WHAT YOUR DR MAY NOT KNOW

The Covid 19 (vaccines) are different from all other vaccines previously on the market.

All previous vaccines were either a dead virus, a part of a dead virus, or a weaken virus.

The Covid 19 (vaccines are likely mRNA viral based gene therapy. Pfizer phase 1 studies were conducted as if it was a viral gene based therapy.

All US Covid 19 vaccines are mRNA viral gene based therapy (vaccine) and have been released due to Emergency Authorization Act

Under the Emergency Use Authorization act (the ~~Popper~~ Prep and the Cures act)

manufactures can do the following:

1) Waive all good manufacturing practices IE no inspections of what is in the vials

   Under the EAU the manufacturers are NOT obligated to tell you what is in it

   UNDER THE CURES ACT SECTION 3024 Dec 13, 2016

   DO NOT HAVE TO PROVIDE A WAIVER OF CONSENT IF IT IS CONTRARY TO THE INTEREST OF SUCH HUMAN BEINGS

   (SO ITS LEGAL FOR THE GOVERNMENT TO INJECT YOU WITH SOMETHING THAT CAN KILL YOU

   LOOK IT UP

2) A pharmaceutical company can stockpile product before approval (never seen this before)
3) Manufactures **Do NOT have to prove efficacy** all they have to say is that it **COULD** be effective
4) Manufacturers **do not have to do animal trials, they can go straight to humans (a violation of the Nuremberg code.**
5) All damages created by the vaccines to patients are the patient's responsibility, manufactures and employers have NO liability

The Emergency Use Authorization Act of mRna viral based gene therapy (vaccines) have not been evaluated like other viral gene therapies.

Normally viral based gene therapies undergo testing for the following:

1) Progeny (antibody enhancement) (where the body can attack itself causing autoimmune diseases). That is why the animal studies are so critical because it takes about 2 years in animal studies to prove that this cannot happen) The FDA originally came out and said the soonest they would consider approval is 2023

2) Shedding the transmission of the spike protein to the unvaccinated (these trials typically require male subjects that are in contact via skin contact or breath contact to make sure that it doesn't pass to unvaccinated people and pregnant mothers. This was followed in Pfizer phase 1 study only

Therefore none of the vaccines on the US market have this data and are not obligated to prove anything until the drug is fully approved

Comirnity, If it is FDA fully approved, should have a full package insert that describes what exactly what is in the vial. Do not let someone tell you it is the same thing as the Pfizer/Biontec. If the vial says Comirnity there may be some liability for damages consult a lawyer before taking the risk.

If it isn't fully approved and remains under Emergency use authorization then patient takes on all injury costs.

CITIZEN FREE PRESS

Fauci finally mentions Vaccine risk for ADE...

Posted by Kane on September 3, 2021 5:03 am

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