IN THE SUPREME COURT OF FLORIDA

Case No.: SC24

JOSEPH SANSONE, M.S., PhD,

Petitioner,

V.

HON. RON DESANTIS, in his

Official Capacity of Governor

of Florida; and

HON. Ashley Moody, in her official capacity

of Attorney General of Florida;

Respondents.

_____/

EMERGENCY PETITION FOR A WRIT OF MANDAMUS

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PETITION FOR A WRIT OF MANDAMUS

(1) This petition for a writ of mandamus is brought under Article V, § 3(b)(8) Florida Constitution, and under Florida Rules of Appellate Procedure 9.03O(a)(3), 9.100 and other relevant authorities to enforce state and federal laws including, and not limited to Biological Weapons 18 USC § 175; Weapons and Firearms § 790.166 Fla. Stat. (2023); Federal Crime of Treason 18 USC § 2381; Treason § 876.32 Fla. Stat. (2023); Domestic Terrorism, 18 USC § 2331, Terrorism § 775.30 Fla. Stat. (2023); Murder § 782.04 (1)(a) Fla. Stat. (2023); and Genocide 18 USC §1091. Petitioner seeks an order of mandamus, requiring the Respondents to immediately prohibit the distribution, promotion, access and administration of COVID-19 injections, mRNA nanoparticle injections, and all mRNA products in the State of Florida.

I. Parties

(2) The petitioner, Joseph Sansone, M.S., PhD, is a citizen and taxpayer in the State of Florida and has been targeted with biological and technological weapons of mass destruction, and has lost and will continue to lose friends and family to these same weapons of mass destruction. The Respondents, the Honorable Ron DeSantis, is Governor of the State of

Florida, and the Honorable Ashley Moody, is Attorney General, of the State of Florida.

II. Jurisdiction

(3) The Court has original jurisdiction under Article V, § 3(b)(8) Florida Constitution. The Florida Supreme Court may issue writs of mandamus and quo warranto to state officers and state agencies. There is concurrent jurisdiction between the Florida Supreme Court, Circuit Courts and the District Courts of Appeal. Article V, §4(b)(3) Florida Constitution and Article V, § 5(b) Florida Constitution. The Florida Supreme Court should clearly exercise its jurisdiction when a writ of mandamus seeks to command the Governor of the State of Florida, the chief constitutional officer of the Executive Branch, to protect Floridians by granting the relief requested herein. It is appropriate that this Court, rather than the lower state courts, should consider such a claim against the Governor. In the case of Wright v. Chiles, 18 Fla. L. Weekly S509 (Fla. Sept. 30, 1993) this Court asserted its jurisdiction under Article V, § 3(b)(8) Florida Constitution regarding a petition for a writ of mandamus directing the Governor to appoint Dr. Wright to a new term as medical examiner for the Seventeenth District. The Florida Medical Examiners Commission nominated only Dr. Wright for the

position and the Governor requested the commission to submit additional nominations.

(4) Based on the interpretation of Article V, § 3(b)(8) Florida Constitution by this Court in Wright v. Chiles, 18 Fla. L. Weekly S509 (Fla. Sept. 30, 1993), this Court has jurisdiction in the present case, as Petitioner seeks to command the Governor of the State of Florida to take the action sought herein to protect himself and other Floridians.

(5) Furthermore, according to the Florida Supreme Court, “A mandamus also must establish that the action being sought is “ministerial.” “An action is ministerial only to the extent that the respondent has no discretion over the matter.” However, mandamus may be used to compel official action that falls within an established legally permissible range if that official fails to act within the range and is required by the law to do so. The petitioner seeks to compel Respondents to take mandatory, required action within a legally permissible range and they are required by law to do so. The fact

that a court may need to interpret a statute to discern the permissible range does not make the legal right any less" clear. 29 Nova L. Rev. 431 (2005) [1]

(6) Article IV, § 1(a) Florida Constitution states, "The governor shall take care that the laws be faithfully executed". Shall is an imperative command, indicating that it is mandatory that the Governor execute the laws, not discretionary. Article IV, § 4(b) Florida Constitution provides that "The attorney general shall be the chief state legal officer. Therefore, the Attorney General is equally bound to enforce the laws of the State of Florida. The Attorney General is also a state officer, and it is proper for this court to assert its jurisdiction.

(7) The gravity of the laws in question makes it clear that the Florida Supreme Court has jurisdiction, and in fact, is the most appropriate Court to hear this case.

[1] Anstead, H. L., Kogan, G., Hall, T. D., & Waters, R. C. (2004). The Operation and Jurisdiction of the Supreme Court of Florida. Nova L. Rev., 29, 431.

III. Statement of Facts

A. Introduction

(8) The distribution of COVID-19 injections (hereafter referred to as ‘COVID-19 nanoparticle injections’ or ‘mRNA nanoparticle injections’ or ‘COVID-19 injections) must be halted immediately in the State of Florida. COVID-19 injections have caused countless deaths, permanent injury, and disability. We intend to demonstrate to the court that; 1- the COVID-19 injections are dangerous, 2 – it is the duty of the Governor and Attorney General to act immediately to prohibit their distribution, 3 – it is incumbent on this Court to compel the Governor and Attorney General to act immediately.

The evidence demonstrates that due to shedding, COVID-19 injections pose a risk of harm, including death and disability, to all Floridians whether ‘vaccinated’ or ‘unvaccinated’. Every Floridian, including members of this Court, and likely the Respondents, were lied to – COVID-19 injections are not safe, nor are they effective.

(9) Due to the extraordinary nature of the crimes committed against the civilian population in the State of Florida, and the threat that continued

distribution of COVID-19 injections will cause further death and disability, and the potential threat posed to the future existence of the human race itself, this Court must act immediately to prevent further harm.

B. EXECUTIVE SUMMARY

(10) A massive mass media and government campaign promoting 'COVID-19 vaccines' as safe and effective 'vaccines' to prevent infection targeted Florida's population of approximately 22 million people. This campaign narrative was false and misleading and has led to numerous deaths, permanent injury, and disability.

(11) It is well-established that the FDA clinical trials for the COVID-19 injections were ***not*** designed to clinically or statistically demonstrate that the COVID-19 nanoparticle injections prevent infection, prevent

transmission, or protect against disease, hospitalizations, and death.[2,3,4,5,6,7,8]

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[2] . Pfizer Inc., BioNTech, initial new drug (IND) application. “A PHASE 1/2/3, PLACEBO-CONTROLLED, RANDOMIZED, OBSERVER-BLIND, DOSE-FINDING STUDY TO EVALUATE THE SAFETY, TOLERABILITY, IMMUNOGENICITY, AND EFFICACY OF SARS-COV-2 RNA VACCINE CANDIDATES AGAINST COVID-19 IN HEALTHY INDIVIDUALS.” PF-07302048 (BNT162 RNA-Based COVID-19 Vaccines) Protocol C4591001; Apr 2020.

[3] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; “Emergency Use Authorization (EUA) for an Unapproved Product.” Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

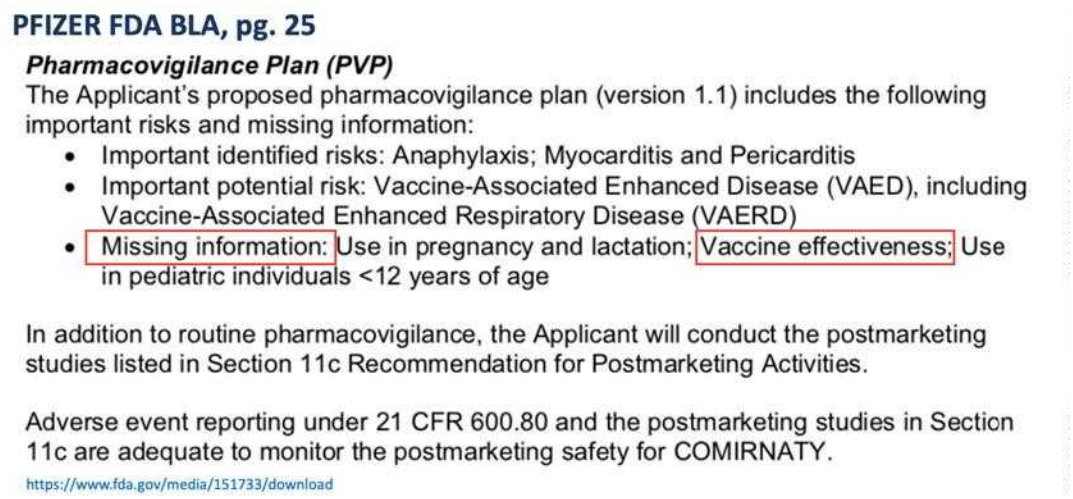
[4] Polack F., Thomas S., Kitchin N., et al. for the C4591001 Clinical Trial Group; “Safety and Efficacy of the BNT162b2 mRNA Covid-19 Vaccine” New Engl J Med; Dec 10, 2020; 383:2603-2615. https://www.nejm.org/doi/full/10.1056/nejmoa2034577

[5] Naik Ramachandra, PhD, Review Committee Chair, DVRPA/OVRR. BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. “COMIRNATY (BNT162b2): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in individuals 16 years of age and older” FDA Approval of the Biological License Application (BLA) for BNT162b2/COMIRNATY; Submitted May 18, 2021. Reviewed November 8, 2021. https://www.fda.gov/media/151733/download

[6] Zaks Tal, ModernaTX, Inc. initial new drug (IND) application. “A Phase 3, Randomized, Stratified, Observer-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of mRNA-1273 SARS-CoV-2 Vaccine in Adults Aged 18 Years and Older, mRNA-1273-P301” Aug 20, 2020. https://covid19crc.org/wp-content/uploads/2020/09/mRNA-1273-P301-Protocol-2020.pdf

[7] Agnihothram Sudhakar, PhD, Review Committee Chair, DVRPA/OVRR. ModernaTX Inc. “SPIKEVAX (mRNA-1273): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in individuals 18 years of age and older” FDA Approval of the Biological License Application (BLA) for COVID-

(12) In Pfizer's August 23, 2021 approval of their biological license application, Pfizer's approved application even states, "**Missing Information: Vaccine Effectiveness.**"



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Pharmacovigilance Plan (PVP)

The Applicant's proposed pharmacovigilance plan (version 1.1) includes the following important risks and missing information:

- Important identified risks: Anaphylaxis; Myocarditis and Pericarditis
- Important potential risk: Vaccine-Associated Enhanced Disease (VAED), including Vaccine-Associated Enhanced Respiratory Disease (VAERD)
- Missing information: Use in pregnancy and lactation; Vaccine effectiveness; Use in pediatric individuals <12 years of age

In addition to routine pharmacovigilance, the Applicant will conduct the postmarketing studies listed in Section 11c Recommendation for Postmarketing Activities.

Adverse event reporting under 21 CFR 600.80 and the postmarketing studies in Section 11c are adequate to monitor the postmarketing safety for COMIRNATY.

https://www.fda.gov/media/151733/download

(13) Moderna's January 4, 2022, FDA approval is also missing "**Vaccine Effectiveness.**"

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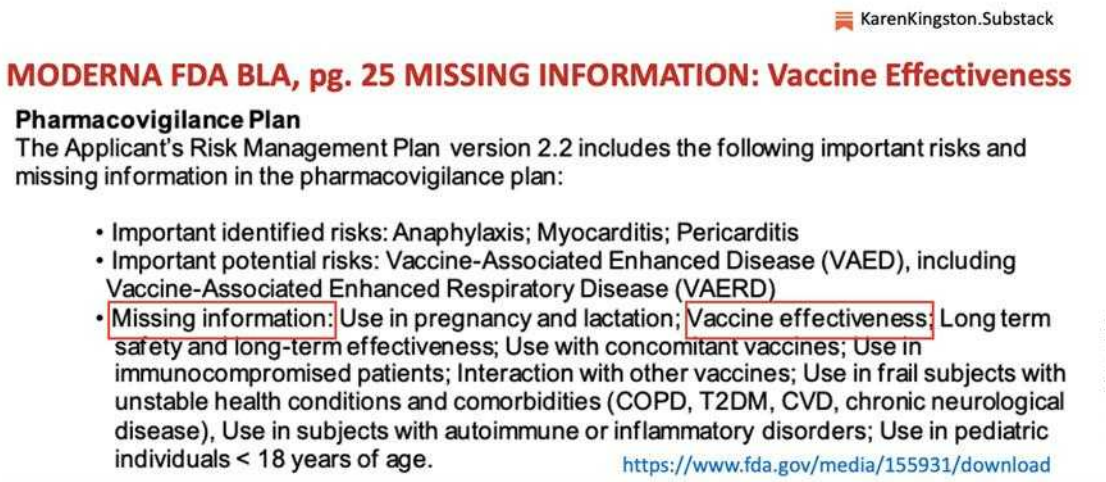
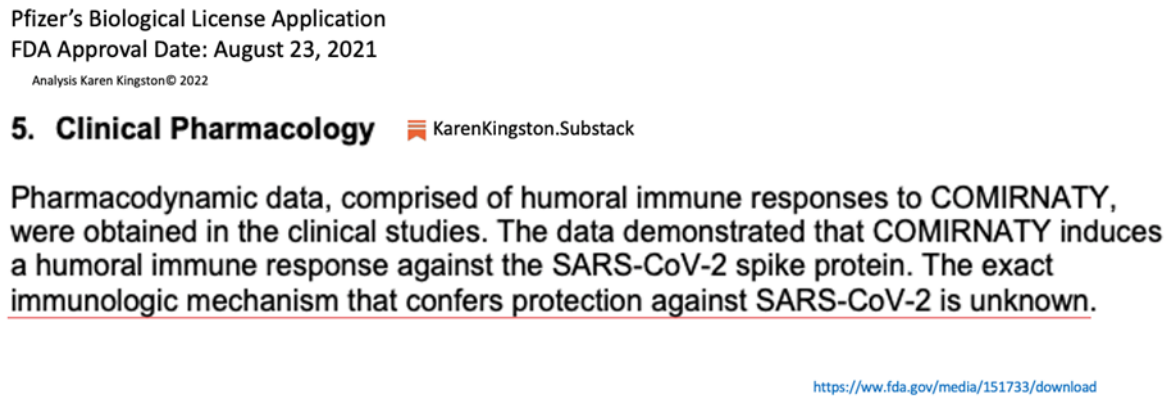
19 Vaccine, mRNA; January 30, 2023. https://www.fda.gov/media/155931/download

[8] ModernaTX, Inc. "FDA Briefing Document Moderna COVID-19 Vaccine" Vaccines and Related Biological Products Advisory Committee (VRBPAC) Meeting. Emergency Use Authorization (EUA); December 17, 2020. https://www.fda.gov/media/144434/download



(14) Pfizer also states in their approved biological license that they don't know how their COVID-19 mRNA vaccine protects against SARS-CoV-2 infection.



(15) FDA clinical trials, US government data, and real-world evidence have demonstrated that mRNA nanoparticle injections cause clinically significant increases in mild-to-moderate disease, serious diseases, disabilities, hospitalizations, and death within days, weeks and/or months of

receiving COVID-19 mRNA nanoparticle injections in formerly healthy infants, children, and adults.[9,10,11,12,13,14,15,16,17,18,19,20]

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[9] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; "Emergency Use Authorization (EUA) for an Unapproved Product." Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

[10] . Agnihothram Sudhakar, PhD, Review Committee Chair, DVRPA/OVRR. ModernaTX Inc. "SPIKEVAX (mRNA-1273): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in individuals 18 years of age and older" FDA Approval of the Biological License Application (BLA) for COVID-19 Vaccine, mRNA; January 30, 2023. https://www.fda.gov/media/155931/download

[11] ModernaTX, Inc. "FDA Briefing Document Moderna COVID-19 Vaccine" Vaccines and Related Biological Products Advisory Committee (VRBPAC) Meeting. Emergency Use Authorization (EUA); December 17, 2020. https://www.fda.gov/media/144434/download

[12] Pfizer, Inc., BioNTech. "Application for licensure of a booster dose for COMIRNATY (COVID-19 Vaccine, mRNA). FDA Briefing Document" Vaccines and Related Biological Products Advisory Committee (VRBPAC) Meeting. September 17, 2021. https://www.fda.gov/media/152176/download

[13] Anderson Steven, PhD, MPP – Dir. Office of Biostats & Epidemiology, CBER. "FDA/CBER Plans for Monitoring COVID-19 Vaccine Safety & Effectiveness Vaccines & Related Biological Products Advisory Committee (VRBPAC) Meeting" October 22, 2020. https://www.fda.gov/media/143557/download?fbclid=IwAR1SooRjTDuhBPqM4TiD3O7vYgX4eAp3CCqB7SzCk04CMve_OzgtMNPfNkc

[14] Senator Ron Johnson, Subcommittee on Investigations, Ranking Member. "United States Senate Letter: The Honorable Lloyd J. Austin III" regarding the Defense Medical Epidemiology Database (DMED); Feb 1, 2022. https://www.ronjohnson.senate.gov/services/files/FB6DDD42-4755-4FDC-BEE9-50E402911E02

[15] Wollersheim Susan, MD, Schwartz Ann, MD. Lee Lucia, MD; Team Leader CRB1/DVRPA/OVRR Allende Maria MD; Chief, CRB1/DVRPA/OVRR, BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. "BLA Clinical Review MEMORANDUM" Biological License

(16) The COVID-19 mRNA nanoparticle injections were administered to civilian adults and children through unlawful human experimentation under the guise of "safe and effective vaccines. Specifically, the FDA and mRNA manufacturers knew that the clinical safety risks outweighed any potential clinical benefits.

(17) Furthermore, the COVID-19 mRNA injections were administered ***without*** informed consent regarding:

Application (BLA); August 23, 2021. https://www.fda.gov/media/152256/download

[16] Shresrha N., Burke P., Nowacki., et al. "Effectiveness of the Coronavirus Disease 2019 Bivalent Vaccine." Open Forum Infect Dis. April 19, 2023. (Cleveland Clinic) https://www.ncbi.nlm.nih.gov/pmc/articles/PMC10234376/

[17] Nafilyan V., Bermingham C., Ward IL., et al. "Risk of death following SARS-CoV-2 infection or COVID-19 vaccination in young people in England: a self-controlled case series study" MedRxiv. March 23, 2022. https://www.medrxiv.org/content/10.1101/2022.03.22.22272775v1

[18] Schmeling M, Manniche V, Hansen PR. "Batch-dependent safety of the BNT162b2 mRNA COVID-19 vaccine" European Journal of Clinical Investigation. March 30, 2023. https://doi.org/10.1111/eci.13998

[19] European Medical Agency. "Safety of COVID-19 vaccines." https://www.ema.europa.eu/en/human-regulatory/overview/public-health-threats/coronavirus-disease-covid-19/treatments-vaccines/vaccines-covid-19/safety-covid-19-vaccines Assessed January 6, 2023.

[20] European Directorate for the Quality of Medicines and Healthcare. "EDQM initiatives in the context of COVID-19 vaccines and therapies." https://www.edqm.eu/en/edqm-initiatives-in-the-context-of-covid-19-vaccines-and-therapies Assessed January 6, 2023.

- the composition and variability of the COVID-19 nanoparticle injections' vials,
- the gene-editing mechanism of action of COVID-19 nanoparticle technologies, and the known harmful, permanently disabling and/or

sometimes deadly clinical outcomes of being injected with engineered COVID-19 mRNA nanoparticle technologies.[21,22,23,24,25,26,27,28,29,]

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[21] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; "Emergency Use Authorization (EUA) for an Unapproved Product." Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

[22] . Naik Ramachandra, PhD, Review Committee Chair, DVRPA/OVRR. BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. "COMIRNATY (BNT162b2): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in individuals 16 years of age and older" FDA Approval of the Biological License Application (BLA) for BNT162b2/COMIRNATY; Submitted May 18, 2021. Reviewed November 8, 2021. https://www.fda.gov/media/151733/download

[23] Anderson Steven, PhD, MPP – Dir. Office of Biostats & Epidemiology, CBER. "FDA/CBER Plans for Monitoring COVID-19 Vaccine Safety & Effectiveness Vaccines & Related Biological Products Advisory Committee (VRBPAC) Meeting" October 22, 2020. https://www.fda.gov/media/143557/download?fbclid=IwAR1SooRjTDuhBPqM4TiD3O7vYgX4eAp3CCqB7SzCk04CMve_OzgtMNPfNkc

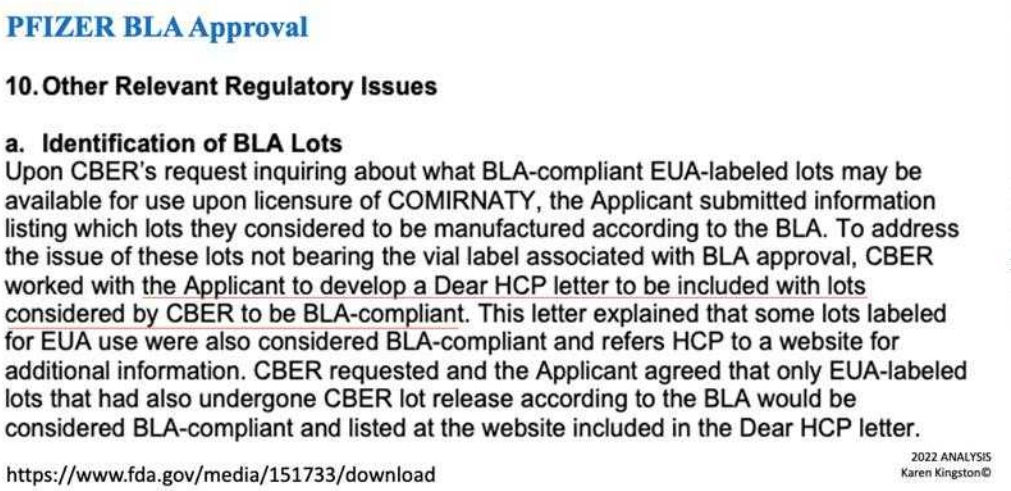
[24] Pfizer-BioNTech COVID-19 Vaccine EUA Amendment for Use in Children 6 Months Through 4 Years of Age. "EUA amendment request for Pfizer-BioNTech COVID-19 Vaccine for use in children 6 months through 4 years of age." VRBPAC Briefing Document. June 15, 2022. https://www.fda.gov/media/159195/download

[25] U.S. Department of Health and Human Services Food and Drug Administration, Office of the Commissioner, Office of the Chief Scientist, Office of Counterterrorism and Emerging Threats. "Emergency Use Authorization of Medical Products and Related Authorities" Guidance for Industry and Other Stakeholders; January 2017. https://www.fda.gov/media/97321/download

[26] Vaccines and Related Biological Products Advisory Committee Meeting. Licensure and Emergency Use Authorization of Vaccines to Prevent COVID-19 for Use in Pediatric Populations" FDA Briefing Document;June 10, 2021. https://www.fda.gov/media/149935/download

[27] Katragadda, CS., Choudhury, KS, and Murthy, PN. "Nanoparticles as Non-Viral Gene Delivery Vectors" IEEE Transactions on Nanobioscience.

- Pfizer states in their FDA application the COVID-19 mRNA vials vary in formulation.



PFIZER BLA Approval

10. Other Relevant Regulatory Issues

a. Identification of BLA Lots

Upon CBER's request inquiring about what BLA-compliant EUA-labeled lots may be available for use upon licensure of COMIRNATY, the Applicant submitted information listing which lots they considered to be manufactured according to the BLA. To address the issue of these lots not bearing the vial label associated with BLA approval, CBER worked with the Applicant to develop a Dear HCP letter to be included with lots considered by CBER to be BLA-compliant. This letter explained that some lots labeled for EUA use were also considered BLA-compliant and refers HCP to a website for additional information. CBER requested and the Applicant agreed that only EUA-labeled lots that had also undergone CBER lot release according to the BLA would be considered BLA-compliant and listed at the website included in the Dear HCP letter.

https://www.fda.gov/media/151733/download

2022 ANALYSIS
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- Pfizer states in their FDA approved license that their mRNA injections edit the genetic make-up of human cells through nucleoside substitution by RNA (modRNA) transcription.

January 2008 6(4):319-30
https://www.researchgate.net/publication/journal/IEEE-transactions-on-nanobioscience-1558-2639

[28] Friedrichs S and Bowman D. "COVID-19 may become nanomedicine's finest hour yet" Nature Nanotechnology;Volume 16, Pages 362–364. April 14, 2021. https://www.nature.com/articles/s41565-021-00901-8

[29] Wang R, Song B, Wu J, et al. "Potential adverse effects of nanoparticles on the reproductive system" Int Jrnl of Nanomedicine: 2018:13 8487–8506 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6294055/pdf/ijn-13-8487.pdf

a. Product Quality

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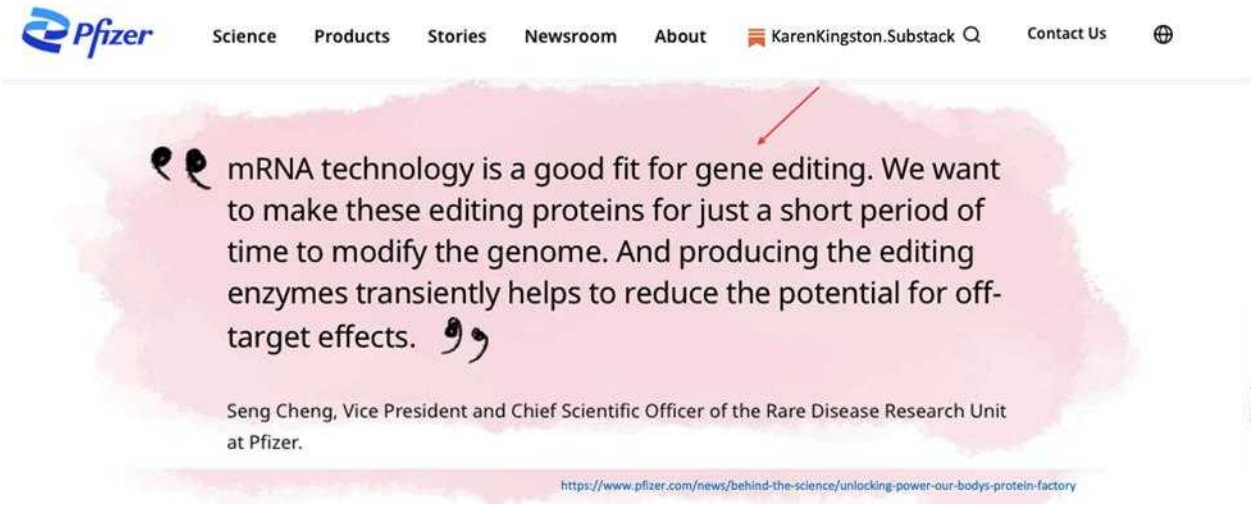
COMIRNATY Manufacturing Overview

The mRNA in COMIRNATY is a single-stranded, 5'-capped mRNA encoding the full-length SARS-CoV-2 spike glycoprotein derived from the Wuhan-Hu-1 isolate (GenBank MN908947.3 and GenBank QHD43416.1). The antigen-coding RNA sequence is codon-optimized and contains two proline mutations ((b) (4) 87P), which ensures an antigenically optimal trimerized pre-fusion confirmation (S-2P). The RNA also contains common structural elements, including 5'-cap, 5'-UTR, 3'-UTR, and poly(A) tail, all of which are designed for mediating high RNA stability and translation efficiency. During RNA transcription, (b) (4) is replaced with the (b) (4) . This nucleoside substitution has been demonstrated to enhance translation of *in vitro* transcribed mRNA while reducing its reactogenicity. Nov 2022 https://www.fda.gov/media/151733/download

- Pfizer states on their corporate website that their mRNA produces 'editing proteins' (aka spike proteins) to modify the genome.



- A recent *in vitro* study demonstrated that Pfizer's COVID-19 mRNA injection (BNT162b2) introduce foreign DNA into human liver cells in as fast as 6 hours upon BNT162b2 exposure.

Intracellular Reverse Transcription of Pfizer BioNTech COVID-19 mRNA Vaccine BNT162b2 In Vitro in Human Liver Cell Line

by Markus Aldén [1], Francisko Olofsson Falla [1], Daowei Yang [1],

Analysis Karen Kingston© 2022

Curr. Issues Mol. Biol. **2022**, *44*(3), 1115-1126; https://doi.org/10.3390/cimb44030073

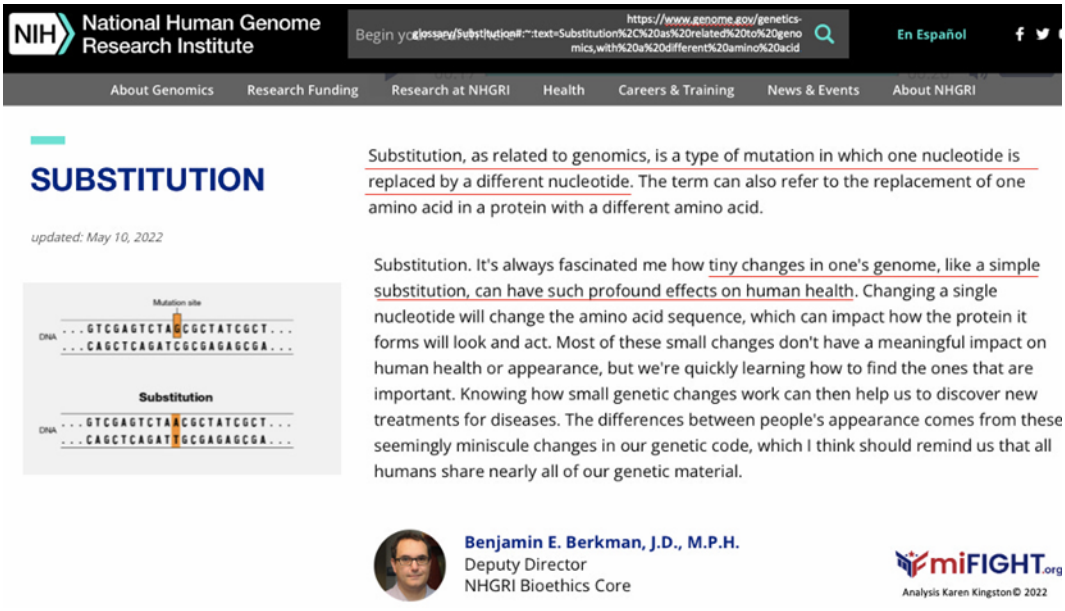
Abstract

Preclinical studies of COVID-19 mRNA vaccine BNT162b2, developed by Pfizer and BioNTech, showed reversible hepatic effects in animals that received the BNT162b2 injection. Furthermore, a recent study showed that SARS-CoV-2 RNA can be reverse-transcribed and integrated into the genome of human cells. In this study, we investigated the effect of BNT162b2 on the human liver cell line Huh7 in vitro. Huh7 cells were exposed to BNT162b2, and quantitative PCR was performed on RNA extracted from the cells. We detected high levels of BNT162b2 in Huh7 cells and changes in gene expression of long interspersed nuclear element-1 (LINE-1), which is an endogenous reverse transcriptase. Immunohistochemistry using antibody binding to LINE-1 open reading frame-1 RNA-binding protein (ORFp1) on Huh7 cells treated with BNT162b2 indicated increased nucleus distribution of LINE-1. PCR on genomic DNA of Huh7 cells exposed to BNT162b2 amplified the DNA sequence unique to BNT162b2. Our results indicate a fast up-take of BNT162b2 into human liver cell line Huh7, leading to changes in LINE-1 expression and distribution. We also show that BNT162b2 mRNA is reverse transcribed intracellularly into DNA in as fast as 6 h upon BNT162b2 exposure.

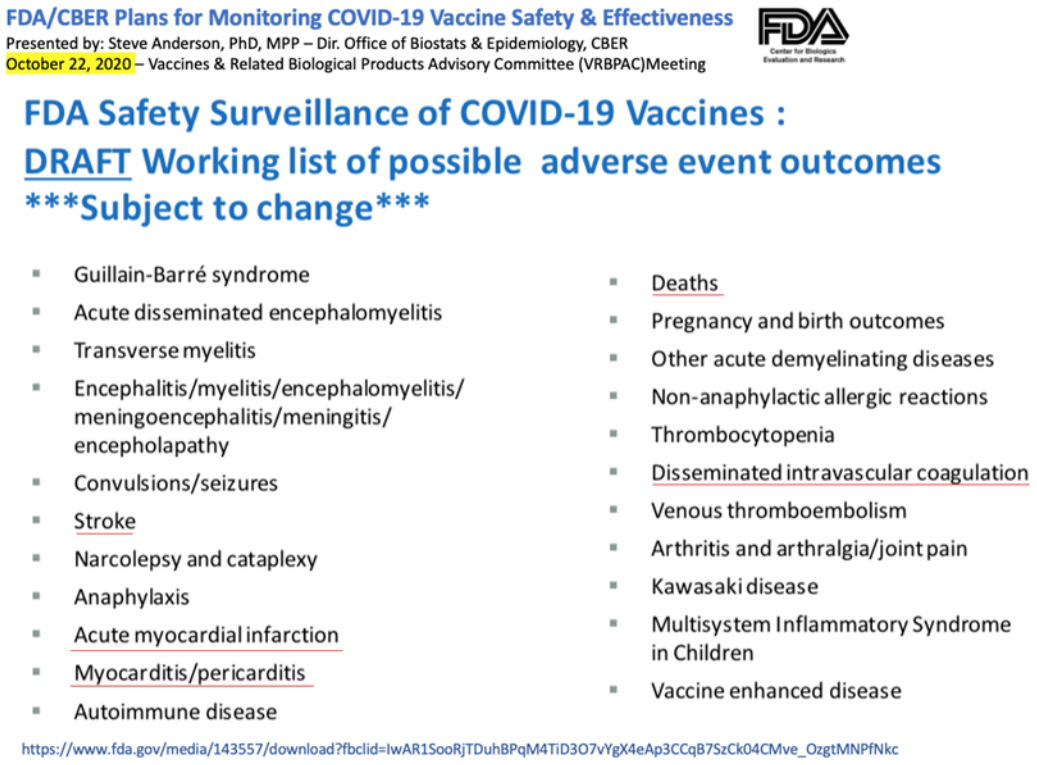
Keywords: COVID-19 mRNA vaccine; BNT162b2; liver; reverse transcription; LINE-1; Huh7

https://www.mdpi.com/1467-3045/44/3/73

- The NIH defines nucleoside substitution as a change to one's genome that can have profound effects on one's health.



- On October 22, 2020, the FDA met with COVID-19 vaccine mRNA manufacturers to discuss the harmful clinical effects of the COVID-19 injections.



FDA/CBER Plans for Monitoring COVID-19 Vaccine Safety & Effectiveness

Presented by: Steve Anderson, PhD, MPP – Dir. Office of Biostats & Epidemiology, CBER

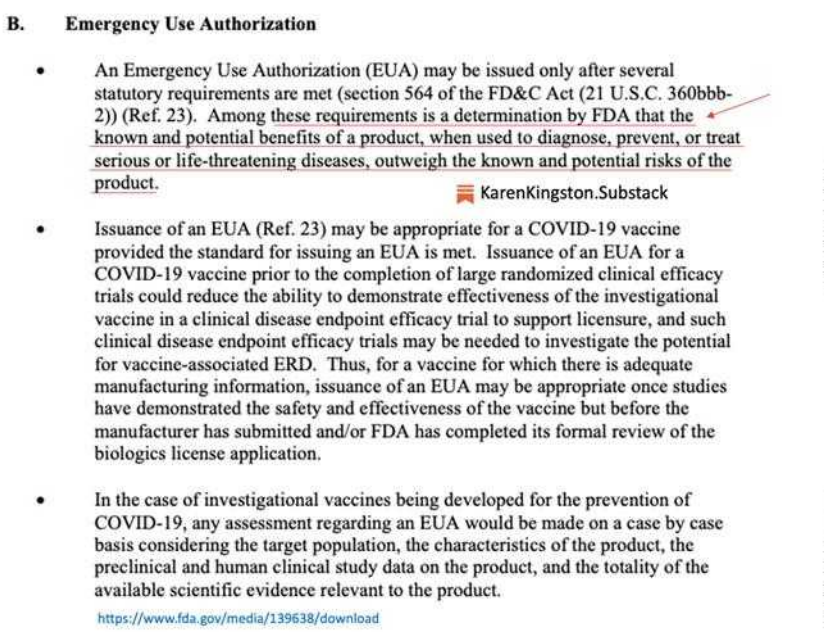
October 22, 2020 – Vaccines & Related Biological Products Advisory Committee (VRBPAC)Meeting

FDA Safety Surveillance of COVID-19 Vaccines : DRAFT Working list of possible adverse event outcomes ***Subject to change***

- Guillain-Barré syndrome
- Acute disseminated encephalomyelitis
- Transverse myelitis
- Encephalitis/myelitis/encephalomyelitis/ meningoencephalitis/meningitis/ encepholapathy
- Convulsions/seizures
- Stroke
- Narcolepsy and cataplexy
- Anaphylaxis
- Acute myocardial infarction
- Myocarditis/pericarditis
- Autoimmune disease
- Deaths
- Pregnancy and birth outcomes
- Other acute demyelinating diseases
- Non-anaphylactic allergic reactions
- Thrombocytopenia
- Disseminated intravascular coagulation
- Venous thromboembolism
- Arthritis and arthralgia/joint pain
- Kawasaki disease
- Multisystem Inflammatory Syndrome in Children
- Vaccine enhanced disease

https://www.fda.gov/media/143557/download?fbclid=IwAR1SooRjTDuhBPqM4TiD3O7vYgX4eAp3CCqB7SzCk04CMve_OzgtMNPfNkc

- Moving forward with the FDA emergency use authorization (EUA) of COVID-19 mRNA LNP injections was in violation of Classification of products 21 USC § 360bbb-2 clearly stating that an EUA product's benefits must outweigh the risks.



B. Emergency Use Authorization

- An Emergency Use Authorization (EUA) may be issued only after several statutory requirements are met (section 564 of the FD&C Act (21 U.S.C. 360bbb-2)) (Ref. 23). Among these requirements is a determination by FDA that the known and potential benefits of a product, when used to diagnose, prevent, or treat serious or life-threatening diseases, outweigh the known and potential risks of the product.

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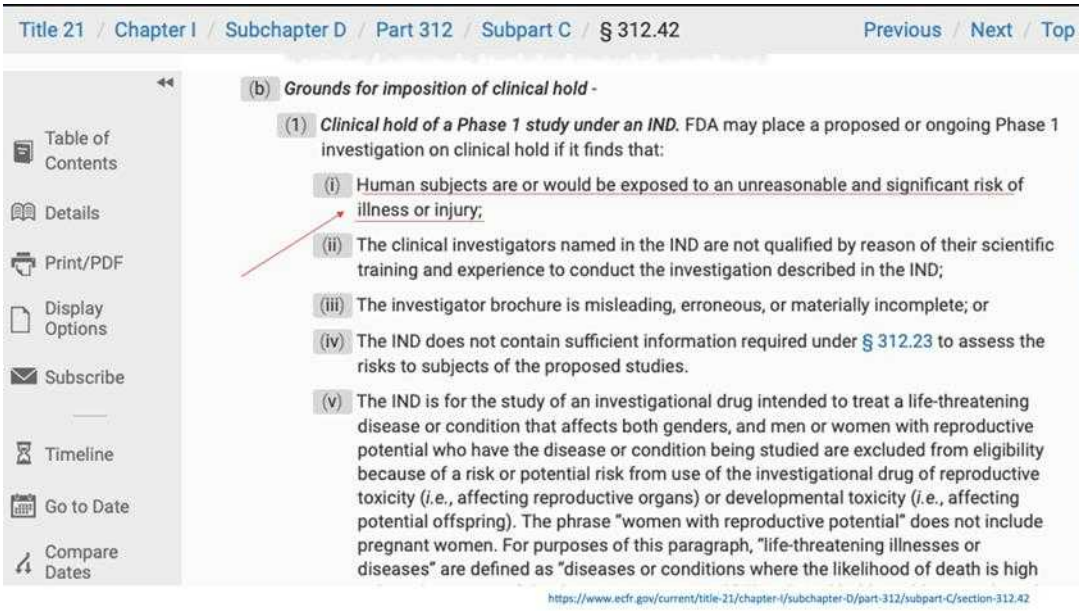
- Issuance of an EUA (Ref. 23) may be appropriate for a COVID-19 vaccine provided the standard for issuing an EUA is met. Issuance of an EUA for a COVID-19 vaccine prior to the completion of large randomized clinical efficacy trials could reduce the ability to demonstrate effectiveness of the investigational vaccine in a clinical disease endpoint efficacy trial to support licensure, and such clinical disease endpoint efficacy trials may be needed to investigate the potential for vaccine-associated ERD. Thus, for a vaccine for which there is adequate manufacturing information, issuance of an EUA may be appropriate once studies have demonstrated the safety and effectiveness of the vaccine but before the manufacturer has submitted and/or FDA has completed its formal review of the biologics license application.

- In the case of investigational vaccines being developed for the prevention of COVID-19, any assessment regarding an EUA would be made on a case by case basis considering the target population, the characteristics of the product, the preclinical and human clinical study data on the product, and the totality of the available scientific evidence relevant to the product.

https://www.fda.gov/media/139638/download

- Moving forward with the FDA emergency use authorization ***and FDA approval*** was in violation of Legal Human Experiments 21 CFR 312.42 (b)(1)(i) for in which subjects are not to be placed at a significant risk for illness or injury.



Important Recent US State Government Events

Texas is Suing Pfizer

(18) On November 30, 2023, State of Texas v. Pfizer, District Court of Lubbuck County, Texas (Pending) Texas Attorney General Ken Paxton filed a lawsuit against Pfizer[30] for:

- Misrepresenting Pfizer's "vaccine efficacy" and relative risk reduction
- Lying about the durability of alleged "vaccine protection"

[30] State of Texas v. Pfizer, District Court of Lubbuck County, Texas, (Pending). https://www.texasattorneygeneral.gov/sites/default/files/images/press/Pfizer%20Vaccine%20Petition%20Filed.pdf

- Lying about Pfizer's mRNA nanoparticle injections' ability to prevent transmission
- Lying about Pfizer's mRNA nanoparticle injections' ability to prevent infection from variants
- Conspiring with social media, specifically Twitter, to suppress the truth regarding the **effects of Pfizer's COVID-19 mRNA nanoparticle injections.**

Florida Dept. of Health Calls for the Halt of the Use of mRNA 'Vaccines'

(19) On January 3, 2024, the Florida Department of Health called for the halt of the use of COVID-19 mRNA vaccines in human beings, with Florida Surgeon General Dr. Ladapo, specifically stating, *"DNA integration poses a unique and elevated risk to human health and to the integrity of the human genome… If the risks of DNA integration have not been assessed*

*for **mRNA COVID-19 vaccines**, **these vaccines are not appropriate for use in human beings**."*[31]

(20) At a minimum this is a clear violation of Florida Drugs and Cosmetic Act § 499.005 (2) Fla. Stat. (2023)—It is unlawful for a person to perform or cause the performance of any of the following acts in this state, "The manufacture, repackaging, sale, delivery, or holding or offering for sale of any drug, device, or cosmetic **that is adulterated** or misbranded or has otherwise been rendered unfit for human or animal use."

(21) Dr. Ladapo went on to rebuke the FDA stating,*"It is my hope that, in regard to COVID-19, the FDA will one day seriously consider its regulatory responsibility to protect human health, including the integrity of the human genome."* In public statements Dr. Ladapo went as far as calling the Covid mRNA injection the *"Anti Christ of drugs"* and stated that they are evil.

[31] Bulletin Florida Department of Health (01/03/2024 08:30 AM EST) "Florida State Surgeon General Calls for Halt in the Use of COVID-19 mRNA Vaccines" (Tallahassee, Florida.) https://content.govdelivery.com/accounts/FLDOH/bulletins/3816863

C. Detailed Facts of This Case

(22) The 'COVID-19 vaccines' contain engineered nanoparticle technologies per the manufacturer's product labeling, FDA submissions, US military contracts, peer-reviewed publications, patents, and manufacturer's websites.[32,33,34,35,36]

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[32] Pfizer Inc., BioNTech, initial new drug (IND) application. "A PHASE 1/2/3, PLACEBO-CONTROLLED, RANDOMIZED, OBSERVER-BLIND, DOSE-FINDING STUDY TO EVALUATE THE SAFETY, TOLERABILITY, IMMUNOGENICITY, AND EFFICACY OF SARS-COV-2 RNA VACCINE CANDIDATES AGAINST COVID-19 IN HEALTHY INDIVIDUALS." PF-07302048 (BNT162 RNA-Based COVID-19 Vaccines) Protocol C4591001; Apr 2020.

[33] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; "Emergency Use Authorization (EUA) for an Unapproved Product." Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

[34] Naik Ramachandra, PhD, Review Committee Chair, DVRPA/OVRR. BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. "COMIRNATY (BNT162b2): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) in individuals 16 years of age and older" FDA Approval of the Biological License Application (BLA) for BNT162b2/COMIRNATY; Submitted May 18, 2021. Reviewed November 8, 2021. https://www.fda.gov/media/151733/download

[35] Zaks Tal, ModernaTX, Inc. initial new drug (IND) application. "A Phase 3, Randomized, Stratified, Observer-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of mRNA-1273 SARS-CoV-2 Vaccine in Adults Aged 18 Years and Older, mRNA-1273-P301" Aug 20, 2020. https://covid19crc.org/wp-content/uploads/2020/09/mRNA-1273-P301-Protocol-2020.pdf

[36] DEPARTMENT OF THE ARMY U.S. ARMY CONTRACTING COMMAND. "COVID-19 PANDEMIC—LARGE SCALE VACCINE

(23) Pfizer ignored and violated, Legal Human Experiments 21 CFR § 312.42 (b)(1)(i), Risk Evaluation and Mitigation Strategies 21 USC § 355-1, Current Good Manufacturing Practice 21 CFR § 225.1. for conducting safe and legal experimentation on humans with the use of FDA-regulated products when Pfizer stated that the formulations of their COVID-19 injections distributed to US adults and children varied by LOT number, per Pfizer's approved August 23, 2021, biological license application (BLA).[37]

(24) Pfizer's criminal experimentation on civilian adults and children with the use of varying biotechnologies in their COVID-19 mRNA nanoparticle formulations by 'vaccine' LOT number (with some lots known to inflict harm, ranging from serious diseases and disabilities to death), combined with lots that are placebos (known to be harmless), was confirmed by a scientific European analysis of 52 different Pfizer mRNA nanoparticle 'vaccine' LOTS, administered to 4,026,575 persons who received 10,793,766 doses

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MANUFACTURING DEMONSTRATION" RPP #: 20-11 Project Identifier: 2011-003. Statement of Work for Pfizer. July 21, 2020. https://www.hhs.gov/sites/default/files/pfizer-inc-covid-19-vaccine-contract.pdf

[37] Wollersheim Susan, MD, Schwartz Ann, MD. Lee Lucia, MD; Team Leader CRB1/DVRPA/OVRR Allende Maria MD; Chief, CRB1/DVRPA/OVRR, BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. "BLA Clinical Review MEMORANDUM" Biological License Application (BLA); August 23, 2021. https://www.fda.gov/media/152256/download

(an average of 2.7 injections/person) between December of 2020 and January of 2022.[38,39,40]

(25) The FDA *and* 'vaccine' manufacturers (i.e. Pfizer) clinically established that the COVID-19 injections would cause an unprecedented incidence of disease, permanent disabilities, and death, when on October 22, 2020 (before the 'COVID-19 vaccine rollout') the FDA met with the manufacturers and reviewed this 'working list' of harmful clinical outcomes caused by the injections; **nervous system disease**(convulsions, seizures, **Guillain-Barre syndrome myelitis encephalitis, encephalopathy, encephalomyelitis, narcolepsy, cataplexy,** meningitis, meningoencephalitis *acute demyelinating diseases*), **cardiac disease** (acute myocardial infarction myocarditis, pericarditis, stroke), **blood and circulatory disease**(disseminated intravascular

[38] Schmeling M, Manniche V, Hansen PR. "Batch-dependent safety of the BNT162b2 mRNA COVID-19 vaccine" European Journal of Clinical Investigation. March 30, 2023. https://doi.org/10.1111/eci.13998
[39] European Medical Agency. "Safety of COVID-19 vaccines." https://www.ema.europa.eu/en/human-regulatory/overview/public-health-threats/coronavirus-disease-covid-19/treatments-vaccines/vaccines-covid-19/safety-covid-19-vaccines Assessed January 6, 2023.
[40] European Directorate for the Quality of Medicines and Healthcare. "EDQM initiatives in the context of COVID-19 vaccines and therapies." https://www.edqm.eu/en/edqm-initiatives-in-the-context-of-covid-19-vaccines-and-therapies Assessed January 6, 2023.

coagulation, thrombocytopenia, venous thromboembolism), **musculoskeletal disease** (arthritis, joint pain), **reproductive and pregnancy disorders** (adverse pregnancy outcomes, adverse birth outcomes), **autoimmune disease** (VAED, multisystem inflammatory syndrome), and **death**.[41]

(26) There were **696,605 nervous system disorders**, **539,299 musculoskeletal** and connective tissue disorders (**92,942 pain in extremities**), and **317,811 gastrointestinal disorders**, **224,633 skin, hair and nail disorders**, **190,720 respiratory** and chest disorders, **178,353 female and male reproductive system disorders** (erectile dysfunction, infertility, heavy menstrual bleeding), **167,382 victims developed bacterial, viral, or parasitic infections (24,9010 herpetic infections), 126,993 cardiac** disorders, **100,970 blood** and lymphatic system disorders, **77,148 psychiatric** disorders, **73,542 vascular** disorders, **61,518 eye** disorders, **47,038 ear a**nd labyrinth disorders (**15,833 tinnitus**), **31,895 autoimmune** disorders, **13,647**

[41] Anderson Steven, PhD, MPP – Dir. Office of Biostats & Epidemiology, CBER. "FDA/CBER Plans for Monitoring COVID-19 Vaccine Safety & Effectiveness Vaccines & Related Biological Products Advisory Committee (VRBPAC) Meeting" October 22, 2020. https://www.fda.gov/media/143557/download?fbclid=IwAR1SooRjTDuhBPqM4TiD3O7vYgX4eAp3CCqB7SzCk04CMve_OzgtMNPfNkc

kidney and urinary disorders, **3,711 cancers** and benign cysts, **4,056 pregnancy complications** (**1,859 spontaneous abortion** complications, **1,143 genetic disorders**, and **3,814 deaths** were documented in an internal Pfizer document as of June 18, 2022.[42]

(27) There were 17,560 deaths, 83,092 hospitalizations, 116,479 urgent care visits,194,594 doctor visits, 36,014 anaphylaxis/severe allergic reactions, 13,515 cardiac events/conditions, 17,076 permanent disabilities, and an additional 14,494 life threatening events have been reported into the CDC's VAERS database as of June 16,2023, with an estimated 100-fold underreporting factor per a Harvard Pilgrim Healthcare Analysis commissioned by HHS.[43,44]

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[42] Pfizer. Confidential. "APPENDIX 2.2: Cumulative and Interval Summary Tabulation of Serious and Non-Serious Adverse Reactions from Post-Marketing Data Sources, BNT162B2." Data collected from December of 2020 through June 18, 2022. https://www.globalresearch.ca/wp-content/uploads/2023/05/pfizer-report.pdf

[43] Center for Disease Control (CDC). "VAERS COVID Vaccine Adverse Event Reports." US Data. June 16, 2023. https://openvaers.com/covid-data

[44] Lazarus Ross, MBBS, MPH, MMed, GDCompSci, Michael Klompas, MD, MPH. "Electronic Support for Public Health–Vaccine Adverse Event Reporting System (ESP:VAERS)." Prepared by Harvard Pilgrim Health Care, Inc. for U.S. Department of Health and Human Services. Inclusive dates: 12/01/07 - 09/30/10 https://digital.ahrq.gov/sites/default/files/docs/publication/r18hs017045-lazarus-final-report-2011.pdf

(28) There were more than one (1) million adverse events were reported in the VAERS database (1,055,219) in the year 2021 from the COVID-19 injections, including; hospitalizations, permanent disabilities, anaphylaxis, heart attacks, miscarriages, adult, child, and newborn deaths which is more than *ALL* reported adverse events from *ALL* childhood and adult vaccines over the past 20 years combined prior to the COVID-19 injection rollout (1990 -2020).[45]

(29) Based on data from the Defense Medical Epidemiology Database (DMED), it was reported that US military men and women experienced a 2,181% increase in hypertension, **1,048% increase in nervous system disorders**, a 894% increase in malignant neoplasms of esophagus, a 680% increase in multiple sclerosis, a 624% increase in malignant neoplasms of digestive organs, 551% increase in Guillain-Barre syndrome (paralysis), a 487% increase in breast cancer, 487% increase in demyelinating disease (damage to the myelin sheath protecting nerve fibers of the brain, optic nerve, and spinal cord), a 474% increase in malignant neoplasms of thyroid and other endocrine glands, a 472% increase in female infertility, a 468% increase in pulmonary embolism, a

[45] Center for Disease Control (CDC). "VAERS COVID Vaccine Adverse Event Reports." US Data. June 16, 2023. https://openvaers.com/covid-data

452% increase in migraines, a 437% increase in ovarian dysfunction, 369% increase in testicular cancer, and a 302% increase in tachycardia.[46]

(30) Data collected by the Joint Artificial Intelligence Center (JAIC) of the U.S. Department of Defense (DoD), demonstrated that among 5.6 million Medicare beneficiaries 65 years and older who received Pfizer's or Moderna's mRNA nanoparticle technology injections or remained uninjected, 71% of COVID-19 cases occurred in fully-vaccinated seniors and 60% of COVID-19 hospitalizations occurred in fully-vaccinated seniors as of August 7, 2021.[47]

(31) Data published by the CDC on June 15, 2023, demonstrated that in adults who were fully vaccinated or fully-vaccinated and boosted, and who

[46] Senator Ron Johnson, Subcommittee on Investigations, Ranking Member. "United States Senate Letter: The Honorable Lloyd J. Austin III" regarding the Defense Medical Epidemiology Database (DMED); Feb 1, 2022. https://www.ronjohnson.senate.gov/services/files/FB6DDD42-4755-4FDC-BEE9-50E402911E02

[47] United States Department of Defense, Joint Artificial Intelligence Center (JAIC), Humetrix data; "Effectiveness of mRNA COVID-19 Vaccines Against the Delta Variant Among 5.6M Medicare Beneficiaries 65 Years and Older." September 28.

were formerly immunocompetent (healthy) experienced an *increased risk* for hospitalization due to COVID-19.[48]

(32) More than 4 million Americans reported a Grade 3 adverse event (as defined as '*unable to perform their daily functions'*) and approximately 200,000 (2%) required admittance to the emergency room or hospital after receiving a COVID-19 injection according to the CDC's V-Safe database of 10 million US residents who were early recipients of COVID-19 injections as of July 31, 2022.[49]

(33) In addition, 403,396 Florida residents who were early recipients of COVID-19 injections, 167,005 (41.1%) reported a Grade 3 adverse event (unable to perform their daily functions) and 8,471 (2.1%) required admittance to the emergency room or hospital after receiving a COVID-19 injection per the CDC's V-Safe database report as of July 31, 2022.[61]

(34) Florida Surgeon General Joseph Ladapo identified 16,406 cardiac deaths from Florida's disease repository (MERLIN), Florida State Health

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[48] Link-Gelles, Ruth, PhD, MPHLCDR, US Public Health Service COVID-19 Vaccine Effectiveness Program Lead Centers for Disease Control and Prevention. "COVID-19 Vaccine Effectiveness Updates." June 15 2023. https://www.fda.gov/media/169536/download

[49] CDC V-SAFE Database. Accessed June 28, 2023. https://vaxxsafedata.com/about-us/

Online Tracking System (FLSHOTS), and death records, in adult Florida residents within 25 weeks of a 1st or 2nd mRNA nanoparticle injection; 3,417 of these cardiac deaths occurred within 28 days of a 1st or 2nd mRNA nanoparticle injection and *none* of these deaths were attributed to COVID-19 infection or a history of heart disease.[62]

(35) A recent systematic review of 100 studies, including case-reports and case studies, demonstrated that the average rate of myocarditis (a formerly rare disease among healthy adults and children) is 1.62% post COVID-19 mRNA nanoparticle injection, as well as demonstrated a clinically significant incidence of cardiomyopathy, pulmonary embolism (PE), and vaccine-induced thrombotic thrombocytopenia post COVID-19 mRNA injection.[63]

(36) It is clinically established that the mRNA 'spike proteins and 'lipid' nanoparticles cross the barrier membranes of the cardiovascular, respiratory, reproductive, and central nervous system (including the brain); causing inflammation that can result in disease, disability, and death, per

peer-reviewed publications and research & development Pfizer documents.[50,51,52,53,]

(37) There were 195% excess mortality claims in the State of Florida made to Group Life Insurance companies in July-September of 2021, during the time period when President Biden's previously announced COVID-19 vaccine mandate was to go into effect by July 4, 2021, for all employed Americans.[54,55] The CDC recorded an excess of 492,851deaths

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[50] Rawati J, Kumar V, Ahlwat P, et al. "Current Trends on the Effects of Metal Based Nanoparticles on Microbial Ecology" Applied Biochemistry and Biotechnology; Feb 17, 2023. https://doi.org/10.1007/s12010-023-04386-0

[51] Wang R, Song B, Wu J, et al. "Potential adverse effects of nanoparticles on the reproductive system" Int Jrnl of Nanomedicine: 2018:13 8487–8506 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6294055/pdf/ijn-13-8487.pdf

[52] Handy RD and Shaw BJ. "Toxic effects of nanoparticles and nanomaterials: Implications for public health, risk assessment and the public perception of nanotechnology" Health, Risk, & Society; Volume 9;2007. https://doi.org/10.1080/13698570701306807

[53] Cupaioli FA, Zucca FA, Boraschi D, Zecca L. Engineered nanoparticles. How brain friendly is this new guest? Prog Neurobiol. 2014 Aug-Sep;119-120:20-38. doi: 10.1016/j.pneurobio.2014.05.002. Epub 2014 May 10. PMID: 24820405. https://pubmed.ncbi.nlm.nih.gov/24820405/

[54] Society of Actuators (SoA) Research Institute. "Group Life COVID-19 Mortality Survey Report" Aug 2022. https://www.soa.org/4a368a/globalassets/assets/files/resources/research-report/2022/group-life-covid-19-mortality-03-2022-report.pdf

[55] The White House. "FACT SHEET: President Biden to Announce Goal to Administer at Least One Vaccine Shot to 70% of the U.S. Adult Population by July 4th." WH Statement made on May 4, 2021. https://www.whitehouse.gov/briefing-room/statements-releases/2021/05/04/fact-sheet-president-biden-to-announce-goal-to-

in the United States in 2022, and an excess of 64,375 deaths in the first 14 weeks (Q1) of 2023.[56]

(38) On November 20, 2020, Pfizer stated *in writing* that the risk-benefit ratio of their COVID-19 mRNA nanoparticle injections were *not favorable* (unfavorable) for children 12 to 15 years of age, based on FDA submitted data from 100 injected children from their Phase 3 trial.[57]

(39) On June 10, 2021, the FDA Vaccine Related and Biological Products Advisory Committee (VRBPAC) stated *in writing* that it would *be infeasible* (it would be *impossible*) to conduct a clinical trial that could clinically and statistically prove that any vaccine could prevent SARS-CoV-2 infection and/or COVID-19 disease in pediatric populations because

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administer-at-least-one-vaccine-shot-to-70-of-the-u-s-adult-population-by-july-4th/

[56] CDC Data submitted to the OECD (Organization for Economic Cooperation and Development) Accessed June 1, 2023. https://stats.oecd.org/index.aspx?queryid=104676#

[57] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; "Emergency Use Authorization (EUA) for an Unapproved Product." Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

teenagers, children, and infants rarely (if ever) become infected or present with symptoms.[58]

(40) Children who received two (2) COVID-19 injections are 1400% (15x) more likely to die of any cause than unvaccinated children and children who received three (3) COVID-19 injections are 4400% (45x) more likely to die of any cause than unvaccinated children per UK Government data.[59,60]

(41) COVID-19 mRNA nanoparticle injections induce anaphylaxis, appendicitis, fevers of greater than 104 degrees Fahrenheit, seizures (with eye rolling), convulsions, status epilepticus (seizures lasting more than 5 minutes and multiple seizures that can lead to permanent brain damage), epilepsy, exanthema subitum (herpes induced fevers and seizures), hypotonia (limp 'lifeless-like baby syndrome'), permanent brain damage confirmed by an EEG, and lissencephaly (genetic-induced brain

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[58] Vaccines and Related Biological Products Advisory Committee Meeting. Licensure and Emergency Use Authorization of Vaccines to Prevent COVID-19 for Use in Pediatric Populations" FDA Briefing Document;June 10, 2021. https://www.fda.gov/media/149935/download

[59] Nafilyan V., Bermingham C., Ward IL., et al. "Risk of death following SARS-CoV-2 infection or COVID-19 vaccination in young people in England: a self-controlled case series study" MedRxiv. March 23, 2022. https://www.medrxiv.org/content/10.1101/2022.03.22.22272775v1

[60] Swart N. "Pfizer documents, official government data confirm: .75m vaccine deaths in 2022, critical infertility" Biz News. Nov 23, 2022. https://www.biznews.com/health/2022/11/23/pfizer-vaccine-deaths

malformation characterized by the absence of convolutions/folds), per Pfizer's June 15, 2022, FDA clinical trial data submission of 6 month old babies through 4 year old toddlers; in which a subgroup of 370 toddlers (2 to 4 year old) only 21 toddlers (5%) made it to their 1-month study follow-up visit after receiving their 3rd COVID-19 mRNA injection, and in a subgroup of 344 babies (6 to 23 months old) only 3 babies (1%) made it to their 1-month study follow-up visit after receiving their 3rd injection of COVID-19 mRNA injection; reasons for discontinuing or withdrawing from the study included adverse events, neurological dysfunctions, ICU admission, hospitalization, and death (but reasons for discontinuation or withdrawal need not be noted by the investigator).[61]

(42) The engineered COVID-19 mRNA nanoparticles can cross the blood brain barrier causing demyelinating disease (deterioration to the protective covering of nerve cells) including permanent changes to nerve cell structures, nerve cell damage, and nerve cell death in the spinal cord and brain leading to permanent brain and neurological disorders and diseases,

[61] Pfizer-BioNTech COVID-19 Vaccine EUA Amendment for Use in Children 6 Months Through 4 Years of Age. "EUA amendment request for Pfizer-BioNTech COVID-19 Vaccine for use in children 6 months through 4 years of age." VRBPAC Briefing Document. June 15, 2022. https://www.fda.gov/media/159195/download

such as the **696,605 neurological disorders** and diseases documented by Pfizer.[62,63]

(43) The engineered mRNA nanoparticles cross the biological barriers of the male reproductive system accumulating in the testis and epididymis adversely affecting sexual health in men, including; sperm quality, quantity, morphology, and motility, and affecting male hormones causing reproductive organ dysfunction such as the **178,353 female and male reproductive system disorders** documented by Pfizer (including male erectile dysfunction, infertility, and testicular pain).[64]

(44) The engineered COVID-19 mRNA nanoparticles cross the biological barriers of the female reproductive system accumulating in the ovaries, placenta, and uterus, causing reproductive dysfunction including damage to

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[62] Cupaioli FA, Zucca FA, Boraschi D, Zecca L. Engineered nanoparticles. How brain friendly is this new guest? Prog Neurobiol. 2014 Aug-Sep;119-120:20-38. doi: 10.1016/j.pneurobio.2014.05.002. Epub 2014 May 10. PMID: 24820405. https://pubmed.ncbi.nlm.nih.gov/24820405/

[63] Pfizer. Confidential. "APPENDIX 2.2: Cumulative and Interval Summary Tabulation of Serious and Non-Serious Adverse Reactions from Post-Marketing Data Sources, BNT162B2." Data collected from December of 2020 through June 18, 2022. https://www.globalresearch.ca/wp-content/uploads/2023/05/pfizer-report.pdf

[64] Wang R, Song B, Wu J, et al. "Potential adverse effects of nanoparticles on the reproductive system" Int Jrnl of Nanomedicine: 2018:13 8487–8506 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6294055/pdf/ijn-13-8487.pdf

eggs and follicle development, and adversely affecting the health of women, unborn babies and newborn babies, as was demonstrated by the **178,353 female and male reproductive system disorders** and **4,056 pregnancy complications** (including heavy menstrual bleeding, irregular menstruation, spontaneous abortions, and infertility). [65]

(45) The engineered mRNA nanoparticle technologies in the COVID-19 injections are classified as electromagnetic devices per Pfizer's Operation Warp Speed contract and Adulterated Drugs and Devices 21 USC § 351**, and the 2017** FDA Guidance on Drugs and Devices**.**[66,67]

[65] Pfizer Inc. Japan. "SARS-CoV-2 mRNA Vaccine (BNT162, PF-07302048) Luciferase RNA-encapsulated LNP administered intravenously to Wistar Han rats at a dose of 1 mg RNA/kg. Plasma and Liver Concentrations of ALC-0315 and ALC-0159"

[66] DEPARTMENT OF THE ARMY U.S. ARMY CONTRACTING COMMAND. "COVID-19 PANDEMIC—LARGE SCALE VACCINE MANUFACTURING DEMONSTRATION" RPP #: 20-11 Project Identifier: 2011-003. Statement of Work for Pfizer. July 21, 2020. https://www.hhs.gov/sites/default/files/pfizer-inc-covid-19-vaccine-contract.pdf

[67] U.S. Department of Health and Human Services Food and Drug Administration, Office of Combination Products, Office of Special Medical Programs Office of the Commissioner "Classification of Products as Drugs and Devices & Additional Product Classification Issues: Guidance for Industry and FDA Staff" September 2017. https://www.fda.gov/media/80384/download

(46) The engineered nanoparticle technologies in COVID-19 mRNA injections are gene-editing technologies per Pfizer's May 18, 2021 FDA-submitted biological license application stating that the COVID-19 mRNA mechanism-of-action is through RNA transcription (nucleoside substitutions) substituting the genetic material of human cells within human bodies with foreign genetic material.[68,69,70]

(47) The engineered nanoparticle technologies in COVID-19 mRNA injections are gene-editing nanotechnologies that use cationic liposome technologies to alter human DNA through RNA transfection; as has been described in Pfizer's biological license application (BLA), on Pfizer's website; and as is demonstrated in multiple scientific papers and Pfizer's

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[68] U.S. Department of Health and Human Services Food and Drug Administration Center for Biologics Evaluation and Research. "Design and Analysis of Shedding Studies for Virus or Bacteria-Based Gene Therapy and Oncolytic Products" Guidance for Industry; Aug 2015. https://www.fda.gov/media/89036/download

[69] Berkman BE, Deputy Director, NHGI Bioethics Core. "SUBSTITUTION" NIH National Human Genome Institute Website visited June 27, 2023. https://www.genome.gov/genetics-glossary/Substitution#:~:text=Substitution%2C%20as%20related%20to%20genomics,with%20a%20different%20amino%20acid

[70] Ganguly P, PhD, Chief Communications Officer, Nucleus Genomics, NHGI Office of Communications. "TRANSLATION" NIH National Human Genome Institute Website https://www.genome.gov/genetics-glossary/Translation

internal report of 1,143 genetic diseases spontaneously reported post-COVID-19 mRNA nanoparticle injection.[71,72]

(48) It is an established scientific fact that the engineered nanoparticle technologies in the COVID-19 mRNA injections are gene-editing technologies with known and unknown risks for; integrating non-human DNA into the human genome, transmission of foreign DNA into the germline (genetic mutations passed from parent to child through sperm or egg), passage foreign genes into sperm, embryo/fetal and perinatal toxicity, genotoxicity (DNA damage that can lead to birth defects and diseases i.e. cancers), and the potential for horizontal transmission (i.e., shedding) is further confirmed in a June 9, 2023 peer-reviewed publication in the *International Journal of Molecular Science.*[73]

(49) The COVID-19 mRNA nanoparticle injections were *NEVER* proven to prevent infection, disease, hospitalization or death, per Pfizer's November

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[71] Pfizer Website. "Unlocking the Power of our Body's Protein Factory" Pfizer Websitevisited June 28, 2023. https://www.pfizer.com/news/behind-the-science/unlocking-power-our-bodys-protein-factory

[72] Pfizer Website. "Turning Your Body into Medicine Factories" Pfizer Website visited June 28, 2023. https://www.pfizer.com/news/behind-the-science/how-once-little-known-molecule-disrupting-medicine

[73] Banoun H. "mRNA: Vaccine or Gene Therapy? The Safety Regulatory Issues." International Journal of Molecular Sciences. 2023; 24(13):10514. https://doi.org/10.3390/ijms241310514

20, 2020, FDA submission, in which Pfizer stated *in writing* that out of 18,198 human subjects originally injected with BNT162b2, 11% or two-thousand and fifty-three (2,053) developed mild, moderate, or severe COVID-19 disease within 2 months of the 1st or 2nd mRNA nanoparticle injection.[74,75,76]

(50) In addition, 19 (0.1%) *deaths were reported by Pfizer within 3 days - 142 days* (less than 4 months) post- Pfizer mRNA nanoparticle injections in previously healthy human subjects per Pfizer's May 18, 2021, post-hoc analysis.[77]

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[74] Pfizer Inc., BioNTech, initial new drug (IND) application. "A PHASE 1/2/3, PLACEBO-CONTROLLED, RANDOMIZED, OBSERVER-BLIND, DOSE-FINDING STUDY TO EVALUATE THE SAFETY, TOLERABILITY, IMMUNOGENICITY, AND EFFICACY OF SARS-COV-2 RNA VACCINE CANDIDATES AGAINST COVID-19 IN HEALTHY INDIVIDUALS." PF-07302048 (BNT162 RNA-Based COVID-19 Vaccines) Protocol C4591001; Apr 2020.

[75] Gruber M. (Dir CBER/OVRR), Naik R., Smith M., Wollersheim S., Huang L., et al. Pfizer Inc. on behalf of Pfizer and BioNTech; "Emergency Use Authorization (EUA) for an Unapproved Product." Review Memorandum; Nov 2020. https://www.fda.gov/media/144416/download

[76] Polack F., Thomas S., Kitchin N., et al. for the C4591001 Clinical Trial Group; "Safety and Efficacy of the BNT162b2 mRNA Covid-19 Vaccine" New Engl J Med; Dec 10, 2020; 383:2603-2615. https://www.nejm.org/doi/full/10.1056/nejmoa2034577

[77] Naik Ramachandra, PhD, Review Committee Chair, DVRPA/OVRR. BioNTech Manufacturing GmbH in partnership with Pfizer, Inc. "COMIRNATY (BNT162b2): Active immunization to prevent coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome

(51) The rotovirus vaccine (RotaShield) was pulled off the US market in 1999 due to five cases (0.05%) of respiratory infection among 10,054 pediatric vaccine recipients.[78]

(52) The manufacturers of the COVID-19 nanoparticle injections *NEVER* submitted clinical trial evidence demonstrating clinically and statistically significant *protection against*; infection, symptomatic illness, medically attended illness, including emergency department and urgent care visits, or severe illness, including hospitalization and death, but did submit clinical data demonstrating an increased risk of heart inflammation, vaccine-related enhanced respiratory disease, and vaccine-related enhanced autoimmune diseases per Pfizer's August 23, 2021 FDA approval and Moderna's January 30, 2022, FDA approval.[79,80,81,]

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coronavirus 2 (SARS-CoV-2) in individuals 16 years of age and older" FDA Approval of the Biological License Application (BLA) for BNT162b2/COMIRNATY; Submitted May 18, 2021. Reviewed November 8, 2021. https://www.fda.gov/media/151733/download

[78] CDC "Monitoring Vaccine Effectiveness." CDC COVID-19 website visited June 29, 2023. https://www.cdc.gov/coronavirus/2019-ncov/vaccines/effectiveness/how-cdc-monitors.html

[79] COMIRNATY Prescribing Information. Purple Cap. Dilute. April 2023. https://labeling.pfizer.com/ShowLabeling.aspx?id=15623

[80] COMIRNATY Prescribing Information. Grey Cap. Do NOT dilute. April 2023. https://labeling.pfizer.com/ShowLabeling.aspx?id=16351

[81] SPIKEVAX Prescribing Information. November 2022. https://www.fda.gov/media/155675/download

(53) The engineered nanoparticles in the COVID-19 injections are nanotechnologies designed to force human cells to produce disease-causing pathogens known as spike proteins, spike proteins that are established lab-made pathogens that cause disease, disabilities, and death per dozens of scientific and clinical publications, abstracts, and patents as well as Pfizer's internal documents and website.[82,83,84,85,86, 87]

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[82] Graham B, McLellan J, and Ward A. "PREFUSION CORONAVIRUS SPIKE PROTEINS AND THEIR USE" US2020/06185A: Feb 27, 2020. https://patentimages.storage.googleapis.com/e9/1f/3a/c72f6a8603a3b5/US20200061185A1.pdf

[83] Arbeitman, C.R., Rojas, P., Ojeda-May, P. et al. The SARS-CoV-2 spike protein is vulnerable to moderate electric fields. Nat Commun 12, 5407 (2021). https://doi.org/10.1038/s41467-021-25478-7

[84] Cross R. "Without these lipid shells, there would be no mRNA vaccines for COVID-19" Chemical & Engineering News. March 6, 2021 | Volume 99, Issue 8; https://cen.acs.org/pharmaceuticals/drug-delivery/Without-lipid-shells-mRNA-vaccines/99/i8

[85] Moderna Patent. Bancel S, Chakraborty T, Fougorelles A, "Modified polynucleotides for the production of secreted proteins" US9828416B2: November 28, 2017. https://patents.google.com/patent/US9828416B2/en Also accessed February 2022 https://www.modernatx.com/sites/default/files/US10703789.pdf

[86] Rawati J, Kumar V, Ahlwat P, et al. "Current Trends on the Effects of Metal Based Nanoparticles on Microbial Ecology" Applied Biochemistry and Biotechnology; Feb 17, 2023. https://doi.org/10.1007/s12010-023-04386-0

[87] Acuitas Therapeutics. A Tissue Distribution Study of a [3H]-Labelled Lipid Nanoparticle-mRNA Formulation Containing ALC-0315 and ALC-0159 Following Intramuscular Administration in Wistar Han Rats. Test Facility Study No. 185350 Sponsor Reference No. ALC-NC-0552. Nov 9, 2020. https://phmpt.org/wp-content/uploads/2022/03/125742_S1_M4_4223_185350.pdf

(54) Dark field live blood analysis demonstrated that not only are hydrogels, which is programable matter, and evidence of nanotechnologies present in the blood of the vaccinated, they are also present in the blood of the unvaccinated due to shedding of this technology. Approximately 73% of embalmers are reporting white fibrous blood clots never seen before. These biosynthetic blood clots are now being seen in children and infants and appear to grow post mortem.[88,89,90]

(55) The engineered nanoparticle technologies (aka vaccine nanotechnology) in the COVID-19 mRNA injections are patented for use as

[88] Evidence of Crimes Against Humanity - Darkfield Blood Microscopy. Video Presentation of evidence by Ana Mihalcea, M.S., PhD. (August 2023) https://rumble.com/v307b3m-evidence-of-crimes-against-humanity-darkfield-blood-microscopy.html?mref=6zof&mrefc=2

[89] Interview by video with Retired Major Thomas Haviland via video (1/31/2024) National Survey of Embalmers. https://josephsansone.substack.com/p/major-thomas-haviland-on-mind-matters

[90] Interview by video with Embalmer Richard Hersman and Ana Mihalcea, M.D., PhD. (2/15/2024). White Fibrous Biosythetic Blood Clots https://anamihalceamdphd.substack.com/p/breaking-news-embalmer-richard-hirschman

a nanocarrier of an ‘agent of biowarfare,’ per US Patent Number 9539210, VACCINE NANOTECHNOLOGY.[91]

(56) COVID-19 injections containing engineered mRNA nanoparticle technologies meet the legal definition of biological weapons see Biological Weapons 18 USC § 175, **“which is a biological agent, toxin and/or delivery device for use *other* *than* prophylactic** (preventative), **protective, *bona fide* research, or other peaceful purpose.”**

(57) COVID-19 injections containing engineered mRNA nanoparticles meet the exact criteria of weapons of mass destruction see Weapons and Firearms § 790.166 Fla. Stat. (2023). A person who manufactures, possesses, sells, delivers, displays, uses, attempts to use, or conspires to use, or who makes readily accessible to others a weapon of mass destruction commits a felony of the first degree see Weapons and Firearms § 790.166 Fla. Stat. (2023).

[91] NIH Patent for “Vaccine Nanotechnology” US 9,539,210 B2 Jan 10, 2017. https://patentimages.storage.googleapis.com/29/d1/ca/18013ced0621f0/US9539210.pdf

(58) Distribution of Covid 19 injections containing mRNA nanoparticles constitute an act of treason see Federal Crime of Treason 18 USC § 2381, see Treason § 876.32 Fla. Stat. (2023).

(59) Distribution of Covid 19 injections containing mRNA nanoparticles qualify as an act of terrorism as see Domestic Terrorism, 18 USC § 2331; and, see Terrorism § 775.30 Fla. Stat. (2023).

(60) Distribution of Covid 19 injections containing mRNA nanoparticles constitute genocide crimes see Genocide 18 USC §1091.

(61) Death via Covid 19 injections containing mRNA nanoparticles is the unlawful killing of a human being and constitutes murder see Murder § 782.04 (1)(a) Fla. Stat. (2023).

(62) Government officials that maintain or assist the principal or an accessory or provide any other aid, knowing that the offender had committed a crime and such crime was a third degree felony with the intent that the offender avoids or escapes detection, arrest, trial, or punishment, is an accessory after the fact see Accessory After the Fact § 777.03 Fla. Stat. (2023).

(63) Studies estimate that COVID 19 injections caused over 17 million dead worldwide, and estimates as high as approximately 600 million dead and injured worldwide.[92,93]

(64) Doctors are reporting that in addition to other diseases and disorder, that the evidence indicates that all those that received COVID injections have developed an autoimmune disease.[94]

IV. Nature of Relief Sought.

(65) Since February 21, 2023, approximately 10 Florida Republican County Political Parties, representing millions of people, have passed resolutions declaring Covid 19 injections and mRNA injections biological

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[92] Rancourt, Denis G., et al. "COVID-19 vaccine-associated mortality in the Southern Hemisphere." Correlation Research in the Public Interest, Ontario, Canada (2023). https://correlation-canada.org/covid-19-vaccine-associated-mortality-in-the-southern-hemisphere/

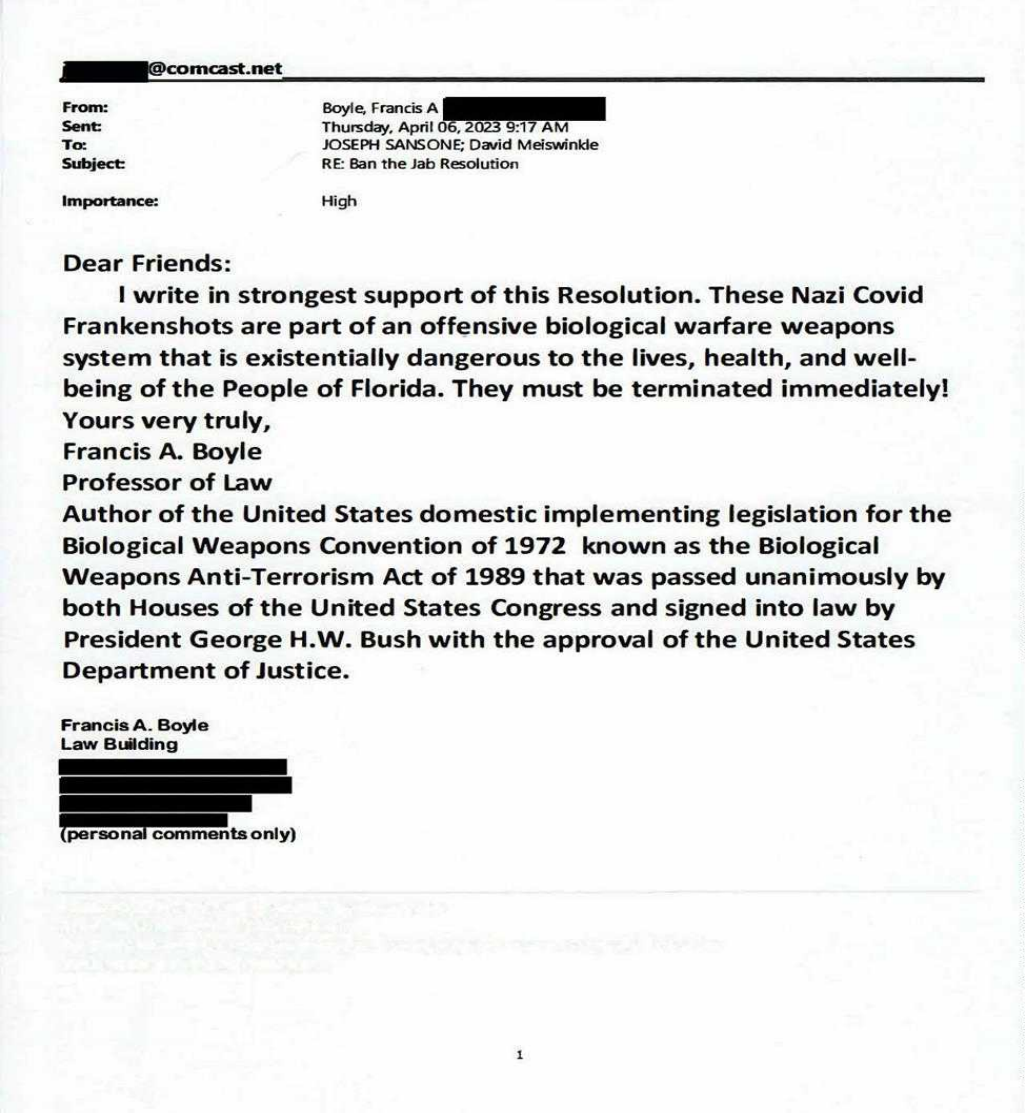
[93] Interview by video with James Thorp, M.D., (December 12, 2023) Estimates, approximately 602 million dead and injured worldwide. https://twitter.com/VigilantFox/status/1734759332804182240?ref_src=twsrc%5Etfw%7Ctwcamp%5Etweetembed%7Ctwterm%5E1734759332804182240%7Ctwgr%5E980b9b38569b4af0897c416131d590360dbbf08d%7Ctwcon%5Es1_&ref_url=https%3A%2F%2Fwww.thegatewaypundit.com%2F2023%2F12%2Fthis-is-unprecedented-injury-kill-ratio-dr-james%2F90

[94] Interview by video with Marivic Villa, M.D. Status of 2000 plus COVID 19 injection patients/victims (12/27/2023) https://josephsansone.substack.com/p/dr-villa-on-mind-matters-and-everything

and technological weapons, calling on the Governor to prohibit their distribution, and the Attorney General to confiscate the vials and conduct a forensic analysis of their contents.[95]

(66) This resolution was endorsed on April 6, 2023, by Dr. Francis Boyle, author of the United States domestic implementing legislation for the Biological Weapons Convention of 1972, known as the Biological Weapons and Anti-Terrorism Act of 1989, that was passed unanimously by both Houses of the United States Congress, and signed into law by President George H.W. Bush, with the approval of the United States Department of Justice.

[95] GOP 'Ban the Jab Resolution'. Originally passed February 21, 2023. Authored by Joseph Sansone, M.S., PhD https://josephsansone.substack.com/p/ban-the-jab-resolution-f6f



[redacted]@comcast.net

From: Boyle, Francis A [redacted]
Sent: Thursday, April 06, 2023 9:17 AM
To: JOSEPH SANSONE; David Meiswinkle
Subject: RE: Ban the Jab Resolution

Importance: High

Dear Friends:

I write in strongest support of this Resolution. These Nazi Covid Frankenshots are part of an offensive biological warfare weapons system that is existentially dangerous to the lives, health, and well-being of the People of Florida. They must be terminated immediately!

Yours very truly,
Francis A. Boyle
Professor of Law
Author of the United States domestic implementing legislation for the Biological Weapons Convention of 1972 known as the Biological Weapons Anti-Terrorism Act of 1989 that was passed unanimously by both Houses of the United States Congress and signed into law by President George H.W. Bush with the approval of the United States Department of Justice.

Francis A. Boyle
Law Building
[redacted]
(personal comments only)

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(67) On approximately October 6, 2023, the Governor Ron DeSantis and Attorney General Ashley Moody received demand letters via certified mail, along with a flash drive, providing ample evidence of a public health hazard and potential law violations to prohibit the distribution of Covid 19 injections and mRNA injections, as well as evidence of potential crimes to be

investigated. In addition, all 20 State Attorneys, and all 67 County Sheriffs were provided this evidence via email.

(68) On February 16, 2024, Governor DeSantis and Attorney General Ashly Moody received demand letters via certified mail requiring immediate action to enforce the aforementioned laws and to prohibit the distribution of these weapons of mass destruction in the State of Florida in order to protect the public.[96]

(69) On behalf of the dead and dying, and those that will die in the future, and for the preservation of the human race, the Petitioner seeks an order of mandamus, requiring the Respondents to immediately prohibit the distribution, promotion, access and administration of COVID-19 injections, mRNA nanoparticle injections, and all mRNA products in the State of Florida. This mandamus also seeks to compel Respondents to seize all vials of Covid 19 injections and all mRNA nanoparticle injections, and conduct a forensic analysis of their contents.

[96] (See Appendix A)

V. **Argument.**

(70) Article V, § 3(b)(8) Florida Constitution authorizes this Court to issue writs of mandamus. The jurisdiction of this Court to issue such writs is limited to actions involving "state officers and state agencies." Art. V, § 3(b)(8), Florida Constitution. The Governor and Attorney General are state officers subject to this Court's jurisdiction under Article V, § 3(b)(8). See, e.g., Flack v. Graham, 453 So. 2d 819 (Fla. 1984). One seeking a writ of mandamus must demonstrate a clear legal right to the performance of a clear legal duty by a public officer, and that no other remedies are available. Hatten v. State, 561 So. 2d 562,563 (Fla. 1990). Mandamus is used to compel the exercise of ministerial duties, which are defined as duties "positively imposed by law to be performed at a time and in a manner or upon conditions which are specifically designated by the law itself absent any authorization of discretion to the agency." See Solomon v. Sanitarians' Registration Bd., 155 So. 2d 353,356 (Fla. 1963).

(71) see Weapons and Firearms § 790.166 Fla. Stat. (2023) a weapon of mass destruction means “any device or object that is designed or intended to cause death or serious bodily injury to any human or animal, or severe emotional or mental harm to any human, through the release,

dissemination, or impact of toxic or poisonous chemicals, or their precursors;" or "Any device or object involving a biological agent", which is defined as "any microorganism, virus, infectious substance, or biological product that may be engineered through biotechnology, or any naturally occurring or bioengineered component of any such microorganism, virus, infectious substance, or biological product, capable of causing death, disease, or other biological malfunction in a human."

(72) The facts of this case evidenced above demonstrate nanotechnology present in the COVID-19 injections which do qualify as a device designed and intended to cause harm, as does the use of such technology, and or a biological agent, resulting in death or harm. Repeatedly distributing a biological agent or device causing harm in mass, especially after it is well known to cause harm, qualifies it as a weapon of mass destruction and a biological weapon. It also meets the threshold of a biological weapon see Biological Weapons 18 USC § 175 those that "knowingly develops, produces, stockpiles, transfers, acquires, retains, or possesses any biological agent, toxin, or delivery system for use as a weapon" "for use as a weapon" includes the development, production, transfer, acquisition, retention, or possession of any biological agent, toxin, or delivery system for other than prophylactic, protective, bona fide research, or other peaceful

purposes.” Continuing to distribute nanoparticle injections despite causing harm qualifies these injections as biological and technological weapons of mass destruction.

(73) This position is indisputable. If the State of Florida, or State Officers, continue to allow the distribution of these biological agents then that makes the State of Florida and State Officers an accessary after the fact. see Accessory After the Fact § 777.03 Fla. Stat. (2023) as such an action would allow the crime to continue and shield perpetrators from prosecution. The Florida Department of Health has called for a halt to these injections because they are known to cause harm, including death, and even pose a threat to the human genome. This adds a tremendous amount of credence to the arguments put forth in this mandamus.

(74) See Treason § 876.32 Fla. Stat. (2023) “Treason against the state shall consist only in levying war against the same, or in adhering to the enemies thereof, or giving them aid and comfort. Whoever commits treason against this state shall be guilty of a felony of the first degree.”

(75) Targeting Floridians with a biological agent and devices that causes harm in mass, including death and disability, meets the threshold of treason. This is biological warfare against the human population residing in

Florida. The facts of this case clearly demonstrate that these injections cause harm and the fact that the Florida Department of Health is calling for a halt to these mRNA injections due to the fact that they do in fact cause harm, means that allowing the carnage to continue is allowing warfare against Floridians to continue. This also violates the federal treason law, see Federal Crime of Treason 18 USC § 2381 as citizens of Florida are also American citizens. Pharmaceutical executives and government officials are not excluded from such laws.

(76) See Terrorism § 775.30 Fla. Stat. (2023) terrorism is defined as "a violent act or an act dangerous to human life which is a violation of the criminal laws of this state or of the United States" and also includes, "Intimidate, injure, or coerce a civilian population" also, "Influence the policy of a government by intimidation or coercion; or "affect the conduct of government through destruction of property, assassination, murder". See Domestic Terrorism, 18 USC § 2331.

(77) Once it was known that these injections cause harm, continuing to promote them and distribute them is an act of terrorism. Pfizer's 5.3.6 Cumulative Analysis of Post-Authorization Adverse Events[97] from the early

[97] 5.3.6 Cumulative Analysis Of Post-Authorization Adverse Event

rollout of the COVID-19 nanoparticle injections, from December 1, 2020, through Feb. 28, 2021, demonstrated approximately 1300 serious adverse reactions, 42,000 adverse cases, 158,000 adverse incidents, and 1223 deaths. This information was made available to the federal government by April of 2021 and to the public via court order in November of 2021.

(78) The body of evidence presented in the facts of this case clearly shows a pattern of intimidation with the intent to harm the civilian population. This Court is well aware that fear and coercion have been used to promote these COVID-19 nanoparticle injections.

(79) See Genocide 18 USC §1091, "Whoever, whether in time of peace or in time of war and with the specific intent to destroy, in whole or in substantial part, a national, ethnic, racial, or religious group" meets the criteria of genocide. The whole of the United States was targeted thus the whole nation was targeted. This qualifies these crimes as acts of genocide.

(80) See Murder § 782.04 (1)(a) Fla. Stat. (2023) "the unlawful killing of a human being" constitutes murder. Every single human being that died or

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Reports Of PF-07302048 (BNT162B2) Received Through 28-FEB-2021 https://phmpt.org/wp-content/uploads/2021/11/5.3.6-postmarketing-experience.pdf

will die as a result of these lethal injections is in fact a murder victim according to Florida law. There was never informed consent, there was coercion, and deception. Once deception was used, and the fact that informed consent was not present, and the fact that once it became clear that these injections were in fact harmful and their distribution continued, there was an intent to harm and resulting deaths constitute murder.

(81) To shield a perpetrator from prosecution or to knowingly allow the crime to continue makes one an accessary after the fact, public officials and state officers are not immune from this law. See Accessory After the Fact § 777.03 Fla. Stat. (2023).

(82) Floridians have been baited like insects. Dark field live blood analysis demonstrated that not only are hydrogels and self assembling nanotechnologies present in the blood of the vaccinated, they are also present in the blood of the unvaccinated due to shedding of this technology.[98]

[98] Evidence of Crimes Against Humanity - Darkfield Blood Microscopy. Video Presentation of evidence by Ana Mihalcea, M.S., PhD. (August 2023) https://rumble.com/v307b3m-evidence-of-crimes-against-humanity-darkfield-blood-microscopy.html?mref=6zof&mrefc=2

(83) This mandamus is not demanding the prosecution of individuals. This mandamus simply seeks to compel the Governor and Attorney General to enforce the law and protect the public from biological and technological weapons of mass destruction and remove them from the market.

(84) According to the Florida Department of Health, based on statements made in an official bulletin by Surgeon Dr. General Ladapo, COVID 19 injections are an adulterated product. See Florida Drugs and Cosmetic Act § 499.005 (2) Fla. Stat. (2023). COVID 19 injections continue to be distributed. COVID 19 injections cause death and permanent disability, and according to the Florida Department of Health, pose an existential threat to the human genome itself. There is a clear duty for the Respondents to enforce this law.

(85) To be entitled to a writ of mandamus the petitioner must have a clear legal right to the requested relief, the respondent must have an indisputable legal duty to perform the requested action, and the petitioner must have no other adequate remedy available. See Turner v. Singletary, 623 So.2d 537, 538 (Fla. 1st DCA 1993).

(86) The Governor is the supreme executive power in the state. Article IV, § 1(a) Florida Constitution provides that “The supreme executive power

shall be vested in a governor, who shall be commander-in-chief of all military forces of the state not in active service of the United States. **The governor shall take care that the laws be faithfully executed**, commission all officers of the state and counties, and transact all necessary business with the officers of government." Furthermore, Article IV, § 4(b) Florida Constitution provides that "The attorney general shall be the chief state legal officer".

(87) It is the sworn duty of both the Governor and the Attorney General to uphold the laws of the State of Florida. Both respondents have an indisputable legal duty to perform the demanded action.

(88) As already demonstrated, the petitioner has presented evidence and demanded action from the Governor and Attorney General and also presented evidence to 67 County Sheriffs and 20 State Attorneys and has not received a reply. The petitioner is left with no other legal remedy.

(89) The petitioner has a clear legal right to the requested relief. The petitioner has a right to be free from being targeted with biological and technological weapons of mass destruction, and to seek relief on behalf of the millions of Floridians that have been targeted, including friends and family members.

(90) Florida's Statewide Grand Jury empaneled by this Court only has the power of indictment and presentment.[99] While the Statewide Grand Jury may indict, this power is directed toward uncovering specific criminal acts and identifying individuals that committed these acts. The Statewide Grand Jury can only issue a recommendation. It does not have the power to prohibit the distribution of Covid 19 injection and mRNA injections, which are weapons of mass destruction.

(91) The body of evidence presented in this mandamus makes it clear that extraordinary crimes have been committed. This mandamus is not seeking an indictment of any individuals or entities or naming specific individuals responsible for these crimes. The intent of this mandamus is to compel the Governor and Attorney General to enforce these laws and stop the aforementioned crimes from continuing, to protect the public.

(92) Due to the extraordinary nature of the crimes perpetrated against the citizens of Florida, this court is obligated to compel the Governor and the Attorney General to enforce the aforementioned laws, and immediately

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[99] FIRST INTERIM REPORT OF THE TWENTY SECOND STATEWIDE GRAND JURY https://www.flgov.com/wp-content/uploads/2024/02/22nd-Statewide-Grand-Jury-First-Interim-Report.pdf

protect the public from further harm by prohibiting the possession or distribution of all COVID-19 injections, COVID-19 nanoparticle injections, mRNA nanoparticle injections, and mRNA products in the state of Florida. On behalf of the preservation of the human race, we plead to the Court to not allow the murder and maiming of Floridians to continue, and not to allow the further contamination of the human genome to continue. We plead to the Court to grant this mandamus immediately.

(93) This petitioner respectfully requests oral arguments and a hearing if necessary to establish the facts of this case.

CERTIFICATE OF SERVICE

I CERTIFY that a copy hereof has been furnished by e-service to Ryan Dean Newman, General Counsel, Executive Office of the Governor, ryan.newman@eog.myflorida.com; gov.legal@eog.myflorida.com, counsel for Respondent Ron DeSantis and Attorney General Ashley Moody, citizenservices@myfloridalegal.com, ashley.moody@myfloridalegal.com on this 3rd day of March, 2024.

CERTIFICATE OF COMPLIANCE WITH RULE 9.045

I CERTIFY that this petition complies with the font (Arial 14-point) and word-count requirements. This filing contains 11,671 words (including sections permitted to be excluded), which is within the 13,000 word-limit prescribed in Fla. R. App. P. 9.100(g)

Dated: March 3, 2024

Respectfully submitted,

Joseph Sansone, Petitioner

27499 Riverview Center Blvd.

Bonita Springs, FL 34134

239-444-1774
DrJosephSansone@Gmail.com

APPENDIX A

Demand Letters and Mail Receipts to Governor and Attorney General

Honorable Governor Ron DeSantis

400 S. Monroe St.,

Tallahassee, FL 32399-0001

Governor DeSantis,

On October 6, 2023, I have previously written to you, providing evidence of the harm, calling for you to immediately halt the distribution of Covid 19/mRNA injections in the state of Florida. I had also previously emailed you a few times, each time providing evidence to support the demand.

As you are aware, since February of 2023, approximately 10 Florida Republican County Parties, representing millions of people, have passed resolutions declaring Covid 19/mRNA injections to be biological and technological weapons, and have called on you to prohibit the distribution of these injections, and for the Attorney General to confiscate the vials and conduct forensic analysis of their contents.

Since then, on December 6, 2023, the Florida Department of Health has called for a halt to mRNA injections in the State of Florida due to multiple health hazards. Surgeon General, Dr. Ladapo, in public statements has went as far as saying these injections are the Anti-Christ of drugs.

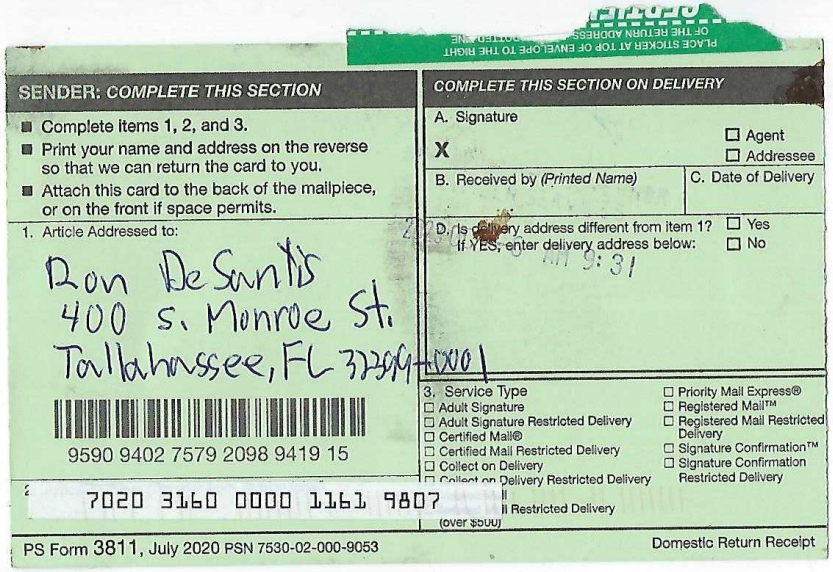
Covid injections and mRNA injections violate State and Federal biological weapons laws, (18 USC 175, Ch. 10: BIOLOGICAL WEAPONS; F.S.790.166); Violate Treason Laws (18 U.S. Code § 238; F.S., 876.32) Terrorism laws, (18 USC 2331 (5), USA Patriot Act; F.S. 775.30) and the federal genocide law (18 USC Chapter 50A).

As Governor you are required to take immediate action to enforce the aforementioned laws, protect the public, and prohibit the distribution of these weapons of mass destruction in the State of Florida. Anything less at this point in time will make you complicit. Consider this a demand for immediate action.

Respectfully, 2/7/2024

Joseph Sansone, M.S., PhD

[redacted]



SENDER: COMPLETE THIS SECTION

- Complete items 1, 2, and 3.
- Print your name and address on the reverse so that we can return the card to you.
- Attach this card to the back of the mailpiece, or on the front if space permits.

1. Article Addressed to:

Ron DeSantis
400 S. Monroe St.
Tallahassee, FL 32399-0001

9590 9402 7579 2098 9419 15

7020 3160 0000 1161 9807

COMPLETE THIS SECTION ON DELIVERY

A. Signature
X
☐ Agent
☐ Addressee

B. Received by (Printed Name)

C. Date of Delivery

D. Is delivery address different from item 1? ☐ Yes
If YES, enter delivery address below: ☐ No

3. Service Type
☐ Adult Signature
☐ Adult Signature Restricted Delivery
☐ Certified Mail®
☐ Certified Mail Restricted Delivery
☐ Collect on Delivery
☐ Collect on Delivery Restricted Delivery
☐ Priority Mail Express®
☐ Registered Mail™
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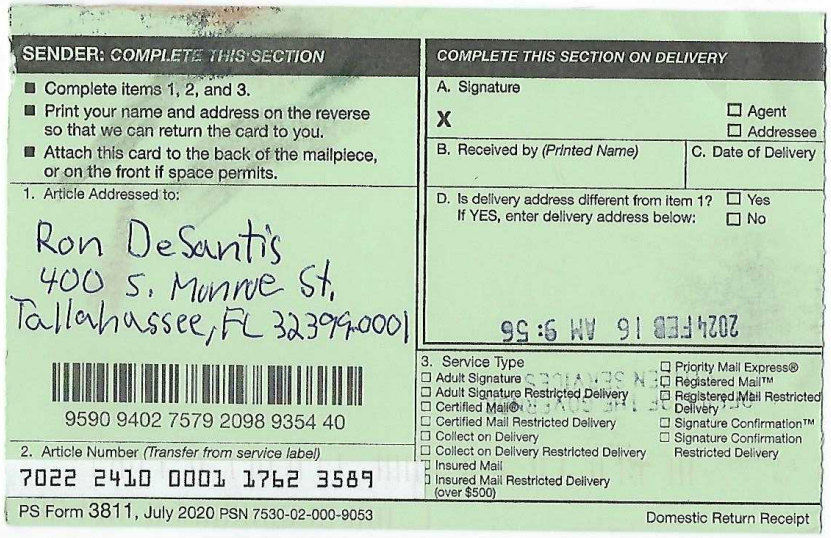
☐ Signature Confirmation™
☐ Signature Confirmation Restricted Delivery

PS Form 3811, July 2020 PSN 7530-02-000-9053 Domestic Return Receipt



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- Complete items 1, 2, and 3.
- Print your name and address on the reverse so that we can return the card to you.
- Attach this card to the back of the mailpiece, or on the front if space permits.

1. Article Addressed to:

Ron DeSantis
400 S. Monroe St.
Tallahassee, FL 32399-0001

9590 9402 7579 2098 9354 40

2. Article Number (Transfer from service label)
7022 2410 0001 1762 3589

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☐ Addressee

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2024 FEB 16 AM 9:56

3. Service Type
☐ Adult Signature
☐ Adult Signature Restricted Delivery
☐ Certified Mail®
☐ Certified Mail Restricted Delivery
☐ Collect on Delivery
☐ Collect on Delivery Restricted Delivery
☐ Insured Mail
☐ Insured Mail Restricted Delivery (over $500)
☐ Priority Mail Express®
☐ Registered Mail™
☐ Registered Mail Restricted Delivery
☐ Signature Confirmation™
☐ Signature Confirmation Restricted Delivery

PS Form 3811, July 2020 PSN 7530-02-000-9053 Domestic Return Receipt

Attorney General Ashley Moody,

PL-01, The Capital

Tallahassee, FL 32399-1050

Attorney General Ashley Moody,

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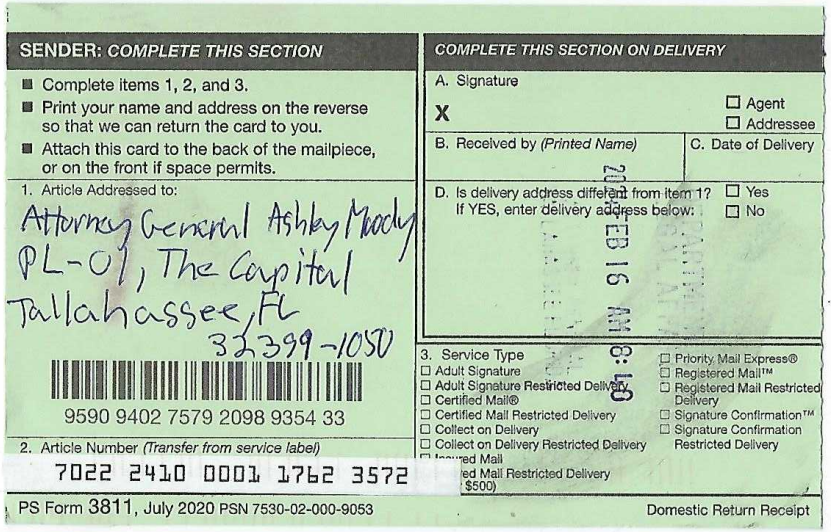
Covid injections and mRNA injections violate State and Federal biological weapons laws, (18 USC 175, Ch. 10: BIOLOGICAL WEAPONS; F.S.790.166); Violate Treason Laws (18 U.S. Code § 238; F.S., 876.32) Terrorism laws, (18 USC 2331 (5), USA Patriot Act; F.S. 775.30) and the federal genocide law (18 USC Chapter 50A).

As Attorney General you are required to take immediate action to enforce the aforementioned laws, protect the public, and prohibit the distribution of these weapons of mass destruction in the State of Florida. Anything less at this point in time will make you complicit. Consider this a demand for immediate action.

Respectfully, 2/7/2024

Joseph Sansone, M.S., PhD

[redacted]



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1. Article Addressed to:

Attorney General Ashley Moody
PL-01, The Capitol
Tallahassee, FL
32399-1050

9590 9402 7579 2098 9354 33

2. Article Number *(Transfer from service label)*

7022 2410 0001 1762 3572

COMPLETE THIS SECTION ON DELIVERY

A. Signature

X ☐ Agent ☐ Addressee

B. Received by *(Printed Name)* | C. Date of Delivery

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3. Service Type
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- ☐ Adult Signature Restricted Delivery
- ☐ Certified Mail®
- ☐ Certified Mail Restricted Delivery
- ☐ Collect on Delivery
- ☐ Collect on Delivery Restricted Delivery
- ☐ Priority Mail Express®
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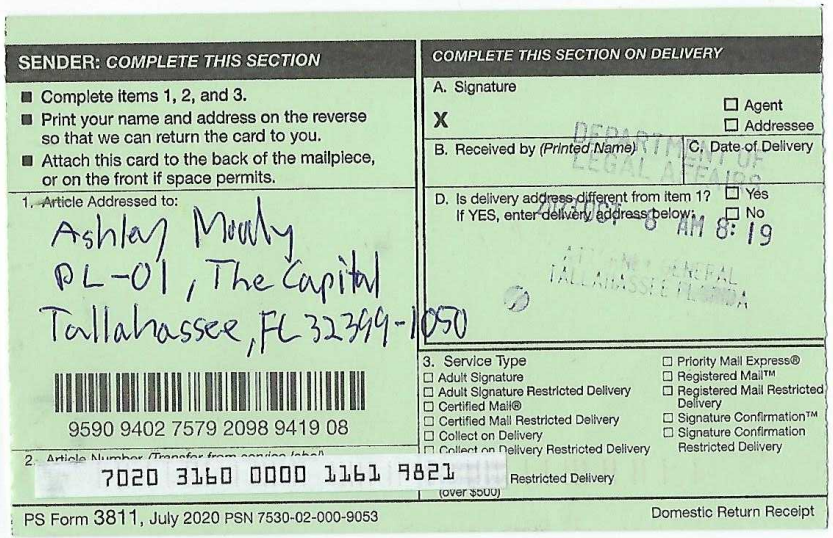
- ☐ Signature Confirmation Restricted Delivery

PS Form 3811, July 2020 PSN 7530-02-000-9053 Domestic Return Receipt



SENDER: *COMPLETE THIS SECTION*

- Complete items 1, 2, and 3.
- Print your name and address on the reverse so that we can return the card to you.
- Attach this card to the back of the mailpiece, or on the front if space permits.

1. Article Addressed to:

Ashley Moody
PL-01, The Capitol
Tallahassee, FL 32399-1050

9590 9402 7579 2098 9419 08

2. Article Number *(Transfer from service label)*

7020 3160 0000 1161 9821

COMPLETE THIS SECTION ON DELIVERY

A. Signature

X ☐ Agent ☐ Addressee

B. Received by *(Printed Name)* | C. Date of Delivery

DEPARTMENT OF LEGAL AFFAIRS

D. Is delivery address different from item 1? ☐ Yes
If YES, enter delivery address below: ☐ No

2020 SEP 8 AM 8:19

ATTORNEY GENERAL TALLAHASSEE FLORIDA

3. Service Type
- ☐ Adult Signature
- ☐ Adult Signature Restricted Delivery
- ☐ Certified Mail®
- ☐ Certified Mail Restricted Delivery
- ☐ Collect on Delivery
- ☐ Collect on Delivery Restricted Delivery
- ☐ Priority Mail Express®
- ☐ Registered Mail™
- ☐ Registered Mail Restricted Delivery
- ☐ Signature Confirmation™
- ☐ Signature Confirmation Restricted Delivery

Restricted Delivery (over $500)

PS Form 3811, July 2020 PSN 7530-02-000-9053 Domestic Return Receipt