PART IV
A. LET'S LOOK AT VAERS

INFORMED CONSENT MATTERS

Cases from VAERS - Vaccine Adverse Event Reporting System

IT IS EVERYONE'S RIGHT TO REVIEW THE FIRST-HAND TESTIMONIES OF THE COVID VACCINE ADVERSE REACTIONS WITHOUT CENSORSHIP BY THE NEWS MEDIA (OR SOCIAL MEDIA).

WITH ACCESS TO THIS DATA, YOU CAN PERFORM A RISK/BENEFIT ANALYSIS FOR YOURSELF.

PART IV

UPDATED MARCH 14, 2021

LET'S LOOK AT

VAERS

THE VACCINE ADVERSE EVENT REPORTING SYSTEM

Did you know that our government records reported vaccine deaths & injuries through VAERS?

VAERS COVID REPORTS

38,444 Reports Through March 12, 2021*

jump to browse reports

1739	3976	6716
DEATHS	HOSPITALIZATIONS	URGENT CARE

5838	324	398
OFFICE VISITS	ANAPHYLAXIS	BELL'S PALSY

Total Heart Attacks	Total Miscarriages	Total Severe Allergic Reaction	Total Thrombocytopenia
386	69	2248	118

* VAERS HHS releases COVID Data weekly, but they release LAST WEEK'S data. So an update will always lag a week behind. When launched, OpenVAERS used the Download date. We have switched to the "data through" date provided by VAERS.

The Vaccine Adverse Event Reporting System (VAERS) Results

Request Form | Results | Map | Chart | Report | About

Dataset Documentation | Other Data Access | Help for Results | Printing Tips | Help with Exports

Save | Export | Reset

Quick Options | More Options

Top | Notes | Citation | Query Criteria

Messages:

- **VAERS data in CDC WONDER are updated every Friday. Hence, results for the same query can change from week to week.**
- **These results are for 38,444 total events.**

Event Category	Events Reported	Percent (of 38,444)
Death	1,739	4.52%
Life Threatening	1,205	3.13%
Permanent Disability	734	1.91%
Congenital Anomaly / Birth Defect *	48	0.12%
Hospitalized	3,976	10.34%
Existing Hospitalization Prolonged	43	0.11%
Emergency Room / Office Visit **	27	0.07%
Emergency Room *	6,689	17.40%
Office Visit *	5,837	15.18%
None of the above	23,754	61.79%
Total	**44,052**	**114.59%**

524 of these deaths were within 24 hours of shot

VAERS is a passive surveillance system in the U.S. which captures adverse events following vaccination.
'Events' are reported by hospital staff, nursing home staff, doctors, etc.
This data is searchable at: https://wonder.cdc.gov/vaers.html

Each VAERS case is assigned a unique ID number and has a description of the incident (death, disability, etc.)

Details for VAERS ID: 0944595-1

Event Information			
Patient Age	56.00	**Sex**	Male
State / Territory	Florida	**Date Report Completed**	2021-01-14
Date Vaccinated	2021-01-12	**Date Report Received**	2021-01-14
Date of Onset	2021-01-12	**Date Died**	2021-01-14
Days to onset	0		
Vaccine Administered By	Senior Living *	**Vaccine Purchased By**	Not Applicable *
Mfr/Imm Project Number	NONE	**Report Form Version**	2
Recovered	No	**Serious**	Yes

* VAERS 2.0 Report Form Only
** VAERS-1 Report Form Only
"Not Applicable" will appear when information is not available on this report form version.

Event Categories	
Death	Yes
Life Threatening	No
Permanent Disability	No
Congenital Anomaly / Birth Defect *	No
Hospitalized	Yes
Days in Hospital	2
Existing Hospitalization Prolonged	No
Emergency Room / Office Visit **	N/A
Emergency Room *	Yes
Office Visit *	No

* VAERS 2.0 Report Form Only
** VAERS-1 Report Form Only
"N/A" will appear when information is not available on this report form version.

Vaccine Type	Vaccine	Manufacturer	Lot	Dose	Route	Site
COVID19 VACCINE	COVID19 (COVID19 (PFIZER-BIONTECH))	PFIZER\BIONTECH	EL0140	1	IM	AR
COVID19 VACCINE	COVID19 (COVID19 (PFIZER-BIONTECH))	PFIZER\BIONTECH	EL3302	2	IM	AR

Adverse Event Description

Cardiac arrest within 1 hour Patient had the second vaccine approximately 2 pm on Tuesday Jan 12th He works at the extended care community and was in good health that morning with no complaints. He waited 10-15 minutes at the vaccine admin site and then told them he felt fine and was ready to get back to work. He then was found unresponsive at 3 pm within an hour of the 2nd vaccine. EMS called immediately worked on him 30 minutes in field then 30 minutes at ER was able to put him on life support yet deemed Brain dead 1-14-21 and pronounced dead an hour or so later

Listed on the following pages are only 70 of the 1,739 deaths recorded in VAERs.

Anyone can view the database via the following steps:

https://wonder.cdc.gov/vaers.html

1. Accept the disclaimer at bottom of page

2. Click on VAERS data search

3. In section 1, click on group results by "VAERS ID" , and by "vaccine type" and by "event category"

4. Under optional measures (still in section 1) tick off "adverse event description"

5. Scroll down to section 3 and under Vaccine Products select "Covid19 vaccine" Make sure it is the only option selected. (You will have to deselect "All vaccine products" from the top of the list.)

6. Scroll down to section 5 and under event category select "death". Make sure it is the only option selected. (Don't forget to deselect "all events" from the top of the list).

Scroll to the very bottom and click send.

VAERS ID # 938118-1 AGE 51. FEMALE Vaccinated 1/5/2021. Died. 1/10/2021. Pfizer vaccine. On 1/8/2021 17:30 patient taken to ER, cerebellar hemorrhage, stroke, aneurysm.

VAERS ID # 946293-1 AGE 51. MALE Vaccinated 1/7/2021. Moderna . Became increasingly hypoxic around 1800hours on 1/7/2021. Transported to ER for acute on chronic hypoxia respiratory failure. Expired on 1/12/2021@2325 at med center.

VAERS ID # 918518-1 AGE 50. FEMALE Vaccinated 12/31/2020. Died 12/31/2020. Moderna

VAERS ID # 930910-1 AGE 52. FEMALE Vaccinated 1/8/2021. Died 1/8/2021. Patient received COVID vaccination 12:15pm. Patient was monitored for the appropriate amount of time by nursing staff. Patient passed away at 2:15pm. Moderna

VAERS ID # 933739-1 AGE 54. FEMALE. Vaccinated 1/8/2021. Died 1/10/2021. 2 days later. Staff member checked on her at 3am and patient stated that she felt like she couldn't breathe. 911 was called and taken to the hospital. While in the ambulance, patient coded. Two EEGs were given to determine that patient had no brain activity. Pfizer

VAERS ID # 923219-1 AGE 41. FEMALE Vaccinated 12/30/2021. Died. 1/1/2021. Pfizer vaccine. The patient did not experience any adverse event at the moment of inoculation with COVID-19 vaccine or the following days. On January 1, 2021, at lunch time, two days after receiving the vaccine, the patient was found unresponsive in her bed by her partner.

VAERS ID # 936805-1 AGE 25. MALE Vaccinated 12/22/2020. Found unresponsive and expired at home on 1/11. Moderna

VAERS ID # 943397-1 AGE 28. MALE Vaccinated 12/23/2020. Died 1/14/2021. Patient was found unresponsive at work in the hospital. Patient pupils were fixed and dilated. Pfizer

VAERS ID # 939050-1 AGE 32. FEMALE Vaccinated 12/28/2020. Died on 1/4/21 at 7:20am. Moderna

VAERS ID # 921667-1 AGE 39. FEMALE Vaccinated 12/29/2020. It was reported that the staff member deceased somewhere between 1/3/2021 and 1/4/2021. Pfizer

VAERS ID # 933578-1 AGE 43. MALE Vaccinated 1/8/2021. Died 1/9/2021. Moderna

VAERS ID # 937527-1 AGE 44. FEMALE Vaccinated 12/23/2020. Died on 1/4/2021. Pfizer

VAERS ID # 929764-1AGE 45. MALE Vaccinated 12/28/2020. Died 12/29/2020. The patient was found deceased at home about 24 hours after immunization. Moderna

VAERS ID # 934968-1AGE 54. MALE Vaccinated 1/4/2021. Died 1/6/2021. Pfizer. The patient received the vaccine on 04Jan2021, after which he started not feeling well. He went right home and went to bed. He woke up and ate a bit but not much and then was kind of pale. The patient then started to vomit, which continued throughout the night. He was having trouble in breathing. Emergency services were called, and they took his vitals and said that everything was okay, but he was very agitated; reported as not like this prior to the vaccine. The patient was taken to urgent care where they gave him an unspecified steroid shot and unspecified medication for vomiting. The patient continued to vomit throughout the day and then he was very agitated again and would fall asleep for may be 15-20 minutes. When the patient woke up, he was very restless (reported as: his body was just amped up and could not calm down). The patient calmed down just a little bit in the evening. When the patient was awoken at 6:00 AM in the morning, he was still agitated. The patient stated that he couldn't breathe, and his mind was racing. The patient's other brother went to him and he was not responsive, passed on 06Jan2021 around 10:15 AM. It was reported that none of the symptoms occurred until the patient received the vaccine.

VAERS ID # 942106-1 AGE 54. MALE Vaccinated 1/8/2021. Died 1/9/2021. Pfizer vaccine. On scene, the patient had a witnessed arrest with EMS starting CPR. He was given 3 rounds of epi without ROSC. Patient's wife, had noted patient had received covid vaccine the prior day.

VAERS ID: 924456-1, AGE 85 vaccination was administered at approx. 10:00 AM and the patient continued throughout day without any complaints or signs of adverse reaction. When the nursing staff went to the room to check on the resident patient was found unresponsive, no chest rises, noted regurgitated small amount of food to mouth left side.

VAERS ID # 996156-1 AGE 58. FEMALE Vaccinated on 02/02/2021. Moderna. Observed by vaccination team for a period of time. She reported shoulder pain radiating into shoulder blade in arm vaccine was received. Vaccination team offered ice pack to her and released back to work. 10pm that evening, she sent text to coworker that her pain was ""off the charts"" and that she had pain covering her whole left side of her body. She did not come to work in the morning and did not contact work. Well being check was performed at approximately 9am on 2/2/2021 and she was found dead in her home.

VAERS ID # 935511-1 AGE 56. FEMALE Vaccinated 1/8/2021. Died 1/9/2021. Moderna. Patient received the 1st dose of Moderna and was found deceased in her home the next day.

VAERS ID # 941811-1 AGE 56. FEMALE Vaccinated 1/4/2021. Died 1/11/2021. Moderna. Resident began having fever on 1/11/21. Resident sent to nearest ER for evaluation. Later in the evening the staff AT Medical Center called to inform staff that resident had expired @ 2230 as a result of Respiratory Failure and Sepsis.

VAERS ID # 944595-1 AGE 56. MALE Vaccinated 1/12/2021. Died 1/14/2021. Pfizer. Cardiac arrest within 1 hour Patient had the second vaccine approximately 2 pm on Tuesday Jan 12th He works at the extended care community and was in good health that morning with no complaints. He waited 10-15 minutes at the vaccine admin site and then told them he felt fine and was ready to get back to work. He then was found unresponsive at 3 pm within an hour of the 2nd vaccine. EMS called immediately worked on him 30 minutes in field then 30 minutes at ER was able to put him on life support yet deemed Brain dead on 1-14-21 and pronounced dead an hour or so later.

VAERS ID # 921768-1 AGE 58. FEMALE Vaccinated 1/4/2021. Died 1/4/2021. Pfizer.

VAERS ID # 930154-1 AGE 60. MALE Vaccinated 1/5/2021. Died 1/8/2021. Moderna.

VAERS ID # 933090-1 AGE 60. MALE Vaccinated 1/5/2021. Died. 1/9/2021. Pfizer.

VAERS ID # 941743-1 AGE 60. FEMALE Vaccinated 1/12/2021. Died 1/13/2021. Moderna. Found deceased at 3am

VAERS ID # 932898-1 AGE 61. MALE Vaccinated 12/17/2020. Died. 12/30/2020. Pfizer vaccine. The patient had an apparent cardiac arrest on 12/23/20 and was admitted to the ICU. He was taken off of life support on 12/30/20.

VAERS ID # 942085-1 AGE 62. FEMALE Vaccinated 1/2/2021. Died 1/8/2021. Pfizer. On 1/8/21 at 0615 resident was shaking. Reported all over pain. At 0850 she was not responsive.

VAERS ID # 940955-1 AGE 66. FEMALE. Vaccinated 1/11/2021. Died 1/11/2021. Pfizer. Cardiac Arrest. Patient was found pulseless and breathless 20 minutes following the vaccine administration. Received the second dose of BNT162B2. Took the first dose on 21Dec2020. MD found no signs of anaphylaxis.

VAERS ID: 926600-1, 65yo Patient did not report any signs or symptoms of adverse reaction to vaccine. Patient reported not feeling well and passed away that day.

VAERS ID: 925154-1, 84yo DEATH within 1 day, no current illness.

VAERS ID: 926797-1, 93yo had a vaccination on 12/31/2020 late morning passed away early morning 01/01/2020.

VAERS ID: 927189-1, 74yo Patient was vaccinated at 11am and was found at the facility in his room deceased at approximately 3:00pm. Nurse did not have cause of death

VAERS ID: 927260-1, 87yo No adverse effects noted after vaccination. Patient found unresponsive at 16:45 on 1/6/21. Abnormal breathing patterns, eyes partially closed SPO2 was 41%, pulseless with no cardiac sounds upon auscultation. CPR, pulse regained, patient breathing. Patient sent to ER had multiple cardiac arrest and severe bradycardia and passed.

VAERS ID: 924664-1, 92yo No current illness. At 1855, I was alerted by caregiver, resident was not responding. Per caregiver, she was doing rounds and found resident in bed, unresponsive, mouth open, observed gurgling noises. Primary caregiver observed resident at baseline and ambulating after dinner at approximately, 1800 less than an hour prior to incident. Resident received the first dose of vaccine on 1/2. Expired at 0615 per Castle RN.

.

VAERS ID: 923993-1, 62yo Patient was vaccinated Dec 30, 2020. Prime dose of Moderna vaccine. Observed for full 15 minutes post-injection. No complaints when asked during observation. Released. Subsequently, vaccine clinic staff learned from the patient's supervisor that patient had expired on Jan 2, 2021at his home.

VAERS ID: 909095-1, 66yo on 12/24/2020 the resident was sleepy and stayed in bed most of the shift. He stated he was doing okay but requested pain medication for his legs at 250PM. At 255AM on 12/25/2020 the resident was observed in bed lying still, pale, eyes half open and foam coming from mouth and unresponsive. He was not breathing and with no pulse.

VAERS ID: 924464-1, 61yo coughing up blood, significant hemoptysis -- > cardiac arrest. started day after vaccine

VAERS ID: 921768-1, 58yo Vaccine received at about 0900 on 01/04/2021 at her place of work, Medical Center, where she was employed as a housekeeper. About one hour after receiving the vaccine she experienced a hot flash, nausea, and feeling like she was going to pass out after she had bent down. Later at about 1500 hours she appeared tired and lethargic, then a short time later, at about 1600 hours, upon arrival to a friends home she complained of feeling hot and having difficulty breathing. She collapsed, when medics arrived, she was still breathing slowly then went into cardiac arrest and was unable to be revived.

VAERS ID: 910363-1, 84yo Patient had mild hypotension, decreased oral intake, somnolence starting 3 days after vaccination and death 5 days after administration.

VAERS ID: 913143-1, 84yo Vaccine administered with no immediate adverse reaction at 11:29am. Vaccine screening questions were completed and resident was not feeling sick and temperature was 98F. At approximately 1:30pm the resident passed away.

VAERS ID: 913733-1, 85yo My grandmother died a few hours after receiving the moderna covid vaccine booster 1. The treating hospital did not acknowledge this and I wanted to be sure a report was made.

VAERS ID: 914604-1, 74yo Spouse awoke 12/20 and found spouse dead. Client was not transferred to hospital.

VAERS ID: 914690-1, 83yo Within 24 hours of receiving the vaccine, fever and respiratory distress, and anxiety developed requiring oxygen, morphine and ativan. My Mom passed away on the evening of 12/26/2020.

VAERS ID: 914805-1, 63yo RESIDENT CODED AND EXPIRED

VAERS ID: 914895-1, 78yo Injection given on 12/28/20 - no adverse events and no issues yesterday; Death today, 12/30/20,

VAERS ID: 914917-1, 63yo Death by massive heart attack. Pfizer-BioNTech COVID-19 Vaccine EUA

VAERS ID: 914961-1, 88yo pt passed away with an hour to hour and 1/2 of receiving vaccine.

VAERS ID: 914994-1, 90yo pt was a nursing home pt. pt received first dose of covid vaccine. pt was monitored for 15 minutes after getting shot. staff reported that pt was 15 days post covid. Pt passed away with in 90 minutes of getting vaccine.

VAERS ID: 915562-1, 88yo pt received vaccine at covid clinic on 12/30 at approximately 3:30, pt vomited 4 minutes after receiving shot--dark brown vomit, Per staff report pt became short of breath between 6 and 7 pm that night. Pt passed away at approximately 10pm.

VAERS ID: 915682-1, 85yo Resident received vaccine per pharmacy at the facility at 5 pm. Approximately 6:45 resident found unresponsive and EMS contacted. Upon EMS arrival at facility, resident went into cardiac arrest, code initiated by EMS and transported to hospital. Resident expired at hospital at approximately 8 pm

VAERS ID: 915920-1, 96yo Resident was living in an assisted living facility. She fell on 11/24/2020 and was admitted to this facility for rehab. Received vaccine 12/28/2020 in am and expired that afternoon.

VAERS ID: 918065-1, 64yo Vaccine 12/30/2020. 1/1/2020: Pronounced deceased 1/1/2020

VAERS ID: 918388-1, 65yo Vaccine 12/30/2020. 1/1/2020: Resident found unresponsive without pulse, respirations at 04:30 CPR performed, expired at 04:52 by Rescue, No acute illness at time of vaccination.

VAERS ID: 918418-1, 65yo Vaccine 12/30/2020. 1/1/2020: Resident became SOB, congested and hypoxic requiring oxygen, respiratory treatments and suctioning. Stabilized after treatment and for the next 72 hours with oxygen saturations in the 90s. On 1/3/2021 was found without pulse and respirations.

VAERS ID: 918487-1, 94yo Two days post vaccine patient went into cardiac arrest and passed away.

VAERS ID: 915880-1, 99yo Patient died within 12 hours of receiving the vaccine.

VAERS ID: 918518-1, 50yo syncopal episode - arrested - CPR - death

VAERS ID: 919108-1, 100yo Fever, Malaise, passed the day after vaccine.

VAERS ID: 919537-1, 96yo Resident exhibited no adverse events during 30 minute monitoring following vaccine administration. Resident found without pulse at 1900.

VAERS ID: 920326-1, 89yo Redness and warmth with edema to right side of neck and under chin. Resident expired on 1.1.21.

VAERS ID: 920545-1, 93yo "The resident received is vaccine around 11:00 am and tolerated it without any difficulty or immediate adverse effects. He was at therapy from 12:36 pm until 1:22 pm when he stated he was too tired and could not do anymore. The therapist took him back to his room at that time and he got into bed himself but stated his legs felt heavy. At 1:50 pm the CNA answered his call light and found he had taken himself to the bathroom. She stated that when he went to get back into the bed it was ""abnormal"" how he was getting into it so she assisted him. At that time he quit breathing and she called a RN into the room immediately. He was found without a pulse, respirations, or blood pressure at 1:54 pm.

VAERS ID: 920815-1, 58yo Found deceased in her home, unknown cause, 6 days after vaccine.

VAERS ID: 920832-1, 104yo Vaccine 12/30/2020 Screening PCR done 12/31/2020 Symptoms 1/1/2021 COVID test result came back positive 1/2/2021 Deceased 1/4/2021

VAERS ID: 921175-1, 77yo Resident received Covid Vaccine, noted after 30 mins with labored breathing BP 161/77, HR 116, R 38, T 101.4,epipen administered, sent to ER, died

VAERS ID: 921481-1, 88yo Vaccine given on 12/29/20 by Pharmacy. On 1/1/21, resident became lethargic and sluggish and developed a rash on forearms. Resident expired on 1/4/2021

VAERS ID: 921547-1, 65yo RESIDENT RECIEVED VACCINE ON 1/2/20. DEATH ON 1/4/2021.

VAERS ID: 921572-1, 87yo Resident had body aches, a low O2 sat and had chills starting on 12/30/20. He had stated that they had slightly improved. On 1/2/21 during the NOC shift his O2 sat dropped again. He later went unresponsive and passed away.

VAERS ID: 921667-1, 39yo LTCF Pfizer Vaccine clinic conducted 12/29/2020 Vaccine lead received a call indicating that a staff member deceased somewhere between 1/3/2021 and 1/4/2021. Cause of death is unknown, and an autopsy is being performed.

VAERS ID: 921880-1, 96yo The resident was found deceased a little less than 12 hours following COVID vaccination.

If you are injured by the COVID vaccine, it will be nearly impossible to receive compensation for lost work days and medical bills

CNBC
Search quotes, news & videos
MARKETS BUSINESS INVESTING TECH POLITICS CNBC TV WATCHLIST PRO
MAKE IT | SELECT

You can't sue Pfizer or Moderna if you have severe Covid vaccine side effects. The government likely won't compensate you for damages either

PUBLISHED THU, DEC 17 2020·8:36 AM EST | UPDATED WED, DEC 23 2020·12:32 AM EST

MacKenzie Sigalos
@KENZIESIGALOS

SHARE

If you experience severe side effects after getting a Covid vaccine, lawyers tell CNBC there is basically no one to blame in a U.S. court of law.

The federal government has granted companies like Pfizer and Moderna immunity from liability if something unintentionally goes wrong with their vaccines.

"It is very rare for a blanket immunity law to be passed," said Rogge Dunn, a Dallas labor and employment attorney. "Pharmaceutical companies typically aren't offered much liability protection under the law."

You also can't sue the Food and Drug Administration for authorizing a vaccine for emergency use, nor can you hold your employer accountable if they mandate inoculation as a condition of employment.

Congress created a fund specifically to help cover lost wages and out-of-pocket medical expenses for people who have been irreparably harmed by a "covered countermeasure," such as a vaccine. But it is difficult to use and rarely pays. Attorneys say it has compensated less than 6% of the claims filed in the last decade.

The FDA has had 10 months to get a safety monitoring system in place, but is no where near implementing one.

02/19/21 • BIG PHARMA › NEWS

One-Third of Deaths Reported to CDC After COVID Vaccines Occurred Within 48 Hours of Vaccination

The numbers reflect the latest data available as of Feb. 12 from the CDC's Vaccine Adverse Event Reporting System website. Of the 929 reported deaths, about one-third occurred within 48 hours.

As is consistent with previous VAERS data reports, 192 of the reported deaths — or 21% — were cardiac-related. As The Defender reported earlier this month, Dr. J. Patrick Whelan, a pediatric rheumatologist, warned the U.S. Food and Drug Administration in December that mRNA vaccines like those developed by Pfizer and Moderna could cause heart attacks and other injuries in ways not assessed in safety trials.

Of the 929 deaths reported since Dec. 14, 2020, the average age of the deceased was 77.8 and the youngest was 23. Fifty-two percent of the reported deaths were among men, 45% were women and 3% are unknown. Fifty-eight percent of the deaths were reported in people who received the Pfizer vaccine, and 41% were related to the Moderna vaccine.

States with the highest reported number of deaths were: California (71); Florida (50); Ohio (38); New York (31); Kentucky (41); Michigan (31); and Texas (31).

According to the latest data, 3,126 "serious" adverse reactions have been reported. Adverse reaction reports from the latest CDC data also include:

- 34 miscarriages and pre-term births
- 917 anaphylactic reactions, 70% of which were reported after a Pfizer vaccine and 30% after the Moderna vaccine
- Bell's palsy (Pfizer 75%; Moderna: 25%)

So far, only Pfizer and Moderna vaccines — approved for emergency use, but not fully licensed — are being used in the U.S.

Meanwhile, the FDA has not yet implemented systems to monitor the safety of the experimental COVID vaccines. FDA officials told The New York Times they don't expect the systems to be up and running before the Biden administration reaches its goal of vaccinating 100 million Americans — nearly one third of the U.S. population.

The U.K. has their own tracking system (Yellow Cards)

BREAKING: UK gov't says over 240 people in Britain died shortly after receiving COVID jab

The report included in its ancillary material the information that a total of 8 miscarriages have been reported

LONDON, England, February 11, 2021 (LifeSiteNews) — A U.K. government vaccine safety review has revealed that 244 people, including 8 unborn babies, have died in Britain shortly after receiving one of two coronavirus vaccines. The government has said that it does not believe the inoculations are responsible.

An extensive report produced by the U.K. government today has detailed all the suspected side effects, including death, reported by medical staff or the people who received at least one Covid-19 vaccine between early December 2020 and January 31, 2021. The two vaccines currently in use in the U.K. are the Pfizer/BioNTech and the Oxford University/AstraZeneca. They were authorized by the country's Medicines and Healthcare products Regulatory Agency (MHRA).

The MHRA monitors the safety of the vaccines as they are used on the public, and to this end they introduced a "Yellow Card Scheme" to collect reports of any suspected side effects. 20,319 Yellow Card reports, including a total of 59,614 suspected adverse reactions, were made regarding the Pfizer vaccine. 11,748 were made regarding the AstraZeneca vaccine, which was rolled out only on January 4, 2021. 72 Yellow Card reports did not specify which vaccine was used.

According to the report:

> The MHRA has received 143 UK reports of suspected ADRs to the Pfizer/BioNTech vaccine in which the patient died shortly after vaccination, 90 reports for the Oxford University/AstraZeneca vaccine and 3 where the brand of vaccine was unspecified.

The report included in its ancillary material the information that 5 miscarriages were reported after the use of the Pfizer vaccine and 3 after the Oxford vaccine.

Suspected adverse reactions after receiving Pfizer/BioNTech vaccine

A total of **59,614 suspected side effects** were recorded after use of the Pfizer/BioNTech vaccine.

The link to the Pfizer vaccine analysis shows clearly that **1,437 blood disorders** have been reported after inoculation, 1,204 of them being lymphadenopathy (enlarged or swollen lymph nodes). **712 cardiac disorders** were reported, notably palpitations (303) and a speeding heart rate (230). **445 ear disorders** were reported, including vertigo (146), ear pain (122), and varying amounts of hearing loss (31).

However, there have been reports of more serious damage associated with the vaccine, notably anaphylaxis and Bell's Palsy. Now 130 "spontaneous adverse reactions associated with anaphylaxis or anaphylactoid reactions" in Britain have been reported for the Pfizer vaccine, and 30 have been reported for the Oxford vaccine. Bell's Palsy, which involves facial paralysis or weakening of the facial muscles, has been reported 99 times for the Pfizer vaccine and 15 times for the Oxford vaccine. The MHRA says that because this ailment "can also occur naturally, an association with the vaccine in not yet confirmed."

There were **224 injuries** reported, and **357 metabolic disorders**, of which 252 were decreased appetite. **8,129 muscle and tissue disorders** were mentioned, of which 2,690 were muscle pain and 1,704 were joint pain. **11,160 nervous disorders** were reported, including 5,506 headaches, 1,546 cases of dizziness, 45 seizures, 15 cases of paralysis, and 1 case of locked-in syndrome. There were **5 miscarriages** reported.

Of the **830 psychiatric disorders** reported, 162 were insomnia, 112 cases of being in a confused state, and 73 bouts of anxiety. Of the **120 renal & urinary disorders**, 35 were renal pain and 12 were urinary incontinence. There were **187 reproductive and breast disorders** mentioned, of which 31 were breast pain and 22 menorrhagia (heavy periods).

There were **2,397 respiratory disorders** reported, including 520 cases of difficulty in breathing, 483 sore throats, and 426 cases of coughing. **3,947 skin disorders** were reported, including at least 796 rashes and 776 cases of itchiness. The **676 vascular disorders** on record included 182 hot flushes.

There were **823 references to eye disorders**, including 145 reports of eye pain, 126 of blurred vision, and 5 of blindness. **6,605 gastrointestinal orders** were reported, among them 2,889 cases of nausea, 924 cases of diarrhea, 775 cases of vomiting, and 283 cases of upper abdominal pain. Of the **19,354 "general disorders"** reported, 4,007 were cases of fatigue, 3,665 were fevers,2,310 were chills, 773 referred to vaccination site pain, 491 were "influenza-like illnesses." There were **308 immune system disorders**, mostly hypersensitivity (146) and anaphylactic reactions (116).

Ironically, there were **1,186 reported infections**, including 308 cases (including "suspected cases") of Covid-19 itself, 231 cases of influenza, and 154 upper respiratory tract infections.

LET'S LOOK AT

THE TRIALS

The 95% effective claim by Pfizer is based on ONLY 170 trial participants.

no one in the trial was tested for the development of antibodies

Everly Report

January 31, 2021

What you need to know about the COVID Vaccine

EFFECTIVENESS:

Claim: 95% effective

What this claim is based on:

Although there were a total of 43,548 trial participants in Pfizer's phase 3 trial, their calculation of effectiveness is based on a total of 170 participants.

These 170 participants were the first to develop symptoms of COVID and test positive for SARS-CoV-2, within the two-month monitoring period starting 7 days after the administration of the second vaccine.

From the chart below, it states that of the first 170 participants to develop symptoms and test positive for SARS-CoV-2, 162 of them were placebo recipients and 8 were vaccine recipients.

This is where the "95% effective" calculation comes from.

Source: Pfizer Vaccine Insert.

However, the rest of the participants were not tested for infection, nor were they tested for the development of antibodies, which is the endpoint typically used to measure vaccine effectiveness.

The published phase 3 clinical trial states the following "Limitations and Remaining Questions":

LIMITATIONS AND REMAINING QUESTIONS

Further study is required to understand the following:

- Safety and efficacy beyond 2 months and in groups not included in this trial (e.g., children, pregnant women, and immunocompromised persons).
- Whether the vaccine protects against asymptomatic infection and transmission to unvaccinated persons.
- How to deal with those who miss the second vaccine dose.

Note: Remaining questions include "whether the vaccine protects against asymptomatic infection and transmission".

Table 6: Vaccine Efficacy – First COVID-19 Occurrence From 7 Days After Dose 2, by Age Subgroup – Participants Without Evidence of Infection and Participants With or Without Evidence of Infection Prior to 7 Days After Dose 2 – Evaluable Efficacy (7 Days) Population

First COVID-19 occurrence from 7 days after Dose 2 in participants without evidence of prior SARS-CoV-2 infection*			
Subgroup	Pfizer-BioNTech COVID-19 Vaccine Na=18,198 Cases n1b Surveillance Timec (n2d)	Placebo Na=18,325 Cases n1b Surveillance Timec (n2d)	Vaccine Efficacy % (95% CI)
All subjects[e]	8 2.214 (17,411)	162 2.222 (17,511)	95.0 (90.3, 97.6)[f]
16 to 64 years	7 1.706 (13,549)	143 1.710 (13,618)	95.1 (89.6, 98.1)[g]
65 years and older	1 0.508 (3848)	19 0.511 (3880)	94.7 (66.7, 99.9)[g]
First COVID-19 occurrence from 7 days after Dose 2 in participants with or without evidence of prior SARS-CoV-2 infection			
Subgroup	Pfizer-BioNTech COVID-19 Vaccine Na=19,965 Cases n1b Surveillance Timec (n2d)	Placebo Na=20,172 Cases n1b Surveillance Timec (n2d)	Vaccine Efficacy % (95% CI)
All subjects[e]	9 2.332 (18,559)	169 2.345 (18,708)	94.6 (89.9, 97.3)[f]
16 to 64 years	8 1.802 (14,501)	150 1.814 (14,627)	94.6 (89.1, 97.7)[g]
65 years and older	1 0.530 (4044)	19 0.532 (4067)	94.7 (66.8, 99.9)[g]

Note: Confirmed cases were determined by Reverse Transcription-Polymerase Chain Reaction (RT-PCR) and at least 1 symptom consistent with COVID-19 (symptoms included: fever; new or increased cough; new or increased shortness of breath; chills; new or increased muscle pain; new loss of taste or smell; sore throat; diarrhea; vomiting).

* Participants who had no evidence of past SARS-CoV-2 infection (i.e., N-binding antibody [serum] negative at Visit 1 and SARS-CoV-2 not detected by NAAT [nasal swab] at Visits 1 and 2), and had negative NAAT (nasal swab) at any unscheduled visit prior to 7 days after Dose 2 were included in the analysis.

a. N = number of participants in the specified group.
b. n1 = Number of participants meeting the endpoint definition.
c. Total surveillance time in 1000 person-years for the given endpoint across all participants within each group at risk for the endpoint. Time period for COVID-19 case accrual is from 7 days after Dose 2 to the end of the surveillance period.
d. n2 = Number of participants at risk for the endpoint.

Up next...testing brand new mRNA technology on children as young as 6 months old...and pregnant women.

Testers are using the meningitis vax as their 'placebo' so they can eliminate know vaccine side effects (like seizures!)

Daily Mail .com News

4.1k shares

Pfizer/BioNTech and Moderna.

Andrew Pollard, professor of paediatric infection and immunity, and chief investigator on the Oxford vaccine trial, said: 'While most children are relatively unaffected by coronavirus and are unlikely to become unwell with the infection, it is important to establish the safety and immune response to the vaccine in children and young people as some children may benefit from vaccination.

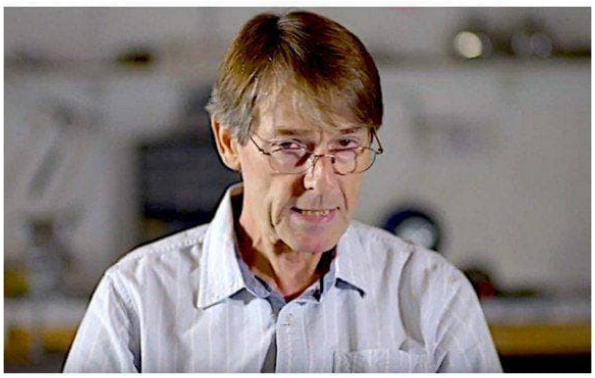
'These new trials will extend our understanding of control of SARS-CoV2 to younger age groups.'

The first vaccinations under the trial will take place this month, with up to 240 children receiving the vaccine and the others receiving a control meningitis jab.

Former Top Pfizer Scientist Michael Yeadon Claims COVID Vaccines May Cause Infertility In Women

DECEMBER 3, 2020 *By* CFT TEAM — LEAVE A COMMENT

Share Tweet Pin Share



Should we assume the risks of this vaccine when we have
NO long term safety studies and
we now know it does NOT prevent transmission of the virus?

THE TIMES OF ISRAEL

Israel's virus czar says 1st dose proving less effective than Pfizer indicated — report

By TOI STAFF
19 January 2021, 1:55 pm | 1

1,311 shares

Coronavirus czar Nachman Ash says the first dose of Pfizer's vaccine is turning out to provide less protection against COVID-19 than the US pharmaceutical firm had initially indicated it would, Army Radio reports.

Many people have gotten infected between the first and second Pfizer shots, he reportedly says, and it appears the protection offered by the firs dose is "less effective than we had thought." The data on the protective effect against the virus of the first dose, it is estimated, is "lower than Pfizer presented," he is quoted saying.

THE EPOCH TIMES

Vaccinated Congressman Tests Positive for COVID-19

BY ZACHARY STIEBER | January 30, 2021 Updated: January 30, 2021

Print

Rep. Stephen Lynch (D-Mass.) tested positive for COVID-19 after receiving two doses of Pfizer's vaccine.

"Congressman Lynch had received the second dose of the Pfizer vaccine and subsequently received a negative COVID-19 test prior to attending President

NEWS

Coronavirus digest: German nursing home sees outbreak after vaccines

Fourteen residents at a German nursing home have tested positive despite receiving two vaccine doses. Markus Söder has warned against lifting Germany's lockdown too soon. All

NEWS

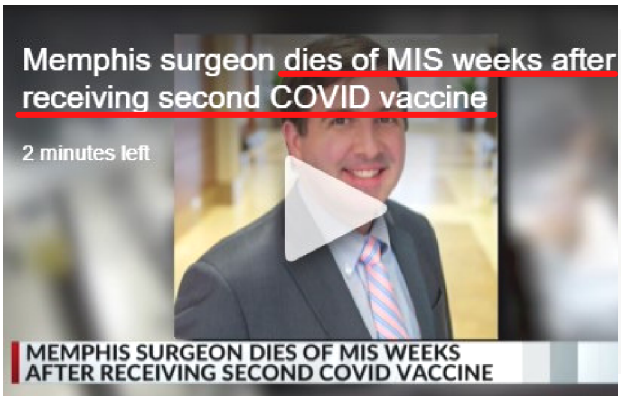
Four people in Oregon who received both doses of vaccine test positive for coronavirus

There are two cases each in Yamhill and Lane counties, the state's Health Authority said.

They all received BOTH vax doses yet still caught Covid?

San Diego County Reports 1st Case of Fully Vaccinated Person Getting COVID-19

"We do expect more [cases]" of vaccinated people getting COVID-19, San Diego County's epidemiologist said. "We expect the number



MEMPHIS, Tenn. — A Mid-South doctor has died of a COVID-related illness. However, he never knew he had the virus and he'd been vaccinated.

Williams, an Orthopedic Surgeon for OrthoSouth, died February 8 of multisystem inflammatory syndrome or MIS, a condition usually affecting children and attacking the immune system.

"The immune system attacks the body in many ways and causes multi-organ system failure," Threlkeld said. "It affects the heart, the gastrointestinal tract and other places."

Threlkeld says Williams tested positive for COVID antibodies, meaning he had COVID at one time, but he never knew it. And he had gotten his second COVID vaccine just weeks before his death.

A typical vaccine safety trial lasts a MINIMUM of 2 years.
Pfizer & Moderna have ENDED their safety trials after only 7 months by allowing placebo recipients to receive the vaccine.
This eliminates all possibilties of capturing long-term adverse effects.

Chicago Tribune

Vaccine trial participants who received placebo now hop the line for the real thing from Pfizer, Moderna

By JUDY PERES
CHICAGO TRIBUNE | JAN 14, 2021

Good news for tens of thousands of volunteers in the COVID-19 vaccine trials: Many of those who received a placebo are now being offered a vaccine — in some cases, earlier than they would otherwise have been eligible.

Participants in Pfizer's vaccine study — some of whom had mounted a noisy campaign on social media — have been advised that anyone who wants one can receive the first of two shots by March 1. Participants in Moderna's vaccine trial already are getting immunized.

That wasn't always the plan, and some experts fear "unblinding" volunteers — that is, letting them know whether they got the vaccine or a placebo — could make it difficult to collect good, long-term data on the experimental vaccines, including how safe they are and when their immunity starts to wear off. But others argue it would be unfair to leave trial participants unprotected from a raging pandemic when an effective vaccine is available.

npr

Long-Term Studies Of COVID-19 Vaccines Hurt By Placebo Recipients Getting Immunized

February 19, 2021 · 5:00 AM ET

Tens of thousands of people who volunteered to be in studies of the Pfizer-BioNTech and Moderna COVID-19 vaccines are still participating in follow-up research. But some key questions won't be easily answered, because many people who had been in the placebo group have now opted to take the vaccine.

Dr. Steven Goodman, a clinical trials specialist at Stanford University, says losing those control groups makes it more difficult to answer some important questions about COVID-19 vaccines.

"We don't know how long protections lasts," he says. "We don't know efficacy against variants — for which we definitely need a good control arm — and we also don't know if there are any differences in any of these parameters by age or race or infirmity."

Below (were) the end dates of the Pfizer & Moderna trials.

Official Title:
A Phase 3, Randomized, Stratified, Observer-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of mRNA-1273 SARS-CoV-2 Vaccine in Adults Aged 18 Years and Older

Actual Study Start Date :
July 27, 2020

Estimated Primary Completion Date :
October 27, 2022

Estimated Study Completion Date :
October 27, 2022

Official Title:
A PHASE 1/2/3, PLACEBO-CONTROLLED, RANDOMIZED, OBSERVER-BLIND, DOSE-FINDING STUDY TO EVALUATE THE SAFETY, TOLERABILITY, IMMUNOGENICITY, AND EFFICACY OF SARS-COV-2 RNA VACCINE CANDIDATES AGAINST COVID-19 IN HEALTHY INDIVIDUALS

Actual Study Start Date :
April 29, 2020

Estimated Primary Completion Date :
August 3, 2021

Estimated Study Completion Date :
January 31, 2023

Also up next, the rollout of the J&J vaccine

Yarah Dalmau
2h ·

"Dec 18th 2020 – My Dad participated in the Johnson & Johnson Trial COVID vaccine shot. They promised all medical expenses paid for any complications related to COVID for the next 2 years. My Dad found security in the medical expense coverage and possibility to have access to a vaccine- that he decided to do the trial.

Dec 25th 2020 - One week later- Christmas Day – Dad was very fatigued. He felt tired right after the shot and when I spoke with him on Christmas day, I recall he sounded exhausted. Which is unlike him, especially on Christmas.

Jan 4th 2021 By the first of the year, he was very sluggish and found it hard to do his exercises that he does weekly. He always likes to stay active.

Jan 11th – 18th Asthma like symptoms, harder to get through his workouts. Very short of breath and started to be winded very easily. Low grade fevers. Hard to make it through calls at work.

Jan 25th – Dad contacted Johnson & Johnson and they did a COVID test. Results came back 3 days later as negative. They advised he has flu like symptoms and to rest and drink lots of fluids.

Jan 29th my Dad could barely walk. His legs became so weak that he shaked when he walked. He thought maybe he was battling a flu and tried to keep resting through the weekend in hopes he would feel better.

Feb 1st - my mom brought him to Lake Mary Emergency room and they immediately transported him via ambulance to Advent Altamonte because of very high troponin levels.

Feb 1st – 4th -My Dad was at Advent Health Altamonte where they ran all kinds of tests and discharged him on the night of Feb 4th . Referring him to an oncologist, infectious disease Dr, cardiologist and PCP.

Feb 5th- Friday I saw my Dad at 11 AM and he could not walk and was very lethargic and sleepy. I called and made all his referral appointments and scheduled him to see his PCP first thing Monday.

Feb 6th- Saturday. I woke up at 6 AM and drove to my parents house and my Dad was completely out of it. He could not get out of bed, pick up his coffee to drink, lift his head. He did not know what year, day, month we were in. We took him to Advent Orlando and within 20 minutes they got him in and found a blood infection and swelling in brain.

Feb 20th We have been here 2 weeks today, and today my Dad started on dialysis.

I am not an anti-vaxxer by any means. But I think there should be some public awareness of the dangers of being part of a trial vaccine. Especially one that is being done in warp speed. And the chance I would recommend someone to get the Johnson & Johnson vaccine, is slim to none. They have denied any type of connection."

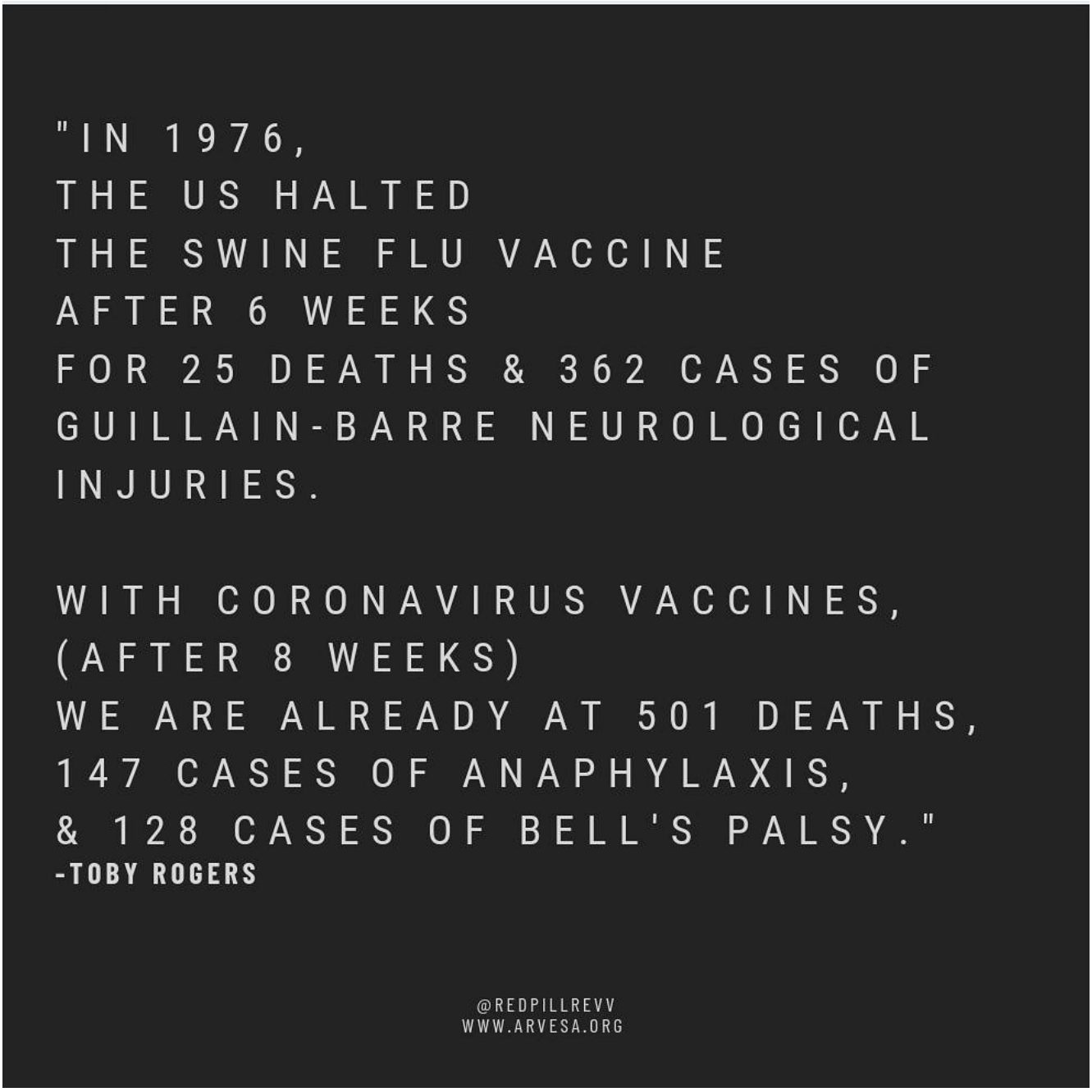
Yarah Dalmau

Yarah Dalmau ···
Feb 27 ·

My Dad took his last breath today on this Earth. I can't wait for our dance in heaven. May he rest with our Lord 🙏

😢🥰💓 423 367 Comments 16 Shares

In 1976 the rushed-to-market Swine Flu vaccine was quickly pulled from the market...after 'only' 25 deaths.



LET'S LOOK AT

THE LAWS

Federal Law states employers, schools, organizations cannot mandate an EAU vaccine.

STAT
The frontiers of health & medicine

Federal law prohibits employers and others from requiring vaccination with a Covid-19 vaccine distributed under an EUA

Ever since the Food and Drug Administration granted emergency use authorization for two new vaccines, employers, schools, and other organizations are grappling with whether to require Covid-19 vaccination.

While organizations are certainly free to encourage their employees, students, and other members to be vaccinated, federal law provides that, at least until the vaccine is licensed, individuals must have the option to accept or decline to be vaccinated.

EUAs are clear: Getting these vaccines is voluntary

The same section of the Federal Food, Drug, and Cosmetic Act that authorizes the FDA to grant emergency use authorization also requires the secretary of Health and Human Services to "ensure that individuals to whom the product is administered are informed ... of the option to accept or refuse administration of the product."

Likewise, the FDA's guidance on emergency use authorization of medical products requires the FDA to "ensure that recipients are informed to the extent practicable given the applicable circumstances ... That they have the option to accept or refuse the EUA product ..."

In the same vein, when Dr. Amanda Cohn, the executive secretary of the CDC's Advisory Committee on Immunization Practices, was asked if Covid-19 vaccination can be required, she responded that under an EUA, "vaccines are not allowed to be mandatory. So, early in this vaccination phase, individuals will have to be consented and they won't be able to be mandatory." Cohn later affirmed that this prohibition on requiring the vaccines applies to organizations, including hospitals.

Much remains unknown about the safety and efficacy of the vaccine

Even though the FDA granted emergency use authorizations for the Pfizer/BioNTech and Moderna vaccines in December 2020, the clinical trials the FDA will rely upon to ultimately decide whether to license these vaccines are still underway and are designed to last for approximately two years to collect adequate data to establish if these vaccines are safe and effective enough for the FDA to license.

The abbreviated timelines for the emergency use applications and authorizations means there is much the FDA does not know about these products even as it authorizes them for emergency use, including their effectiveness against asymptomatic infection, death, and transmission of SARS-CoV-2, the virus that causes the disease.

Given the uncertainty about the two vaccines, their EUAs are explicit that each is "an investigational vaccine not licensed for any indication" and require that all "promotional material relating to the Covid-19 Vaccine clearly and conspicuously ... state that **this product has not been approved or licensed by the FDA**, but has been authorized for emergency use by FDA" (emphasis added).

Do you know what Ohio's Constitution says about forced participation in Health Care?

as it applies to Lake Erie or the navigable waters of the state.

(G) Nothing in Section 1e of Article II, Section 36 of Article II, Article VIII, Section 1 of Article X, Section 3 of Article XVIII, or Section 7 of Article XVIII of the Constitution shall impair or limit the rights established in this section.

(2008)

Powers reserved to the people.

§20 This enumeration of rights shall not be construed to impair or deny others retained by the people; and all powers, not herein delegated, remain with the people.

(1851)

Preservation of the freedom to choose health care and health care coverage

§21 (A) No federal, state, or local law or rule shall compel, directly or indirectly, any person, employer, or health care provider to participate in a health care system.

(B) No federal, state, or local law or rule shall prohibit the purchase or sale of health care or health insurance.

(C) No federal, state, or local law or rule shall impose a penalty or fine for the sale or purchase of health care or health insurance.

(D) This section does not affect laws or rules in effect as of March 19, 2010; affect which services a health care provider or hospital is required to perform or provide; affect terms and conditions of government employment; or affect any laws calculated to deter fraud or punish wrongdoing in the health care industry.

(E) As used in this Section,

(1) "Compel" includes the levying of penalties or fines.

(2) "Health care system" means any public or private entity or program whose function or purpose includes the management of, processing of, enrollment of individuals for, or payment for, in full or in part, health care services, health care data, or health care information for its participants.

(3) "Penalty or fine" means any civil or criminal penalty or fine, tax, salary or wage withholding or surcharge or any named fee established by law or rule by a government established, created, or controlled agency that is used to punish or discourage the exercise of rights protected under this section.

(2011)

Article II: Legislative

In whom power vested.

§1 The legislative power of the state shall be vested in a General Assembly consisting of a Senate and House of Representatives but the people reserve to themselves the power to propose to the General Assembly laws and amendments to the constitution, and to adopt or reject the same at the polls on a referendum vote as hereinafter provided. They also reserve the power to adopt or reject any law, section of any

Forcing people to submit to a pharmaceutical injection as a condition to attend school, church, travel or go into a store violates our God-given, inalienable rights protected under the Common Law and the U.S. Constitution.

Is forcing someone to take the COVID vaccine in violation of the Nuremberg Code?

What Are The Nuremberg Code's Ethical Guidelines For Research?

The Nuremberg Code aimed to protect human subjects from enduring the kind of cruelty and exploitation the prisoners endured at concentration camps. The 10 elements of the code are:

1. Voluntary consent is essential
2. The results of any experiment must be for the greater good of society
3. Human experiments should be based on previous animal experimentation
4. Experiments should be conducted by avoiding physical/mental suffering and injury
5. No experiments should be conducted if it is believed to cause death/disability
6. The risks should never exceed the benefits
7. Adequate facilities should be used to protect subjects
8. Experiments should be conducted only by qualified scientists
9. Subjects should be able to end their participation at any time
10. The scientist in charge must be prepared to terminate the experiment when injury, disability, or death is likely to occur

As the COVID vaccine was approved as an emergency measure, it is still classified as 'experimental.' It is not FDA approved.
Therefore it violates 6 elements of the Nuremberg Code including:

Number 1 - Requiring an experimental vaccine to participate in society is in violation of the Code.
Number 3 - The Covid vaccine skipped required animal safety trials
Number 4 - 36,000 adverse reactions (including PERMANENT DISABILITY) have been recorded (in the US alone) in the first 90 days of rollout
Number 5 - 1,524 deaths from vaccine are recorded in VAERS in first 90 days of rollout (USA only)
Number 6 - The vast majority of Americans have a 99.96% chance of surviving Covid ***without*** a vaccine
Number 10 - No one is pulling the plug on this experimental-status vaccine even though we have thousands of deaths and permanent disabilities.