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VAERS data released Friday by the Centers for Disease Control and Prevention included a total of 1,168,894 reports of adverse events from all age groups following COVID vaccines, including 25,158 deaths and 203,888 serious injuries between Dec. 14, 2020, and March 4, 2022.

by <u>Megan Redshaw</u>, <u>The Defender</u> March 11, 2022

The Centers for Disease Control and Prevention (CDC) today released new data showing a total of 1,168,894 reports of adverse events following COVID vaccines were submitted between Dec. 14, 2020, and March 4, 2022, to the Vaccine Adverse Event Reporting System (VAERS). VAERS is the primary government-funded system for reporting adverse vaccine reactions in the U.S.

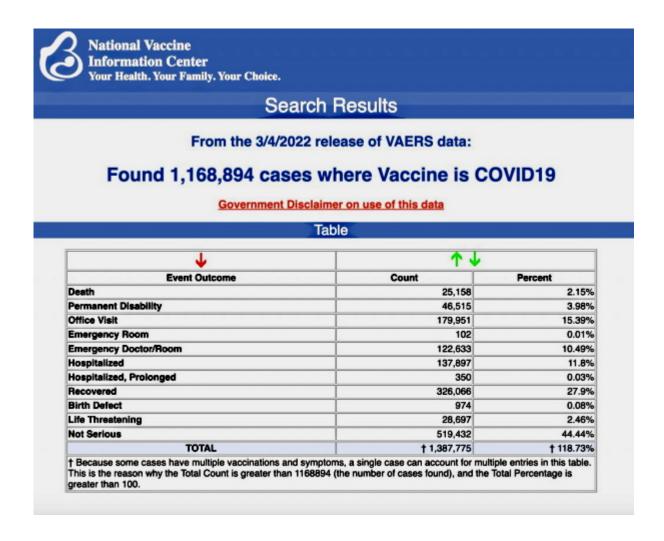
The data included a total of <u>25,158 reports of deaths</u> — an increase of 331 over the previous week — and <u>203,888 reports</u> of serious injuries, including deaths, during the same time period — up 3,557 compared with the previous week.

Excluding "foreign reports" to VAERS, <u>783,282 adverse events</u>, including <u>11,505 deaths</u> and <u>75,286 serious injuries</u>, were reported in the U.S. between Dec. 14, 2020, and March 4, 2022.

Foreign reports are reports foreign subsidiaries send to U.S. vaccine manufacturers. Under U.S. Food and Drug Administration (FDA) regulations, if a manufacturer is notified of a foreign case report that describes an event that is both serious and does not appear on the product's labeling, the manufacturer is required to submit the report to VAERS.

Of the 11,505 U.S. <u>deaths reported</u> as of March 4, 17% occurred within 24 hours of vaccination, 22% occurred within 48 hours of vaccination and 60% occurred in people who experienced an <u>onset of symptoms</u> within 48 hours of being vaccinated.

In the U.S., 554 million COVID vaccine doses had been administered as of March 4, <u>including</u> 327 million doses of Pfizer, 209 million doses of Moderna and 18 million doses of Johnson & Johnson (J&J).



Every Friday, <u>VAERS</u> publishes vaccine injury reports received

as of a specified date. Reports submitted to VAERS require further investigation before a causal relationship can be confirmed. Historically, VAERS has been shown to report only 1% of actual vaccine adverse events.

U.S. VAERS data from Dec. 14, 2020, to March 4, 2022, for 5- to 11-year-olds show:

• <u>9,009 adverse events</u>, including <u>213 rated as serious</u> and <u>5 reported deaths</u>.

The most recent death involves a 7-year-old boy (VAERS I.D. 2152560) from Washington who died 13 days after receiving his first dose of Pfizer's COVID vaccine when he went into shock and suffered cardiac arrest. He was unable to be resuscitated and died in the emergency department.

• 17 reports of myocarditis and pericarditis (heart inflammation).

The CDC uses a <u>narrowed case definition</u> of "myocarditis," which <u>excludes cases</u> of cardiac arrest, ischemic strokes and deaths due to heart problems that occur before one has the chance to go to the emergency department.

• 32 reports of blood clotting disorders.

U.S. VAERS data from Dec. 14, 2020, to March 4, 2022, for 12- to 17-year-olds show:

• <u>30,193 adverse events</u>, including <u>1,734 rated as serious</u> and <u>40 reported deaths</u>.

The most recent death involves a 14-year-old boy (VAERS I.D. 2148498) who experienced a cerebral aneurysm leading to death one day after receiving his first dose of Pfizer's COVID vaccine.

• 69 reports of anaphylaxis among 12- to 17-year-olds where the reaction was life-threatening, required treatment or resulted in death — with 96% of cases

- attributed to Pfizer's vaccine.
- <u>650 reports</u> of myocarditis and pericarditis with <u>631</u> cases attributed to Pfizer's vaccine.
- <u>161 reports</u> of blood clotting disorders, with all cases attributed to Pfizer.

U.S. VAERS data from Dec. 14, 2020, to March 4, 2022, for all age groups combined, show:

- 19% of deaths were related to cardiac disorders.
- 54% of those who died were male, 41% were female and the remaining death reports did not include the gender of the deceased.
- The <u>average age</u> of death was 72.7.
- As of March 4, <u>5,233 pregnant women</u> reported adverse events related to COVID vaccines, including 1,664 reports of <u>miscarriage or premature birth</u>.
- Of the <u>3,596 cases of Bell's Palsy</u> reported, 51% were attributed to <u>Pfizer</u> vaccinations, 40% to <u>Moderna</u> and 8% to <u>J&J</u>.
- ■859 reports of <u>Guillain-Barré syndrome</u>, with 41% of cases <u>attributed to Pfizer</u>, 30% to <u>Moderna</u> and 28% to <u>J&J</u>.
- 2,358 reports of anaphylaxis where the reaction was life-threatening, required treatment or resulted in death.
- 1,632 reports of myocardial infarction.
- <u>13,428 reports</u> of blood clotting disorders in the U.S. Of those, <u>5,992 reports</u> were attributed to Pfizer, <u>4,784 reports</u> to Moderna and <u>2,607 reports</u> to J&J.
- 4,065 cases of myocarditis and pericarditis with 2,499 cases attributed to Pfizer, 1,381 cases to Moderna and 174 cases to J&J's COVID vaccine.

CDC study concludes COVID vaccine adverse events 'mild'

A <u>study</u> funded by the CDC and published Monday in The Lancet <u>concluded</u> most COVID vaccine-related adverse events

reported during the first six months of the rollout in the U.S were "mild and short in duration," despite thousands of deaths reported to VAERS.

For the study, researchers analyzed data captured between Dec. 14, 2020, and June 14, 2021, by VAERS and v-safe, both of which are overseen by the CDC. Nearly 300 million doses of COVID vaccines were administered during the study period.

The authors found that of the 340,522 adverse events reported to VAERS, 27,023 (8%) were serious, 4,496 were deaths. The authors said the cause of the increased reporting of deaths during the first few days after vaccination might represent "reporting bias."

The authors suggested that deaths occurring soon after vaccination were more likely to be reported than deaths that occurred later. This, they believe, is why the number of deaths asymptotically approaches zero as more time elapses since vaccination.

<u>Jessica Rose</u>, Ph.D., attempted to duplicate the Lancet authors' findings through her independent <u>analysis</u> of the VAERS data. Despite filtering the database using three different date stamps, Rose was unable to duplicate the Lancet study's results.

Florida surgeon general breaks with CDC, recommends against shots for healthy kids

Florida's surgeon general on Monday said he will issue guidance formally recommending against COVID vaccines for healthy children. Florida is the first state to break with official guidance from the CDC, which <u>recommends</u> all children over age 5 get the vaccine.

Dr. Joseph Ladapo made the announcement at a <u>roundtable</u>, hosted by Gov. Ron DeSantis, featuring physicians and other medical experts who criticized CDC and government policies,

including mask mandates and lockdowns, which they said were ineffective and harmful.

Ladapo and DeSantis said the new guidance had to do with lingering questions about the vaccines' potential health risks for young people and the fact that children are in a low-risk category for severe COVID.

COVID vaccines may be enhancing disease

COVID vaccines may be causing enhanced disease because they target an old version of the coronavirus, Dr. Robert Malone told the Epoch Times in a recent interview.

"The data are showing that vaccination can actually increase the risk of being infected with the <u>Omicron</u> version of this virus," Malone said, referring to how in some areas, including Scotland and New Zealand, patients hospitalized with COVID are more likely to have received a COVID vaccine.

U.S. drug regulators identified vaccine-associated enhanced diseases (VAED) as an "important potential risk" of COVID vaccines, along with enhanced respiratory disease.

Some adverse events reported following COVID vaccination "could indicate" VAED, according to a CDC team.

VAED refers to disease "resulting from infection in individuals primed with non-protective immune responses against the respective wild-type viruses," researchers <u>said</u> <u>last year</u>.

"Given that these enhanced responses are triggered by failed attempts to control the infecting virus, VAED typically presents with symptoms related to the target organ of the infection pathogen," they added.

Vaccine researcher develops tinnitus after COVID shot, calls for further study

Dr. Gregory Poland, director of the Mayo Clinic's Vaccine Research Group in Rochester, Minnesota, <u>developed lifealtering tinnitus</u>, or ringing in the ear, after receiving his second dose of a COVID vaccine.

"It was like someone suddenly blew a dog whistle in my ear," Poland told MedPage Today. "It has been pretty much unrelenting."

Poland then received a booster, after which his tinnitus briefly disappeared but then returned at a slightly higher pitch. Poland realized his life may never be the same and says he has received emails from people across the world struggling with the same condition to the point they're considering taking their own lives.

Poland, who said he supports COVID vaccines, believes there may be tens of thousands of people affected in the U.S. and is calling for more research to be done to provide help to people desperate for relief.

Michigan woman files claim over mom's COVID vaccinerelated death

Tatum Strieter-Byron is asking the federal government to compensate her for the death in April of her mother Sandra Jacobs. An <u>autopsy confirmed</u> Jacobs died from a blood-clotting disorder caused by J&J's COVID vaccine.

Strieter-Byron received confirmation Monday her claim to the Countermeasures Injury Compensation Program (CICP) had been received. The program was established to give pharmaceutical companies blanket liability protection from harm caused by their COVID vaccines.

In 2020, the U.S. Health and Human Services secretary invoked the <u>Public Readiness and Emergency Preparedness (PREP) Act</u> and declared COVID-19 a public health emergency, providing J&J other COVID vaccine makers immunity from lawsuits.

The only exception under the PREP Act is if a plaintiff can prove a vaccine-related death or serious physical injury was caused by "willful misconduct." The protections, unless amended or rescinded, extend through Oct. 1, 2024.

Claims to the CICP must be made within one year of the date the vaccine was received.

Jacobs, 60, <u>received</u> the single-shot vaccine at a CVS pharmacy on April 8, 2021, just five days before federal health agencies temporarily <u>paused the vaccine</u> to examine numerous reports of a serious and potentially fatal blood-clotting disorder.

Jacobs died on April 21 of "complications of cerebral venous sinus thrombosis," a type of stroke caused by the vaccine.

<u>Children's Health Defense</u> asks anyone who has experienced an adverse reaction, to any vaccine, to file a report following <u>these three steps</u>.

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