

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.

By [Children's Health Defense](#)

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According to new data released today, as of Feb. 12, 15,923 adverse reactions to COVID vaccines, including 929 deaths, [have been reported](#) to the Centers for Disease Control and Prevention's (CDC) [Vaccine Adverse Event Reporting System](#) (VAERS) since Dec. 14, 2020.

VAERS is the primary mechanism in the U.S. for reporting adverse vaccine reactions. Reports submitted to VAERS require further investigation before a determination can be made as to whether the reported adverse event was directly or indirectly caused by the vaccine.

From the 2/12/2021 release of VAERS data:

Found 15,923 cases where Vaccine is COVID19

Table

 Event Outcome	 Count	 Percent
Death	929	5.83%
Permanent Disability	318	1.99%
Office Visit	2,193	13.75%
Emergency Room	21	0.13%
Emergency Room/Room	3,433	21.54%
Hospitalized	1,889	11.74%
Recovered	5,709	35.85%
Birth Defect	25	0.16%
Life Threatening	616	3.87%
Not Serious	5,681	35.69%
TOTAL	↑ 20,788	↑ 130.42%

† Because some cases have multiple vaccinations and symptoms, a single case can account for multiple entries in this table. This is the reason why the Total Count is greater than 15,923 (the number of cases found), and the Total Percentage is greater than 100.

The latest VAERS data show that 799 of the deaths were reported in the U.S., and that about

one-third of those deaths occurred within 48 hours of the individual receiving the vaccination.

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)

[CBS Detroit reported](#) this week that a 68-year old news anchor died one day after being vaccinated for COVID of a [suspected stroke](#).

Reports of deaths among elderly people after being vaccinated for COVID continue to surface, including [the article](#) published this week by The Defender about 46 nursing home residents in Spain who died within one month of receiving the Pfizer vaccine.

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.

[AstraZeneca’s COVID vaccine](#), which does not use mRNA technology, was approved for emergency use this week by the World Health Organization, paving the way for some countries to start using it. However, as [The Defender reported](#) this week, some nations have said they won’t use it, citing safety and efficacy concerns.

[FiercePharma reported today](#) that the FDA may reject the AstraZeneca vaccine over concerns relating to efficacy, especially against new COVID variants, and manufacturing issues.

News reports indicate that a growing number of people, including [nearly 30% of healthcare workers](#), now say they don't want the COVID vaccine, citing safety concerns.

[The Washington Post reported](#) this week that nearly a third of military personnel are opting out of the vaccines, and [ESPN reported](#) that top NBA players are reluctant to promote the vaccine.

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.

As of Feb. 19, about [56.3 million people](#) in the U.S. had received one or both doses of a COVID vaccine.

While the VAERS database numbers may seem sobering, according to a U.S. Department of Health and Human Services [study](#), the actual number of [adverse events](#) is likely significantly higher. VAERS is a [passive surveillance system](#) that relies on the willingness of individuals to submit reports voluntarily.

According to the VAERS website, healthcare providers are required by law to report to VAERS:

- Any adverse event listed in the VAERS Table of Reportable Events Following Vaccination that occurs within the specified time period after vaccination
- An adverse event listed by the vaccine manufacturer as a contraindication to further doses of the vaccine

[The CDC says](#) healthcare providers are strongly encouraged to report:

- Any adverse event that occurs after the administration of a vaccine licensed in the United States, whether or not it is clear that a vaccine caused the adverse event
- Vaccine administration errors

However, “within the specified time” means that reactions occurring outside that timeframe may not be reported, in addition to reactions suffered hours or days later by people who don't report those reactions to their healthcare provider.

Vaccine manufacturers are required to report to VAERS “all adverse events that come to their attention.”

Historically, however, fewer than [fewer than 1%](#) of adverse events have ever been reported to VAERS, a system that Children's Health Defense has previously referred to as an “abject [failure](#),” including in a December 2020 [letter](#) to Dr. David Kessler, former FDA director and now co-chair of the COVID-19 Advisory Board and President Biden's [version](#) of Operation Warp Speed.

A critic familiar with VAERS' shortcomings bluntly condemned VAERS in [The BMJ](#) as “nothing more than window dressing, and a part of U.S. authorities' systematic effort to reassure/deceive us about vaccine safety.”

CHD is calling for complete transparency. The children's health organization is asking Kessler and the federal government to release all of the data from the clinical trials and suspend COVID-19 vaccine use in any group not adequately represented in the clinical trials, including the elderly, frail and anyone with comorbidities.

CHD is also asking for full transparency in post-marketing data that reports all health outcomes, including new diagnoses of autoimmune disorders, adverse events and deaths from COVID vaccines.

Children's Health Defense asks anyone who has experienced an adverse reaction, to any vaccine, to file a report following [these three steps](#).

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