

Top 10 Things You Should Know About COVID-19 Vaccines

1 VACCINE MAKERS ARE IMMUNE FROM LIABILITY. Vaccine manufacturers have no incentive to ensure their vaccines are as safe as possible. With legal immunity established in 1986 with the National Childhood Vaccine Injury Act and reinforced by the 2005 PREP Act, vaccine makers cannot be sued even if they are shown to be grossly negligent.

<https://www.congress.gov/bill/99th-congress/house-bill/5546>

<https://www.phe.gov/Preparedness/legal/prepact/Pages/default.aspx>



2 VACCINE COMPANIES HAVE LONG RAP SHEETS. Vaccine makers have paid out tens of billions of dollars for crimes, including fraud and violating the False Claims Act, when they knew products would cause injuries and death. (Think Vioxx, opioids and more.) If they knowingly put harmful products on the market even when they *can* be sued, why should we trust them to make safe vaccines when they have no liability and rushed clinical trials?

<https://violationtracker.goodjobsfirst.org/industry/pharmaceuticals>

3 PREVIOUS ATTEMPTS TO MAKE SIMILAR VACCINES HAVE FAILED. In one study, vaccinated infants got much sicker than the unvaccinated infants when exposed to the respiratory syncytial virus (RSV) naturally, with 80% of the vaccinated infants requiring hospitalization. Two died. In subsequent studies, vaccinated animals became very sick when they later became infected with the actual virus. Many died. This phenomenon is called Antibody Dependent Enhancement (ADE).

<https://www.nature.com/articles/s41579-020-00462-y#Sec11>

4 COVID VACCINES HAVE NO LONG-TERM SAFETY TESTING. There is no way to determine what these experimental vaccines will do to humans in the medium- to long-term. Not all vaccine injuries manifest immediately. Additionally, given that all COVID vaccines currently available in the U.S. have Emergency Use Authorization status only, people cannot be subject to mandates under federal and international law.

<https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization>

5 SERIOUS ADVERSE VACCINE REACTIONS ARE REAL AND ARE UNDERREPORTED. As of 9/24/21, over three quarters of a million (752,803) adverse events following COVID vaccines had been reported to the national Vaccine Adverse Event Reporting System (VAERS), including 15,937 deaths. According to a government-funded study at Harvard, less than 1% of all vaccine adverse reactions are actually submitted to VAERS.

<https://digital.ahrq.gov/sites/default/files/docs/publication/r18hs017045-lazarus-final-report-2011.pdf>

<https://www.medalerts.org/vaersdb/findfield.php?TABLE=ON&GROUP1=CAT&EVENTS=ON&VAX=COVID19>

<https://www.medalerts.org/vaersdb/findfield.php?TABLE=ON&GROUP1=AGE&EVENTS=ON&VAX=COVID19&DIED=Yes>

6 COVID VACCINES DO NOT STOP TRANSMISSION. The clinical trial study designs for COVID vaccines did not address transmission but merely assessed reduced symptoms, as explained in the materials manufacturers submitted to the FDA to obtain Emergency Use Authorization.

<https://www.fda.gov/media/144245/download>

<https://www.fda.gov/media/144434/download>

<https://www.fda.gov/media/146217/download>

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YOUNG ADULTS HAVE UNDER .01% RISK OF DEATH FROM COVID-19.

According to the CDC, COVID overall has a 99.74% survival rate. Among young people, that number is even higher. For people aged 0-19, the survival rate is 99.997%, and for those in their 20s, it is 99.986%. Consider this low risk from COVID when deciding whether to take an experimental vaccine that causes significant side effects, including death.

<https://data.cdc.gov/NCHS/Provisional-COVID-19-Death-Counts-by-Sex-Age-and-S/9bhg-hcku/data>
<https://www.medrxiv.org/content/10.1101/2021.07.08.21260210v1.full.pdf>



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GOVERNMENT EMPLOYEES OWN PATENTS ON THE MODERNA VACCINE.

The Bayh-Dole Act allows government employees to file patents on their research, even though they have government salaries. Moderna's research and development partner is the National Institute of Allergy and Infectious Diseases (NIAID), directed by Dr. Anthony Fauci. Moderna benefited from \$2.5 billion in federal government funding when developing its vaccine technology, and the company shares joint ownership of vaccine patents with NIAID scientists, prompting Public Citizen to rename Moderna's COVID vaccine "the NIH vaccine." This is a troubling conflict of interest.

<https://www.documentcloud.org/documents/6935295-NIH-Moderna-Confidential-Agreements.html>
<https://www.statnews.com/pharma/2020/08/28/moderna-covid19-vaccine-coronavirus-patents-darpa/>

Dr. Fauci provided U.S. taxpayer funding for illegal gain-of-function research conducted in China. "Gain-of-function" research makes viruses more virulent and transmissible, and promotes research on vaccines against the lab-created viruses. When the President imposed a moratorium on this research in December 2014 in the U.S., Dr. Fauci's NIAID outsourced its coronavirus bat research to Wuhan, China, spending \$7.4 million in six years. Members of Congress have called for an investigation into the Wuhan Institute of Virology's role in the SARS-CoV-2 epidemic.

<https://www.dailysignal.com/2021/04/06/fauci-must-explain-why-oversight-bypassed-for-funding-to-wuhan-lab-congressman-says/>

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COVID VACCINES CONTAIN PROBLEMATIC INGREDIENTS.

Both mRNA vaccines (Pfizer's and Moderna's) contain polyethylene glycol (PEG), and J&J's vaccine contains polysorbate 80—structurally similar ingredients associated with hypersensitivity reactions and anaphylaxis. Although the unlicensed mRNA vaccines are the first in widespread use to feature PEG, there are a number of approved vaccines that include polysorbate 80—all of which document anaphylaxis in their package inserts.

<https://childrenshealthdefense.org/defender/pfizer-covid-vaccine-reaction-fda-peg/>
<https://childrenshealthdefense.org/defender/inactive-ingredients-covid-vaccines-allergic-reactions/>
<https://www.nejm.org/doi/full/10.1056/NEJMra2035343>
<https://childrenshealthdefense.org/news/read-the-fine-print-part-two-nearly-400-adverse-reactions-listed-in-vaccine-package-inserts/>

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THE J & J VACCINE CONTAINS ABORTED HUMAN FETAL CELL LINES.

The viral vector that forms the backbone of the J&J vaccine is grown in a continuous ("immortalized") human embryonic cell line (PER.C6) derived from the abortion of a healthy 18-week-old fetus, leading some Catholic leaders to describe the vaccine as "morally compromised." FDA officials have acknowledged for over two decades that such cell lines are a "major safety concern." Pfizer's COVID vaccine manufacturing process includes the use of the HEK293T human embryonic cell line.

<https://childrenshealthdefense.org/defender/media-ignores-jj-pharma-giants-checkered-past/>
https://religionnews.com/2021/03/01/new-orleans-archdiocese-urges-catholics-to-avoid-new-johnson-johnson-vaccine/?fbclid=IwAR0O-bTGv7WKyiNK5_thILH_z9PK53A-eHb3oIh3u81qeJGkCuY8_25jYw
<https://childrenshealthdefense.org/wp-content/uploads/FDA-Pink-Sheets-99.pdf>
<https://www.fda.gov/vaccines-blood-biologics/biologics-research-projects/investigating-viruses-cells-used-make-vaccines-and-evaluating-potential-threat-posed-transmission>
<https://childrenshealthdefense.org/defender/project-veritas-melissa-strickler-pfizer-covid-vaccine-fetal-cells/>