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The real data behind the new COVID vaccines the White House is pushing

By Marty Makary and Tracy Beth Høeg

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I I nurse quilty in \$1.5M COVID vay scam blames 'government mandates'

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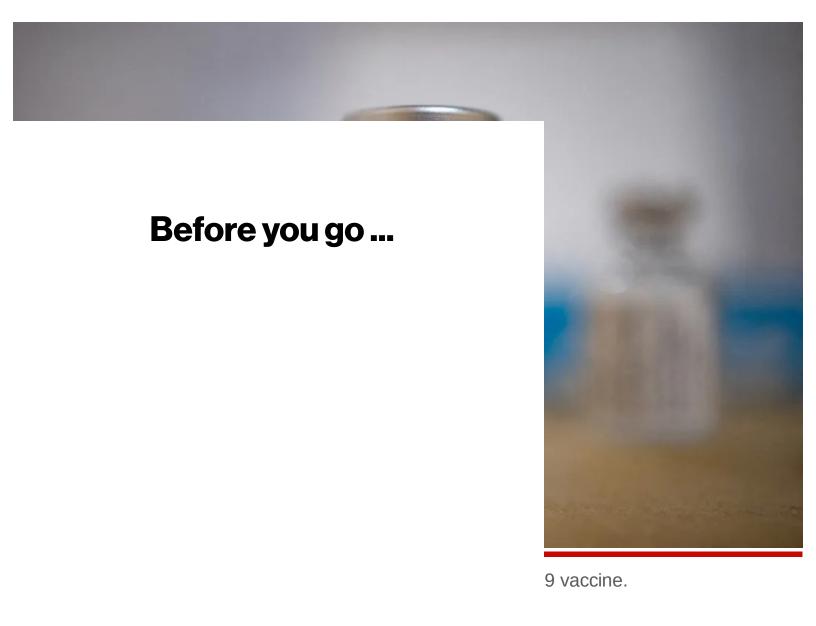
stration cleared it without s are not universally of and Prevention is?

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D coordinator Dr. Ashish pported claims the new the likelihood you will None of those claims has a shred of scientific support.

In fact, if the manufacturers said that, they could be fined for making false marketing claims beyond an FDA-approved indication.

The questions surrounding Moderna's new COVID vaccine approved this week are still looming.



as zero efficacy data have data about

The FDA, or Moderna (frankly, it's hard to tell the difference sometimes), should disclose what happened to the patient who took the new vaccine and had a complication that required medical attention.

The public has a right to know.

The last time the Biden administration approved and recommended a novel COVID bivalent booster, last fall, with no human-outcomes data, it was an epic fail.

Only 17% of Americans took it (and some of those were forced to do so by

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it infects the body could

oster last year, the Biden .71 million doses —

orders for the pediatric on doses at \$1.3 billion to four times as many There clearly seems to be a special push this time to give it to children — the same group European regulators are not supporting.

In fact, the original Moderna vaccine was banned in parts of Europe for people under age 30.

European doctors are not alone.

Dr. Paul Offit, a vaccine-mandate supporter and FDA adviser from the University of Pennsylvania, told The Atlantic this week that he's not going to take the new COVID vaccine.

spite being 72 years



ently confessed, "Yes, he ."

tant news

America is tired of political apologists as medical experts. They want the truth.

Offit is at least more honest than most experts who put their heads in the sand and parroted whatever public health officials said.

Pfizer made \$100 billion during the pandemic. It can afford to fund a randomized trial to demonstrate to the American people the new booster is effective.

That's the scientific process.

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It's time for the FDA to resume its role as a regulator and not the marketing department for Pfizer and Moderna.

e severity of COVID all the more reason a

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cluding severe and lifeolus-year safety record whereas COVID vaccines have been associated with a serious adverse event rate of one in 5,000 doses, according to a German study by the Paul-Ehrlich-Institut.

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urnal Vaccine, estimated one in 556 COVID

is six to 28 times higher les, according to a 2022

ral national colleagues Jates appear to have

ncerned about what is ple booster doses can

escribed a reduced received three COVID If public health officials get their way, a healthy 5-year-old boy will get 72 COVID vaccine shots over the course of his lifetime, if he has an average lifespan, with a risk of myocarditis after each one.

Inexplicably and defying science, the CDC is saying even if a child had COVID three weeks ago, he or she should still get the new COVID shot.

Two of the FDA's best vaccine experts are gone. Dr. Marion Gruber, who was director of the FDA's vaccine office, and her deputy director, Dr. Philip Krause, both quit the agency in 2021 in protest over political pressure to authorize vaccine boosters for young people.

e agency's vaccine cozy relationship

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Public health leaders cannot afford to squander any more credibility and money on interventions with no scientific support.

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