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-19 vaccine. The most recent COVID-19 vaccine was administered in a pharmacy or drug store facility. On 02Apr2021, the patient experienced right lip swelling and paralysis. The events were considered non-serious. No treatment was received for the adverse events. Since the vaccination, the patient has not been tested for COVID-19. The outcome of the events was unknown. Information about lot number and expiration date for the suspect product will be requested in follow-up attempts.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1234842-1	40 hours after vaccination severe light sensitivity causing paralysis and fainting in daylight.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1239036-1	Blood clotting, hospitalized, emergency, paralyzed, thrombosis, body not responding, therapy, treatment.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1246196-1	Eye strain, possibly conjunctivitis. Some very slight flu like symptoms (sneezing, congestion)
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1247276-1	March 15th COVID shot. March 16th started to get severe dizziness and fatigue. Went to bed early. March 17th speech was impaired and difficulty with gait and balance. Admitted to MAMC ER, which confirmed a stroke and released. March 18th early AM admitted again to ER due to unable to move/walk. Due to lacking proper equipment and space was transferred via ambulance to hospital in March 18th until March 24 Obeservations Inpatient at Hospital. March 24 to April 16th Inpatient rehab.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1248132-1	She got the vaccine at 2:00, around 4:00 she started with fatigue with headache, dizziness, somewhat blurry vision (not extreme), chills up her full spine to her head from the bottom. She also at that time noticed a lump on her right arm and difficulty raising her arm straight out due to severe intensity of pain. She had difficulty of neurological function and paralysis and reaction of the right leg to just above the knee into her toes. She had paralysis feeling and also heavy swelling that was noticed by 6:00 that day. She also noticed that night that no position gave her comfort to be able to sleep, which kept going and she still has the paralysis, numbness, chills, fatigue. The headaches are getting sharper, more intense in her right temple area which is very sharp, which is not normal for her. The tumor she had gave her squeezing headache pain across her eyes and the to of her head but not the temples. The 2nd shot brought about 3 days of wheezing, which she does not have any lung or heart issues. She has been having difficulty with just having normal breathing patter as she does not feel it is quite right for getting oxygen as it had been prior. She went to her PCP last Friday, 4/16/21 who has ordered an MRI. He has consulted with the neuro-specialist that she had with her tumor. He will follow through with more blood work more than likely. He did mention to her about having a COVID test to double check that end of that. She did not have this done due to the fact that she had the brain tumor surgery and that is how they did that approach. She was just staring to have resolution of some of her symptoms from the 1st vaccine from the right leg when she got this one. When she got the 2nd vaccine she had intense swelling in the left leg from the knee down. There is a definite difference in the size of the legs. Her PCP said that there is definite damage from his testing at her recent appointment.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1248200-1	day 1 sore arm day 2 temp 101, headache day 3 lower temp day 4 no temp day 7 some left had paralysis, woke up with it, resolved in 20 minutes day 9 herpes cold sore day 2-21 still have headache

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Shaky; Muscles and nerves seemed to be locking up like almost paralyzed; swollen lymph nodes; injection site redness; severe allergic reactions; slight dizziness; severe headache; nausea; weak; tired; feverish; sweat; had pain at the Pfizer Covid 19 Vaccine injection site; right arm started to swell; chills; extreme muscle pain; certainly extreme and joint pain; Heart beat seemed to speed up; Has not been able to walk or move body in normal fashion; This is a spontaneous report from a contactable consumer (patient herself) A 67-year-old female consumer received first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Formulation: Solution for injection, lot number: EP7533) a week ago via an unspecified route of administration on 02Apr2021 at 13:45 hrs on the right upper arm as a single dose for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported.When caller was 18 they required a booster for Small pox and caller received the small pox booster on the leg in the same site where it was given at age 5. Caller's leg swelled up to 5 times the normal size, at the time caller was 18 years old.Took a long time for swelling to go away, a couple weeks. Caller requires it hindered sports activities during this time. On 02Apr2021, the patient started reported slight dizziness, severe headache, Shakiness, nausea, weakness, tiredness, have been feverish, pain at the injection site and stated that her right arm started to swell. The patient stated that through out the night of 02Apr2021, she had extreme muscle pain, certainly extreme and joint pain that was also extreme and heart beat seemed to speed up within the first hour and by the end of the hour. On 03Apr2021, the patient stated that her muscles and nerves seemed to be locking up like almost paralyzed. The patient described the events she experienced slight dizziness and didn't worry because it wasn't major and caller was driving herself and with in an hour the dizziness got worse. Caller had severe headache and had to drive home a significant distance and the time the caller got home, she reports being shaky, dizzy, had headache, nausea, was weak and tired and must have been feverish. Caller reports she must have been feverish because she was drenched in sweat by the time she got home and it was a 2 hour drive home from the vaccine administering facility. Caller had pain at the Pfizer Covid 19 Vaccine injection site, and caller's right arm started to swell.Caller clarified that the injection site was in the right upper arm, doesn't know what the muscle is called. Caller says she couldn't see if her arm was getting red, caller has some discs out in her neck and is unable to look at the arm. Caller could feel the injection site on her right arm was swollen and swelling. Caller reports fever was gone and fever got replaced with chills later on. Caller reports there was progression of the symptoms after the first hour. Caller states the information provided thus far was the first hour of her experience and through the night a lot of other things changed and added.Through the night caller had extreme muscle pain, certainly extreme and joint pain that was also extreme. Muscles and nerves seemed to be locking up like almost paralyzed: By the next morning, after caller had the Pfizer Covid 19 Vaccination, states she was not a doctor but all of her muscles and nerves seemed to be locking up like caller was almost paralyzed. Caller reports she couldn't move, couldn't get out of bed. It took a long time to get caller's fingers and toes moving and the rest of the body, but fingers and toes were the first ones. Caller was able to kind of release from muscle cramping. Whatever was going on, caller says can't tell exactly what was going on, does not have a Doctor in the home. States the information given should cover that part of the experience. Through the night had extreme muscle pain, certainly extreme and joint pain that was also extreme: Reports since the morning after receiving the Pfizer Covid 19 Vaccine, caller has had continued muscle pain, although not as extreme, its pretty extreme, and join pain. The patient has not been able to walk or move body in normal fashion. The patient stated that she had every side effect listed on the fact sheet at the top minus the swollen lymph nodes and injection site redness and she had two of the side effects listed under the severe allergic reactions. The patient wanted to know whether it is safe to receive the second dose of the vaccine. The patient lives in an isolated area where all through COVID-19 she tried to get information from the local medical clinics and they did not seem to stay informed. Caller states she has a doctor's appointment, and she will let them know the side effects she experienced after the first dose but she is not confident in the recommendations they might provide. Patient was given a list of side effects to look for, ones they were assuming, doesn't know if the hand out was published by Pfizer or not.The hand out listed emergency kind of side effects people were experiencing.Caller reports they are over age 65 range and knew a lot of people before her who had the Pfizer Covid 19 Vaccine and said they only had injection site soreness. Caller was expecting the same reaction, caller was told by the Doctor she was in pretty good shape. Caller reports she had most all of the reactions except anaphylactic reaction of not breathing after receiving first dose Pfizer Covid 19 Vaccine. Caller is nervous about going for the second round of the Pfizer Covid Vaccine, is not sure how to proceed and would like that information. The case was assessed as serious medically significant for the event paralysis. It was reported that patient took Aspirin as treatment medication for pain. The outcome of the events was resolved for fever and nausea, resolving for swelling arm, unknown for swollen lymph nodes, Chills , Vaccination site pain, injection site redness, severe allergic reactions, sweat, shaky feelings, heartbeat increased, movement disorder and not resolved for all other events.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1255121-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255434-1	entire left area where the patient received the vaccine had fallen asleep and paralyzed/it moved to the right area; throat problems such as throbbing; shoulder pain became unbearable; panicked; feel it in the throat and I could not breathe; the patient took an Advil to be able to sleep; This is a spontaneous report from a contactable consumer (patient). A patient of unspecified age and gender received the second dose of BNT162B2 (Pfizer-BioNTech COVID-19 vaccine, lot number and expiry date unknown), dose 2 via an unspecified route of administration on 08Apr2021 as a single dose for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient previously received the first dose of the BNT162B2 for COVID-19 immunization. The patient received the Pfizer vaccine in the morning of 08Apr2021 and she had no reaction and it was perfect at 2 in the afternoon however at 4 the patient began to have throat problems such as throbbing and the shoulder pain became unbearable, the patient could not move it. During that night, the patient took an Advil to be able to sleep because the pain in the shoulder was unbearable, the patient put an ice pack and tried to sleep but got up at 12 and the entire left area where the patient received the vaccine had fallen asleep and paralyzed. The patient then panicked because it moved to the right area and the patient began to feel it in the throat and could not breathe. The patient was treated in the ER. It was also reported that the patient was confident that the effects will be normal and at most have a fever or any of the 7 side effects that were listed on the sheet that was given during vaccination but the patient did not think that he/she was going to be put in these circumstances because the patient could not breathe, or speak, or move for a period of time that also mixed with his/her nerves because the patient was not prepared and did not know what to do and they had to see the patient in the ER to be certain that he/she was fine and happily, the patient was stable. The outcome of the events was recovering. Information on the lot/batch number has been requested.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255726-1	she could not feel her body from the neck down- She believes the vaccine paralyzed most of her body.; she could not feel her body from the neck down- She believes the vaccine paralyzed most of her body.; This is a spontaneous report from a contactable consumer. A 33-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Solution for injection, Lot number-UNKNOWN) via an unspecified route of administration on unspecified date as single dose for covid-19 immunization. The patient medical history and concomitant medications were not reported. The patient experienced she could not feel her body from the neck down- she believes the vaccine paralyzed most of her body on an unspecified date. The patient underwent lab tests and procedures which included blood test: clear and/or negative, lumbar puncture: clear and/or negative, magnetic resonance imaging: clear and/or negative. The outcome of events was unknown. Follow-up (19Apr2021): This follow-up contains no new safety information. Follow-up attempts have been completed and no further information is expected.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1255735-1	new numbness and pins & needles in left leg; It causes paralysis in muscles; CPK increased; This is a spontaneous report from a contactable consumer (patient). This 60-year-old male consumer received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 26Mar2021 at single dose in right arm for COVID-19 immunisation at the age of 60-year-old. Medical history and concomitant medications were unknown. Historical vaccine included fatigue with BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 05Mar2021 for Covid-19 immunisation at the age of 60-year-old (1st dose, at 09:30, leftarm). The patient got up one day and legs where paralyzed on 08Apr2021. The patient kept trying to stand and walk thinking it was just legs fallen asleep, but the patient was falling every time she would try to walk it off. On 08Apr2021 at 07:30, new numbness and pins and needles in left leg. The patient was hospitalized due to the events and they were considered as disabiliting and life-threatening. The patient then had a bad moment when on his way to leave for pcp appt (as reported), a steel cabinet fell on him. The patient was taken by ems to trauma unit in Hospital. Creatine phosphokinase (CPK) levels were over 2,000 on unknown date. The outcome of Creatine phosphokinase increased was unknown, the patient was recovering from other events. Information on the Lot/Batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1266001-1	loss of consciousness plus all the normal symptoms; paralysis/partial paralysis; Fever of 103.7; diarrhea; nausea; dizzy; joint pain; migraine; throat swelling; nose swelling; eyes swelling; This is a spontaneous report received a contactable consumer, the patient. A 33-year-old adult [non-pregnant] female received the second dose of BNT162b2 (solution for injection; Lot ER8729 and expiry information not reported) in the right arm via unspecified route on 06Apr2021 at 09:45 (at 33 years-old) for COVID-19 immunisation. Relevant medical history included asthma, allergies, penicillin and adhesive tape allergies, and high blood pressure. Concomitant medications included lisinopril. The patient previously received COVID-19 immunisation with the first dose of BNT162b2 (Lot ER7531) in the right arm on 16Mar2021 at 09:45 (at 32-years-old). The patient previously received treatment for an unspecified indication with hydrocodone bitartrate/paracetamol (NORCO) and experienced an allergy. The denied being diagnosed with COVID-19 prior to the vaccine. The patient was not pregnant at time of vaccination, and the patient denied receiving any other vaccine within four weeks of this vaccine. On 06Apr2021 at 20:00 the patient experienced fever of 103.7, diarrhea, nausea, dizzy, paralysis, partial paralysis, joint pain, migraine, throat swelling, nose swelling, eyes swelling, loss of consciousness plus all the normal symptoms. The patient reported as a result of the events the patient sought care with a doctor or other healthcare professional office/clinic visit, emergency room/department or urgent care. The patient was tested for COVID-19 since the vaccine with nasal swab test on 07Apr2021 and was negative. Treatment for the events received in the emergency room (ER) included diphenhydramine HCl (BENADRYL), ibuprofen, and acetaminophen (TYLENOL). The outcome of the events, fever of 103.7, diarrhea, nausea, dizzy, paralysis, partial paralysis, joint pain, migraine, throat swelling, nose swelling, eyes swelling, loss of consciousness plus all the normal symptoms, was not recovered.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1268633-1	Woke up middle of the night with bilateral arm numbness/paralysis and then shooting pain on both arms and neck. Lasted up to 5 minutes. Next day -- pain in both hands and periods of bilateral arm numbness. Following evening woke up in the middle of the night with bilateral arm numbness/paralysis and pain shooting up fingers to shoulders in both arms and neck. Pins and needles on spine and arm and bilateral hand pain and numbness followed for 27 days.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1269765-1	paralysis; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received bnt162b2 (BNT162B2, Solution for injection, lot number was not reported), via an unspecified route of administration on 16Apr2021 (Batch/Lot number was not reported) as SINGLE DOSE for COVID-19 immunisation. The patient had partial paralysis since getting the Pfizer vaccine last week (unspecified date). Her claim is that the paralysis was due to her getting the shot last Friday (unspecified date). The doctors that are tending to her say they are indeed looking into the cause and that a possible one could be the vaccine. The patient was hospitalized due the event. The event was assessed as life-threatening. Outcome of the event was unknown. Information about the Lot/batch number has been requested.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1270827-1	"On or about April 14, 2021 patient yelled out to a friend to come help him. Patient stated that he ""could not see or move"". The friend called the ambulance and patient was taken to hospital. One of the doctors tending to patient told me patient had several strokes (over 5). These strokes have left him blind and incapacitated. He is still currently at hospital."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1273210-1	Complete paralysis of arms, hands and face for more than 1 hour. Tingling and some numbness in legs and feet. Extreme pain in the arms. Rapid heart rate and sweating accompanied the paralysis.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1279279-1	"I had a very bad headache not in front of my head, at the back of my head where my neck was from where I was paralyzed and my shoulder where my dystonia was; I got a slight headache at the back of my head and neck and my shoulder and when I woke up, I was completely paralyzed, I could not move; I got a slight headache at the back of my head and neck and my shoulder and when I woke up, I was completely paralyzed, I could not move; Suffer from dystonia; I was paralyzed for at least 24 hours, I could not move; I could not move or anything for 24 hours; This is a spontaneous report from a contactable consumer (patient). A 55-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), second 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 reported that ""I am a 'post op asexual' (not clarified), I had a surgery but I am a female now, I don't know if is there anything to do with it, I had a surgery 7 years ago."" The patient also had dystonia. The patient's concomitant medications were not reported. The patient previously took bnt162b2 first dose on an unspecified date for COVID-19 immunization and experienced sore arm. The patient stated that: ""I had a very severe (sentence incomplete), I took my shot and my first shot was fine, I only had a sore arm from my first shot but from my second shot, I suffered a very severe side effect. I suffer from dystonia (further clarification unknown) and I was paralyzed for at least 24 hours, I could not move and I could not call 911 or anything, I was just by myself, I was just by myself with my dog, I could not move or anything for 24 hours. I had some medicine and I was able to get some sleep and the following morning (sentence incomplete) but I was technically paralyzed, I could not move or anything. So, anyone with dystonia should not take the second shot without somebody being there because it would be too dangerous to take it, it is way too dangerous."" The patient further stated, ""The side effect was, I had a very bad headache not in front of my head, at the back of my head where my neck was from where I was paralyzed and my shoulder where my dystonia was."" The patient added: ""I felt great, I felt good, I took my shot at 10 o' clock in the morning, I felt good, I felt good. I took it on the 15th at 10:15 in the morning (unknown exact date) and I felt good all day but at the middle of that night, I got a slight headache at the back of my head and neck and my shoulder and when I woke up, I was completely paralyzed, I could not move and I suffer from dystonia (history)."" For the treatment the patient stated: ""I took Tylenol and things like that, basic stuff. I took Clonazepam for my dystonia because this Clonazepam helps my dystonia and that did not work it all. Clonazepam is the only medication which helps with dystonia and that did not helped, that is the only medication that would work."" The outcome of the event dystonia was not recovered; and unknown for all other events. Information on the Lot/Batch number has been requested."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1284845-1	Neurologist agreed that the vaccine has attacked nervous system; They thought he was having a stroke because the left side of his body was twitching and numbness all the way from the arm to the feet; He had some kind of palsy where his face was not reacting bilaterally; Got really sick 18 hours after it and it lasted for like two days where he was really sick, all sorts of symptoms; He hasn't been able to work since 25th; He cannot coordinate his thoughts and fantasies; Dizziness; He felt his head was spinning; Nausea; He couldn't stand up without falling; Pressure in his brain; Pain in the rear of his head between the neck and the cranium; Inability to speak/Slurring his speech; twitching and numbness all the way from the arm to the feet; twitching and numbness all the way from the arm to the feet; This is a spontaneous report from a contactable consumer or other non hcp. A 66-years-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Formulation: Solution for injection, lot number: Unknown), dose 2 via an unspecified route of administration on 25Mar2021 as single dose for covid-19 immunization. The patient medical history was not reported. The patient's concomitant medications were not reported. The Reporter stated, her husband had the Pfizer vaccine and the second dosage he got really sick 18 hours after it and it lasted for like two days where he was really, really sick, all sorts of symptoms. He got very sick. the Reporter stated, there was a problem and it's recurring and the vaccine was affecting people neurologically and if they have had something like concussion or any neurological damaging the path was attacking their nervous system. So, this was major, and this was happening not just 3 people we know and the doctors were agreeing that this was happening from the Pfizer vaccine. The Reporter stated, the patient had been in emergency 3 times, patient thought he was going to die. He hasn't been able to work since 25th and he has been in emergency 3 times they have done MRI's and CT scans. And he was very sick, he cannot coordinate his thoughts and fantasies. He was not dead; he was very sick. On Mar2021, the Reporter stated that, Everything CT scans, MRI, ear exams, audiology test was done it. On Mar2021, the reporter stated that, the patient was hospitalized 1 night in emergency for observation. He was hospitalized about three days ago. He was discharged the next day, but he was in emergency overnight. He was hospitalized for the vaccine he took, and the neurologist agreed that the vaccine has attacked nervous system. The Reporter again stated that the patient had some of the symptom's dizziness, he felt his head was spinning, they thought he was having a stroke because the left side of his body was twitching and numbness all the way from the arm to the feet, he thought he was having a stroke. He had some kind of palsy where his face was not reacting bilaterally, he had nausea, he couldn't stand up without falling, pressure in his brain, pain in the rear of his head between the neck and the cranium and inability to speak, he was slurring his speech. The outcome of the events was reported as unknown No follow-up attempts are needed; information about lot/batch number cannot be obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1291309-1	felt a little woozy/ woozy and dizzy; I began violently shaking/ tremble; I couldn't lift my foot up I had to crawl up the steps; convulsions/ seizure like shaking it is terrifying; couldn't talk; she can't walk; palsy; not being able to function like I did/ am unable to be a minister now; stuttering; This is a spontaneous report from a non-contactable consumer (patient). A 68-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, solution for injection), via an unspecified route of administration on 02Mar2021 (Batch/Lot number was not reported) as single dose (dose 1) and then via an unspecified route of administration on 25Mar2021 (Batch/Lot number was not reported) as single dose (dose 2) for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient took the first shot on 02Mar2021, a few days, maybe on 2nd day or day or so after she took the shot, she went to the store and I felt a little woozy. She thought maybe she needed something to eat, but when she went to restaurant, she began to tremble and needed to get home. The patient drove slowly, when she got home, her husband was outside and she got out of the car and told him she was woozy and dizzy. She went in side and when she got to steps, she couldn't lift her foot up and had to crawl up the steps. She began violently shaking, she sat in a chair, it got worse. Her husband called the number, because she couldn't talk, like convulsions, just shaking, when Emergency people came, they took her to the hospital and she was there for 2 days. They did 2 computerized tomography (CT) scans, Magnetic resonance imaging (MRI) couldn't detect anything, but she continued with shaking and trembling. They weren't doing anything to find out what it was. One doctor said one thing the other said they didn't see anything, they sent me home. The patient was in hospital for 2 days, may have been 3 days, on 3rd day she went home (as reported). The patient went to the doctor and he sent her for other tests, but no one can determine or figure out what was going on. She didn't stutter, she was speaking normal, she was a minister. She was unable to be a minister now. She was well and doing well, then after that she can't walk, some times in bed at night, she go into shaking all over like she has palsy. She had to use a cane to walk. Some days she was feeling fine. She was still using a cane to walk and was stuttering, did not have her normal walk, she was afraid that she will fall. She was really trembly and shaky. Some days she can be fine, but then it comes right back. The patient stated that before she took the shot, she was fine. She is now 68 years old, she owns her own business sowing, active, walks and had no issues at all. Now, she was not able to go down stairs to sowing area and do anything. She had to use the buggy at the grocery store, because she can't walk and at night, some nights, she has a bell, her husband had to hold her from seizure like shaking it was terrifying. She had her second dose on 25Mar2021, the next day, she experienced worsening effects, she never had this shaking and not being able to function like she did before. She was concerned Ithat she was having seizure and palsy, because she had never been like this. It didn't start until after the first shot was bad and after the second one, it intensified even more. The patient stated that ever since the shot, her life had changed, she had never been in this position before. She can't think of anything else, checked her medications, never had any problems. She had been to the doctor, they can't detect anything else that could have caused her to get in this condition. Nothing like this. The patient stuttering, having to think of what she has to say, she had never ever had anything like this. She wanted to reach out because it was life changing and she was afraid and have never experienced anything like this at all. It has taken away any opportunity to get a job or anything. The outcome of the events was not recovered. Information about the Lot/batch number has been requested.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1293024-1	Patient is a 74 y/o female with PMH significant for meningitis at age 2 resulting in developmental delay (non-verbal at baseline and seizure disorder) who presented for poor oral intake, dysphagia, and inability walk due to increased lower extremity weakness that has been occurring since April 9th. The patient is typically ambulatory but needs significant assistance with ADLs. She is able to feed herself (tends to eat baby food). Eventually, the patient was admitted for sepsis 2/2 UTI and now with concerns for remote stroke with persistent dysphagia and acute encephalopathy. Since April 9th, the family has noted that the patient has had difficulty swallowing baby food and seems to be uncomfortable; she is able to make a couple of audible sounds or gestures that suggest pain and that is why they brought her. The family also reports that the patient received her second dose of the Pfizer-BioNTech COVID-19 vaccine on 04/07/2021. On 04/08/2021, the patient had one staring episode, but then returned to baseline. She was at baseline during breakfast on the morning on 04/09/2021, but around lunchtime, she had another staring spell and was unable to swallow food as well as unable to walk, stand, or move her lower extremities. The patient has limited language at baseline but also has not been able to say the few words she used to communicate with her family, only moaning and grunting since that time. Since 04/09, she has not been able to consume any solid foods, only eating a very small amount of smashed/liquified fruit and drinking Pedialyte. Sister has been massaging her legs regularly in hopes of re-establishing movement but saw nothing until yesterday (05/05/2021) when the patient moved toes bilaterally and briefly lifted her right leg. Neurology Consult on 05/05/2021: The patient had mildly bradycardiac and hypertensive (170s/80s). Labs reviewed and showed mildly elevated WBC (i.e., 12.9 K/MM3) and urinalysis was consistent with UTI. Imaging was reviewed and consistent with chronic CVA, likely occurring on 04/09/2021 while the patient was in hypercoagulable post-COVID vaccine state. Neurology recommends CT angio head and neck, TTE w/ bubble, A1c, fasting lipid panel, DVT prophylaxis via lovenox 40mg SC daily, D-dimer levels, EKG, PT/OT?Speech therapy evals, BP control via lisinopril 10mg daily w/ goal of < 140/90, and ASA 81mg daily

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1293231-1	awoke 4/26/2021 with double vision and slight dizziness; saw eye doctor 4/29 and was told I was experiencing palsy in my 3rd Cranial nerve; saw eye surgeon 5/4/2021 after my right eyelid went completely shut, he confirmed issues with 3rd Cranial nerve. Now wearing eye patch on right eye to prevent double vision and I have follow up appointment on Tuesday, May 11.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294699-1	Severe pain and weakness in extremities that progressed upwards through legs, hips, arms, back, neck and head; new and sudden onset of popping joints; Temporary paralysis and numbness upon waking up in the morning; Temporary paralysis and numbness upon waking up in the morning; severe muscle fatigue; experiencing pain, weakness, and tingling in upper back, both arms and hands; Heart rate spike to 180; dizziness; sweating; sudden aching and weakness in both hands, progressed upwards through legs, hips, arms, back, neck and head; sudden aching and weakness in both hands; Severe head and facial pain; Severe head and facial pain; Pain, tingling, burning, and weakness in extremities and limbs; Pain, tingling, burning, and weakness in extremities and limbs; Vibrations/tremor-like sensation in upper back; Tingling tongue; This is a spontaneous report from a contactable consumer, the patient. A 33-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, lot EN6200, first dose) solution for injection intramuscular in the left arm on 27Feb2021 at 10:00 (at the age of 33-years-old) as a single dose for COVID-19 vaccination. Medical history included SVT (supraventricular tachycardia). Concomitant medications were not reported. Prior to the vaccination, the patient was not diagnosed with COVID-19. On 27Feb2021, the patient experienced heart rate spike to 180 within first 2-3 minutes of vaccine with dizziness, sweating and sudden aching and weakness in both hands; severe head and facial pain, pain, tingling, burning, and weakness in extremities and limbs; vibrations/tremor-like sensation in upper back; tingling tongue; severe muscle fatigue; experiencing pain, weakness, and tingling in upper back, both arms and hands, sent to ER for monitoring. Continued heart rate spikes for approx. 7 days afterwards. Severe pain and weakness in extremities that progressed upwards through legs, hips, arms, back, neck and head over 1 week period. On 09Mar2021, 10 days after vaccination; two more ER visits. Severe head and facial pain. Pain, tingling, burning, and weakness in extremities and limbs. Vibrations/tremor-like sensation in upper back. Tingling tongue, new and sudden onset of popping joints, temporary paralysis, and numbness upon waking up in the morning. Multiple lab work orders, X-rays, MRIs - all ruled out other causes/conditions in Mar2021. The events resulted in doctor or other healthcare professional office/clinic visit, Emergency room/department or urgent care, with disability or permanent damage due severity of symptoms required 4 weeks of medical leave from work. Treatment for the events included medication and physical therapy. The outcome of the events was not recovered. Post vaccination, a Nasal Swab, Covid test result was negative in Mar2021.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1294738-1	"Significant weight loss; Chronic back pain; terrible pain in the back; Osteolysis; Terrible pain in her extremities; It was her first dose and she was afraid to get the second one now; patient was supposed to get her second one last week and she didn't; Paralyzed from her neck down to her arm, legs, and everything; weakness in extremities; could not move; Received BNT162B2 via subcutaneous route; This is a spontaneous report from a contactable nurse (patient who is a retired nurse who did anesthesia). A 66-year-old female patient received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; Lot Number and Expiration Date were not reported), subcutaneous in right deltoid of the upper arm, on 09Mar2021, at a single dose, for COVID-19 immunization. The patient's medical history included pain, use of medication to help with sleep, gastroesophageal reflux disease (GERD), and thyroid. The patient's concomitant medications included levothyroxine for thyroid, pantoprazole sodium sesquihydrate (PROTONIX) for GERD, trazodone to help with sleep, zolpidem tartrate (AMBIEN) to help with sleep, morphine (extended relief) for pain, and morphine sulfate (immediate relief) for pain. They also gave the patient diazepam (VALIUM) if she needs to take it but she did plan; she didn't take it. The patient just didn't take diazepam and then pain management doctor put her on morphine extended release. Then, morphine sulphate immediate relief, so she can take up to 4 for day and she was not using now. The COVID-19 vaccine was administered in a pharmacy. The patient would like to report a reaction that she received that she had approximately for an hour and a half, that basically started in an hour after the injection. The patient's first COVID-19 shot was given by Pfizer on 09Mar2021 and what happens in next hour was she became totally paralyzed from her neck down to her arm, legs, and everything. It was approximately about 30 minutes after the shot that she began to notice some weakness in extremities (also reported as eternities) and then they got home an hour later, and her husband left the patient then she got to know she could not move. The patient can breathe, and she was totally conscious, but she could not move, could not pull herself along the floor, could not pull from the sofa, could not move her knee, and could not move her body. So, the patient thought maybe she should report this. On an unspecified date, the patient had significant weight loss. The patient was eating but she was rapidly losing. The patient didn't have COVID-19; she had been tested negative, so she didn't have COVID-19. The patient was tested 4 times for her to be around her grandchildren and make sure everything is okay. The patient had been tested against COVID-19 because of her grandchildren. The patient tried to take them, and she just now received the paper it was negative. So, it was her third one to try to watch her grandchildren because they were infant, and all test results were negative. It was also reported that the patient had no other problem, just the pain clarified as chronic back pain since an unspecified date. The patient checked this as persistent for chronic back pain; it is positive, definitely it's totally osteolysis on an unspecified date. Treatment was received for all events (unspecified medications). The patient went to her primary healthcare doctor as a result of the events and he said it was no big deal and she told him that she disagrees with that. Trazodone was added to zolpidem tartrate to help w with the sleep because she had terrible pain in the back (right here) and her extremities after this incident (unspecified date). It was her first dose and she was afraid to get the second one now. The patient was supposed to get her second one last week and she didn't. So, they told her to come back in 3 weeks or 4 weeks. The patient underwent laboratory test back in December (unspecified year). The patient recovered from ""Paralyzed from her neck down to her arm, legs, and everything; weakness in extremities; could not move"" on 09Mar2021. The patient had not recovered from ""Chronic back pain; terrible pain in the back"". The outcome of the other events was unknown. Information about lot number and expiration date for the suspect product will be requested in follow-up attempts.; Sender's Comments: A contributory role of BNT162B2 to event paralyzed from her neck down to her arm, legs, and everything cannot be excluded based on close 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 Regulatory Authorities, Ethics Committees and Investigators, as appropriate."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1299120-1	the neurologist agreed that the vaccine has attacked nervous system; developed signs of stroke, he can't stand up and walk up straight; feeling unwell and was unable to work for a few days; he had tingling on the left side that moved to his right side; he has difficulty speaking/ word finding/Slurring his speech; tinnitus; nausea; pain in the rear of his neck, between the neck and skull; His symptoms continue to be severe and is unable to work; he cannot coordinate his thoughts and fantasies; dizziness, he felt his head was spinning; He had some kind of palsy where his face was not reacting bilaterally; pressure in his brain; Got really sick 18 hours after it and it lasted for like two days where he was really sick, all sorts of symptoms; This is a spontaneous report from a contactable consumer (patients wife). A 65-years-old male patient received bnt162b2 (BNT162B2), dose 2 via an unspecified route of administration on 25Mar2021 (Batch/Lot number was not reported) as SINGLE DOSE for covid-19 immunisation . The patient medical history was not reported. The patient's concomitant medications were not reported. The patient experienced the neurologist agreed that the vaccine has attacked nervous system (nervous system disorder) (hospitalization) on an unspecified date with outcome of unknown , developed signs of stroke, he can't stand up and walk up straight (cerebrovascular accident) (medically significant) on an unspecified date with outcome of unknown , feeling unwell and was unable to work for a few days (malaise) (medically significant) on an unspecified date with outcome of unknown , he had tingling on the left side that moved to his right side (paraesthesia) (medically significant) on an unspecified date with outcome of unknown , he has difficulty speaking/ word finding/slurring his speech (dysarthria) (medically significant) on an unspecified date with outcome of unknown , tinnitus (tinnitus) (non-serious) on an unspecified date with outcome of unknown , nausea (nausea) (non-serious) on an unspecified date with outcome of unknown , pain in the rear of his neck, between the neck and skull (neck pain) (non-serious) on an unspecified date with outcome of unknown , his symptoms continue to be severe and is unable to work (loss of personal independence in daily activities) (medically significant) on an unspecified date with outcome of unknown , he cannot coordinate his thoughts and fantasies (coordination abnormal) (medically significant) on an unspecified date with outcome of unknown , dizziness, he felt his head was spinning (dizziness) (medically significant) on an unspecified date with outcome of unknown , he had some kind of palsy where his face was not reacting bilaterally (paralysis) (medically significant) on an unspecified date with outcome of unknown , pressure in his brain (head discomfort) (non-serious) on an unspecified date with outcome of unknown , got really sick 18 hours after it and it lasted for like two days where he was really sick, all sorts of symptoms (illness) (non-serious) on an unspecified date with outcome of unknown. Information on the lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1299130-1	a woman who was paralyzed after getting her doses of the Pfizer vaccine.; This is a spontaneous report from a contactable consumer. A female patient of an unspecified age received bnt162b2 (BNT162B2), first dose via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose, second dose via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as single dose for COVID-19 immunization. The patient medical and concomitant medications were not reported. The reporter reported that they are doing a story with a woman who was paralyzed on an unspecified date after getting her doses of the Pfizer vaccine. Outcome of the event was unknown. Information on the lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1299193-1	paralysis; rapid heartbeat; heart palpitations; cold sweats; chills; muscle pain; in arm; nausea; difficulty breathing; This is a spontaneous report from a Contactable Consumer (patient). An 88-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 mRNA VACCINE, Lot Number: ER2613), first dose via an unspecified route of administration, administered in Arm Left on 05Apr2021 (at 88 years old) for covid-19 immunisation. Medical history included breast cancer. She had breast cancer surgery on that side previously. She had chills on and off before the Covid vaccine also. Says it started after her Pneumonia and Flu shots that she believes were in Nov2020; possibly 12Nov2020. Concomitant medications included spironolactone and other unspecified medicines. She had them go over her medications to see if any of them would interact with the vaccine. They said that all of her medications were okay. On Apr2021, patient experienced heart palpitations, paralysis, cold sweats, chills, muscle pain, in arm, nausea and difficulty breathing about three weeks after the vaccine. She noticed that it mentioned concerns if you experience a rapid heartbeat. States she did, and is wondering about getting her second vaccine. Her rapid heartbeat became so bad this past Tuesday night she thought she was having a stroke. It came over her and she could not move. She went to an urgent care center. They did a Covid 19 Test and it was negative on Apr2021. Also states she has been having lots of chills. The outcome of the events Heartbeats increased, palpitations, Chills was not recovered, other events was unknown.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1299274-1	"legs were heavy and numb; weakness in legs; bladder was now the problem/brain doesn't tell the body to go to the bathroom she is leaking because the bladder is filling up filling over; reason for hospitalization;her kidneys; totally became paralyzed from the waist down; legs were heavy and numb; Patient was a little nervous; painful tingling feeling she have that kind of feeling in her legs; This is a spontaneous report from a contactable consumer (patient's mother) from a Pfizer-sponsored program,. A 39-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 20Apr2021 at 19:30 (Batch/Lot number was not reported) as single dose for covid-19 immunization and pregabalin (LYRICA), via an unspecified route of administration from an unspecified date at 100 mg, 3x/day for neuropathy. Medical history included neuropathy from 2020 and ongoing. The patient's concomitant medications were not reported. It was reported that the patient got the vaccine and she was taking pregabalin as well and after vaccination, she experienced an adverse event, her legs were heavy and numb and also her bladder was now the problem she was actually now at the emergency room. Patient was a little nervous. The patient was still in the hospital and she was not getting any care. What happened at the beginning was that she totally became paralyzed from the waist down, when your foot fall asleep you get painful tingling feeling she have that kind of feeling in her legs not able to move any part of her anywhere from her waist down later on that night, after she took pregabalin, went to sleep about an hour she was able to move her left toes and able to drag one leg a little bit. She was supposed to take pregabalin 3 times a day and they were giving her a difficult time about it, she took one at 12 O' clock on Wednesday and the hospital they did not give another one until 11 o clock that evening because she was supposed to take 3 times a day. After she no longer feel from the waist down, she took a pregabalin and it didn't help her it didn't work and her physician told her to take 3 times a day and then at 10 O' clock yesterday morning she was still asking the hospital to give her another one. When asked to repeat the concern, patient stated ""weakness in her legs that felt very heavy like she was dragging. The date of hospitalization was reported as 21Apr2021. The reason for hospitalization was reported as numbness in her legs and weakness in legs, became paralyzed from the waist down, and her kidneys. So her brain doesn't tell the body to go to the bathroom she is leaking because the bladder is filling up filling over and the body doesn't know to get it out its leaking because it is filling up it is so full. The patient underwent lab test in the hospital. The action taken with pregabalin in response to the events was unknown. Outcome of the events was unknown."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1299354-1	had a stroke that left him paralyzed; had a stroke that left him paralyzed; This is a spontaneous report from a contactable consumer (patient). A 60-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 24Apr2021 at 09:45 (lot number: EW0158) as SINGLE DOSE for COVID-19 immunization. The patient medical history and concomitant medications were not reported. The patient had no allergies. 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. On 27Apr2021 at 11:15, the patient had a stroke that left him paralyzed. The events resulted in emergency room/department or urgent care and doctor or other healthcare professional office/clinic visit. The events were considered serious (hospitalization, disability, life threatening). Therapeutic measures were taken as a result of the events which included full stroke treatment (as reported). The patient had Covid-19 nasal swab test with unknown results on 27Apr2021. The outcome of the events was recovered with sequel. No follow-up attempts are needed. No further information is expected.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1304459-1	Weak when woke up. Fell 2 times was then paralyzed on that Sunday A. M. went to ER was Dx with GBS . Guillan Barre Syndrome. Hospitalized. Now in rehab hospital
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1306231-1	paralyzed; This is a spontaneous report from a contactable consumer reporting for a patient. A female patient of an unspecified age received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as 2nd dose, single for COVID-19 immunisation. The patient medical history and concomitant medications were not reported. The reporter interviewed a woman who said she was paralyzed on an unspecified date after receiving the second dose of the Pfizer vaccine. The reporter mentioned they understood this is incredibly rare, and this has not been confirmed to have a connection to the vaccine, but would like to include a statement from Pfizer on these claims. The event outcome was unknown. Information on the lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1309612-1	Paralysis; This is a spontaneous report from a non-contactable consumer. A 73-year-old male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on an unspecified date (Batch/Lot number was not reported) as 1st dose, single for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient suffered from paralysis after he was vaccinated. He has been having issues with the left side of his body from head to toe since receiving the first shot of the Pfizer vaccine. He went to local hospitals where doctors ran every test and can't find anything wrong. He went to a local chiropractor who said he's seen similar issues from the vaccine in about 12-20 patients. The outcome of the event was unknown. The event was assessed as serious-medically significant. No follow-up attempts are possible, information about lot/batch number cannot be obtained.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1310979-1	At five mins Began with dizziness, then rash then diff breathing. Emt called to store and during ambulance transport to emergency room, extremities grew tingly then numbe then contracted stiffly (they called it posturing) in ER became stiff and entirely paralyzed. Spiked fever. After unknown amount of time gradually loosened up, did not have full mobility for over four hours. Following day still weak and trouble walking, bad leg pain from when they were postured. Brain fog, trouble thinking of right words, off and on dizzy.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1313002-1	"complete paralysis/paralyzed/can't move anywhere/can't move her arms and legs; muscle weakness; This is a spontaneous report from two contactable consumers (patient herself and her husband). A 45-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number and expiry date not reported), via an unspecified route of administration, administered in the right arm (also reported as right shoulder), on 13Apr2021, as 1st dose, single, for COVID-19 immunization, at a pharmacy/drug store. Medical history included allergies to latex, bulimia, depression, and COVID-19 (prior to vaccination). The patient previously took codeine and experienced allergies. The patient was not pregnant at the time of vaccination. The patient was not tested for COVID-19 since the vaccination. Concomitant medications were unspecified (taking other medications). The patient did not receive any other vaccines within 4 weeks prior to BNT162B2. On 16Apr2021, the patient experienced complete paralysis which started with muscle weakness 3 days after the vaccine. The events resulted in emergency room visit, hospitalization (also reported as prolonged hospitalization, pending clarification), disability or permanent damage. The patient was still in the hospital at the time of the report. Treatment for the events included spinal tap, IV fluids, and potassium intake treatment. On 29Apr2021, it was reported that the patient was having a really bad reaction (unspecified), was paralyzed, could not move anywhere and in the hospital from 25Apr2021 and ongoing. The patient underwent lab tests in Apr2021 with unknown results. On 08May201, it was reported that the patient could not move her arms and legs and is ""nursing home"" (pending clarification). The reporter assessed the events as serious (hospitalization and disability). The outcome of the events was not recovered."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1316147-1	"he had an episode which he states is similar to a TIA (transient ischemic attack); Reports experiencing something similar last may except for 'the paralysis'.; numbness, pain and tightness in his lower half of his body which was more prominently felt on his left side.; numbness, pain and tightness in his lower half of his body which was more prominently felt on his left side.; numbness, pain and tightness in his lower half of his body which was more prominently felt on his left side.; States he felt as though his knee had a tear in it- it felt it was on fire; tightness in left knee; shortness of breath; feeling foggy; headache; balance issue; coordination off; having a hard time to get a sentence out (felt like his motor skills were affected); having a hard time to get a sentence out (felt like his motor skills were affected); confusion; blurred vision; impending sense of doom; post fatigue; his energy doesnt feel up to par; This is a spontaneous report from a contactable consumer (patient) reported for himself. A 28-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 1 via an unspecified route of administration on 19Apr2021 at 28 years old (Batch/Lot number was not reported) as single dose for covid-19 immunisation. The patient's medical history and concomitant medications were not reported. Patient stated that on 19Apr, he had the first shot of pfizer covid vaccine. This past sunday (25Apr2021), he had an episode which he stated was similar to a TIA (transient ischemic attack). Reported numbness, pain and tightness in his lower half of his body which was more prominently felt on his left side. Stated he felt as though his knee had a tear in it- it felt it was on fire; he experienced burning and tightness in left knee. He also reported having shortness of breath, headache, feeling foggy, balance issue, coordination off, having a hard time to get a sentence out (felt like his motor skills were affected), confusion, blurred vision and an impending sense of doom. Stated this episode persisted for 15 min and then it resolved. Currently he was experiencing a 'post fatigue' and his energy didn't feel up to par. Reported experiencing something similar last may except for 'the paralysis'. Had not followed up with his provider. Patient was scheduled for second shot of Pfizer covid vaccine on 10May, he was asking for information if he decided to delay his second shot past may 10th. He was asking for information on getting the second shot given his recent experience possibly related to the vaccine shot. Was not sure if he should get the second shot. He was asking about side effects for the second shot- if it would be worse. Asking if there were any reports of blood clots with Pfizer covid vaccine. The outcome of the events 'post fatigue' and ""his energy didn't feel up to par"" was unknown, of other events was recovered on 25Apr2021. Information about lot/batch number has been requested."

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Paralysis; Sore for four days\Arm was very sore; Mild headache; Mild nausea; throbbing pain temple; tongue swells; lips felt numb/her tongue went numb; Heaviness to her tongue; flu like symptoms; her face a has a feeling of numbness on her right side; asthma; allergies/Food allergy/completed for initial report (PRD 30Apr2021, SRD 01May2021) and FU#1 (PRD 29Apr2021, SRD 01May2021); the second shot hurts; This is a spontaneous report received from a contactable consumer (patient reported for herself). A 71-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), dose 2 via an unspecified route of administration on 25Mar2021 at the age of 71 years old (Batch/Lot number: ER8732, expiration date: 31Jul2021) as 2nd dose, single for covid-19 immunisation. Medical history included asthma, some allergy, seasonal allergies, allergy to mold, high blood pressure, high cholesterol. Concomitant medications were not reported. The patient previous received the first dose of bnt162b2 on 04Mar2021 (Lot Number: En6199, Expiration Date:30Jun2021) at the age of 71 years old for covid-19 immunisation and what happened was just this injection site was hurting for about 4 days and she just had some fever and nausea. The patient asked if Pfizer had provided a statement on the need for a booster dose (3rd dose), if Pfizer was investigating the need for a booster dose, if she would receive a Pfizer dose. The patient reported that usual her arm did not even swell. Sore for four days. She did not take anything. Mild headache. She could not stand the smell of chicken for instance. Mild nausea. Second dose, throbbing pain temple. She had asthma, allergies. Took Tylenol. Five and half hours later she craved salad. She had had Haig tabbouleh. Fresh lemon. She did react to walnuts, tomatoes. Her tongue swells. She took one taste of the tabbouleh. Her lips felt numb. Heaviness to her tongue. She called the nurse. On-call Dr Food allergy. Suggested she went to bed. Arm was very sore. She took a Ibuprofen. By that night, flu like symptoms. Just went to bed. Her own doctor. That was the vaccine. Immune system was working. Paralysis. eggplant. Do you think it could cause food allergies? The patient hesitant to take a booster dose. As of 29Apr2021, patient reported she got her first shot on 04Mar2021 and her 2nd shot on 25Mar2021. She was surprised at herself that was not sacred. She had been sheltering at home for a year as she came under the critical category. Patient further stated she had reaction she thought to the vaccine from second shot so she wanted to report it and she needed some clarification on that she did reported this last week about it (further clarification unknown). She just wanted some reassurance and because what happened was, from the first shot she was fine she had been tested there for 30minutes of 15 for her allergies and asthma was fier. So, what happened was just this injection site was hurting for about 4 days and she just had some fever and nausea but the second shot hurts and hours after the second shot she wanted to eat something and so she ate some salad. For Weight, Consumer stated was about 165 pounds, she had not weighed herself lately. For Treatment: Consumer stated she came back she did take Tylenol for the pain and for the headache. Consumer further stated she had been waiting to talk to you not for the headache but because she had reaction when she ate something. She ate salad and she had eaten the salad before and it had some lemon juice, olive oil and it was cracked with tomatoes and cucumber and parsley and she had eaten the one my whole life and these particular take out that, she took from grocery for the salad. So, when she ate it, it bothered her tongue just a little bit because her tongue reacted when she eat citrus fruits like tomatoes and spicy things. So, what happened was as soon as she took a taste of this salad, her tongue went numb like she had a Novocaine shot (further clarification unknown), her lips went numb, her face a has a feeling of numbness on her right side and it lasted for 8 hours and she talked to her doctor and he said, it was food allergy or it may due to be a vaccine. Investigation Assessment was no. The outcome of the events was unknown.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1320123-1	
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1320187-1	Pins/Needles; Paralysis; Full body weakness; Numbness; This is a spontaneous report from a contactable consumer (patient) A 43-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 07Apr2021 15:15 as single dose, for covid-19 immunisation. Medical history included Raynaud's phenomenon, cystitis interstitial, motion sickness, allergy to sulfa and macrobid. The patient's concomitant medications were not reported. 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 paralysis on 07Apr2021 15:15, full body weakness on 07Apr2021 15:15, numbness on 07Apr2021 15:15, pins/needles on an unspecified date. The patient was hospitalized for the events for 4 days (start date not reported). No treatment received. The patient underwent lab tests and procedures which included sars-cov-2 test: negative on 17Apr2021, sars-cov-2 test: negative on 29Apr2021. The events outcome was not recovered. Information on the lot/batch number has been requested

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1320243-1	Severe muscle soreness (paralyzing); Severe muscle soreness (paralyzing); soreness in tissue and bones; soreness in tissue and bones; unable to walk straight; numbness in hands, arms, and feet.; This is a spontaneous report from a contactable consumer (patient). A 49-year-old female patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), dose 2 via an unspecified route of administration, administered in arm left on 15Apr2021 11:00 (Batch/Lot Number: EW0162) at the age of 49 years, as 0.3 ml single for covid-19 immunisation. Previously the patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) Lot: EN6207, on 25Mar2021 at 11:00 AM at 0.3 ml single in left arm. Medical history included calcium deposits and mitral valve prolapse, from an unknown date and unknown if ongoing. The patient was not diagnosed with COVID-19 prior to vaccination and has not been tested for COVID-19 since the vaccination. The patient did not receive any other vaccines within 4 weeks prior to the COVID vaccine. The patient's concomitant medications were not reported. On 15Apr2021 at 18:00 the patient experienced severe muscle soreness (paralyzing), soreness in tissue and bones, unable to walk straight, numbness in hands, arms, and feet (severe symptoms as such for 4 days). The outcome of the events was resolving.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1329523-1	bacterial meningitis; paralysis; he ended up in a coma.; This is a spontaneous report from a contactable consumer. A male patient of an unspecified age received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE lot/batch number and expiration date not reported) via an unspecified route of administration on an unspecified date as 1st dose, single for COVID-19 immunization. The patient's medical history and concomitant medications were not reported. The patient experienced paralysis after getting the first dose of the vaccine, then was later diagnosed with bacterial meningitis. The patient said his doctors said it was a possibility that it was linked to the vaccine because it was the only thing he did differently that day. Symptoms came on quickly and he ended up in a coma. Outcome of the events was unknown. Information on the lot/batch number has been requested.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332777-1	"Transverse myelitis; Paralyzed; A young pregnant woman (Patient); 6 months pregnant; This is a spontaneous report from a contactable consumer. This is the second of 2 reports for the second dose. A 37-year-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) via an unspecified route of administration on 01Apr2021 (Lot number was not reported) as single dose for covid-19 immunisation. The patient medical history and concomitant medications were not reported. Historical Vaccine included the first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE) on an unspecified date as single dose for covid-19 immunisation and experienced myelitis transverse. The patient received the second dose of bnt162b2 on 01Apr2021 and by 03Apr2021 she was ""paralyzed"". The patient was 6 months pregnant and had a ""transverse myelitis"" attack after the vaccine. Events outcome was unknown. Information about lot/batch number has been requested."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1332780-1	Leg spasms; She has a patient who is pregnant with her first child. She had the second dose of the vaccine on April 1st.; Just 2 days after she had leg spasms then it led to being paralized; she had leg spasms then it led to being paralized. It spread through her whole body, even her stomach; She is now a transverse/ have Transverse mylighttis as well; she had leg spasms then it led to being paralized. It spread through her whole body, even her stomach; This is a spontaneous report from a Pfizer sponsored program support from a non contactable consumer. A 37-year-old female patient received second dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE) on 01Apr2021 at single dose for COVID-19 immunization. Medical history and concomitant medications were not reported. The patient previously received first dose of BNT162B2 on unknown date. The patient was pregnant with her first child. She had the second dose of the vaccine on 01Apr2021. Just 2 days after she had leg spasms then it led to being paralized. It spread through her whole body, even her stomach. She was put into the hospital and she was now a transverse and was going to hospice. She was only 37 years old and the caller wanted to know what happened to other people who have transverse mylighttis as well. The outcome of all events was unknown. No follow-up attempts are possible; information about lot/batch number cannot be obtained.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1334472-1	Shot given 11:58am. Within 2 minutes symptoms began with feeling symptoms: nauseous, hot forehead. I asked CVS for the nurse to help. By 12:01 patient was feeling extreme pain in her stomach, was sweating profusely, felt tingling sensation from head down to her stomach (including her arms and hands). Patient began screaming in pain. Her hands became rigid and paralyzed. She was completely unable to move both hands. Patient was able to feel if someone was touching her hands. Her face began tingling and feeling numb. Patient was lying down and almost passed our. I asked CVS to call 911 and if they should administer an epi pen. The epi pen was not administered. I (mother) went w/ child in the ambulance to Hospital. EMT monitored her vitals, which were normal. Patient was still unable to move her hands until after she was admitted to Hospital. Patient did not receive any IV r medication at Hospital. The sensation in her hands slowly came back with the numbness subsiding after about 1.5 hours in the ER. Patient was discharged at about 2:00pm on 5/16/2021.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1344160-1	Are swelling and paralysis, fever, disorientation, headaches,

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1349564-1	I was suffering from it, I was not able to move my mouth so, I tell my husband I couldn't move; tired; had a chills; it also made my back (incomplete sentence), I have problem with my back; My back pain was worsen; I was getting out of breath too; I can hardly walk; Bell's Palsy/she can move the rest of her body except for her face; paralyzed; This is a spontaneous report from a contactable consumer (patient) reported for herself. A 64-years-old female patient received the second dose of bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 06May2021 at 10:15 (Batch/Lot Number: ER8736) as single dose for covid-19 immunisation. The medical history and concomitant medications were not reported. On 06May2021 night the patient woke up paralyzed which was what a little bit she can find out was Bell's palsy. The patient was not able to move her mouth so, she tell her husband she couldn't move. The patient was also tired and had a chills too and other than that it also made her back (incomplete sentence), the patient have problem with her back. Her back pain was worsen for some reason. The patient do not know if it was from the shot or what. She have concerns with paralysis. The patient can hardly walk and it was outside and it will take a little while. She got a bad back, a bad her and a bad shoulder. The patient was falling apart. Hopefully she can get a new her and a new shoulder. The patient was getting out of breath too. Other things are tiredness and my back hurting a lot. Yesterday she was like getting tired and out of breath, usually she do not get out of breath like this. The outcome of the events was unknown.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1357367-1	Severe sore throat; severe pain; could not swallow; Ramsay Hunt syndrome; herpes zoster/shingles; cannot speak; half face paralyzed; This is a spontaneous report from a contactable consumer (patient). A 74-year-old male patient received bnt162b2 (PFIZER-BIONTECH COVID-19 VACCINE, lot number: EN6199), dose 1 via an unspecified route of administration, administered in Arm Left on 04Mar2021 18:30 as 1st dose, single for covid-19 immunisation. Medical history was reported as none. Concomitant medication included irbesartan. On 18Mar2021 16:00, a severe sore throat began, on 21Mar2021, the patient entered for emergency to hospital, severe pain and could not swallow, on 22Mar2021 he was hospitalized, for 14 days, the first 3 in the intensive care unit. Doctors have given his ailment different names: Ramsay Hunt syndrome, herpes zoster, shingles. On 04Mar2021 they sent me home, but under medical supervision from the hospital. To this date 22May2021 the patient continue with the adverse effects: he cannot swallow, tube feeding into the stomach, he cannot speak, half face paralyzed. The patient received treatment for ae included steroids, antivirals, analgesics. AE_treatment Lupus Erythematosus, Rheumatoid Arthritis, Fibromyalgia (pending confirmation). The adverse event result in visit to the doctor's office or other healthcare professional, Emergency room / department, Hospitalization, Life-threatening illness (immediate risk of death from the event. 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 outcome of the events was resolving.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1362833-1	"Patient states he got the covid vaccine in the morning on 4/30/2021 and by afternoon he was having difficulty walking - states he was having trouble picking his feet up. Patient states he was feeling weak and the next day had to start using a cane (walked without assistance prior to vaccination). Patient states he started losing feeling/strength/movement in feet and legs 4/30/2021, with the weakness/paralysis moving up the legs over the course of a month. Patient states he went to primary doctor approximately 3 weeks after his vaccination and reported his symptoms but his doctor ""wasn't concerned"". On arrival to the ER 5/31/2021 patient was totally unable to sit up unassisted and stated his decreased sensation and loss of movement had risen to bilateral hips. Patient had to be lifted to the stretcher by four staff as he was unable to help at all, with legs dangling/flaccid paralysis obvious. Patient also reports loss of bowel/bladder control since vaccination - states started losing that control as decreased sensation/movement ascended to hips/pelvic region."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1370488-1	Paralyzed can't feel/move my legs; feel tired; Paralyzed can't feel/move my legs; This is a spontaneous report received from a contactable consumer (patient). An elderly male patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection, Batch/Lot number and Expiration date was not reported), via an unspecified route of administration on an unspecified date, as 2nd dose, single dose for COVID-19 immunization. The patient medical history and concomitant medications were not reported. Patient previously took 1st dose of BNT162B2 for COVID-19 immunization. Patients wife and patient had the second dose, in the middle of the night paralyzed can't feel/move his legs. 03:00 am felt better. 10 days since the shot, feel tired. Mentioned it to the doctor. Rapid COVID Test. Covid have the second dose. Then patient had the five day thing. Feeling very tired. Doctor stated could endure 6 months. Patient and his wife are gonna take another rapid test. Patient saying have your stock. The outcome of the events was unknow. Information on lot number/batch number has been requested.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1370586-1	Nightly sleep seizures/spastic seizure lasting 5 minutes of uncontrollable myoclonus only of the left leg; Temporary, passing paralysis; spastic seizure lasting 5 minutes of uncontrollable myoclonus only of the left leg; longer aphasia (inability to speak); severe migraines; This is a spontaneous report from a contactable consumer. This 32-year-old female consumer reported that she received 2nd dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE; lot Ew0179) on 13May2021 at 02:45 PM into left arm for COVID-19 immunisation. Medical history included Hypermobile Ehlers-danlos syndrome, non-epileptic seizures, migraines, mild traumatic brain injury, intractable pain, severe food allergies. Known allergies: Wheat, gluten, eggs, soy, estrogens. Historical vaccine included 1st dose of BNT162B2 (lot Ew0171) on 22Apr2021 at 02:45 PM into left arm for COVID-19 immunisation. Concomitant drugs included Morphine, hydrocodone bitartrate, paracetamol (NORCO), diazepam (VALIUM), Propranolol, Hydroxyzine. The patient had a non-epileptic seizure disorder that was well controlled. After the second dose she began to have nightly sleep seizures, severe migraines, but am writing because of a spastic seizure lasting 5 minutes of uncontrollable myoclonus only of the left leg, with longer aphasia (inability to speak) and temporary, passing paralysis. The symptoms passed in an hour but an epileptic patient could have died. Event onset time was 19May2021 06:30 PM. No treatment was received. The outcome of the event was resolving.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1370634-1	Within 30 minutes of receiving the 1st dose of the covid19 vaccine in my left shoulder my left hand and wrist became completely paralyzed.; Ulnar Palsy; This is a spontaneous report from a contactable consumer (patient). A 43-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Lot Number: EN6207, expiration date not provided), 1st dose via an unspecified route of administration, administered in arm left on 16May2021 15:30 at age of 43 years old as a single dose for COVID-19 immunisation. Medical history included latex sensitivity. The patient's concomitant medications were not reported. The patient experienced within 30 minutes of receiving the 1st dose of the COVID-19 vaccine in her left shoulder her left hand and wrist became completely paralyzed on 16May2021 16:00. The ER said it was random coincidence and diagnosed it as Ulnar Palsy on 16May2021 16:00. The events were reported as being disability. Therapeutic measures were taken as a result of all events. The outcome of the events was not recovered. The patient had not received any other vaccines within 4 weeks prior to the COVID vaccine. The patient had not received any other medications within 2 weeks of vaccination. The adverse event resulted in visiting doctor or other healthcare professional office/clinic. The adverse event result in visiting Emergency room/department or urgent care. Treatment for the adverse event was included ER (emergency room) Exam. The patient was not pregnancy at time of vaccination. Prior to vaccination, the patient was not diagnosed with COVID-19. Since the vaccination, the patient had not been tested for COVID-19. Information on the lot/batch number has been requested.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1373739-1	"Collapsed day after vaccine first dose/Collapsed day after vaccine second dose; His left side still feels paralyzed and can't walk; legs wouldn't move, couldn't turn around; He had compression fractures L1 to L3 that were noticed on his x-ray; he collapsed and had a hard time getting up/felt bad; General malaise; His left side still feels paralyzed and can't walk; bedridden; Pain; Half of back is black from incident; Mobility got worse after first shot and then after second dose his mobility got even worse; felt like what I would call the flu; Collapsed day after vaccine first dose/Collapsed day after vaccine second dose; This is a spontaneous report received from a Pfizer sponsored program. A 72-years-old male patient received first dose of bnt162b2 (PFIZER-BIONTECH COVID-19 MRNA VACCINE, Solution for injection), via an unspecified route of administration, administered in Arm Left on 16Apr2021 (Batch/Lot Number: EP7533) as 1st dose, single dose and second dose via an unspecified route of administration, administered in Arm Left on 07May2021 14:00 (Batch/Lot Number: EW0182) as 2nd dose, single dose for covid-19 immunisation. Medical history included L1 to L3 compression fractures somewhere around 2008 to 2010, ongoing gout (from about 35 years ago) (pain in both hips and right shoulder, left shoulder some times), gouty arthritis (30 years constant pain), ongoing moderate emphysema (since 20 years ago), stroke (happened in 2010). States he was taking Warfarin at the time of the stroke and the stroke was permanent, cholesterol, smoker (20 years), excess uric acid, allergy in right sinus, cough uncontrollably, perirectal abscess and low blood pressure. Family medical history was none. States he is adopted. Concomitant medications included atorvastatin taken for blood cholesterol from an unspecified start date to unspecified date in 2021 (Taking for about 5 years, not sure if the dose is 30mg, 50mg, or 300mg, stopped after adverse event with covid vaccine), zolpidem tartrate (AMBIEN) taken for sleeping from an unspecified start date to unspecified date in 2021 at 10mg (taking for couple of decades, stopped after adverse event with covid vaccine, stated it may cause drowsiness) and allopurinol taken for gout at a dose of 300 mg, (taking for 6 or 7 years) and azelastine hydrochloride nasal spray taken for uncontrollable coughing via nasal route as 2 shots each nostril twice a day (uncontrollable coughing caused perirectal abscess, had scope done, diagnosed as allergy. Azelastine works for cough). The patient previously took pneumonia shot and stated that it hurt like hell for a week, flu shot and got sick and also took warfarin. The patient reported that he received his first dose on 16Apr2021, then the next day he collapsed and had a hard time getting up. He also reports general malaise, ""just felt bad"", and ""felt like what I would call the flu"" for the next 5 days. On 07May2021 he received his second dose, and the same thing happened where he collapsed the next day. This time, he was not able to get up on his own and had to call 911. He was admitted to the hospital and ""had 2 X-rays and a CT scan"". CT of torso he reports his CT scan

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				and "had 2 X-rays and a CT scan", CT of torso he reports his CT scan showed compression fractures at L1, L2, and L3 on his x-ray. He is taking a lot of Norco to treat his pain. States the music on hold is so uplifting that he wants to go cut his throat. Caller stated he was bedridden. He reported ""I can't walk, I used to be able to, and now I can't"". He spent 5 years in a computer testing laboratory and he knows a little about testing. He states a lot of the rules apply to conductors and automobiles. Caller stated he wants to know what happened to him. He says if he dies as a result of this, there will be mountains of paperwork. He also said if his condition is permanent, there will also be mountains of paperwork. He can't go to the bathroom and this is an issue. He was falling down and collapsing several times with both shots. He says this is a major problem. He declined being admitted to assisted living saying ""if I'm going to die I'm going to die at home"". Caller would like to know if anything like this has been reported before, if they got better, and what treatment they received to get better? He also asked if there were any reported deaths after vaccination? Events paralyzed, bedridden, can't walk, legs wouldn't move, couldn't turn around was reported as worsened, half of back is black from incident. The patient was taking Pfizer Covid Vaccine as needs to show he had vaccine to travel. He hasn't measured his height since compression fractures. Lost 20% on 3 vertebrae, so he might only be 5 feet 10 inches. He figures 20% would be an inch lost with his calculation. He can't reach around 180 degrees to measure himself. When asked to provide NDC, lot number and expiration date from vaccine card, caller stated he needed to use magnifying glass. He then stated that his vision is good. He stated that he collapsed several times after his second shot. He says this happened approximately 2 to 3 days after the second shot. He called his doctor and was told to call 911. He went to the hospital. He was in the ER, but confirmed he was not admitted to the hospital. Outcome for collapsing after both doses, had worsened as mentioned by patient. Caller stated after the first vaccine, he noticed his left leg did not work very well. After first dose, he had trouble walking after 5 days. When he would try to move his left leg forward it would shake, not move and he had to slide it sideways across the floor. Right now he can get up and turn around before he pees all over himself and could not get to the bathroom before. Now he pees in a cooking pot. He asked to be excused while on call and stated he had to urinate again. Mobility got worse after first shot and then after second dose his mobility got even worse. Confirmed his compression fractures are an existing condition from 10 to 12 years ago. Reported he has gouty arthritis and it has been going on over 30 years and has constant pain for over 30 years but has always been able to walk and feed himself. The pain is not our fault. Reported that the malaise was better after 5 days, but his mobility went down. He has trouble walking. Sometimes knees would give out and he collapses. Caller states he now uses a walker, cane, and a wheelchair. If he has to go more than 10 feet, he has to have assistance. Treatment for Event: Norco 10-325: Takes up to 3 times per day. States the other day he had 45mg but was still in pain. This is discrepant but details were documented as provided. Caller states he is still in pain, but he can't get more without going to pain specialist. His doctor stopped atorvastatin and Ambien after adverse events with the covid vaccine. ER doctor wanted his primary doctor to review his medications since the ER doctor thought he was on too many medications. Caller tried to explain the reasons he was on the different medications. When talking about his gout: He has a picture after incident and half of his back is black, not blue, it is black. He was not in pain for about 2 years. Then he had the test to determine what was going on and that is when the pain got really bad. He has had bad pain for 35 years with gout in his hips especially, and in right shoulder. He cannot reach if something is in his right back pocket. Left shoulder sometimes. All the time with both hips. Also in right shoulder but if he does not use right shoulder then there is no problem. He was even thinking about getting a hip transplant due to the pain. His doctor did blood tests to find out that he had excess uric acid. He was on a medication called Uloric and it worked wonders but it is no longer made. Then it was found out that some people had bad reactions with it harming their heart and that was the end of Uloric. Then he started taking Allopurinol, states it does work but it works so little that it is almost worth not taking. When the pain is at 10 or 11 on pain scale, states what is the difference. He has an allergy in right sinus that caused to cough uncontrollably and his uncontrollable cough let to a perirectal abscess. He suggested to his doctor to stop his coughing and not worry about the abscess. He then went to ENT and stuck a scope down his nose and told him his sinus is inflamed and there is no infection, there is an allergy and cannot tell what the allergy is and will probably never find out what the allergy is, and prescribed him the Azelastine hydrochloride nasal spray. The pharmacist thinks the nasal spray is too much but states the pharmacist is not a physician and ENT doctor prescribed it. The two sprays a day works and he has not had uncontrollable coughing since then. Out of an abundance of caution he was sent to a lung doctor to get breathing test. Testing: He had two x-rays and CT scan of the torso. He does not know what x-rays showed. CT scan shows kidneys, liver, pancreas, and gall bladder are normal. Everything is normal except for the heart. It did not explain what was wrong with him. Clarifies he had the same CT scan at another facility a couple of years ago and he was told that calcium around the heart is normal. He knows that he has low blood pressure normally. Attempted to clarify if x-rays were off his chest, stated they stuck something under his back. States nothing was mentioned about his x-rays and if they looked at the x-rays they could see he has a broken back. No other vaccinations was done within four weeks prior to the first administration date of the suspect vaccine. Got sick the next day after the flu shot twice. Will never take again (unspecified flu shot). Patient reported about 4 years ago he let his doctor give him a pneumonia shot and his arm hurt like hell for a week, states he is never doing that again. When discussing Allopurinol, caller stated it was not working very well, but it had positive effect and he will continue taking it as long as his doctor will let him

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				States when taking it, the gout does not get better, it gets worse. Allopurinol, states it does work but it works so little that it is almost worth not taking. When the pain is at 10 or 11 on pain scale, states what is the difference. Caller states he was on a different medication before that worked better, but he could not continue to take it. Caller stated that microprint was difficult to read. He was able to give NDC number using magnifying glass. Product strength and count size dispensed: 300mg, 90 count. Expiration Date: Dec 2021. The sample of the product was available to be returned, if requested. Packaging was sealed and intact. Treatment for event was Norco 10-325: Takes up to 3 times per day. Caller states he is still in pain, but he can't get more without going to pain specialist. Confirms he has been in chronic pain for over 30 years due to gouty arthritis. Advised caller to retain sample of product in the event he receives Mailer within 3-10 business days for the return. Swap. Caller took 1st dose of Covid vaccine 16Apr2021 and collapsed the next day and then for the next 5 days he felt general malaise and weakness. On 07May2021 he took 2nd dose, 21 days after 1st dose, and again collapsed the next day. Stated that he needed to call 911 to help him get up and they took him to hospital where they did a scan of his torso and said he was fine. Doctor also told him that he did not have stroke, blood clot or aneurysm. Caller thinks it is neurological as he is still having trouble standing and walking and his left side feels paralyzed. He is wondering if these side effects were documented during clinical trials. Therapeutic measures were taken as a result of pain. The outcome of the events experienced collapsed day after vaccine first dose/collapsed day after vaccine second dose, legs wouldn't move, couldn't turn around, he had compression fractures l1 to l3 that were noticed on his x-ray, pain, half of back is black from incident, mobility got worse after first shot and then after second dose his mobility got even worse, felt like what i would call the flu was unknown, for the events his left side still feels paralyzed and can't walk, he collapsed and had a hard time getting up/felt bad, his left side still feels paralyzed and can't walk, bedridden the outcome was not resolved, for the event general malaise the outcome was resolving. Follow-up attempts are needed. Further information is expected."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1374313-1	"On the third day after the injection, I had a sudden return of sx previously experienced with medications cleared 3A4/5 and 2C19 including rigid paralysis for >11 minutes, multiple contractions, numbness/tingling in legs. I had not experienced any of these symptoms for > 3 months. Also fibromyalgia sx exacerbated to include burning pain in L arm (injection site) For 1 week, my arm turned red, like sunburn and burned. After week 1, there was no redness but there was more pain than burning and that continued for about 4 weeks. This has since abated. I currently have extreme fatigue, fibromyalgia pain in multiple sites, esp breast, chest and legs. Numbness and tingling continues in my feet and legs. I am also experiencing transient numbness in my lower face, periods of severe ""brain fog"" (this seems to be improving) and GERD sx relieved by antacids."
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	1381174-1	felt like she could not move it and then she felt like paralyzed for less than three second; right arm little bit hurting/pain in her hand like she couldn't grab something; pain in her hand like she couldn't grab something, and it's twitching; felt short breath/felt like she need to grab air to breathe, she can't breathe/it was hard to breathe; chest kept hurting/short pain in chest; felt like she could not move it; body felt super tired; Chills; little fever; This is a spontaneous report from a contactable consumer (patient). A 36-year-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE), via an unspecified route of administration on 18May2021 (at the age of 36 years old; Lot Number: EW0182, unknown expiration) as 1st dose, single for COVID-19 immunisation. Medical history and concomitant medications reported as none. The patient reported that she received her first shot on 18May2021. She came home with her right arm little bit hurting, felt pain in her hand like she couldn't grab something, and it's twitching. As soon as she received the vaccine, she felt short breath, she called the place where she received the vaccine, she was there for 20 minutes. In her aid, she walked out as she needed air and her chest kept hurting and hurting on 18May2021. Every time she breathes, she felt like she need to grab air to breathe, she can't breathe/it was hard to breathe. Since 18May2021 to today, she felt short pain in chest and right hand and in the morning it was like twitch, she felt like she could not move it and then she felt like paralyzed for less than three second and then the shortening went to exactly to her chest and felt like she was losing air. She mentioned she need air to breathe and her body felt super tired on 18May2021. She is a very active person; she feels so tired. She also experienced chills, little fever, and her chest was hurting more on 18May2021. Due date of second shot is on 08Jun2021. The patient did not receive treatment for the events. The patient had not yet recovered from the events. No follow-up attempts are possible. No further information is expected.

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
				Anaphylactic reaction; left cheek started swelling; looking woozy; throat and tongue started swelling; throat and tongue started swelling; having a hard time breathing; from the neck down, she had a temporary paralysis and she could not move./she could not feel anything from the waist down.; whole body was shaking; numbness and tingling all over.; numbness and tingling all over.; extreme pain in her back; When they would touch her hips, toes, spine, ankles, and knees she would complain of extreme pain, like joint pain.; When they would touch her hips, toes, spine, ankles, and knees she would complain of extreme pain, like joint pain.; When they would touch her hips, toes, spine, ankles, and knees she would complain of extreme pain, like joint pain.; very weak; chest was tight.; cannot walk without assistance; itching all over and in her throat; itching all over and in her throat; This is a spontaneous report from a contactable consumer (parent). A 13-years-old female patient received first dose of bnt162b2 (Pfizer-BioNTech COVID-19 Vaccine, 0.3 ML), at the age of 13-years-old , via an unspecified route of administration, administered in Arm Right on 21May2021 11:55 (Lot Number: EW0179; Expiration Date: Aug2021) as 0.3 ML SINGLE for covid-19 immunisation. Medical history included ongoing asthma, Diagnosed as a toddler. Had RSV as a baby. ongoing seasonal allergy, To like grass and pollen. ongoing constipation, Diagnosed about a year ago. allergic to shellfish because she gets sick. She said that it happened when her daughter was 4-5 years old. Whenever she has contrast for a MRI she throws it up. The patient's concomitant medications were not reported. The patient previously took Azithromycin and Amoxicillin and were allergic. The patient experienced anaphylactic reaction on 21May2021 with outcome of recovering, left cheek started swelling on 21May2021 with outcome of unknown, looking woozy on 21May2021 with outcome of unknown, throat and tongue started swelling on 21May2021 with outcome of unknown, having a hard time breathing on 21May2021 with outcome of unknown, from the neck down, she had a temporary paralysis and she could not move./she could not feel anything from the waist down on 21May2021 with outcome of unknown, whole body was shaking on 21May2021 with outcome of unknown, numbness and tingling all over on 21May2021 with outcome of unknown, extreme pain in her back on 21May2021 with outcome of unknown , when they would touch her hips, toes, spine, ankles, and knees she would complain of extreme pain, like joint pain on 21May2021 with outcome of unknown, very weak on 21May2021 with outcome of unknown , chest was tight on 21May2021 with outcome of unknown , cannot walk without assistance on 21May2021 with outcome of unknown , itching all over and in her throat on 21May2021 with outcome of unknown. Patient was in hospital from 21May2021 to 23May2021. AE(s) require a visit to Emergency Room. No AE(s) following prior vaccinations. No Family Medical History Relevant to AE(s). No Relevant Tests. Clinical courses as follows: It was reported they wanted her to wait for 30 minutes after her daughter received the vaccine. She said that after about 20 minutes, her daughter said that it felt like someone punched her in the face and her left cheek started swelling. She said that her husband said that their daughter started looking woozy and her eyes were closing. The staff laid her down on a mat on the floor. Then her daughter's throat and tongue started swelling and she was having a hard time breathing. Then her daughter complained that from the neck down, she had a temporary paralysis and she could not move. Her daughter asked if she asked if she was going to die. The nurse gave her an EpiPen and called a ambulance. Her whole body was shaking, probably from the EpiPen, like almost convulsing. She said that they got her onto the ambulance and her daughter was having trouble breathing. They gave her another dose of an EpiPen in the ambulance. They also gave her daughter a steroid and some Benadryl through an IV. Her daughter was rushed to the ER, having a hard time breathing and complaining she could not feel anything from the waist down. Caller said that her daughter was also complaining of numbness and tingling all over. She also started to complain about extreme pain in her back. When they would touch her hips, toes, spine, ankles, and knees she would complain of extreme pain, like joint pain. Her daughter had numbness from waist down. If they touched her foot she could not feel it. Her daughter was admitted to the hospital and got released yesterday 23May2021. She is still experiencing pain and numbness. The numbness is from the waist down and the tingling was up and down her arms. She said that her daughter was very weak and could not feed herself. They had to give her several albuterol breathing treatments in the hospital because her chest was tight. She was not wheezing, but they could tell it was constricted and treated her with Albuterol. She said that her daughter still cannot walk without assistance this morning. She was treated with Motrin, Benadryl and Atarax, and Prednisone in the hospital. The caller said that her daughter also complained about itching all over and in her throat. The caller said that her daughter did not have a rash though.
PARALYSIS	**COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)**	**PFIZER\BIONTECH**	**1384603-1**	

Symptoms	Vaccine	Vaccine Manufacturer	VAERS ID	Adverse Event Description
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1387759-1	Paralyzed; This is a spontaneous report from a contactable consumer (patient). A 47-years-old female patient received BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), dose 2 via an unspecified route of administration in the right high shoulder (right deltoid) on 27Mar2021 at 10:03 at the age of 47 years old (Batch/Lot Number: ERZ613 OR ER2613) as 2ND DOSE, SINGLE DOSE for COVID-19 immunisation. The patient's medical history and concomitant medications were not reported. The patient previously received the first dose of BNT162B2 (PFIZER-BIONTECH COVID-19 VACCINE, Solution for injection), at the age of 47 years old, batch number: EN6205 on 06Mar2021 at 10:11AM or 09:11AM given in her right high shoulder as an injection for COVID-19 immunisation. The patient was calling about the COVID 19 vaccination that she received. She stated that with the first dose, she did not feel anything, and she is calling to report on the second dose. She has spoken with Pfizer before, but she had questions and she was told how to go about it. She stated she received the vaccine and the symptom started exactly 8 hours after. She got the vaccine at 10:05AM or 10:03AM and it was about 9:03PM when the symptom started. She stated she got the vaccine probably at 10:03AM on 27Mar2021 and she was paralyzed. She stated picture someone laying there, it went into the right arm and then went into left and paralyzed there then it came down from the top to across her chest and shot straight down and missed the breast area and went into the ribs then it paralyzed her left leg and shot right across and paralyzed her whole right thigh and all she knows was that she could not move. She was paralyzed, and it shot into her and came across and then straight down. She stated it just did it at the moment exactly 8 hours and it started at around 9:08pm. She clarified, well it started actually 11 hours after the vaccine and not 8 hours. She wanted to say it lasted for 2 minutes where she could not move for a while and it was strange. The exact time was maybe 2 minutes after it started was when it stopped. She clarified and stated, well actually it lasted a couple of hours and stopped at 11:08PM. She stated she is not sure if she has recovered completely from this. The event outcome was unknown.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1394241-1	"Systemic: Weakness-Severe, Additional Details: Patient reports neurological induced paralysis from date of injection until and through current date. Has consulted with dr who thinks vaccine is likely cause to to current medical history. Reports inability to walk and ""needing to be carried""."
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1396012-1	4/18/21 pfizer shot in left arm. 8 hours later nausea, heart race, dizzy. go to sleep. wake normal. no more symptoms for 10 days. Day 10, arm pain returns at the injection site. Day 20 arm pain has spread in size and pain, i cancel the second shot. elbow and wrist become inflamed, tight, pain restricted. become sick for 5 days, headaches exhausted. prescribed 40mg prednisone for 5 days. first day it took the swelling down, then swelling came right back 2nd day. pinky and ring finder go numb, paresthesia/paresis. nerve pain in forearm to hand. ulnar nerve at first then all fingers. arm injection pain throbbing. rash shows up on stomach (now at 3.5 to 4 weeks since shot) cannot grasp a cup, weakness and numbness in hand. At this time its confined to the injection arm. Over the next week nerve pain and numbness spread to left side of face, both arms, hands, fingers, legs, feet, toes. physical therapist confirms elbow lymph node on injection arm swelling, hands turning red extreme pain and swelling, loss of feeling. nerve pain in the entire body, nerve and muscle spasms in all parts of body. Fingers and does burning at the tips feel like a string is cutting off circulation. Visibly red. Around the 4 week mark, fully unable to lift legs or arms, left half of face numb, zap/jots seizure like hitting my body. go to ER seemingly becoming paralyzed, within hours strength of legs returns but with numb feet/toes. no diagnosis other than peripheral neuropathy. neck begins to swell, hurts to swallow, throat closing. trying to sleep, throat closed, wake up, completely unable to open throat cant breath. throat episodes lasted about 5 days and resolved. complete insomnia and 24/7tinitus. any attempt to sleep or become tired results in heavy body zaps like hooked to a tenz unit. can't use hands or type, arm to hand seized nerves, feel like shrunken or contracted nerves. PCP prescribes benzos to sleep. this allows some sleep with waking every hour. over the following weeks symptoms seem to improve slowly but still debilitating. today at week 8+ since one shot only, I am able to type again, but with nerve pain and numbness in all fingers. loud tinnitus, numbness in all limbs, episodes of extreme debilitating muscle fatigue come in waves, almost unable to stand agonizing muscle fatigue and nerve zapping. These full body muscle episodes only seem to come in waves of 24 hours, then seem to feel recovering, only to relapse again. injection site pain seems to have wrapped around to the bicep 24/7 pain. unsure if this is nerve or muscle. 3 ER visits, PCP visits, blood tests, no help. I cannot work, constant nerve firing muscle spasms and nerve pain and episodes of weakness so bad you cannot sit without agonizing pain. I am disabled from this and still searching for help
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1400228-1	Flacid arm several hours after shot. Had not shown much improvement 60 days after original vaccine shot.
PARALYSIS	COVID19 (COVID19 (PFIZER-BIONTECH)) (1200)	PFIZER\BIONTECH	1407825-1	After Dose 1 (given 03/14/2021): EXTREME fatigue/exhaustion; sleeping 5-6 hours during the day Daily migraine-like headaches Brain Fog Memory Loss After Dose 2 (given 4/11/2021): EXTREME fatigue/exhaustion; sleeping 3-4 hours during the day Frequent headaches Brain Fog Memory Loss Joint Pain Extreme Lower Back Pain Numbness and tingling in arms and hands One bout of paralyzation for 6-7 hours (seen in ER)

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 5:03:40 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 5:03:40 PM

Query Criteria:

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