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FDA Advisers Are Angry at Moderna for Hiding Data

Are they pretending to be independent to save themselves?

By Igor Chudov
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Region: USA
Theme: Science and Medicine

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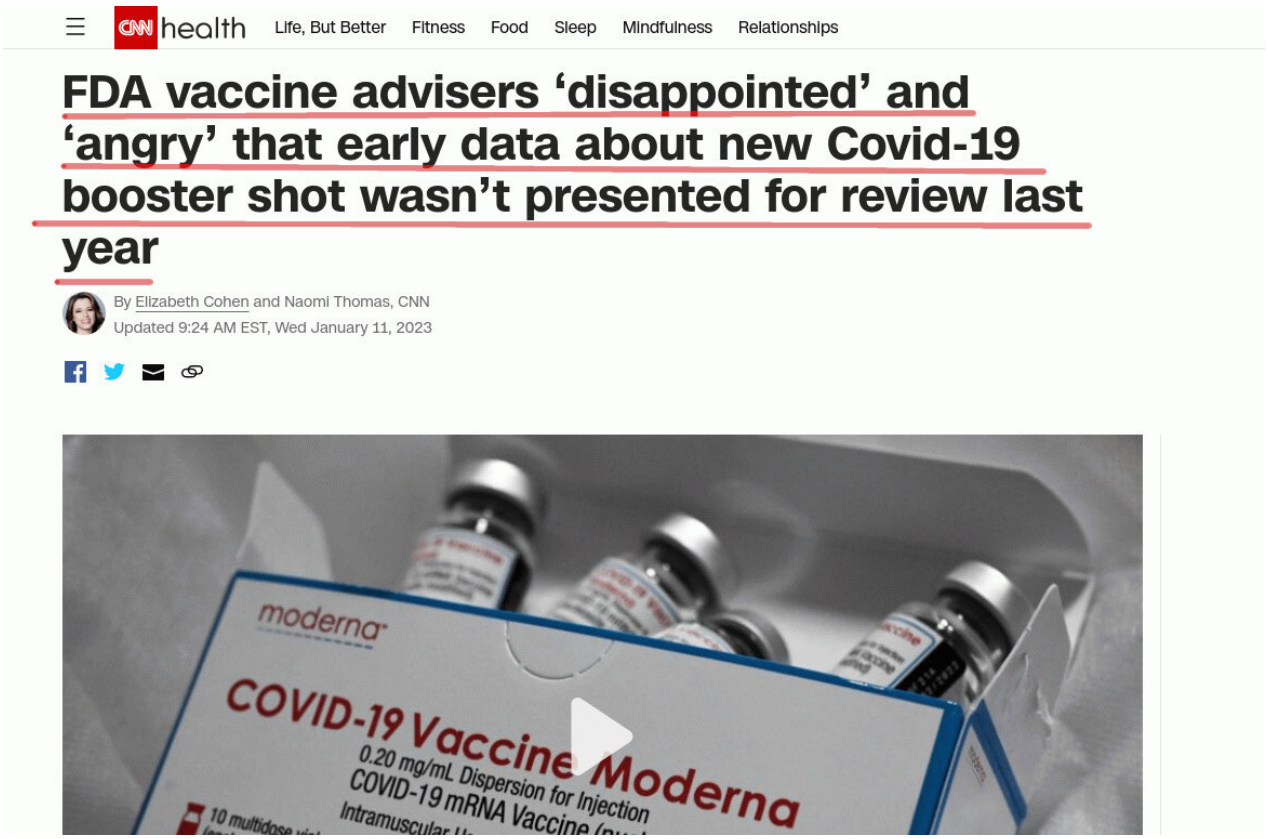
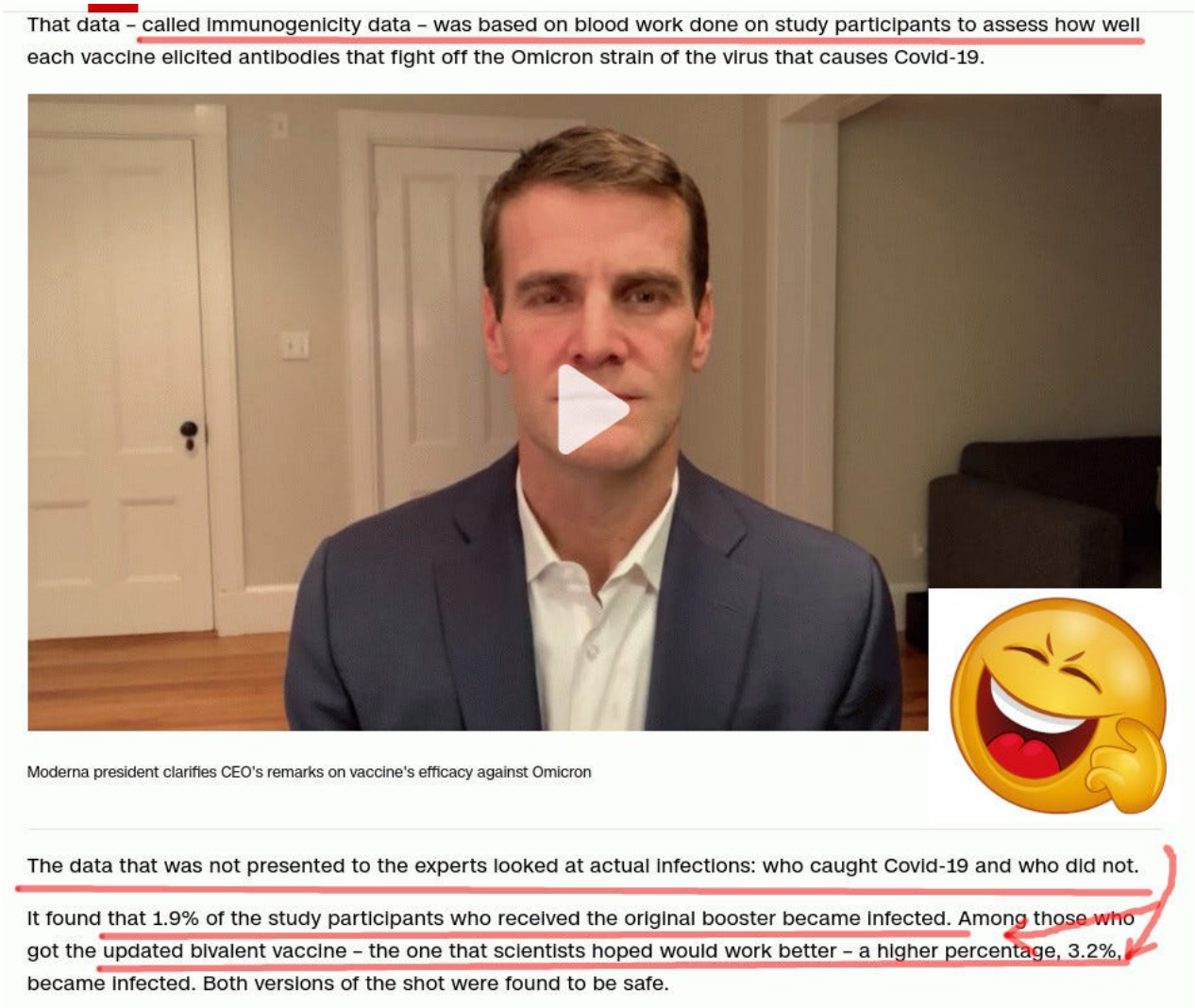
The times are changing!

The FDA's "Independent advisers" are reportedly disappointed and angry at Moderna.



We, the antivaxxers, have been disappointed and angry at Moderna for about two or three years based on mountains of data we unearthed, some of which the FDA tried to hide for 75 years. So, may we ask, what disappointed the FDA advisers so late in the game? This is

what it was:



It turns out that the FDA advisers approved Moderna booster shots based on "antibody counting," a *quack medicine* approach called *immunobridging*.

The reason immunobridging is medical quackery is pointed at by the "laughing emoji" above. Despite having "*more antibodies,*" MORE people in the *bivalent group* caught Covid compared to people in the *monovalent* (old booster) group.

Despite being there in larger numbers, the antibodies *facilitated* the infection instead of *preventing* it. Thus, reliance on antibody counts is medical quackery, as the real-life experimental data proves them useless at best.

FDA charlatans approved the Moderna bivalent Covid booster for millions of people based on antibodies in TEN MICE (who all got sick with Covid when challenged with the virus).

And now, the FDA advisers are "angry" that Moderna did not present real-life data showing that bivalent-boosted people are 68% (3.2/1.9) more likely to get Covid.

Why did they not get angry several months before?

The data that was not presented was available in plain sight!

> Despite these imperfections, the data was included in a preprint study that was posted

> online in June, again in September in an FDA document and then later that month in a top medical journal – and advisers to the FDA and the CDC said the data should have been shared with them, too.

The FDA advisers are not reading the literature and top medical journals. What do they do in their spare time?

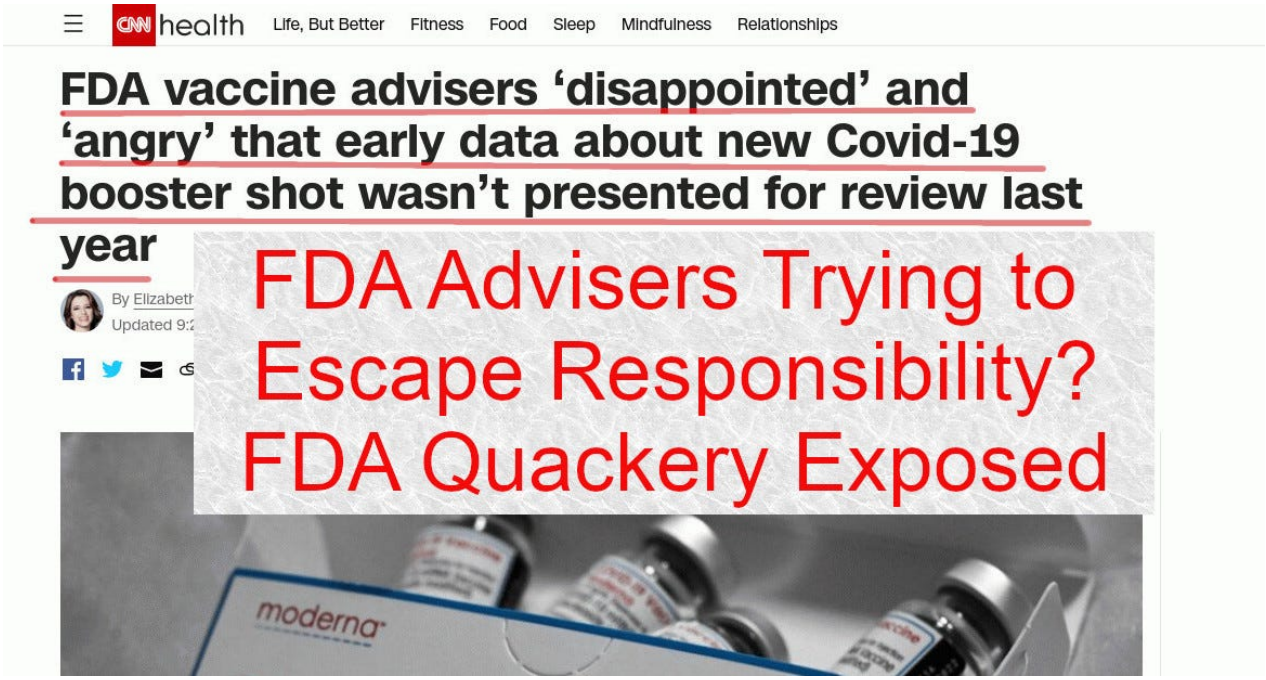
> Michael Felberbaum, an FDA spokesman, told CNN in an email that "the FDA received the preprint less than a day prior to the advisory committee meeting," and "the information was therefore not provided in an adequate timeframe for it to be included in the agency's meeting materials, and generally the FDA only discusses data at advisory committee meetings that the agency has had the opportunity to substantively review."

Hm. The FDA only discusses data at advisory committee meetings that the agency has had the opportunity to substantively review??? That's a lie. The FDA approved the "Covid vaccine for babies" just TWO DAYS after receiving the submission.

The FDA was good at rubber stamping stuff while they considered themselves invincible.

Now that the public is asking pointed questions, FDA advisers are suddenly "angry" at Moderna. Are they trying to save themselves from repercussions and future prosecution?

Should we offer them "pandemic amnesty"?



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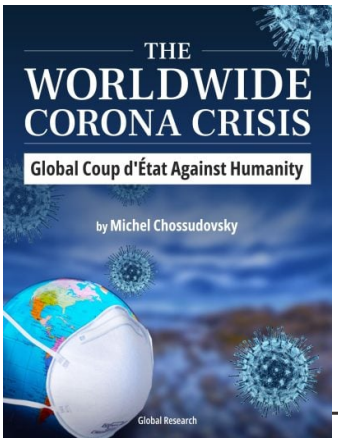

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