



Written by [Rebecca Terrell](#) on October 1, 2021

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Correction, Please!

FDA Sleight of Hand: Does the Pfizer Vaccine Really Have Full Approval?



It's official: The FDA approved Pfizer's Comirnaty vaccine for ages 16 and up. Pfizer's BioNTech product is still not approved. Could this approval be a way to avoid liability in the United States? (*Photo credit: Capuski/E+/GettyImagesPlus*)

Item: *The British Medical Journal reported August 24, "The U.S. Food and Drug Administration has granted full approval to the two dose Pfizer-BioNTech vaccine to protect against covid-19 for people aged 16 and above.... The vaccine has been used under emergency authorisation since December 2020, with Pfizer applying for full approval on 7 May 2021 and approval being granted in under four months, the fastest time in FDA history.... Anthony Fauci, director of the National Institute of Allergy and Infectious Diseases, told the national television programme PBS Newshour that the FDA ... 'worked double time fast. They just did not cut any corners.'"*

Item: *According to a September 9 article in USA Today, "Biden announced a new rule by the Labor Department that will require all businesses with 100 or more employees to be vaccinated or submit to testing at least once a week.... 'What more is there to wait for? What more do you need to see?' he said. 'We've made vaccinations free, safe and convenient. The vaccine is FDA approved. Over 200 million Americans have gotten at least one shot. We've been patient, but our patience is wearing thin, and your refusal has cost all of us.'"*

Item: *Air Force News Service noted September 22, "The COVID-19 vaccine has been mandated across the Department of Defense and despite its demonstrated effectiveness and safety, a host of myths have left some Airmen and Guardians hesitant to receive it. While social media posts and some news outlets may make it harder to keep up with what is fact or fiction, the science is clear ... approved COVID-19 vaccines work."*

Correction: The FDA has seemingly removed any objection to the Pfizer COVID vaccine by approving the company's biologics license application. That gives the gene-altering serum general use approval for individuals age 16 and over, as opposed to the more restrictive Emergency Use Authorization (EUA)



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assigned to other COVID vaccines.

EUA products are experimental, meaning that under U.S. law, everyone has a right to refuse them. Forcing someone to get the experimental jab so they can work or go to school is tantamount to Nazi human experimentation during World War II. Those atrocities gave birth to the Nuremburg Code, which dictates that voluntary consent is “absolutely essential” when it comes to medical experiments involving humans, and that those involved should be protected “against *even remote* possibilities of injury, disability or death.” (Emphasis added.)

FDA approval seemingly removed Nuremburg objections.

But there’s a catch: The FDA only licensed Pfizer’s Comirnaty vaccine, not its BioNTech product, which remains under EUA restriction. Confusing the matter further, in its approval documentation, FDA acknowledges “insufficient stocks” of the Comirnaty vials, but “sufficient amounts” of BioNTech, claiming the products can be used “interchangeably.” FDA chief scientist Denise Hinton wrote, “The licensed vaccine has the same formulation as the EUA-authorized vaccine and the products can be used interchangeably to provide the vaccination series without presenting any safety or effectiveness concerns. The products are legally distinct with certain differences that do not impact safety or effectiveness.”

Did the FDA just equate an EUA vaccine to one that is fully licensed? Let’s look at the “legally distinct” differences. First, U.S. law permits employers and schools to require licensed vaccines. Second, while manufacturers of EUA-approved vaccines are immune from product liability (as are those who produce vaccines placed on the FDA’s mandatory childhood immunization schedule), licensed adult vaccines do not enjoy liability protection. They are subject to the same product liability as any consumer goods in the U.S. market. In other words, if you were injured by the Comirnaty vaccine, you could sue. Injured by BioNTech? No chance.

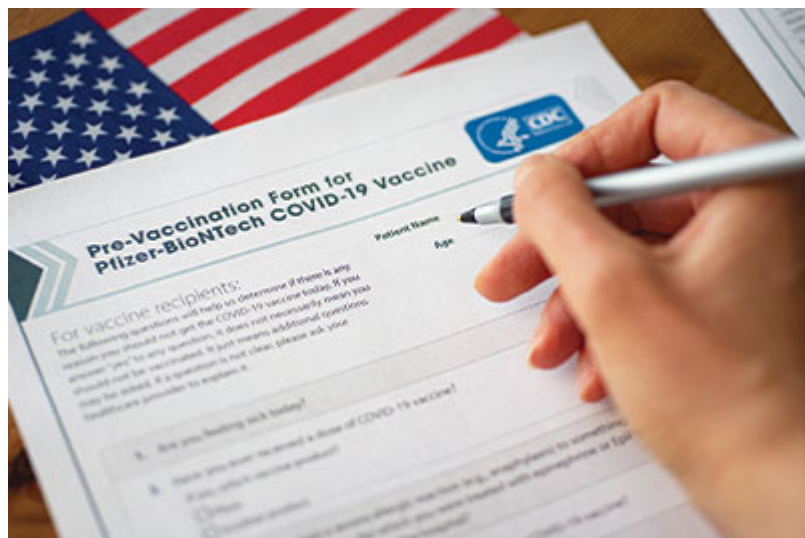
So why would Pfizer market Comirnaty in the United States, where it can end up in court when patients are injured? With “significant amounts” of BioNTech available, written FDA go-ahead, media silence on this criminal sleight-of-hand, and general public ignorance of what amounts to illegal mandates resulting from FDA approval, Pfizer can unload all its liability-immune product on an unsuspecting public while President Biden, in Hitleresque fashion, orders private companies nationwide to mandate the jab.

What’s particularly disconcerting is that there are more reported deaths attributable to Pfizer vaccines than to the others combined. The Vaccine Adverse Event Reporting System (VAERS), operated jointly by the FDA and the U.S. Centers for Disease Control and Prevention (CDC), is a voluntary reporting system designed to detect possible early warnings in regard to vaccine safety. As of mid-September, VAERS recorded more than 10,000 deaths associate with Pfizer’s product, whereas deaths reportedly caused by other manufacturers’ vaccines amounted to under 5,000. Overall, VAERS has received more than 700,000 reports of adverse events, including nearly 20,000 that involve permanent disability and tens of thousands of life-threatening events. How does that measure up to the Nuremburg mandate “against *even remote* possibilities of injury, disability or death”?



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Not buying it: The FDA approval of Comirnaty did not cause the increase in vaccination rates the government and Pfizer hoped for. In fact, weekly vaccination rates in the United States have declined since the approval. *(Photo credit: EvgeniaParajanian/GettyImagesPlus)*

Contrast this, too, with past VAERS experience. “A typical new drug ... at about 50 unexplained deaths for a new product, it’s pulled off the market,” Dr. Peter McCullough told *The New American*. “In the 1976 swine flu pandemic we attempted to vaccinate 55 million Americans. At 25 deaths, the program was killed.” Dr. McCullough is a principal faculty in internal medicine at Texas A&M University Health Sciences Center and developed a successful COVID outpatient treatment early in the pandemic.

Want more undiluted proof of Pfizer’s track record? Let’s look at Israel, where the government cut an early exclusivity deal with Pfizer, paying above-market value for premium access to its vaccine. The Pfizer CEO blatantly called Israel “the world’s lab.”

What are the lab results? Journalist Sharyl Attkisson has kept a running tally of COVID-19 vaccine concerns on her blog, where Israel is heavily featured. For example, a recent case-control study from that country found an increased risk of myocarditis (heart inflammation), appendicitis, lymphadenopathy (inflammation of lymph nodes), and herpes zoster infection (shingles). Israel’s Ministry of Health has admitted that nearly half of adults infected with the COVID-19 virus are fully vaccinated, and that 60 percent of hospitalizations involve fully vaccinated individuals. A spokesman for Israel’s largest health maintenance organization called their Pfizer experience “a very clear warning sign for the rest of the world.” So Israel has lifted mandates, right? On the contrary, its reaction is unbelievable: The country now only awards vaccine passports to those who have had three shots, not a mere two, and the passport is only good for six months.

The good news is that stateside, we’re not fooled. FDA’s thumbs-up “didn’t cause the uptick in vaccinations that people had hoped, data shows,” announced MSN Business Insider on September 9. “The US’s average weekly vaccination rate has declined 38% since the FDA approved Pfizer’s COVID-19 vaccine.”

And more good news is what you can do to protect yourself. Writing for Children’s Health Defense, attorney Robert F. Kennedy, Jr. gives this advice: “When someone orders you to get the vaccine: Ask to see the vial. If it says ‘Comirnaty,’ it’s a licensed product. If it says ‘Pfizer BioNTech,’ it’s an experimental product, and under 21 U.S. Code 360bbb, you have the right to refuse.” Of course, all



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other so-called COVID vaccines are still under EUA restrictions, so you have the right to refuse them as well.



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