Top 10 Things You Should Know About COVID-19 Vaccines

VACCINE MAKERS ARE IMMUNE FROM LIABILITY
Vaccine manufacturers have no incentive to ensure their
vaccines are as safe as possible. Established in 1986 with the
National Childhood Vaccine Injury Act and reinforced by the
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



WACCINE 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 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

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

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 current COVID vaccines 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

SERIOUS ADVERSE VACCINE REACTIONS ARE REAL AND ARE UNDERREPORTED As of 4/1/21, 55,869 adverse events following COVID vaccines have been reported to VAERS, including 2,342 deaths. According to a government-funded study at Harvard, less than 1% of all adverse reactions to vaccines are actually submitted to the National Vaccine Adverse Events Reports System (VAERS).

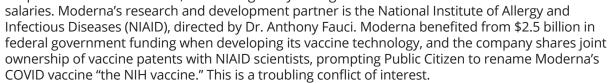
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COVID VACCINES DO NOT STOP TRANSMISSION The clinical trial study designs for COVID vaccines did not address transmission, but merely addressed reducing symptoms, as explained in the materials they submitted to the FDA to obtain Emergency Use Authorization.

https://www.fda.gov/media/144245/download https://www.fda.gov/media/14434/download https://www.fda.gov/media/146217/download YOUNG ADULTS HAVE UNDER .04% 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 18 to 29, the survival rate is 99.97%. 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.documentcloud.org/documents/6935295-NIH-Moderna-Confidential-Agreements.html https://www.statnews.com/pharmalot/2020/08/28/moderna-covid19-vaccine-coronavirus-patents-darpa/

Additionally, 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 them. When the U.S. President put 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 SARS-CoV-2 epidemic.

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

mRNA 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/NEJMra2035343https://childrenshealthdefense.org/defender/inactive-ingredients-covid-vaccines-allergic-reactions/

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."

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=lwAR0O-bTGv7WKyiNK5_thILH_z9PK53A-eHb3olh3u81qeJGkCuY8_25jYwhttps://childrenshealthdefense.org/wp-content/uploads/FDA-Pink-Sheets-99.pdfhttps://www.fda.gov/vaccines-blood-biologics/biologics-research-projects/investigating-viruses-cells-used-make-vaccines-and-evaluating-potential-threat-posed-transmission



For more information on COVID-19, go to CHD's website, www.childrenshealthdefense.org where we have reliable, up-to-date, science-based information on COVID-19 and other issues affecting health.