CERTIFIED MAIL-RETURN RECEIPT REQUESTED

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Attorney General Andrea Campbell,

The pharmaceutical industry moved quickly in 2020, in conjunction with the United States government's "Operation Warp Speed," to develop vaccines in response to the COVID-19 pandemic. Billions were paid to facilitate the development and distribution of the vaccines. Due to the seriousness of the pandemic, the Pfizer-BioNTech vaccine quickly received its initial Emergency Use Authorization on December 11, 2020. https://www.fda.gov/news-events/press-announcements/fda-takes-key-action-fight-against-covid-19-issuing-emergency-use-authorization-first-covid-19. Almost immediately, millions of people lined up to receive these injections. These same millions of people trusted that the FDA and the CDC had done their due diligence to ensure the vaccines were safe for human use. In fact, celebrities and politicians alike were engaged to encourage people to take the vaccines. Even President Biden did so.

But according to a recent study, two months after the COVID-19 vaccines rollout began, the FDA and CDC Vaccine Adverse Event Reporting System (VAERS) had received reports that myocarditis was occurring in young males at a rate that indicated it was a statistically significant event. The study, Jablonowski, K., & Hooker, B. S. (2022). Delayed Vigilance: A Comment on Myocarditis in Association with the COVID-19 Injections. *International Journal of Vaccine Theory, Practice, and Research*, *2*(2), 651.1–651.4. https://doi.org/10.56098/ijvtpr.v2i2.61 shows that a potent, statistically significant vaccine adverse event 'signal' for myocarditis in males 8 to 21 years of age was perceptible as early as February 19, 2021. Furthermore, the VAERS data already demonstrated that myocarditis was a risk after only 14.23% of the U.S. population had been administered injections.

Worse, in an October 2020 meeting, two months before approval, the "Vaccines, and Related Biological Products Advisory Committee," at a slideshow presentation, showed a slide entitled "FDA Safety Surveillance of Covid-19 Vaccines: DRAFT Working list of possible adverse event outcomes***Subject to change***" https://www.fda.gov/media/143557/download and acute myocardial infarction and myocarditis were on the list. The slide, however, was quickly glossed over and barely seen by those in attendance. https://www.youtube.com/watch?v=1XTiL9rUpkg In May 2021, the CDC finally confirmed the possible connection between the mRNA vaccines acknowledging cases of myocarditis or pericarditis in people who received a second dose, typically in patients under the age of 30 https://www.biospace.com/article/likely-link-between-

mrna-vaccines-and-rare-cases-of-heart-inflammation-cdc-panel-says/The labels for the vaccines now include these warnings.

Even now, we are learning that reports presented to the CDC's Advisory Committee on Immunization Practices on September 1, 2022, show for the 12- to 15-year-old group, there was an incidence rate of myocarditis of 150.5 per million—or about 1 in 6,600—and 137.1 per million for the 16- to the 17-year-old group—or about 1 in 7,262. Among 16–17-year-old males, the rate jumps to 188 per million following the first booster. These rates are 3-5 times higher than the CDC reported for young men last year.
https://www.cdc.gov/vaccines/acip/meetings/downloads/slides-2022-09-01/05-covid-shimabukuro-508.pdf

The above information demonstrates that the Federal Government, under the auspices of the FDA and the CDC, hid pertinent details on the side effects of the Pfizer vaccine before they issued the EUA and again within two months after issuing the EUA.
Had the FDA and the CDC released the side effects in a timely manner, the public would have had the information they needed to make an informed decision on the vaccine. But they did not, the public was left in the dark, and many people were harmed or died.

This information is still not well known to the Massachusetts public. Given the established informed consent laws in this state, the citizens are entitled to this knowledge so that they can make an informed decision as to whether they will continue to take the vaccine, especially if they are a young male. We are asking you to require the information regarding the myocarditis side effects to be attached to every public and paid advertisement for the vaccines.
Furthermore, the CDC and the FDA, having caused serious injury to many people by suppressing the vaccine side effects, may have violated several Massachusetts criminal statutes. As the Chief Law Enforcement Officer of Massachusetts, we ask you to investigate this matter further and determine what actions, if any, must be taken against the CDC and the FDA.

Sincerely,

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