US COVID-19 RESPONSE – THE COUNTERMEASURES

• The US DoD is the Executive branch of Operation Warp Speed (OWS) and they own the "mRNA COVID-19 vaccines" until they are injected in arms.

• The National Security Council (NSC) is in charge of the COVID Policy and not HHS as they will have everyone to believe.

• The Federal Emergency Management Agency (FEMA) is in charge of Operations for the public health emergency COVID-19 response.

• HHS and their Public Health Emergency Medical Countermeasures Enterprise (PHEMCE) only act as scientific advisory group to Operation Warp Speed (OWS), they are not executive but US DoD is. Their role was only to issue recommendations and rubber stamp approvals whilst financing the R&D effort.

• COVID-19 Vaccine are in fact Security Covered Countermeasures not regulated by the FDA under Emergency Use Authorisation (EUA). The US Heath Secretary and the United States Secretary of Homeland Security must secure presidential approval to develop and manufacture security covered countermeasures (not available in the stockpile) to respond to a biological (CBRN) national security threat on the strength of a Public Health Emergency and when an Emergency Use Authorisation (EUA) has been authorised by the FDA Commissioner.

• The Biomedical Advanced Research and Development Authority (BARDA) under the dome of HHS, is in charge of financing the development and manufacturing of these countermeasures. They also seems to have taken on the role of a regulator which they are not.

• Advanced Technology International (ATI) micro-manages Big Pharma OTA contracts for the US DoD and for BARDA. They also have an oversight over the contracts allocated to pharma companies members of the Medical Technology Enterprise Consortium (MTEC) and the Medical CBRN Defence Consortium (MCDC).

• FDA mandate does not provide them with the legal authority to regulate Security Covered Countermeasures under EUA. The law is clear… Development & Manufacturing of countermeasures under EUA cannot be considered to constitute a clinical investigation, meaning none of the clinical trials including subjects, investigators and trials outcomes exist within the context of the US code (as Pfizer demonstrated during the Jackson Brook law suit).

• FDA reviewed the Pfizer clinical trials data for the Pfizer BNT162b2 mRNA shot before trying to bury it for 70+ years as the “Cumulative Analysis of Post-authorization Adverse Event Reports” were confirming an inconvenient 1,223 fatal cases and 42,000 adverse reactions as part of Pfizer clinical trial outcome” (source section 5.3.6)

• Pfizer and big pharma were contracted by US DoD and the Biomedical Advanced Research and Development Authority (BARDA) using Other Transaction Authorities (OTA) a form of contract which by nature drastically reduces vital oversight opportunities and protect IP meaning the US taxpayers never get to know what’s actually in the jab and same goes for US Government agencies which too will find it very hard if not impossible to get clarity or access.

• OTA contract confirms that Pfizer was contracted for a demonstration (demo) of a prototype mRNA vaccine in view of large scale manufacturing. This does not make their product a pharmaceutical product as demo and prototype are not subjected to strict manufacturing regulatory measures and protocols but only require “reasonable efforts” on the manufacturers part . Clinical trials data can be incomplete and it is not required for the vaccine companies to prove safety or efficacy when SOW is a demo and a prototype.

• FDA / CDC and mass media were deceitful and misleading by promoting efficacy and safety of these injections as we now know immunity studies were never conducted prior to big pharma receiving their initial marketing approval. (this was confirmed by Janine Small from Pfizer). People were told sometimes even forced to get the jab to keep their job and to allegedly protect others when FDA and CDC had no idea whatsoever if the jab could stop infection and transmission.

• We know NIH have been sponsoring Game of Function for a very long time, in and out of the United States and even when Obama’s moratorium forbidding GOF research was in place

• SARS CoV alleged isolated virus (for the virus therory believers) was patented by the CDC hence it cannot be fully natural and if it was fully natural, it was illegally patented. US7220852B1 - US7776521B1

• COVID-19 so called vaccines are not pharmaceutical products as they are not strictly regulated, they are not prophylactic, they do not stop infection nor transmission and they do harm people. This experimental products and delivery systems were not developed in good faith since we can establish from available early data that Big Pharma and FDA had prior knowledge of their dangerousness which establish intent to harm. The above information pertaining to these injectables qualifies and characterises this COVID-19 injections as a “Military grade Security Covered Countermeasure” and by definition as a “Bioweapon” or “Synthetic Bioweapon”