FDA Director Peter Marks and the Ever-Shifting COVID Vaccine Narrative

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by **Robert F, Kennedy, Jr.**, Chairman, *Children's Health Defense*
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After suckering us into ruinous lockdown awaiting rescue by vaccine, the Pharma grifters are frantically dialing back the expectations they inflated. During an FDA teleconference on July 8, CBER's Director Peter Marks said FDA is now willing to license COVID vaccines with a dismal 50%–and as low as 30%–efficacy, a humiliating retreat from the Gates/Fauci promise of a vaccine they intend to give to seven billion people in order for society to get back to "normal". Equally deflating, NIH's Tony Fauci conceded that vaccine immunity may only last a few months …

https://youtu.be/HFP1yaQiR3A

… and joined Gates hinting that the vaccine may not even prevent transmission:

https://youtu.be/CjRTIcf4Tk8

Astra Zeneca is making two billion doses of its "Oxford" jab (Gates is heavily invested) despite proof that monkeys vaccinated transmit COVID. Paul Offit told CBS that the jab may not stop transmission and may only weaken not prevent symptoms.

https://youtu.be/QpiaQA2ifZs

Marks conceded that a vaccine with 50% efficacy will not stop the virus. “We’re going to need a vaccine that’s probably in the order of 70% effective and 70%, at least, of the population is going to need to take it.”

Marks’s justified FDA’s willingness to license jabs with a pathetic 30-50% efficacy citing industry convenience. “Can we show you some calculation of how we got there? No,” he confessed, adding “If you go much lower than 50% then the lower bounds of things start to get to a place where vaccines may have very little efficacy…On the other hand, if we held that number at 70% to 80% … we may not have a vaccine until there’s herd immunity that’s occurred naturally.”

Fear that the wild virus might vanish before a vaccine is ready for human trials has prompted FDA to scrap its traditional ethical revulsion for “challenge trials” in which drug companies deliberately expose humans to wild viruses. FDA has issued new protocols for “controlled human infection models” wherein drug companies intentionally expose vaccinated volunteers to a pathogen. FDA says such trials could be necessary if COVID becomes so rare that “it is no longer possible to demonstrate vaccine effectiveness by way of conducting clinical disease endpoint efficacy studies.” When will the press call fraud on these quacks?

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Marks said that the thing that "scares me more than anything else is that a third or half of Americans are hesitant about taking a vaccine [for COVID-19]."

Marks stressed that FDA is determined to restore public faith in vaccines by insuring that COVID vaccines are safe and high quality.Recent tests by Corvelva have found every vaccine tested to be loaded with extraneous contaminants not listed as ingredients including particles of mercury, steel tungsten, lead, organic chemicals and pesticides and human and animal DNA & pharmaceutical drugs including amphetamines. Corvelva speculated that these dangerous substances might be manufacturing debris. 👉

Marks promised improved quality control for COVID vaccines. "For any of these vaccines targeting SARS-CoV-2, important things for us from the standpoint of our guidance... will be things like consistency of manufacturing, and the need for manufacturing processes and controls that have appropriate steps in them, the need to have facilities inspected to produce vaccines under good manufacturing practices, that's important because we really do need to make sure that these are going to be high quality products that when we say they're safe, they really are."