

# 1.4 Million Adverse Events After COVID Vaccines Reported to VAERS, Canadian Woman Dies 7 Minutes After Bivalent Booster Shot

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*The Centers for Disease Control and Prevention (CDC) today released new data showing a total of [1,418,220 reports of adverse events](#) following [COVID-19](#) vaccines were submitted between Dec. 14, 2020, and Sept. 16, 2022, to the Vaccine Adverse Event Reporting System (VAERS). That’s an increase of 10,811 adverse events over the previous week.*

VAERS is the primary government-funded system for reporting adverse vaccine reactions in the U.S.

The data included a total of [31,074 reports of deaths](#) — an increase of 139 over the previous week — and [258,480 serious injuries](#), including deaths, during the same time period — up 1,253 compared with the previous week.

There were a total of [563 reports of adverse events](#) following the new bivalent booster COVID-19 vaccine as of Sept. 16, 2022. The data included a total of [5 deaths](#) and [31 serious injuries](#).

Of the 31,074 reported deaths, [19,934 cases](#) are attributed to Pfizer’s COVID-19 vaccine, [8,385 cases](#) to Moderna, [2,704 cases](#) to Johnson & Johnson (J&J) and [no cases](#) yet reported for Novavax.

Excluding “[foreign reports](#)” to VAERS, [868,175 adverse events](#), including [14,531 deaths](#) and [90,422 serious injuries](#), were reported in the U.S. between Dec. 14, 2020, and Sept. 16, 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 14,531 [deaths reported](#) as of Sept. 16, 7% occurred within 24 hours of vaccination and 15% occurred within 48 hours of vaccination.

In the U.S., 600 million COVID-19 vaccine doses had been administered as of Sept. 14, [including](#) 363 million doses of Pfizer, 231 million doses of Moderna and 19 million doses of Johnson & Johnson (J&J).






From the 9/16/2022 release of VAERS data:

**Found 1,418,220 cases where Vaccine is COVID19 or COVID19-2**

[Government Disclaimer on use of this data](#)

Table

 Age	Count	 	Percent
< 6 Months		176	0.01%
6-11 Months		230	0.02%
1-2 Years		726	0.05%
3-5 Years		2,890	0.2%
6-17 Years		51,745	3.65%
18-29 Years		121,598	8.57%
30-39 Years		161,886	11.41%
40-49 Years		161,665	11.4%
50-59 Years		166,897	11.77%
60-64 Years		78,939	5.57%
65-79 Years		184,815	13.03%
80+ Years		55,660	3.92%
Unknown		430,993	30.39%
<b>TOTAL</b>		<b>1,418,220</b>	<b>100%</b>

Every Friday, [VAERS](#) 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](#).

VAERS data from Dec. 14, 2020, to Sept. 16, 2022, for 6-month-olds to 5-year-olds show:

- [3,843 adverse events](#), including [169 cases rated as serious](#) and [6 reported deaths](#).
- [5 reports](#) of myocarditis and pericarditis (heart inflammation).
  - The CDC uses a [narrowed case definition](#) of “myocarditis,” which [excludes cases](#) of cardiac arrest, [ischemic strokes](#) and deaths due to heart problems that occur before one has the chance to go to the emergency department.
- [26 reports](#) of blood clotting disorders.
- [43 reports](#) of seizures.

VAERS data from Dec. 14, 2020, to Sept. 16, 2022, for 5- to 11-year-olds show:

- [14,115 adverse events](#), including [665 rated as serious](#) and [28 reported deaths](#).
- [45 reports](#) of myocarditis and pericarditis.
- [67 reports](#) of blood clotting disorders.
- [180 reports](#) of seizures.

VAERS data from Dec. 14, 2020, to Sept. 16, 2022, for 12- to 17-year-olds show:

- [39,472 adverse events](#), including [4,277 rated as serious](#) and [120 reported deaths](#).
  - According to the CDC, “VAERS data [available to the public](#) include only the initial report data to VAERS. Updated data which contains data from medical records and corrections reported during follow up are used by the government for analysis. However, for numerous reasons including data consistency, these amended data are not available to the public.”
- [269 reports](#) of anaphylaxis among 12- to 17-year-olds where the reaction was life-threatening, required treatment or resulted in death.
- [1,310 reports](#) of myocarditis and pericarditis with [1,158 cases](#) attributed to Pfizer’s vaccine.
- [300 reports](#) of blood clotting disorders with [277 cases](#) attributed to Pfizer.
- [26 cases](#) of postural orthostatic tachycardia syndrome (POTS) with all cases attributed to Pfizer’s vaccine.

VAERS data from Dec. 14, 2020, to Sept. 16, 2022, for all age groups combined, show:

- 16% of deaths were related to cardiac disorders.
- 53% of those who [died were male](#), 42% [were female](#) and the remaining death reports did not include the gender of the deceased.
- The [average age](#) of death was 72.
- As of Sept. 16, [8,471 pregnant women](#) reported adverse events related to COVID-19 vaccines, including [4,898 reports of miscarriage or premature birth](#).
- Of the [16,353 cases of Bell’s Palsy](#) reported, 73% were [attributed to Pfizer](#) vaccinations, [22% to Moderna](#) and [5% to J&J](#).
- [2,968 reports](#) of Guillain-Barré syndrome, with 64% of cases attributed to Pfizer, 19% to Moderna and 17% to J&J.
- [10,015 reports](#) of anaphylaxis where the reaction was life-threatening, required treatment or resulted in death.
- [4,756 reports](#) of myocardial infarction.
- [43,030 reports](#) of blood-clotting disorders. Of those, [29,522 reports](#) were attributed to Pfizer, [9,659 reports](#) to Moderna and [3,789 reports](#) to J&J.
- [23,926 cases](#) of myocarditis and pericarditis with [18,186 cases](#) attributed to Pfizer, [5,304 cases](#) to Moderna and [410 cases](#) to J&J.
- [67 cases](#) of Creutzfeldt-Jakob disease with [54 cases](#) attributed to Pfizer, [12 cases](#) to Moderna and [1 case](#) to J&J.
- [534 cases](#) of POTS with [395 cases](#) attributed to Pfizer, [119 cases](#) to Moderna and [21 cases](#) to J&J.

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

## Canadian woman dies 7 minutes after booster shot, coroner rules 'natural causes'

A Canadian woman on Sept. 14 [died suddenly](#) in a drug store within minutes of receiving the new bivalent COVID-19 booster shot, [according to her daughter](#).

Carol Pearce [texted her daughter](#), Stephanie Foster, at 12:31 p.m. to say she received her booster shot and was waiting the recommended 15 minutes at the drug store. At 12:38 p.m., Pearce was unconscious.

An eyewitness [told SASKTODAY.ca](#) she was in the store when she heard screaming and crying. "They were saying in the store that it was about seven minutes," after Pearce got the booster shot that she collapsed on the floor, the witness said.

A spokesperson from Saskatchewan Health told SASKTODAY.ca on Thursday Pearce died from natural causes.

"The Saskatchewan coroner's service has investigated this instance and determined that the person died from natural causes," Dale Hunter, a communications consultant from the Saskatchewan Ministry of Health said in an email.

"I do not believe this was caused from natural causes," Foster said. "My Mom had no health conditions. I believe had she not gotten that Covid shot then she would be here with us today!"

The family said when Pearce left for her appointment, she was happy and healthy.

Reportedly 15 minutes after getting a shot in Saskatoon @ Shoppers Drug Mart, Stephanie Foster's mother dies. <https://t.co/iZoBTeikclpic.twitter.com/ZwlwnVvjge>

— wlctv.ca (@wlctv\_ca) [September 17, 2022](#)

## CDC predicts COVID boosters for kids by mid-October

The CDC expects [new bivalent COVID-19 boosters](#) to be ready for kids ages 5 to 11 by mid-October, [Reuters reported](#).

The agency said in a document released Tuesday it [will issue a recommendation](#) in early- to mid-October if the FDA authorizes the modified booster for this age group.

The CDC said it expects Pfizer-BioNTech's bivalent booster to be available for children ages 5 to 11 and Moderna's bivalent booster to be available for children ages 6 to 17, pending authorization by the FDA.

The bivalent booster targets both the original Wuhan strain [no longer in circulation](#) and the Omicron BA.4/BA.5 subvariants. The CDC [expects pre-orders](#) of the Pfizer booster to begin next week even though the FDA has not yet signed off on the vaccine for this age group.

Moderna's booster is the same formula used for children and adults and will not require a separate pre-order period, [the agency said](#).

## Woman's brain inflammation caused by Moderna COVID vaccine, case study confirms

A [healthy 35-year-old woman](#) who experienced a seizure two days after her second dose of Moderna's COVID-19 vaccine was diagnosed with [limbic encephalitis](#), a rare form of brain inflammation.

The authors of the report said the vaccine caused the woman's condition.

According to the case report, when the woman arrived at the hospital she had a fever, followed by generalized [tonic-clonic seizures](#) that lasted for approximately five minutes and a [postictal \(post-seizure\) phase](#) of confusion that lasted approximately 30 minutes.

While in the ER, she had two more generalized seizures and was stable. Upon admission to the hospital, a lumbar puncture was performed revealing significant [lymphocytosis](#).

She then developed [anisocoria](#), a condition in which the pupil of one eye differs in size from the pupil of the other eye. An urgent CT scan of the head showed possible temporal lobe [hypodensities](#) (darker areas on the CT scan). A head MRI showed features suggestive of limbic encephalitis. A subsequent electroencephalogram showed severe post-seizure encephalitic changes.

While hospitalized, the patient [developed abnormal behavior](#) associated with limbic encephalitis and was diagnosed with autoimmune limbic encephalitis.

Researchers concluded the patient's limbic encephalitis — which occurred quickly after COVID-19 vaccination — is an uncommon side effect of the vaccine based on the timing of symptoms and absence of other identifiable causes.

## U.S. Supreme Court to consider case against COVID vaccine mandate for New York City employees

The U.S. Supreme Court on Oct. 7 will consider a [New York Police Department detective's challenge](#) to the city's COVID-19 vaccine mandate for municipal employees.

Supreme Court Justice Clarence Thomas granted the request by the detective, Anthony Marciano, after Justice Sonia Sotomayor initially rejected it. Marciano's challenge seeks to strike down the mandate for all city workers — including firefighters, police and teachers.

According to data from the mayor's office, as of Aug. 30, [1,761 city workers](#) have been fired for not getting the vaccine. That number is likely higher after the New York City Department of Education this month [fired another 850 teachers](#) and other staff, bringing the total to nearly 2,000 school employees fired for failure to comply with the mandate, according to the New York Post.

The Supreme Court's decision to [consider Marciano's request](#) came one day after New York Mayor Eric Adams rescinded the city's COVID-19 vaccine mandate for private employers and student athletes — but not municipal employees.

## Woman develops blood clots, hallucinations, mini-strokes after Pfizer shots

In April 2021, Cathy Nagy's 84-year-old mother-in-law received her first dose of the [Pfizer](#)

[vaccine](#).

According to Nagy, “about a week-and-a-half later, Marilyn experienced an [episode of hallucinations](#), seeing people outside who were not there.”

Nagy told [The Defender](#):

“Marilyn’s family attributed [the episode] to her dementia, even though she had not had any prior episodes ... [They] did not make a connection to the vaccination.”

But the family’s opinion changed a few weeks later, after Marilyn got her second Pfizer shot and “had a more dramatic reaction,” experiencing “another episode of hallucinations that same day, extreme weakness and pain in her leg,” Nagy said.

Marilyn seemed to recover, but over the coming weeks, she continued to experience occasional hallucinations, and an increased decline in her cognition, Nagy said, prompting a visit to a neurologist.

An MRI “revealed she was experiencing blood clots, bleeding and mini-strokes in her brain, and she was sent directly to the ER for observation and treatment,” Nagy said.

At the hospital, Marilyn was administered blood-thinning medications and appointments were scheduled “with several specialists.”

About a month later, she developed [deep vein thrombosis](#) in her left leg, which according to Nagy, was “the same leg that had been experiencing pain since her second vaccine.”

“Her right leg also became painfully swollen, although no blood clot was found,” Nagy said.

## Biden says ‘pandemic is over’ but mandates, federal health emergency remain in effect

President Joe Biden’s [Sept. 18 statement](#) that the COVID-19 pandemic is “over” prompted questions about why his administration hasn’t ended the federal public health emergency and how the government or employers can continue to mandate Emergency Use Authorized vaccines.

In an interview aired Sunday on “60 Minutes,” President Joe Biden declared the COVID-19 “pandemic is over,” in the first such statement by a prominent political figure in the U.S. Biden made the comment from the North American International Auto Show in Detroit, which was held for the first time since 2019.

In his remarks, Biden, using the auto show as a point of reference, stated:

“We still have a problem with COVID. We’re still doing a lotta work on it ... but the pandemic is over.

“If you notice, no one’s wearing masks. Everybody seems to be in pretty good shape. And so I think it’s changing. And I think this is a perfect example of it.”

[Sarah Lovenheim](#), a spokesperson for the U.S. Department of Health and Human Services (HHS), said the federal [public health emergency](#) “remains in effect” and that a 60-day notice

would be provided by the agency before it is rescinded.

The earliest possible date when the emergency declaration would be lifted is in late November — well past the current Oct. 15 expiration date.

HHS first declared the public health emergency on Jan. 31, 2020, and renewed it 10 times since, most recently on [July 15](#). In order for COVID-19 vaccines to be authorized for emergency use, a “public health emergency” [must remain in effect](#).

## Woman with heart condition injured by Pfizer shot after being forced to choose between job and vaccine

When her employer in 2021 mandated all employees get the COVID-19 vaccine, Heather Elkins, who had a pre-existing heart condition, [requested a medical exemption](#).

But the Oregon Department of Human Services turned down her application — even though Elkins had transitioned, in 2020, to a permanent work-from-home employee.

Elkins, 45, told The Defender she was reluctant to get the vaccine in part because of the lack of long-term safety data but also because she saw reports of people developing [heart issues](#).

Elkins got the [Pfizer](#) vaccine on Sept. 12, 2021, and “within about an hour,” she said, began experiencing [adverse reactions](#).

The reactions she experienced after the first shot didn’t qualify her for a medical exemption from the second shot, she said.

So despite continued efforts to secure a medical exemption, Elkins received her second Pfizer shot on Oct. 3, 2021, and experienced “the same severe reaction as with the first dose.”

A year later, Elkins continues to experience symptoms that affect her ability to work.

Elkins told The Defender:

“To date, I’m experiencing all the same issues in addition to other symptoms like brain fog, horrible joint pain, pins and needles sensations, painful nerve zaps, shakiness and tremors in my hands and feet, and vision issues.

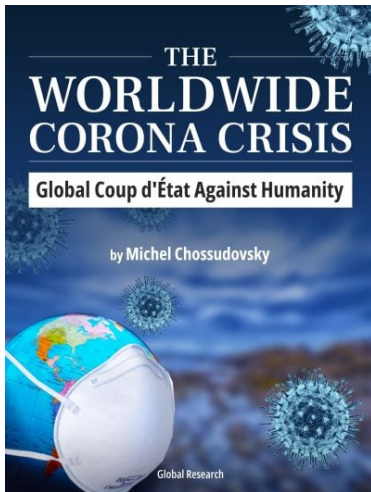
“I’ve now been mostly confined to my bed for the last 11 months, unable to do minimal activities because of my debilitating symptoms.”

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*Megan Redshaw is a staff attorney for Children’s Health Defense and a reporter for The Defender.*

*Featured image is from CHD*



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by Michel Chossudovsky

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