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Intended Consequences: mRNA Vaccines were Designed to Cause Severe Disease and Be Resistant to Antibodies

Karen Kingston

11-13 minutes

Data published by spike protein inventors, Barney Graham & Jason

McLellan, COVID-19 mRNA vaccines were purposefully designed to not protect against SARS-CoV-2 or variants and cause disease

Barney Graham of the NIAID and Jason McClellan of the University of Texas are the inventors of the S-2P spike proteins produced by the COVID-19 mRNA vaccines.

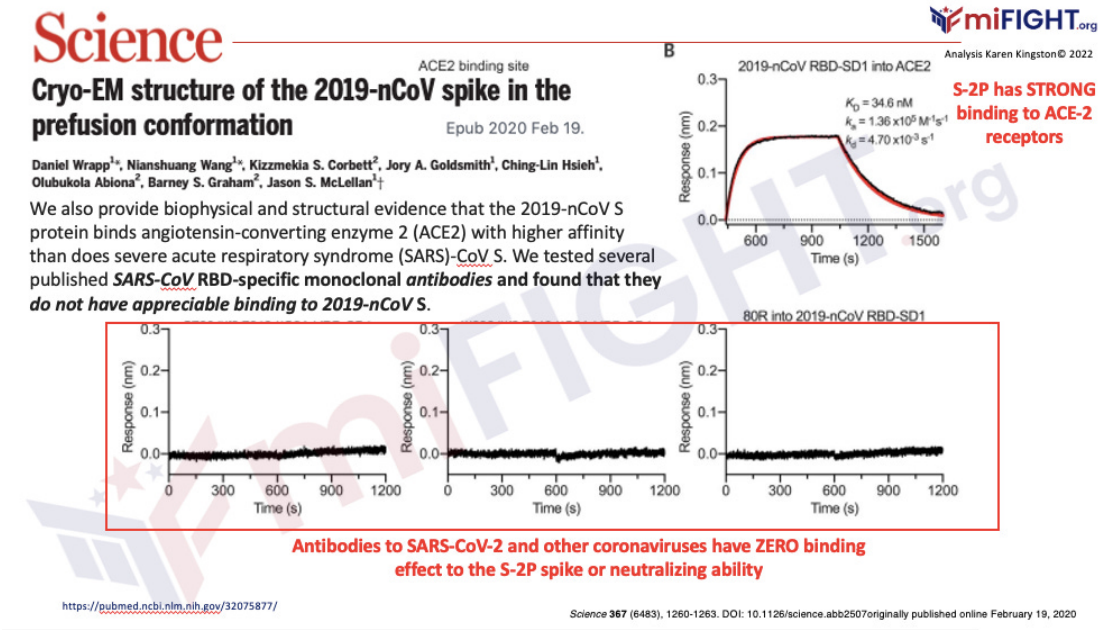
Barney Graham, NIAID/NIH

Jason McLellan, Univ of Texas

In a February 19, 2020 article in *Science*, authored by the inventors of coronavirus S-2P spike proteins, **Barney Graham and Jason McClellan**, the authors **state that the S-2P 'spike protein' has *stronger binding affinity to the ACE-2 receptors* (in the hearts, lungs, kidneys, and endothelial cell line of blood vessels) *than the original SARS-CoV-2 (S)* spike protein.**

Graham and McLellan also tested synthetically recreated antibodies for coronaviruses (SARS-CoV-2) against the S-2P spike proteins. Their research showed that ***none of the antibodies***

for coronaviruses bound to the new trimeric two-proline spike (S-2P) proteins and ***no coronavirus antibodies were able to neutralize it.***



This scientific evidence (authored by the inventors of the spike proteins) **confirms that the *COVID-19 mRNA vaccines*;**

- ***do not produce antibodies to SARS-CoV-2*, and**

- ***are resistant to the antibodies for SARS-CoV-2*, and therefore -**

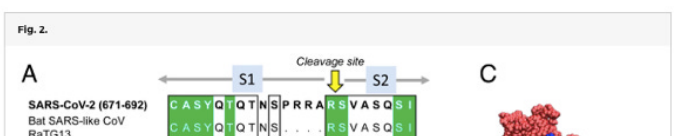
  It is scientifically and clinically *IMPOSSIBLE* for Pfizer's FDA-approved mRNA vaccines *TO PROVIDE ANY PROTECTION AGAINST SARS-CoV-2 infection, or any coronavirus, including 'VARIANTS'.*

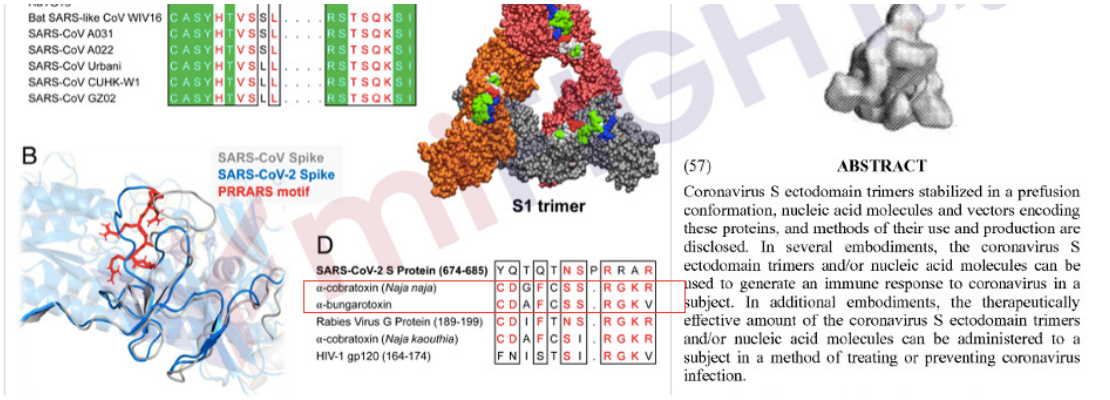
  This is why Dr. Fauci discouraged people to test for SARS-CoV-2 antibodies, vaccinated or not. The COVID-19 'public health' vaccine campaign was never about 'preventing SARS-CoV-2 infection', it was about getting a needle in every arm of every American; from adult seniors to

teenage adults and even pregnant women and six-month old babies, in order to force their bodies to produce the pathogenic spike proteins, *not* SARS-CoV-2 antibodies.

The strong binding of the S-2P spike proteins to the ACE-2 receptors (***produced by the mRNA vaccines***), is further scientific proof that the *COVID-19 mRNA vaccines were designed to cause disease (pathogenic)*.

Per the inventors, the S-2P two-proline trimeric spike is different and more pathogenic than the (S) wild type spike produced by the SARS-CoV-2 virus.





Perhaps one can logically conclude that the FDA authorized and approved Pfizer's mRNA vaccines because the mRNA injections elicited a 'robust immune response.' But the FDA had no evidence that the 'robust immune response' provides any clinical benefit in reducing the risk of SAR-CoV-2 infection, severe disease, hospitalization, or death. But don't take my word for it, take the FDA's.

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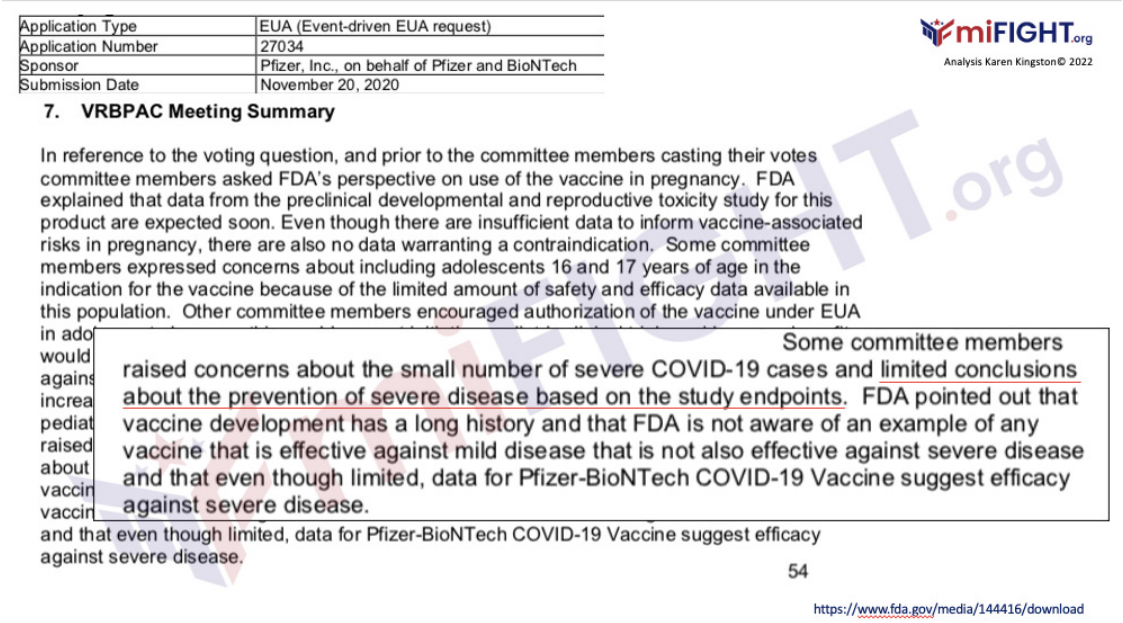
During the FDA's December 11, 2020, emergency use authorization meeting for the Pfizer COVID-19 mRNA vaccines, **FDA committee members *pointed out*** that the ***mRNA vaccines appear to provide no clinical benefit, specifically regarding severe disease.***

"Some committee members raised concerns about the small number of severe COVID-19 cases and **limited conclusions about the prevention of severe disease** based on the study endpoints. **FDA pointed out that vaccine development has a long**

history and that FDA is not aware of an example of any vaccine that is effective against mild disease that is not also effective against severe disease and that even though limited, data for Pfizer-BioNTech COVID-19 Vaccine suggest efficacy against severe disease." - FDA EUA Review Committee for Pfizer EUA COVID-19 mRNA vaccines, Dec 11, 2020



Application Type	EUA (Event-driven EUA request)
Application Number	27034
Sponsor	Pfizer, Inc., on behalf of Pfizer and BioNTech
Submission Date	November 20, 2020

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7. VRBPAC Meeting Summary

In reference to the voting question, and prior to the committee members casting their votes committee members asked FDA's perspective on use of the vaccine in pregnancy. FDA explained that data from the preclinical developmental and reproductive toxicity study for this product are expected soon. Even though there are insufficient data to inform vaccine-associated risks in pregnancy, there are also no data warranting a contraindication. Some committee members expressed concerns about including adolescents 16 and 17 years of age in the indication for the vaccine because of the limited amount of safety and efficacy data available in this population. Other committee members encouraged authorization of the vaccine under EUA in ado
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54

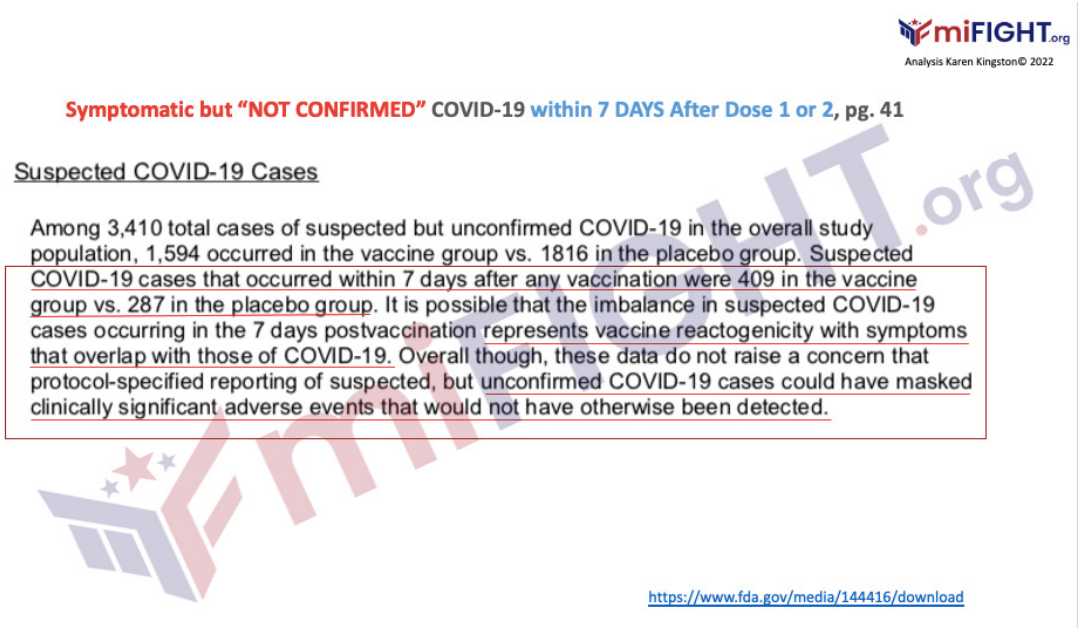
https://www.fda.gov/media/144416/download

Per the FDA's own words, in the history of the FDA, the *FDA has never approved a vaccine that does not protect against severe disease* (until the agency approved the COVID-19 mRNA vaccines).

This is where it gets interesting though (and by interesting, I mean 'evidence of criminal intent' interesting). Per the data discussed during that same FDA meeting, not only do ***Pfizer's COVID-19 mRNA vaccines NOT prevent severe disease***, ***Pfizer's mRNA vaccines cause SEVERE COVID-19 within 7 days of being injected.***

Page 41 of PFIZER's EUA submission,

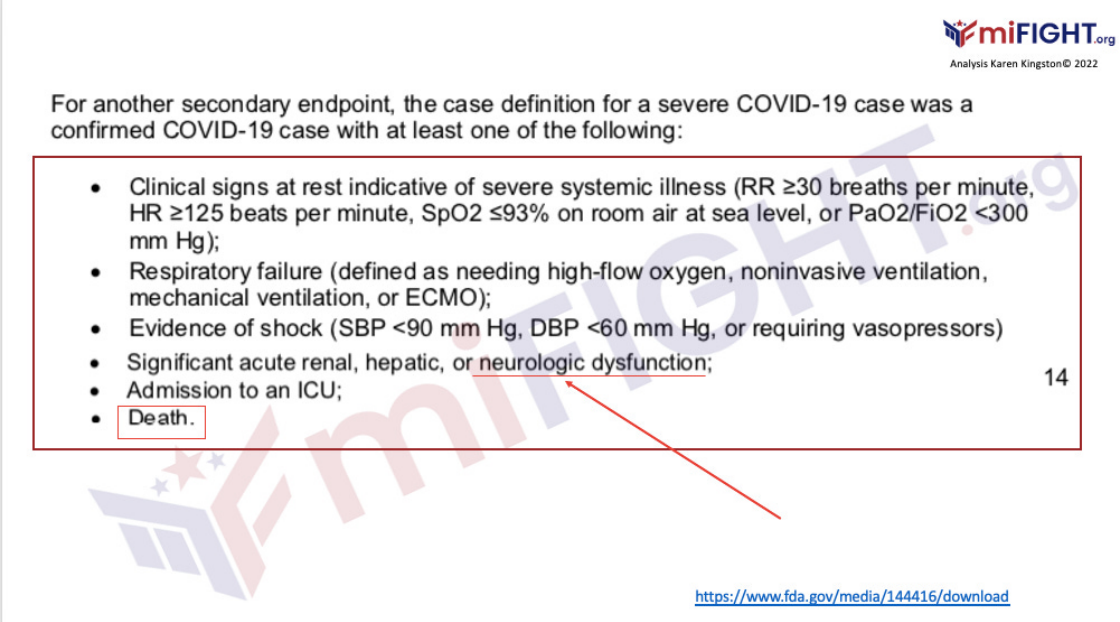
states that there were 409 patients who had COVID-19 symptoms within 7 days of getting their first (1st) or second (2nd) PFIZER shot, ***BUT these patients did not have a positive PCR-test*** for SARS-CoV-2.



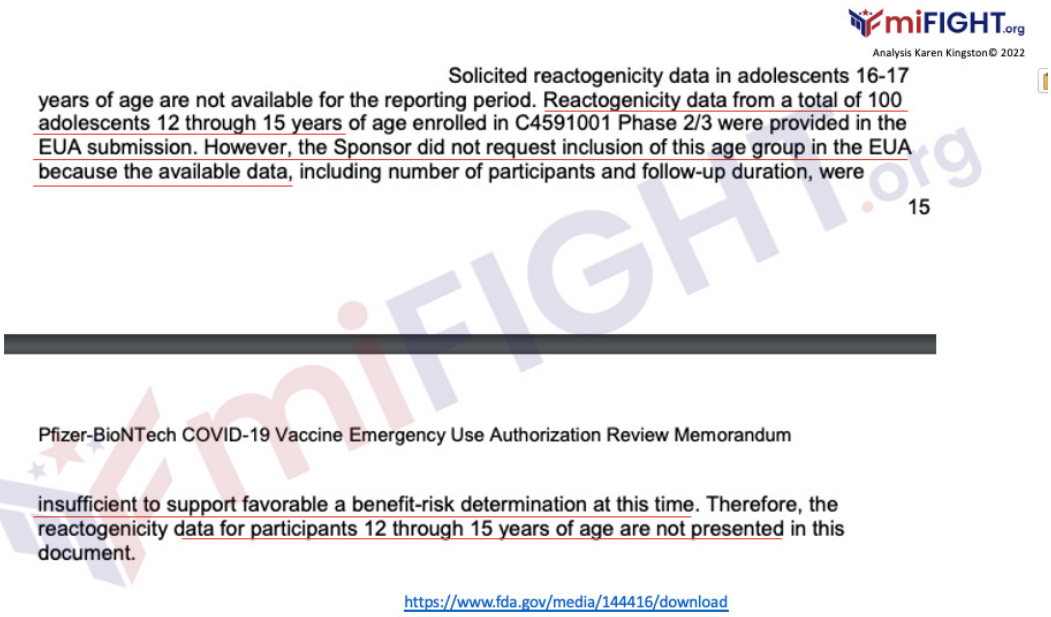
Per PFIZER's own document, "unconfirmed COVID-19 cases could have masked clinically significant adverse events that

would have otherwise been detected."

Per PFIZER's EUA submission, clinically significant or severe COVID-19 cases were defined in the Phase 3 Study as kidney, liver, or ***neurological dysfunction****, low oxygen levels, respiratory failure, mechanical ventilation, systemic shock, admission to the intensive care unit (ICU), or ***death***.



Pages 15-16 of PFIZER's EUA submission, states **that the incidence of expected *serious adverse events* (reactogenicity) in 100 children aged *12-15 years*, was *so incriminating* that Pfizer and the FDA agreed to not release (*cover-up*) the data.**



"**Reactogenicity data** (including severe adverse events from the

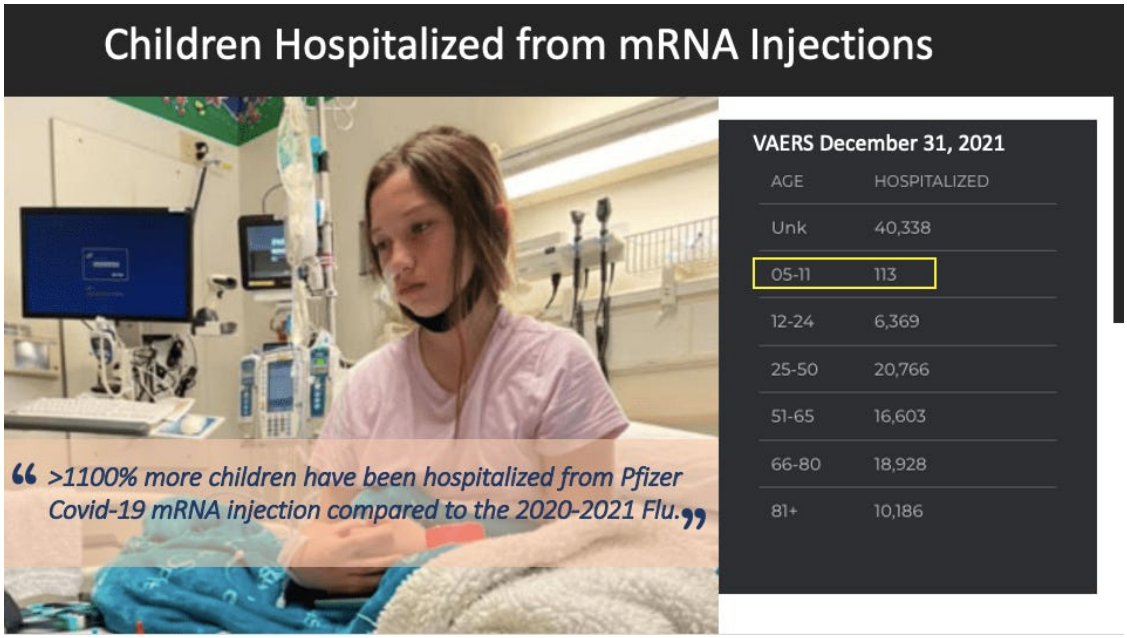
October 22, 2020 FDA/Industry meeting) from a total of **100 adolescents 12 through 15 years of age**…were provided in the EUA submission. **However, the Sponsor** (**Pfizer**) d**id not request inclusion of this age group** in the EUA because the available data…were **insufficient to support favorable a benefit-risk determination** at this time. Therefore, **the reactogenicity data for participants (children) 12 through 15 years of age are not presented in this document**." - FDA EUA Review Committee for Pfizer EUA COVID-19 mRNA Vaccines, 12/11/20

The FDA knew Pfizer's mRNA

vaccines would permanently injure and harm children. Immediately after this meeting, the FDA should have stopped all trials, especially the trial for children. Allowing the pediatric trials to move forward was a criminal act by the FDA resulting in the unnecessary battery (bodily injury) of children.

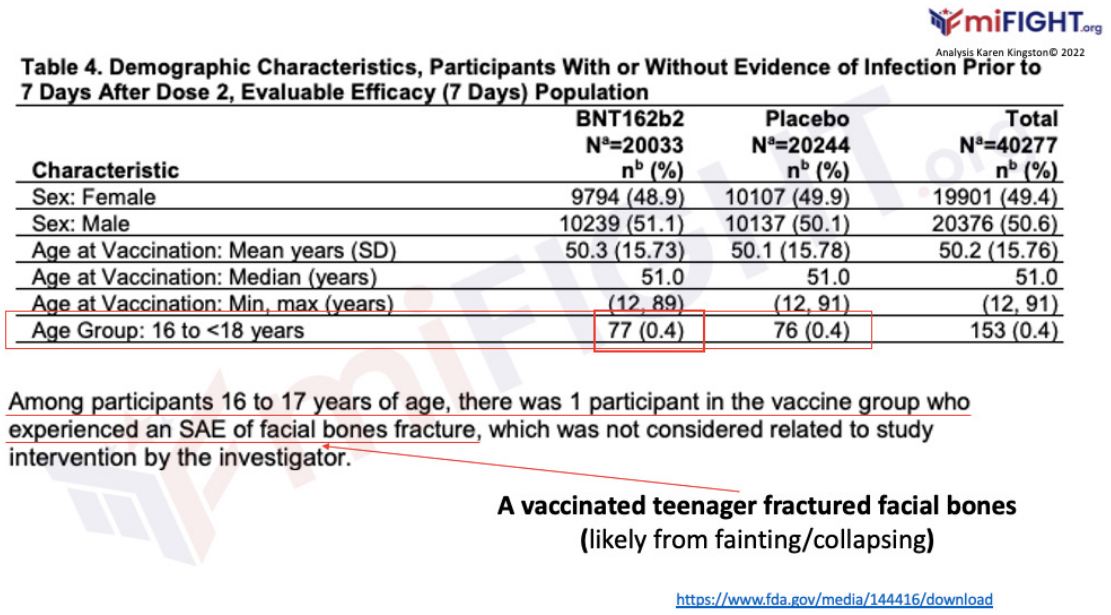
> Maddie de Garay was a healthy, happy 12-year old girl prior to being a study participant in Pfizer's 12-15 year old Phase 3 trial. Within 24 hours of receiving her 2nd mRNA Pfizer vaccine, Maddie suffered severe crippling nerve pain, memory loss, vision disturbances, and irregular menstrual cycles. Maddie is suffers

particle paralysis and requires a wheel chair and NG tube. Her serious adverse events were recorded as a stomach ache in the Pfizer trial.

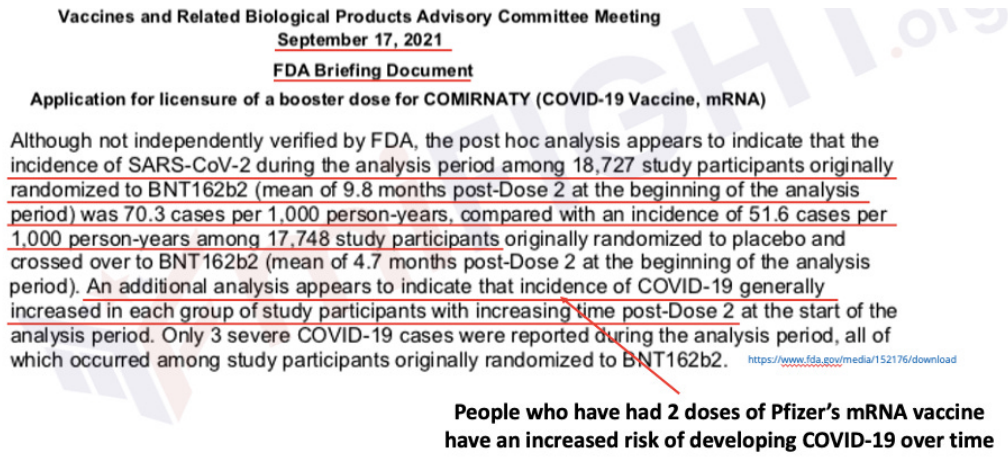


During the December 11, 2020 FDA meeting, data was disclosed of a teenager from the 16-17 year old group (n=77) who was vaccinated with the Pfizer mRNA vaccine. The vaccinated teenager fractured his facial bone

structures, likely from the fainting caused by the mRNA vaccines.



Per the September 17, 2021, FDA committee meeting, the FDA has full knowledge that two (2) injections of Pfizer's mRNA vaccines increases a person risk for developing COVID-19 over time versus a person who remains unvaccinated.



Vaccines and Related Biological Products Advisory Committee Meeting
September 17, 2021
FDA Briefing Document
Application for licensure of a booster dose for COMIRNATY (COVID-19 Vaccine, mRNA)

Although not independently verified by FDA, the post hoc analysis appears to indicate that the incidence of SARS-CoV-2 during the analysis period among 18,727 study participants originally randomized to BNT162b2 (mean of 9.8 months post-Dose 2 at the beginning of the analysis period) was 70.3 cases per 1,000 person-years, compared with an incidence of 51.6 cases per 1,000 person-years among 17,748 study participants originally randomized to placebo and crossed over to BNT162b2 (mean of 4.7 months post-Dose 2 at the beginning of the analysis period). An additional analysis appears to indicate that incidence of COVID-19 generally increased in each group of study participants with increasing time post-Dose 2 at the start of the analysis period. Only 3 severe COVID-19 cases were reported during the analysis period, all of which occurred among study participants originally randomized to BNT162b2. https://www.fda.gov/media/152176/download

People who have had 2 doses of Pfizer's mRNA vaccine have an increased risk of developing COVID-19 over time

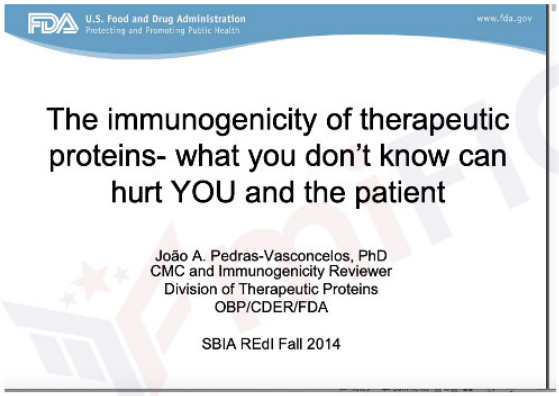
In other words, Pfizer's COVID-19 mRNA vaccines cause COVID-19.

So there you have it. The FDA authorized and approved the COVID-19 mRNA vaccines knowing the injections would cause injury, disease, disabilities, and death at a higher rate than the 'SARS-CoV-2 virus' (much, much higher rate).

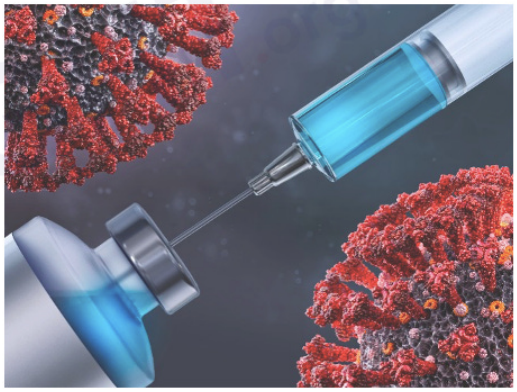
The FDA mRNA vaccine committee is also well-versed in the immunogenicity

and lethality of 'mRNA vaccines' based on over a decade of analysis by the agency on protein-based therapies.

"A major problem with protein-based therapeutics is their immunogenicity, that is, their tendency to trigger an unwanted immune response against themselves . . . Such antibodies can cause complications that can be life threatening."



https://www.fda.gov/files/drugs/published/The-immunogenicity-of-therapeutic-proteins--what-you-don%E2%80%99t-know-can-hurt-YOU-and-the-patient--Fall-2014.pdf



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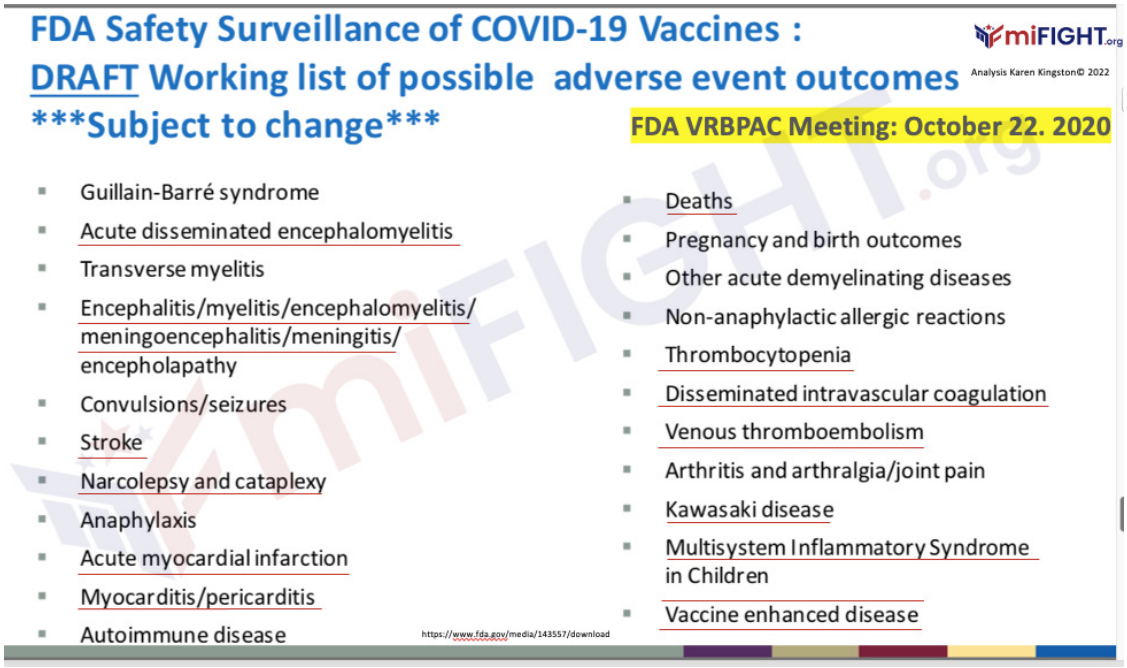
https://www.fda.gov/vaccines-blood-biologics/biologics-research-projects/immunogenicity-protein-based-therapeutics#:~:text=A%20major%20problem%20with%20protein,or%20eliminate%20their%20therapeutic%20effects.

Lastly, on October 22, 2020, (prior to any mRNA vaccine authorizations) the FDA had a meeting with industry, where a list of the ***expected severe diseases and outcomes caused by the mRNA vaccines was disclosed***. **When the FDA** authorized and *then*

approved **the COVID-19 mRNA vaccines, the agency knew that the mRNA vaccines *do not prevent* severe disease, but rather *cause* severe disease and death.**



The above list of diseases and outcomes are not side effects. The above list of diseases and outcomes, including death, are the *INTENDED CONSEQUENCES* of the

COVID-19 mRNA vaccines.

Based on the body of evidence submitted to the FDA by Pfizer, the FDA is fully knowledgeable that Pfizer's COVID-19 mRNA vaccines *cause* mild-moderate disease, severe disease, injuries, disabilities, and even death in adults and children. Th FDA is fully aware that the mRNA vaccines put American adults and children at significant risk for hospitalizations **and death while the '**SARS-CoV-2 virus**' does not.**

The evidence in this article is primarily from Pfizer's documents submitted to the FDA. The other evidence is from the FDA and a peer-reviewed journal

authored by the inventors of the 2-SP spike protein produced by COVID-19 mRNA vaccines. Each statement made can be verified with the link provided to a government website (either NIH.gov or FDA.gov) where you will find the documents that are cited.

This article is based on evidence that can stand up in a court of law. Please share with local and national government officials, school boards, churches, and your health care providers. Please share with your friends and family. Our loved ones, our communities, and our country cannot be healed if they are unaware of how they have been harmed.

[Share The Kingston Report]

The FDA's approval of COVID-19 mRNA vaccines is not only fraudulent, it is conspiracy to commit premeditated battery and murder of Americans.

The Kingston Report. TRUTH WINS.

Psalm 140: 1-3

"Rescue me, Lord, from evildoers; protect me from the violent who devise evil plans in their hearts and stir up war every day. They make their tongues as sharp as a serpent's; the poison of vipers is on their lips."

[Take Down COVID-19]

The uncomfortable truth is that we all have been lied too and deeply

betrayed by leaders we trust, and some we even adored. All offices of power across our nation; from our President and Federal Healthcare agencies to our local governors, mayors, city counsel members, and even our health care service providers, employers, and school boards members. If you're questioning on how to know if a government, public official or even if your employer or school is an ally of the American people of our children, there is a simple challenge to give them;

Demand Local Officials Take Down COVID-19.

Call for governors, mayors, school

boards, colleges and universities, health care officials, health care centers, businesses, *and churches* to;

- **Make a public declaration that COVID-19 mRNA vaccines cause disease and death and must be banned and recalled immediately**
- Immediately STOP ALL COVID-19 testing, treatments and mRNA vaccines
- REJECT and STOP ALL FUNDING for all COVID-19 programs
- *CALL FOR GOVERNORS to CRIMINALIZE the promotion and administration of mRNA VACCINES

*Governors have the power to reject the HHS declaration that SARS-CoV-2

is a threat to public health and national security and to criminalize the use of all EUA designated COVID-19 products, tests, and mRNA vaccines. Demand that they do.

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