THE NEW MAGA DEAL

The Unofficial Deplorables Guide to Donald Trump's 2024 Policy Platform

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CHAPTER 22

Eradicating Forced Vaccinations and Holding Mainstream Media Liable for Deadly Misreporting

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In the 1950s, President Eisenhower issued a warning to Americans about the dangers of an unjustified influence exerted by the U.S. military-industrial complex. This concept referred to an interconnected network of interests, commonly known as the "iron triangle," in which Congress passed legislation favoring defense contractors who, in turn, provided support for reelection campaigns. These contractors would then lobby Congress on behalf of the U.S. military, resulting in special treatment for themselves. Ultimately, the bureaucracy frequently received increased funding to administer federal policies.

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Recently, the COVID-19 pandemic has exposed a second and more sinister variant of the iron triangle that has remained concealed within the federal bureaucracy for the past three decades. This insidious entity takes the form of the "Big Pharma -federal health agency industrial complex," which has now come to light due to relentless Freedom of Information Act (FOIA) requests and lawsuits filed by organizations like Judicial Watch and America's First Legal.

Throughout the COVID-19 pandemic, the CDC, the NIH, and the FDA have repeatedly assured all Americans that the COVID-19 mRNA "vaccines" were highly effective, well-tested, and completely safe. In actuality, the mRNA vaccines have never been safe, effective, or cornpletely tested.

Before any new FDA approval, pharmacokinetic studies are necessary to establish product effcacy and safety in humans. Yet FOIA documents recovered by Judicial Watch show that no absorption studies, no metabolism studies, no excretion studies, and no pharmacokinetic studies were ever conducted with the Moderna experimental mRNA 1273 COVID-19 vaccine. We also now know that Pfizer never released all of its animal testing data after the FDA issued it an Emergency Use Authorization (EUA)• A Japanese government lawsuit forced the release of Pfizer's biodistribution study, which showed that the mRNA vaccines did not remain localized upon injection but instead quickly spread throughout many tissues. We now know that the vaccine mRNA can stay active in the body for days if not weeks.

We also know that repeated vaccination with mRNA boosters can lead to toxic viral spike protein damage to the human liver' heart, testes, ovaries, and spleen, with widespread damage to the

microcirculation and the deposition of the abnormal spike protein amyloid in the brains of repeated vaccine/booster recipients. This abnormal amyloid deposition is accompanied by biomarkers of neurodegeneration formerly associated with Alzheimer's, the Parkinson's disease spectrum, and dementia. The long-term effects of this are unknown at this time.

In addition, further FOIA documents show that a "statistically significant" number of baby rats were born with skeletal deformations after their mothers were injected with the Moderna vaccine. The FDA has dismissed this animal data, saying that the skeletal anomalies were "not considered adverse." However, leading experts in this field agree that the highly experimental mRNA "vaccines" should never have been authorizedfor use during pregnancy. Lax and inaccurate vaccine adverse event surveillance makes the true incidence of human miscarriages associated with the mRNA vaccines uncertain.

FOIA records also detail internal discussions about mRNA vaccine-induced heart inflammation (myocarditis). Publicly, the CDC and FDA have tried to downplay this serious event. However, leading cardiologists maintain that all cases of myocarditis are serious. The mRNA vaccines should never have been authorizedfor Young adults, children, and babies who lack any benefit—risk ratio. In late 2020, the Brighton Collaboration created a priority list of adverse events that should be assessed for the COVID-19 mRNA vaccines. In 2022, a small group of prominent clinical data Scientists used this list to reanalyze the placebo-controlled, phase Ill randomized clinical trials of the Pfizer and Moderna mRNA COVID-19 "vaccines" in adults (clinical trials NCT04368728 and NCT04470427). This reanalyzed manufacturer clinical trial data

clearly demonstrates a negative benefit-to-risk ratio in individuals who took the Moderna or Pfizer mRNA COVID-19 "vaccines."

The injected mRNA "vaccine" instructs the body to produce a highly toxic, harmful molecule called the spike protein for weeks after injection. Both the Moderna and Pfizer manufacturer's clinical trials show these mRNA vaccines cause a higher risk of postvaccine hospitalization, disability, and a life-changing event that is greater than the risk ofbeing unvaccinated and naturally catching a COVID-19 infection and being hospitalized with it. These mRNA products should never have been given to humans.

Instead of an immediate halt to mRNA mass vaccination in January 2022, the program continued to run, poorly supervised, for another sixteen months until the pandemic emergency ended.

Throughout 2021, 2022, and 2023, a majority of the American population has essentially served as uninformed, often coerced or mandated medical test subjects for the ineffective, poorly tested, unsafe, and highly experimental mRNA COVID-19 "pseudo-vaccines." This coercion violated the 1947 Nuremberg Code, which provided ethical guidelines for human research, the later 1964 Declaration of Helsinki that reaffrrned the need for informed consent in human research, the 1978 Belmont Report that framed these issues into "broader" ethical principles, and the International Covenant on Civil and Political Rights (ICCPR), which provides that "no one shall be subjected without his free consent to medical or scientific experimentation."

During the U.S. mass vaccination program, the highly experimental COVID-19 mRNA vaccines were administered withoUt informed consent. Even worse, all federal employees, the U•S• military, healthcare workers, students, and airline pilots were

mandated to receive the problematic vaccines with repeated federal health agency recommendations for the vaccination of young adults, children, and infants. These were groups without any demonstrable benefit-to-risk ratio.

As observed with the mRNA vaccine debacle, in spite of international agreements outlining ethical guidelines for human experimentation, there is still a needfor a definitive, legal, enforceable norm to protect the rights of all Americans against experimental vaccine mandates. As of this writing, the American people have little to no trust in the federal health agencies.

President Trump will restore this trust through urgent action. Recent FOIA documents obtained by America First Legal clearly outline that the CDC was actively suppressing the free speech of U.S. citizens on the social media platforms of Twitter, Facebook, and Instagram, intentionally blocking and actively deplatforming highly educated physicians and scientists who were trying to warn the public of the proven safety and effectiveness of hydroxychloroquine and Ivermectin as early outpatient treatments for COVID- 19 and the dangers of the experimental mass mRNA vaccination program.

- By executive order, President Trump should explicitly prohibit all federal agencies from interfering with the practice Of medicine and the freedom of speech exercised by doctors and scientists on social media platforms or any other public "town hall" communications.
- President Trump should immediately put into place mechanisms to hold the commercial mainstream media accountable for any prolonged and incorrect reporting on medical topics

that result in injury or death. This accountability will help safeguard the public from misinformation and ensure that accurate and reliable information is disseminated. Liability measures with no statute of limitations should be the consequences for spreading false or misleading medical information.

In the face of the overwhelming statistics, the histologically proven vaccine tissue damage, and the lack of effcacy, the Biden vaccination mandates severely tested the U.S. Constitution and Bill of Rights. Technically, many state governors, schools, universities, hospitals, companies, and corporations all violated international ethics considerations by coercing and subjecting their state residents to experimental medical procedures [sic, mRNA vaccines] that for months carried a significant demonstrable risk without any benefit.

- By executive order, President Trump should immediately roll back all forced vaccination policies and ban medical coercion and mandates for medical treatments and procedures. will protect individuals' rights to make autonomous decisions about their own healthcare and prevent undue pressure or force to undergo medical interventions.
- President Trump should also propose an amendment to the Bill of Rights that explicitly requires fully informed consent for all clinical trial participants and the general public.

The carnage caused by Anthony Fauci, individual companies like Pfizer, and, most broadly, the Big Pharma—federal health agency industrial complex must never be allowed to happen again.