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The Vaccinated Can Be Patented (Owned)

In a court case in 2013 Pathology v Myriad Genetics, inc, in the United States the Supreme Court ruled that you cannot patent human DNA as it was "a product of nature". But at the end of the ruling the Supreme Court did rule that if you were to change a humans genome by mRNA vaccines (which are being used currently) then the genome can be patented.

This means that everyone who has had the vaccine is now technically 'patented' and something that is patented is 'owned' and will come under the definition of 'trans human'.

Those people that are legally identified as 'trans human' do not have access to Human Rights or any rights provided by the State. This is because they are not classed as 100% organic or human.

Therefore, technically anyone having this vaccine could no longer have any access to human rights. There have been a few legal papers discussing this recently, so clarification should be available on this soon.

https://www.supremecourt.gov/opinions/12pdf/12-398_1b7d.pdf

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(12) **United States Patent**
De Fougerolles et al.

(10) **Patent No.: US 10,703,789 B2**
(45) **Date of Patent: *Jul. 7, 2020**

(54) **MODIFIED POLYNUCLEOTIDES FOR THE PRODUCTION OF SECRETED PROTEINS**

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.

This patent is subject to a terminal disclaimer.

(21) Appl. No.: **16/438,978**

(22) Filed: **Jun. 12, 2019**

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Related U.S. Application Data

(63) Continuation of application No. 14/987,328, filed on Jan. 4, 2016, now Pat. No. 10,385,106, which is a (Continued)

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A61K 9/127	(2006.01)
C07K 14/535	(2006.01)
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C07K 14/47	(2006.01)
A61K 31/7088	(2006.01)
C07K 19/00	(2006.01)
C12N 15/85	(2006.01)
A61K 38/18	(2006.01)
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A61K 47/10	(2017.01)
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CPC ***C07K 14/535*** (2013.01); ***A61K 9/1271*** (2013.01); ***A61K 9/1272*** (2013.01); ***A61K 9/1277*** (2013.01); ***A61K 9/14*** (2013.01); ***A61K 9/5031*** (2013.01); ***A61K 31/7088*** (2013.01); ***A61K 38/1767*** (2013.01); ***A61K 38/1816*** (2013.01); ***A61K 38/1866*** (2013.01); ***A61K 38/191*** (2013.01); ***A61K 38/193*** (2013.01); ***A61K 38/212*** (2013.01); ***A61K 38/215*** (2013.01); ***A61K 38/36*** (2013.01); ***A61K 38/363*** (2013.01); ***A61K 38/44*** (2013.01); ***A61K 38/4833*** (2013.01); ***A61K 38/4846*** (2013.01); ***A61K 39/3955*** (2013.01); ***A61K 47/10*** (2013.01); ***A61K 47/54*** (2017.08); ***A61K 47/542*** (2017.08); ***A61K 48/0033*** (2013.01); ***A61K 48/0066*** (2013.01); ***A61K 48/0075*** (2013.01); ***C07K 14/47*** (2013.01); ***C07K 14/475*** (2013.01); ***C07K 14/505*** (2013.01); ***C07K 14/525*** (2013.01); ***C07K 14/56*** (2013.01); ***C07K 14/565*** (2013.01); ***C07K 14/745*** (2013.01); ***C07K 14/75*** (2013.01); ***C07K 16/2887*** (2013.01); ***C07K 16/32*** (2013.01); ***C07K 19/00*** (2013.01); ***C12N 9/0069*** (2013.01); ***C12N 9/644*** (2013.01); ***C12N 15/85*** (2013.01); ***C12N 15/88*** (2013.01); ***C12Y 113/12007*** (2013.01); ***C12Y 304/21005*** (2013.01); ***C12Y 304/21022*** (2013.01); ***A61K 9/0019*** (2013.01); ***A61K 48/00*** (2013.01); *C12N 2840/00* (2013.01)

(58) **Field of Classification Search**

CPC C07H 21/02; C12N 15/67; C12N 15/11

See application file for complete search history.

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(57) **ABSTRACT**

A pharmaceutical composition which has a plurality of lipid nanoparticles that has a mean particle size of between 80 nm and 160 nm and contains a modified mRNA encoding a polypeptide. The lipid nanoparticles include a cationic lipid, a neutral lipid, a cholesterol, and a PEG lipid. The mRNA contains a 5'-cap, 5'-UTR, N1-methyl-pseudouridine, a 3'-UTR, and a poly-A region with at least 100 nucleotides.

14 Claims, 14 Drawing Sheets

Specification includes a Sequence Listing.

tured herein have morpholino backbone structures of the above-referenced U.S. Pat. No. 5,034,506.

Modifications at the 2' position may also aid in delivery. Preferably, modifications at the 2' position are not located in a polypeptide-coding sequence, i.e., not in a translatable region. Modifications at the 2' position may be located in a 5'UTR, a 3'UTR and/or a tailing region. Modifications at the 2' position can include one of the following at the 2' position: H (i.e., 2'-deoxy); F; O—, S—, or N-alkyl; O—, S—, or N-alkenyl; O—, S- or N-alkynyl; or O-alkyl-O-alkyl, wherein the alkyl, alkenyl and alkynyl may be substituted or unsubstituted C_1 to C_{10} alkyl or C_2 to C_{10} alkenyl and alkynyl. Exemplary suitable modifications include $O[(CH_2)_nO]_mCH_3$, $O(CH_2)_nOCH_3$, $O(CH_2)_nNH_2$, $O(CH_2)_nCH_3$, $O(CH_2)_nONH_2$, and $O(CH_2)_nON[(CH_2)_nCH_3)]_2$, where n and m are from 1 to about 10. In other embodiments, the polynucleotides, primary constructs or mmRNA include one of the following at the 2' position: C_1 to C_{10} lower alkyl, substituted lower alkyl, alkaryl, aralkyl, O-alkaryl or O-aralkyl, SH, SCH_3, OCN, Cl, Br, CN, CF_3, OCF_3, $SOCH_3$, SO_2CH_3, ONO_2, NO_2, N_3, NH_2, heterocycloalkyl, heterocycloalkaryl, aminoalkylamino, polyalkylamino, substituted silyl, an RNA cleaving group, a reporter group, an intercalator, a group for improving the pharmacokinetic properties, or a group for improving the pharmacodynamic properties, and other substituents having similar properties. In some embodiments, the modification includes a 2'-methoxyethoxy (2'-O—$CH_2CH_2OCH_3$, also known as 2'-O-(2-methoxyethyl) or 2'-MOE) (Martin et al., *Helv. Chin. Acta*, 1995, 78:486-504) i.e., an alkoxy-alkoxy group. Another exemplary modification is 2'-dimethylaminooxyethoxy, i.e., a $O(CH_2)_2ON(CH_3)_2$ group, also known as 2'-DMAOE, as described in examples herein below, and 2'-dimethylaminoethoxyethoxy (also known in the art as 2'-O-dimethylaminoethoxyethyl or 2'-DMAEOE), i.e., 2'-O—CH_2—O—CH_2—$N(CH_2)_2$, also described in examples herein below. Other modifications include 2'-methoxy (2'-OCH_3), 2'-aminopropoxy (2'-$OCH_2CH_2CH_2NH_2$) and 2'-fluoro (2'-F). Similar modifications may also be made at other positions, particularly the 3' position of the sugar on the 3' terminal nucleotide or in 2'-5' linked dsRNAs and the 5' position of 5' terminal nucleotide. Polynucleotides of the invention may also have sugar mimetics such as cyclobutyl moieties in place of the pentofuranosyl sugar. Representative U.S. patents that teach the preparation of such modified sugar structures include, but are not limited to, U.S. Pat. Nos. 4,981,957; 5,118,800; 5,319,080; 5,359,044; 5,393,878; 5,446,137; 5,466,786; 5,514,785; 5,519,134; 5,567,811; 5,576,427; 5,591,722; 5,597,909; 5,610,300; 5,627,053; 5,639,873; 5,646,265; 5,658,873; 5,670,633; and 5,700,920 and each of which is herein incorporated by reference.

In still other embodiments, the polynucleotide, primary construct, or mmRNA is covalently conjugated to a cell penetrating polypeptide. The cell-penetrating peptide may also include a signal sequence. The conjugates of the invention can be designed to have increased stability; increased cell transfection; and/or altered the biodistribution (e.g., targeted to specific tissues or cell types).

In one embodiment, the polynucleotides, primary constructs or mmRNA may be conjugated to an agent to enhance delivery. As a non-limiting example, the agent may be a monomer or polymer such as a targeting monomer or a polymer having targeting blocks as described in International Publication No. WO2011062965, herein incorporated by reference in its entirety. In another non-limiting example, the agent may be a transport agent covalently coupled to the polynucleotides, primary constructs or mmRNA of the present invention (See e.g., U.S. Pat. Nos. 6,835,393 and 7,374,778, each of which is herein incorporated by reference in its entirety). In yet another non-limiting example, the agent may be a membrane barrier transport enhancing agent such as those described in U.S. Pat. Nos. 7,737,108 and 8,003,129, each of which is herein incorporated by reference in its entirety.

In another embodiment, polynucleotides, primary constructs or mmRNA may be conjugated to SMARTT POLYMER TECHNOLOGY® (PHASERX®, Inc. Seattle, Wash.).

Self-Assembled Nanoparticles

Nucleic Acid Self-Assembled Nanoparticles

Self-assembled nanoparticles have a well-defined size which may be precisely controlled as the nucleic acid strands may be easily reprogrammable. For example, the optimal particle size for a cancer-targeting nanodelivery carrier is 20-100 nm as a diameter greater than 20 nm avoids renal clearance and enhances delivery to certain tumors through enhanced permeability and retention effect. Using self-assembled nucleic acid nanoparticles a single uniform population in size and shape having a precisely controlled spatial orientation and density of cancer-targeting ligands for enhanced delivery. As a non-limiting example, oligonucleotide nanoparticles were prepared using programmable self-assembly of short DNA fragments and therapeutic siRNAs. These nanoparticles are molecularly identical with controllable particle size and target ligand location and density. The DNA fragments and siRNAs self-assembled into a one-step reaction to generate DNA/siRNA tetrahedral nanoparticles for targeted in vivo delivery. (Lee et al., Nature Nanotechnology 2012 7:389-393; herein incorporated by reference in its entirety).

In one embodiment, the polynucleotides, primary constructs and/or mmRNA disclosed herein may be formulated as self-assembled nanoparticles. As a non-limiting example, nucleic acids may be used to make nanoparticles which may be used in a delivery system for the polynucleotides, primary constructs and/or mmRNA of the present invention (See e.g., International Pub. No. WO2012125987; herein incorporated by reference in its entirety).

In one embodiment, the nucleic acid self-assembled nanoparticles may comprise a core of the polynucleotides, primary constructs or mmRNA disclosed herein and a polymer shell. The polymer shell may be any of the polymers described herein and are known in the art. In an additional embodiment, the polymer shell may be used to protect the polynucleotides, primary constructs and mmRNA in the core.

Polymer-Based Self-Assembled Nanoparticles

Polymers may be used to form sheets which self-assembled into nanoparticles. These nanoparticles may be used to deliver the polynucleotides, primary constructs and mmRNA of the present invention. In one embodiment, these self-assembled nanoparticles may be microsponges formed of long polymers of RNA hairpins which form into crystalline 'pleated' sheets before self-assembling into microsponges. These microsponges are densely-packed sponge like microparticles which may function as an efficient carrier and may be able to deliver cargo to a cell. The microsponges may be from 1 um to 300 nm in diameter. The microsponges may be complexed with other agents known in the art to form larger microsponges. As a non-limiting example, the microsponge may be complexed with an agent to form an outer layer to promote cellular uptake such as polycation polyethyleneime (PEI). This complex can form a 250-nm

DO YOU KNOW WHAT'S IN A VACCINE?

NONE OF THESE SHOULD BE INJECTED INTO YOUR BODY

Aluminum
Known to cause brain damage at all doses, linked to ALZHEIMER'S DISEASE, dementia, seizures, autoimmune issues, SIDs and cancer. This toxin accumulates in the brain and causes more damage with each dose.

Beta-Propiolactone
Known to cause CANCER. Suspected gastrointestinal, liver, nerve and respiratory, skin and sense organ POISON.

Gentamicin Sulphate & Polymyxin B [antibiotics]
ALLERGIC reactions can range from mild to life-threatening.

Genetically Modified Yeast, Animal, Bacterial and Viral DNA
Can be incorporated into the recipient's DNA and cause unknown GENETIC MUTATIONS.

Glutaraldehyde
Poisonous if ingested. Causes BIRTH DEFECTS in animals.

Formaldehyde [formalin]
Known to cause CANCER in humans. Probable gastrointestinal, liver, respiratory, immune, nerve and reproductive system POISON. Banned from injectables in most European countries.

Human and Animal Cells
Human DNA from aborted BABIES. Pig blood, horse blood, rabbit brains, dog kidneys, cow hearts, monkey kidneys, chick embryos, calf serum, sheep blood & more. Linked to childhood leukemia and diabetes.

Mercury [thimerosal]
One of the most toxic substances known. Even if a thermometer breaks, the building is cleared and HAZMAT is called. Tiny doses cause damage to the brain, gut, liver, bone marrow, nervous system and/or kidneys. Linked to autoimmune disorders, and neurological disorders like AUTISM.

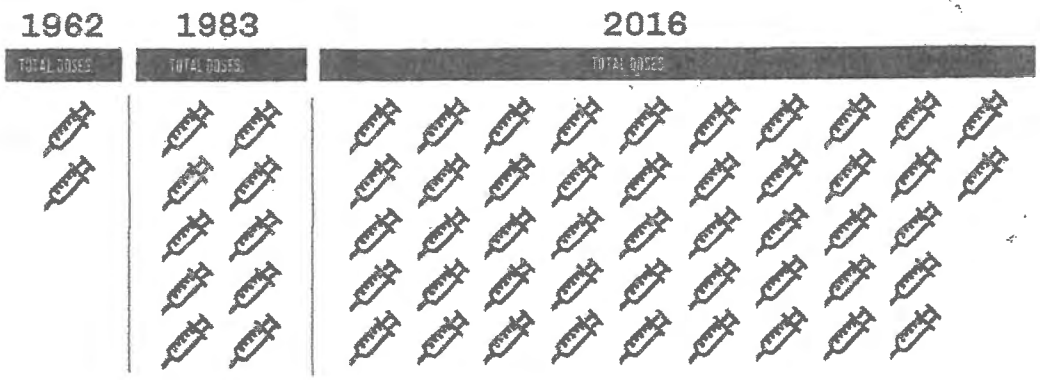
Monosodium Glutamate [MSG]
A toxic chemical that is linked to birth defects, developmental delays and infertility. Banned in Europe.

Neomycin Sulphate [antibiotic]
Interferes with vitamin B6 absorption which can lead to epilepsy and brain damage. Allergic reactions can range from mild to life-threatening.

Phenol/Phenoxyethanol [2-PE]
Used as anti-freeze. TOXIC to all cells and capable of destroying the immune system.

Polysorbate 80 & 20
Known to cause CANCER in animals and linked to numerous autoimmune issues and infertility.

VACCINES DOSES for U.S. CHILDREN



The US gives 2-3x more vaccines than most developed countries, yet we have the sickest population -- with skyrocketing rates of health issues like asthma, childhood diabetes, food allergies, leukemia, developmental delays, ADHD, autism, lupus, arthritis, eczema, epilepsy, brain tumors, Alzheimer's and more. ***It's NOT a coincidence.***

In 1986, Pharmaceutical manufacturers producing vaccines were freed from ALL liability resulting from vaccine injury or death by the Childhood Vaccine Injury Act. With this, vaccines became HIGHLY profitable. There are 271 vaccines in development and mandatory vaccine laws for children — and ADULTS — being pushed in most states.

Help us raise awareness by supporting the Learn The Risk campaign.

Learn more at **LearnTheRisk.org**

Vaccine Excipient & Media Summary

Excipients Included in U.S. Vaccines, by Vaccine

This table includes not only vaccine ingredients (e.g., adjuvants and preservatives), but also substances used during the manufacturing process, including vaccine-production media, that are removed from the final product and present only in trace quantities.
In addition to the substances listed, most vaccines contain Sodium Chloride (table salt).

Last Updated February 2015

All reasonable efforts have been made to ensure the accuracy of this information, but manufacturers may change product contents before that information is reflected here. If in doubt, check the manufacturer's package insert.

Vaccine	Contains	Source: Manufacturer's P.I. Dated
Adenovirus	sucrose, D-mannose, D-fructose, dextrose, potassium phosphate, plasdone C, anhydrous lactose, micro crystalline cellulose, polacrilin potassium, magnesium stearate, cellulose acetate phthalate, alcohol, acetone, castor oil, FD&C Yellow #6 aluminum lake dye, human serum albumin, fetal bovine serum, sodium bicarbonate, human-diploid fibroblast cell cultures (WI-38), Dulbecco's Modified Eagle's Medium, monosodium glutamate	March 2011
Anthrax (Biothrax)	aluminum hydroxide, benzethonium chloride, formaldehyde, amino acids, vitamins, inorganic salts and sugars	May 2012
BCG (Tice)	glycerin, asparagine, citric acid, potassium phosphate, magnesium sulfate, Iron ammonium citrate, lactose	February 2009
DT (Sanofi)	aluminum potassium sulfate, peptone, bovine extract, formaldehyde, thimerosal (trace), modified Mueller and Miller medium, ammonium sulfate	December 2005
DTaP (Daptacel)	aluminum phosphate, formaldehyde, glutaraldehyde, 2-Phenoxyethanol, Stainer-Scholte medium, modified Mueller's growth medium, modified Mueller-Miller casamino acid medium (without beef heart infusion), dimethyl 1-beta-cyclodextrin, ammonium sulfate	October 2013
DTaP (Infanrix)	formaldehyde, glutaraldehyde, aluminum hydroxide, polysorbate 80, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium	November 2013
DTaP-IPV (Kinrix)	formaldehyde, glutaraldehyde, aluminum hydroxide, Vero (monkey kidney) cells, calf serum, lactalbumin hydrolysate, polysorbate 80, neomycin sulfate, polymyxin B, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium	November 2013
DTaP-HepB-IPV (Pediarix)	formaldehyde, gluteraldehyde, aluminum hydroxide, aluminum phosphate, lactalbumin hydrolysate, polysorbate 80, neomycin sulfate, polymyxin B, yeast protein, calf serum, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium, Vero (monkey kidney) cells	November 2013
DTaP-IPV/Hib (Pentacel)	aluminum phosphate, polysorbate 80, formaldehyde, sucrose, gutaraldehyde, bovine serum albumin, 2-phenoxethanol, neomycin, polymyxin B sulfate, Mueller's Growth Medium, Mueller-Miller casamino acid medium (without beef heart infusion), Stainer-Scholte medium (modified by the addition of casamino acids and dimethyl-beta-cyclodextrin), MRC-5 (human diploid) cells, CMRL 1969 medium (supplemented with calf serum), ammonium sulfate, and medium 199	October 2013
Hib (ActHIB)	ammonium sulfate, formalin, sucrose, Modified Mueller and Miller medium	January 2014
Hib (Hiberix)	formaldehyde, lactose, semi-synthetic medium	March 2012
Hib (PedvaxHIB)	aluminum hydroxphosphate sulfate, ethanol, enzymes, phenol, detergent, complex fermentation medium	December 2010

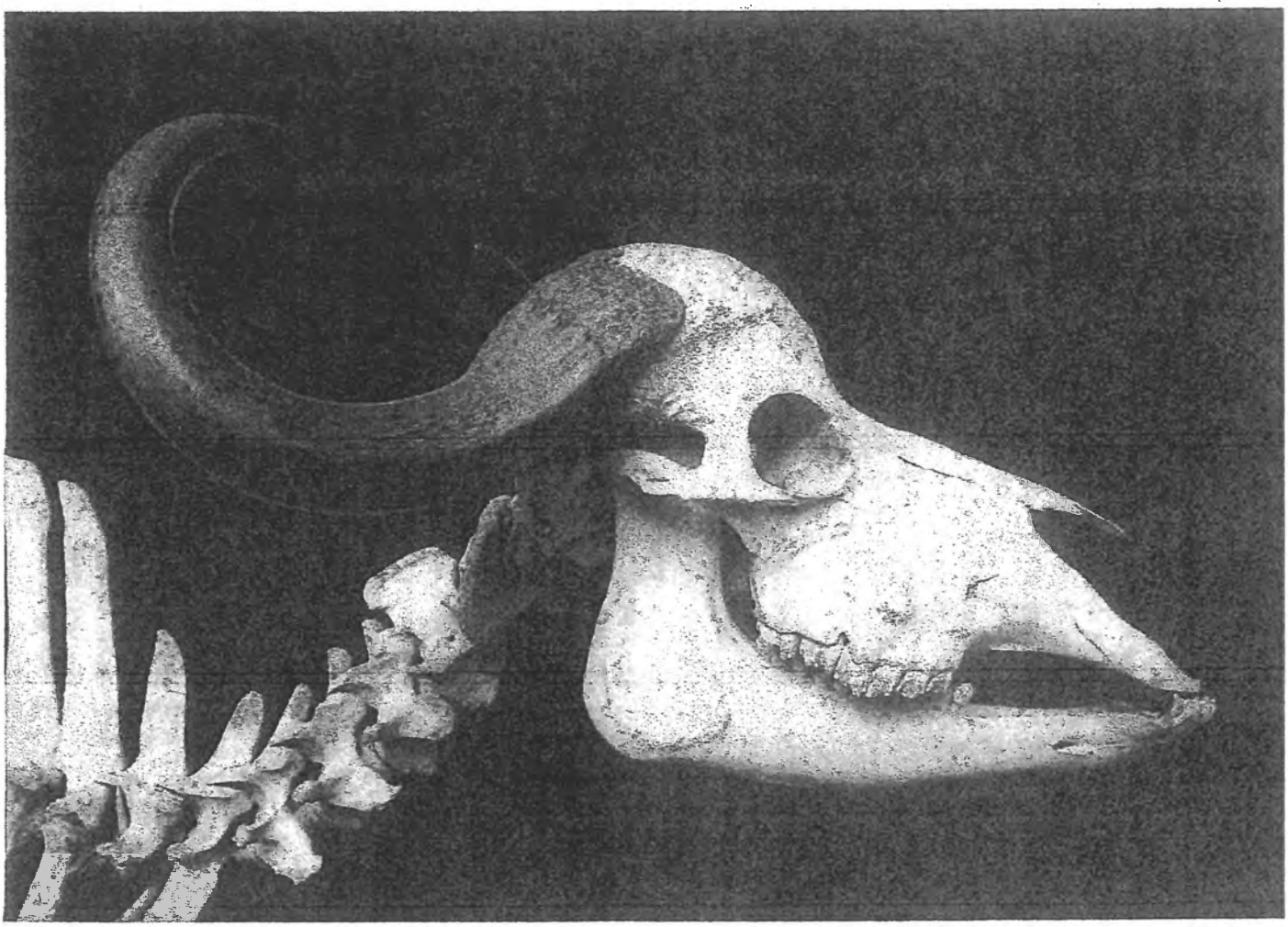
Vaccine	Contains	Source: Manufacturer's P.I. Dated
Influenza (FluMist) Quadrivalent	ethylene diamine tetraacetic acid (EDTA), monosodium glutamate, hydrolyzed porcine gelatin, arginine, sucrose, dibasic potassium phosphate, monobasic potassium phosphate, gentamicin sulfate, egg protein	July 2013
Japanese Encephalitis (Ixiaro)	aluminum hydroxide, Vero cells, protamine sulfate, formaldehyde, bovine serum albumin, sodium metabisulphite, sucrose	May 2013
Meningococcal (MCV4-Menactra)	formaldehyde, phosphate buffers, Mueller Hinton agar, Watson Scherp media, Modified Mueller and Miller medium, detergent, alcohol, ammonium sulfate	April 2013
Meningococcal (MCV4-Menveo)	formaldehyde, amino acids, yeast extract, Franz complete medium, CY medium	August 2013
Meningococcal (MPSV4-Menomune)	thimerosal (multi-dose vial only), lactose, Mueller Hinton casein agar, Watson Scherp media, detergent, alcohol	April 2013
Meningococcal (MenB – Bexsero)	aluminum hydroxide, *E. coli*, histidine, sucrose, deoxycholate, kanomycin	2015
Meningococcal (MenB – Trumenba)	polysorbate 80, histodine, *E. coli*, fermentation growth media	October 2015
MMR (MMR-II)	Medium 199 (vitamins, amino acids, fetal bovine serum, sucrose, glutamate) , Minimum Essential Medium, phosphate, recombinant human albumin, neomycin, sorbitol, hydrolyzed gelatin, chick embryo cell culture, WI-38 human diploid lung fibroblasts	June 2014
MMRV (ProQuad)	sucrose, hydrolyzed gelatin, sorbitol, monosodium L-glutamate, sodium phosphate dibasic, human albumin, sodium bicarbonate, potassium phosphate monobasic, potassium chloride, potassium phosphate dibasic, neomycin, bovine calf serum, chick embryo cell culture, WI-38 human diploid lung fibroblasts, MRC-5 cells	March 2014
Pneumococcal (PCV13 – Prevnar 13)	casamino acids, yeast, ammonium sulfate, Polysorbate 80, succinate buffer, aluminum phosphate, soy peptone broth	January 2014
Pneumococcal (PPSV-23 – Pneumovax)	phenol	May 2014
Polio (IPV – Ipol)	2-phenoxyethanol, formaldehyde, neomycin, streptomycin, polymyxin B, monkey kidney cells, Eagle MEM modified medium, calf serum protein, Medium 199	May 2013
Rabies (Imovax)	Human albumin, neomycin sulfate, phenol red indicator, MRC-5 human diploid cells, beta-propriolactone	April 2013
Rabies (RabAvert)	β-propiolactone, potassium glutamate, chicken protein, egg protein, neomycin, chlortetracycline, amphotericin B, human serum albumin, polygeline (processed bovine gelatin), sodium EDTA, bovine serum	March 2012
Rotavirus (RotaTeq)	sucrose, sodium citrate, sodium phosphate monobasic monohydrate, sodium hydroxide, polysorbate 80, cell culture media, fetal bovine serum, vero cells *[DNA from porcine circoviruses (PCV) 1 and 2 has been detected in RotaTeq. PCV-1 and PCV-2 are not known to cause disease in humans.]*	June 2013
Rotavirus (Rotarix)	amino acids, dextran, sorbitol, sucrose, calcium carbonate, xanthan, Dulbecco's Modified Eagle Medium (potassium chloride, magnesium sulfate, ferric (III) nitrate, sodium phosphate, sodium pyruvate, D-glucose, concentrated vitamin solution, L-cystine, L-tyrosine, amino acids solution, L-glutamine, calcium chloride, sodium hydrogenocarbonate, and phenol red). *[Porcine circovirus type 1 (PCV-1) is present in Rotarix. PCV-1 is not known to cause disease in humans.]*	May 2014
Smallpox (Vaccinia – ACAM2000)	human serum albumin, mannitol, neomycin, glycerin, polymyxin B, phenol, Vero cells, HEPES	September 2009

B

LETHAL INJECTIONS: 18% of cattle DIE immediately following mRNA "vaccination"

Friday, October 21, 2022 by: Ethan Huff
Tags: animal health, Cattle, chemical violence, COVID, death, food supply, mRNA, new vaccines, vaccines, world agriculture

This article may contain statements that reflect the opinion of the author



(Natural News) Much of the conversation surrounding mRNA (messenger RNA) "vaccines" centers around their impact on humans how about all the animals that are being injected with it?

Believe it or not, cattle are reportedly now getting jabbed with the stuff, which in a recent mass "vaccination" campaign of an Austra herd resulted in 35 of the 200 animals dying *immediately*.

We are told that dairy farmers and others are now being *forced* to inject their animals for the Fauci Flu in order to remain in busine and that the animals are not responding well to it.

Just like in humans, the shots are causing such profound damage that many of the animals are succumbing to *instant death*, while others are getting sick and dying over a longer period of time. (Related: mRNA spike proteins linger in the heart and brain long afte injection.)

For the animals that survive, one wonders what is becoming of their milk, which gets passed on as food for other animals as well a humans. Is it safe to consume mRNA-tainted milk and cheese from a "fully vaccinated" dairy cow? The answer is *probably not*.

"Dairy herd DNA is altered," one report explains. "Milk is altered and you consume it! Butter constitution, yoghurt, and cheese is alt meat is altered – will chicken and other meats be next?

the great worldwide BABY BOOM. It was the culmination of all man's efforts to survive through history. It was modern medicine, better diets, heat in winter, pure running water, and proper disposal of sewage. It was the point in history when the birth rate so exceeded the death rate that the world's population doubled between 1957 and 1990. It was the most wonderful time in the history of the world, but it was also the worst. It signaled the end of man's most precious achievement. An alliance of all of the powers on earth, open and hidden, decided that individual freedoms could no longer be tolerated in the interest of the preservation of the human race. They believed the common man could not be trusted.

What had been the unfulfilled dream of many individual groups became reality by the concentration of power in the alliance known as the Bilderberg Group. What had been impossible before was now promised. The New World Order that so many had envisioned was now a certainty.

The first study was made during World War II to determine the impact of the returning soldiers upon the economy. The results mobilized the ruling elite. A second secret study was conducted in 1957 by scientists meeting in Huntsville, Alabama. It confirmed the results of the first. The conclusion was that civilization as we know it would collapse shortly after the year 2000 unless the population was seriously curtailed. The study expressed a concern that since atomic weapons existed they would ultimately be used. Total worldwide disarmament was urged. Congress adopted the disarmament plan and created the U.S. Disarmament Agency. President Dwight David Eisenhower had this to say in 1957: "As a result of lowered infant mortality, longer lives, and the accelerating conquest of famine there is under way a population explosion so incredibly great that in little more than another generation the population of the world is expected to double."

A third study was made by the Club of Rome ending in 1968 to determine the limits to growth. The result was the same. The Club of Rome was commissioned to develop a computer model of the world so as to predict the outcome of corrections made to social and economic structures by the elect. The Club of Rome was also asked to develop a computer model of a New World Order. Both tasks were accomplished.

Studies were done to determine a method to arrest the population explosion before the point of no return would be reached. It was determined that an immediate attack on the problem would involve two points of intervention. The first was to lower the birth rate and the second was to increase the death rate.

To lower the birth rate several programs were put into motion. The first was the development of positive birth-control methods using

mechanical (diaphragm and condom), chemical (foam and birth-control pills), and medical (sterilization, abortion, and hysterectomy) procedures. These were developed and implemented. The Women's Liberation movement was started with the demand for free abortions, using "pro choice" as its rallying cry. Homosexuality was encouraged and Gay Liberation was born. Homosexuals do not have children. Zero population growth became a hot subject at cocktail parties. Individual freedom, "the heat of the moment," religion, and the old blue laws sabotaged these efforts, and while zero population growth became a reality in some areas, population increased rapidly in others.

The only alternative left to the world's ruling elite was to increase the death rate. This was a difficult thing to do, as no one wanted to pick people out of a crowd and line them up for execution. Neither did they relish the possible consequences of an enraged public upon discovering that they were being systematically murdered. Of course, a very short but very deadly global war using nuclear weapons upon select population concentrations was contemplated and, to tell you the truth, was not ruled out. The fact that such a population control was even contemplated confirmed the worst fears of those who had participated in the 1957 study. War was put on the back burner to simmer, but may become a reality. In the meantime something else had to be done that would absolve the decision makers of guilt and place the blame on those who did not lead clean lives. Something that could be blamed upon Mother Nature. What was needed was the bubonic plague or some other horrible but natural disease. The answer came from Rome.

Several Top Secret recommendations were made by Dr. Aurelio Peccei of the Club of Rome. He advocated that a plague be introduced that would have the same effect as the famous Black Death of history. The chief recommendation was to develop a microbe which would attack the auto immune system and thus render the development of a vaccine impossible. The orders were given to develop the microbe and to develop a prophylactic and a cure. The microbe would be used against the general population and would be introduced by vaccine. The prophylactic was to be used by the ruling elite. The cure will be administered to the survivors when it is decided that enough people have died. The cure will be announced as newly developed when in fact it has existed from the beginning. This plan is a part of Global 2000. The prophylactic and the cure are suppressed.

"Man has skyrocketed from a defensive position, largely subordinate to Nature's alternatives, to a new and dominant one. From it he not only can and does influence everything in the world but, voluntarily or unwittingly, can and indeed does determine the alternatives of his own future –

Appendix B

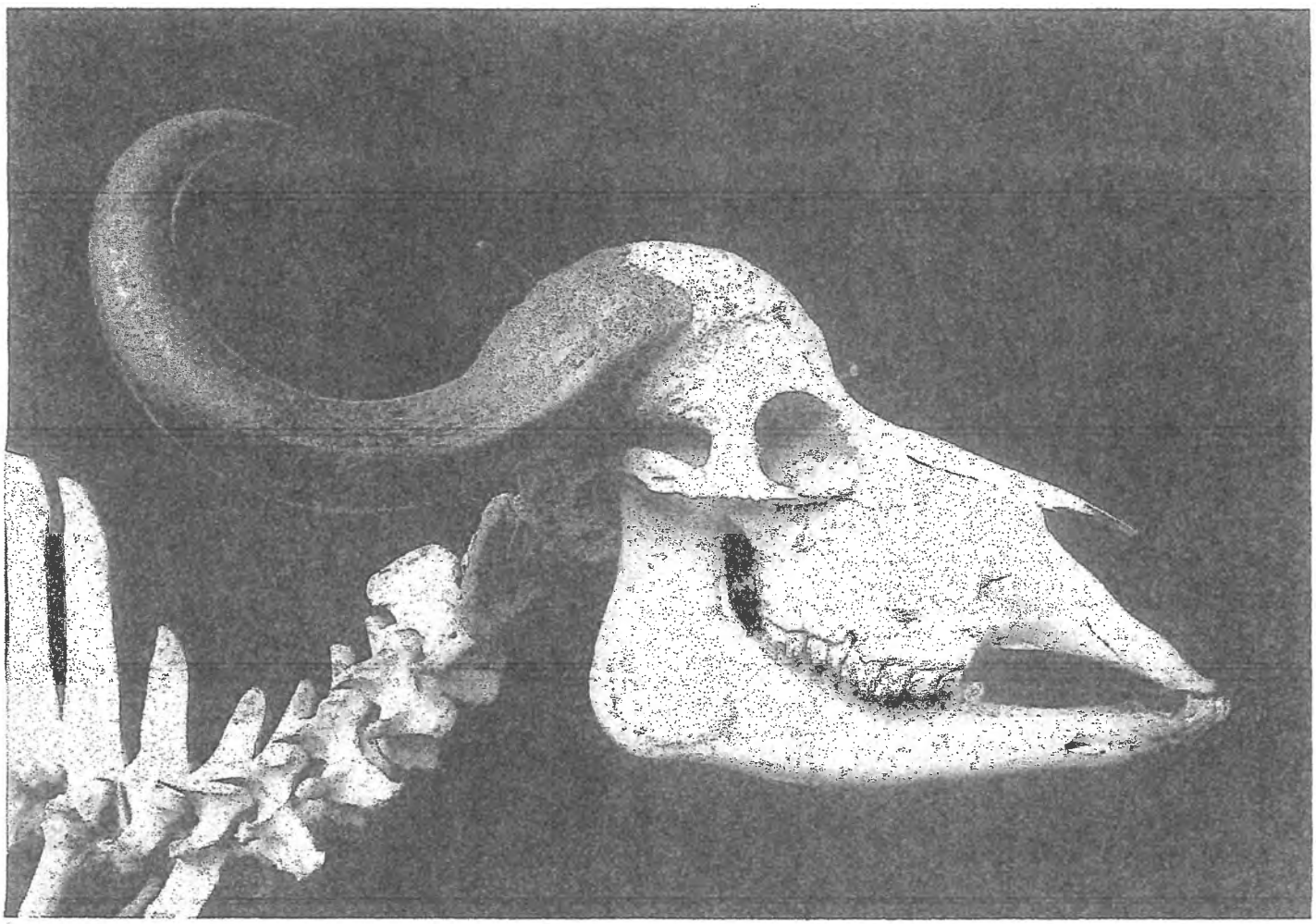
Vaccine	Contains	Source: Manufacturer's P.I. Dated
Hib/Hep B (Comvax)	yeast (vaccine contains no detectable yeast DNA), nicotinamide adenine dinucleotide, hemin chloride, soy peptone, dextrose, mineral salts, amino acids, formaldehyde, potassium aluminum sulfate, amorphous aluminum hydroxyphosphate sulfate, sodium borate, phenol, ethanol, enzymes, detergent	December 2010
Hib/Mening. CY (MenHibrix)	tris (trometamol)-HCl, sucrose, formaldehyde, synthetic medium, semi-synthetic medium	2012
Hep A (Havrix)	aluminum hydroxide, amino acid supplement, polysorbate 20, formalin, neomycin sulfate, MRC-5 cellular proteins	December 2013
Hep A (Vaqta)	amorphous aluminum hydroxyphosphate sulfate, bovine albumin, formaldehyde, neomycin, sodium borate, MRC-5 (human diploid) cells	February 2014
Hep B (Engerix-B)	aluminum hydroxide, yeast protein, phosphate buffers, sodium dihydrogen phosphate dihydrate	December 2013
Hep B (Recombivax)	yeast protein, soy peptone, dextrose, amino acids, mineral salts, potassium aluminum sulfate, amorphous aluminum hydroxyphosphate sulfate, formaldehyde, phosphate buffer	May 2014
Hep A/Hep B (Twinrix)	formalin, yeast protein, aluminum phosphate, aluminum hydroxide, amino acids, phosphate buffer, polysorbate 20, neomycin sulfate, MRC-5 human diploid cells	August 2012
Human Papillomavirus (HPV) (Cerverix)	vitamins, amino acids, lipids, mineral salts, aluminum hydroxide, sodium dihydrogen phosphate dehydrate, 3-O-desacyl-4' Monophosphoryl lipid A, insect cell, bacterial, and viral protein	November 2013
Human Papillomavirus (HPV) (Gardasil)	yeast protein, vitamins, amino acids, mineral salts, carbohydrates, amorphous aluminum hydroxyphosphate sulfate, L-histidine, polysorbate 80, sodium borate	June 2014
Human Papillomavirus (HPV) (Gardasil 9)	yeast protein, vitamins, amino acids, mineral salts, carbohydrates, amorphous aluminum hydroxyphosphate sulfate, L-histidine, polysorbate 80, sodium borate	December 2014
Influenza (Afluria)	beta-propiolactone, thimerosol (multi-dose vials only), monobasic sodium phosphate, dibasic sodium phosphate, monobasic potassium phosphate, potassium chloride, calcium chloride, sodium taurodeoxycholate, neomycin sulfate, polymyxin B, egg protein, sucrose	December 2013
Influenza (Agriflu)	egg proteins, formaldehyde, polysorbate 80, cetyltrimethylammonium bromide, neomycin sulfate, kanamycin, barium	2013
Influenza (Fluarix) Trivalent and Quadrivalent	octoxynol-10 (Triton X-100), α-tocopheryl hydrogen succinate, polysorbate 80 (Tween 80), hydrocortisone, gentamicin sulfate, ovalbumin, formaldehyde, sodium deoxycholate, sucrose, phosphate buffer	June 2014
Influenza (Flublok)	monobasic sodium phosphate, dibasic sodium phosphate, polysorbate 20, baculovirus and host cell proteins, baculovirus and cellular DNA, Triton X-100, lipids, vitamins, amino acids, mineral salts	March 2014
Influenza (Flucelvax)	Madin Darby Canine Kidney (MDCK) cell protein, MDCK cell DNA, polysorbate 80, cetyltrimethlyammonium bromide, β-propiolactone, phosphate buffer	March 2014
Influenza (Fluvirin)	nonylphenol ethoxylate, thimerosal (multidose vial–trace only in prefilled syringe), polymyxin, neomycin, beta-propiolactone, egg proteins, phosphate buffer	February 2014
Influenza (Flulaval) Trivalent and Quadrivalent	thimerosal, formaldehyde, sodium deoxycholate, egg proteins, phosphate buffer	February 2013
Influenza (Fluzone: Standard (Trivalent and Quadrivalent), High-Dose, & Intradermal)	formaldehyde, octylphenol ethoxylate (Triton X-100), gelatin (standard trivalent formulation only), thimerosal (multi-dose vial only) , egg protein, phosphate buffers, sucrose	2014

B

LETHAL INJECTIONS: 18% of cattle DIE immediately following mRNA "vaccination"

Friday, October 21, 2022 by: Ethan Huff
Tags: animal health, Cattle, chemical violence, COVID, death, food supply, mRNA, new vaccines, vaccines, world agriculture

This article may contain statements that reflect the opinion of the author



(Natural News) Much of the conversation surrounding mRNA (messenger RNA) "vaccines" centers around their impact on humans, how about all the animals that are being injected with it?

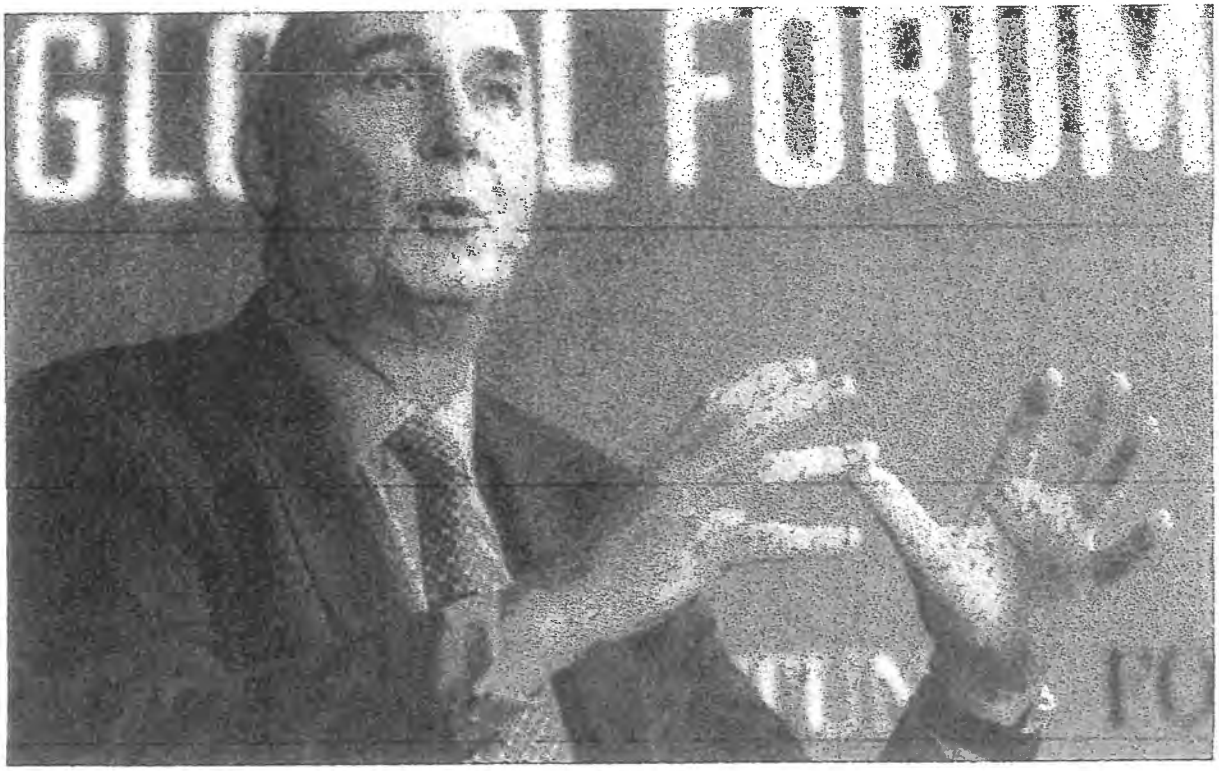
Believe it or not, cattle are reportedly now getting jabbed with the stuff, which in a recent mass "vaccination" campaign of an Austral herd resulted in 35 of the 200 animals dying *immediately*.

We are told that dairy farmers and others are now being *forced* to inject their animals for the Fauci Flu in order to remain in busines and that the animals are not responding well to it.

Just like in humans, the shots are causing such profound damage that many of the animals are succumbing to *instant death*, while others are getting sick and dying over a longer period of time. (Related: mRNA spike proteins linger in the heart and brain long after injection.)

For the animals that survive, one wonders what is becoming of their milk, which gets passed on as food for other animals as well as humans. Is it safe to consume mRNA-tainted milk and cheese from a "fully vaccinated" dairy cow? The answer is *probably not.*

"Dairy herd DNA is altered," one report explains. "Milk is altered and you consume it! Butter constitution, yoghurt, and cheese is alte meat is altered – will chicken and other meats be next?



While we all fixate on glyphosate, Monsanto prepares its next GM trick: RNA pesticides

JP Sottile | 11th April 2016

The global pesticide and bioscience giant Monsanto is a byword for evil for millions of campaigners and concerned citizens, writes JP Sottile. But that has never stopped it getting its way with the people that matter - politicians and regulators. And now the company is on the verge of biggest victory ever - winning clearance to spray biologically active RNA sequences on US crops.

12/09/22 • BIG PHARMA › VIEWS

'All These Vaccines Need to Be Withdrawn From the Market': COVID Roundtable Part 2

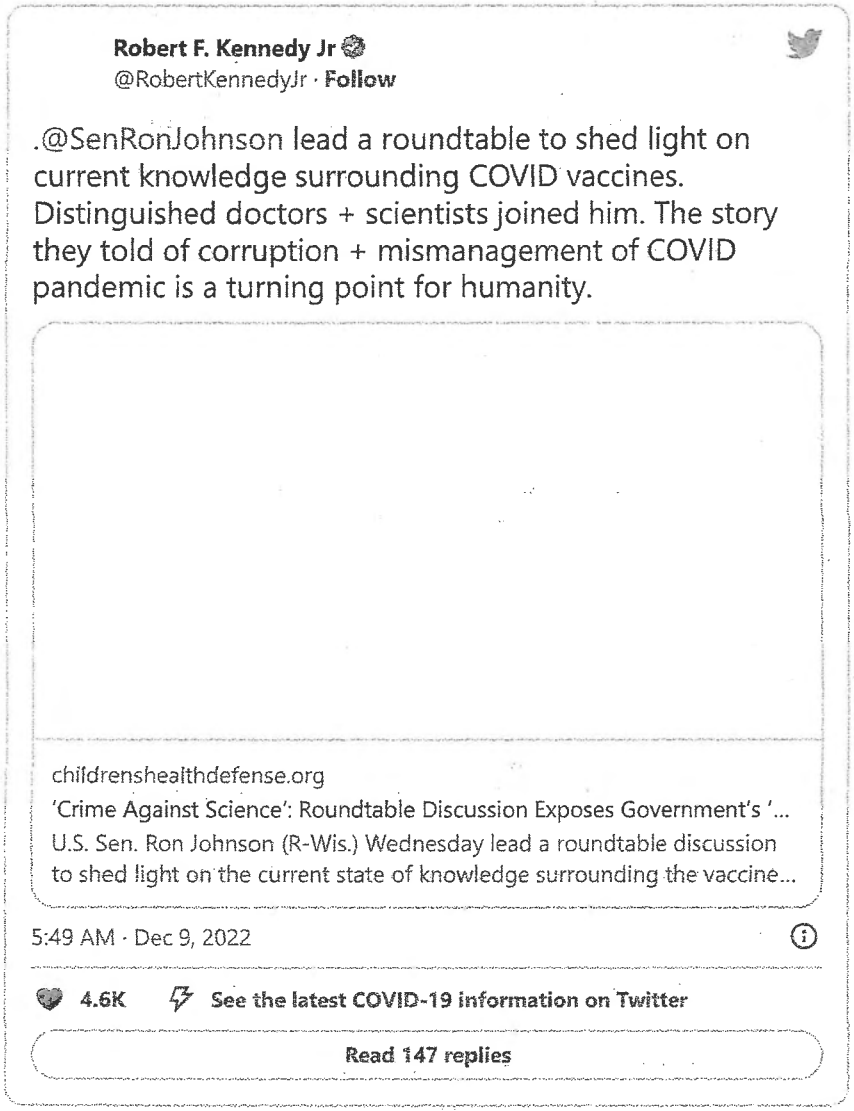
During a roundtable discussion on COVID-19, led by U.S. Sen. Ron Johnson (R-Wis.) on Wednesday, Dr. Peter McCullough concluded that in order to prevent future harm, all COVID-19 vaccines need to be immediately withdrawn from the market.

By Josh Mitteldorf, Ph.D.

U.S. Sen. Ron Johnson (R-Wis.) on Wednesday led a roundtable discussion — "COVID-19 Vaccines: What They Are, How They Work and Possible Causes of Injuries" — to shed light on the current state of knowledge surrounding the vaccine and the path forward.

Distinguished doctors and scientists who specialize in COVID-19 vaccine research and treatment participated in the three-hour event.

This is my summary of the second part of the roundtable discussion. You can read the first part here.



Before recapping more of the roundtable discussion, I'm taking the liberty of inserting my personal perspective on a topic that was avoided during the event, advisedly I am sure: the origin of SARS-CoV-2 as a bioweapon.

One reason the laboratory origin of COVID-19 is important is that it implies there are people who designed the SARS-CoV-2 virus, who know what it does and does not do, and probably know how to treat it.

Why weren't these scientists located, offered protection and subpoenaed to tell public health officials what they knew?

Natural viruses don't want to hurt you. Fitness, for a virus, is the ability to spread from host to host.

Viruses are evolved to make copies of themselves and maybe to make you sneeze to disperse the virus through aerosols, but they don't have any interest in causing permanent harm, in infecting your heart, or causing blood clots or infertility.

For this reason, the vast majority of viruses are harmless. There are some that get too enthusiastic about making copies and hijack enough of the body's machinery that they can make us very sick, but always this is an evolutionary miscalculation and it doesn't last long.

A bioweapon is engineered, not evolved. It may have features that are gratuitously toxic to the host (that's us) even though they confer no adaptive benefit for the virus.

In the case of the SARS-CoV-2 virus, evidence points to the spike protein as the part of the virus that is different from other coronaviruses — this is the part of the virus that was likely engineered and is highly toxic.

The spike is designed to break off into the bloodstream when it comes in contact with a common human enzyme (furin). The spike is neurotoxic and damages heart muscles and attacks the epithelium of our blood vessels.

Consider now that all the COVID-19 vaccines were designed to deliver spike protein to the body.

During the roundtable, Dr. Robert Malone cited a Stanford study demonstrating that the average vaccinated individual gets larger doses of spike protein from the vaccine than the average COVID-19 patient gets from the virus.

In an evolved virus, the spike protein would be optimized for connecting to the human receptor (in this case ACE-2) that allows it to enter the cell. It would likely be harmless in itself. But this is not an evolved spike protein. It has been engineered and weaponized.

Thousands of scientists worked on the COVID-19 vaccines and they innocently played their parts. They did not know that the spike protein that they were delivering was designed to do damage.

But a few people at the top knew this and made the decision nevertheless to base their vaccine development on the spike protein.

This was a heinous crime.

The first publications I know that gave evidence that the spike protein was toxic came out in June 2020, when vaccines were still in the early stage of warp-speed development.

Malone said there are studies on toxicity of coronavirus spike proteins going back to 1992.

The people who made the decisions to proceed nevertheless with a vaccine for universal distribution that was based on the spike protein were burying the first red flag, and starting their companies and their governments down a path of dissembling a great many more safety signals over the ensuing 2 years.

Roundtable, part II

David Gortler, Ph.D., former faculty member at Yale School of Pharmacology, recently from the Center for Ethics and Public Policy within the U.S. Food and Drug Administration (FDA), told us he was fired for being one of the few persons within the FDA who called out the leadership and asked them to follow their own procedures for evaluating vaccine safety.

Usually, the FDA starts by getting an ingredient list, knowing contents and dosage quantities for any medication before even considering approval. In the case of the COVID-19 vaccines, this rudimentary information was withheld by the manufacturers, and they still maintain ingredients lists as a trade secret.

Del Bigtree showed a short video, a composite of journalists and public officials who stated unequivocally that the COVID-19 vaccines would prevent a vaccinated person from passing the virus to anyone else. But assessing transmission of the virus was not part of the protocol when the vaccines were being tested. The question was not even asked.

Despite protests about the "speed of science," all they would have had to do was collect PCR samples from family members when they tested the experimental subjects.

Johnson listed some of the officials who were invited to the roundtable to give balance and present their perspectives on the story.

The list included Dr. Anthony Fauci, Dr. Rochelle Walensky, Albert Bourla (Pfizer), Stéphane Bancel (Moderna), Dr. Robert Califf (FDA) and Peter Marks, M.D., Ph.D. (FDA) and Robin Baily (CDC).

All of them declined to appear or to send a representative. Johnson indicated that this is consistent with his experience in the past. The people who are making COVID-19 policies avoid debating the issues with other scientists.

Dr. Peter McCullough outlined normal procedures for establishing safety monitoring boards and critical event committees, ethics committees and institutional review boards whenever a new drug is released. None of this was done with the COVID-19 vaccines, despite the fact that these vaccines were released to the public with far less testing than any approved product in the past.

David Wiseman, Ph.D. reported on the gap between Pfizer's vaccine insert, which says there is no data establishing safety for pregnant women, and the CDC's guarantee that the vaccines are safe for pregnant women in their public promotions. (The vaccine insert was

added last summer. For the first 19 months of the vaccines, they were distributed with a large, folded package insert saying only "This page intentionally left blank.")

Gortler went into the history of the testing procedures that were established in the 1930s through the '60s to back up the words "safe and effective" with explicit tests that a product would have to pass. This has always included long-term studies in a diverse population, but these were bypassed when COVID-19 vaccines were approved under "emergency" rules.

He described the dangers of our country sourcing most of our drugs from India and China, where they are produced cheaply. We should be inspecting these foreign manufacturing facilities and randomly sampling their products for analytic testing to assure that the vials contain what the label says.

Not enough of this is currently done with respect to common drugs, no inspection or testing is being done for the mRNA vaccines. There is no excuse for not collecting data on quality control and making it public.

Malone added that the mRNA vaccines are made from lipid nanoparticles that require exacting conditions far beyond the normal organic chemistry that is involved in making a drug. There are dozens of sites around the world where the vaccines are being manufactured, and FDA is not inspecting any of them.

How-bad-is-my-batch is a website that compares safety results from different vaccine batches, and there is wide variation by lot number in the number of vaccine injuries reported.

In the next segment of the hearings, a few people who were injured by the COVID-19 vaccines described their experience. (Malone mentioned in passing that his diastolic blood pressure rose to a life-threatening level of 230 mm after vaccination.)

Brianne Dressen, a 42-year-old mother and classroom teacher, had been healthy before volunteering for the AstraZeneca vaccine trials. After vaccination, her immediate symptoms included double vision, tingling and numbness in her arms, followed by paralysis in her left leg the next morning.

She is now unable to work or engage in normal household activities because of nerve damage. Doctors tell her that the damage is progressive. In the reported results of AstraZeneca trials, her case was not even mentioned as a safety concern. This, unfortunately, was typical.

Adverse reactions to the vaccines during the test phase were routinely understated or ignored. A woman in the Moderna trials developed lymphoma (blood cancer), but in Moderna's write-up, she was listed as fully recovered.

A 12-year-old from the Pfizer trials who has been confined to a wheelchair and feeding tube the rest of her life was coded as a "stomach ache." Dressen became aware of these cases and many others when she and her husband organized an online support group for the vaccine injured, which quickly grew to 20,000 members.

The National Institutes of Health (NIH) has protocols for the vaccine-injured that have been disclosed privately to a few of those injured by vaccines, but these remedies are not being shared with the public.

Dr. Joel Wallscog was an orthopedic surgeon in Madison, Wisconsin before nerve damage from a Moderna shot ended his career. Symptoms include balance problems, dizziness and weakness in both legs. His diagnosis was transverse myelitis, [an injury to the spinal cord that is related to multiple sclerosis].

Wallscog has created an advocacy network, support group and funding source for people who are vaccine-injured. Of course, Federal agencies should be providing medical care for people who are injured by vaccines, but this would require recognition of the vast scope of the project, which is politically inexpedient.

One result of this is that most GP doctors don't know about vaccine injuries, don't recognize them when they appear and don't believe their patients when injuries are reported. 90% of vaccine-injured patients report being gaslighted when they came to their family doctors for medical care. They were diagnosed as psychosomatic disorders.

People who are called "anti-vaxxers" were not born that way.

Sen. Roger Marshall (R-Kan.) made a cameo appearance and expressed concern that the government agencies which people trusted to protect them were withholding information and lying outright during the pandemic.

Dr. Kirk Milhoan, a pediatric cardiologist, defined myocarditis as inflammation of the heart muscle. The spike protein manufactured by our bodies in response to vaccination has been found to cause myocarditis. Rates of myocarditis following vaccination are not being measured or reported in this country.

A study from Thailand indicates a rate of 2-3% of adolescent males with myocarditis following vaccination. For college students, the risk of hospitalization after COVID-19 is about 0.002%. The risk of myocarditis from vaccination is about a thousand times higher.

Yet our governments and our universities are demanding that students be vaccinated as a condition of enrollment. Ninety days after vaccination, damage to the heart was still detectable in more than half of those who suffer myocarditis.

We do not know for a portion of myocarditis patients the injury will be permanent, or eventually fatal.

Dr. Renata Moon, a pediatrician, had seen only three cases of myocarditis in her 20-year career. Now she sees myocarditis routinely.

Dr. James Thorp is an ob-gyn in St Louis. He reported a "substantial, massive, unprecedented increase in menstrual abnormalities, infertility, miscarriage, fetal death and fetal malformation. We have published many studies over the last two years based on data from VAERS and CDC."

Lt. Col. Theresa Long, M.D., M.S. listed some of the injuries she has seen personally, including strokes, clotting in the spleen and liver, spinal tumors, brain tumors, sarcoidosis [a once rare condition, involving warts that grow on internal organs], lupus, cognitive impairment, myocarditis, pericarditis, avascular necrosis [dying bone that has been deprived of blood supply] that required hip replacement and a "shocking, pervasive" suppression of the immune system.

Dr. Ryan Cole said that the nanolipid particles that deliver mRNA in the COVID-19 vaccines were designed to bypass the body's barriers, including cell walls and the blood-brain barrier.

The toxic spike protein goes to the heart, the brain, the reproductive organs and all the most sensitive areas of the body, where it causes inflammation and autoimmunity.

We were told that the vaccine stays in the deltoid muscle of the arm, and this is just wrong. Sen. Johnson then pointed out that "stays in the arm" was not a mistake but a deception. Companies that produced the vaccines had results from animal tests that showed the presence of the spike protein in sensitive areas of the body.

A discussion of the toxicity of the spike protein ensued. Malone reported being censored when he mentioned this fact in a September 2001 podcast. McCullough cited a 1992 paper by Ralph Baric [virologist and bioweapons expert from the University of North Carolina] that detected heart damage from a coronavirus spike protein.

The virus and the vaccine both dose the body with spike protein and the damage compounds. For this reason, COVID-19-recovered people were excluded from the vaccine trials — and yet, the vaccines are now being recommended, even mandated, for people who have recovered from COVID-19.

Malone cited a paper in which spike protein levels were measured in people who had the COVID-19 virus and others who had been vaccinated. Higher levels of spike protein were reported in the vaccinated.

Johnson asked, where do we go from here?

Malone said there are 200 trials listed on ClinicalTrials.gov for mRNA-based vaccines. The original plan was to demonstrate that the mRNA platform is intrinsically safe, and then new products that are based on plugging different sequences into the RNA could be rapidly approved without testing.

Of course, the mRNA platform has not been proven safe — quite the opposite, the current mRNA vaccines have by far the highest rates of associated complications and death of any vaccine product in history.

Still, the medical establishment is so committed to their paradigm and a new, profitable industry that they are conducting accelerated trials of over 200 new products (not just vaccines), for which they will cite the record of the COVID-19 vaccines as a safety precedent.

With their patents, Moderna and Pfizer-BioNTech hope to lock in a duopoly on a lucrative pipeline of future products.

The National Institute of Allergy and Infectious Diseases gets royalties on these patents and is motivated to help them, safety be damned.

McCullough concluded:

> "In order to prevent future harm, all these vaccines need to be withdrawn from the market. That needs to happen immediately. All the vaccine mandates should be dropped. We need requests for applications and immediate funding for vaccine injury. Centers of excellence across the United States for screening, detection, and diagnosing of vaccine injuries. We need a massive shift in our healthcare system towards managing this large number of vaccine-injured people. What is at stake here is increased risk of death."

Amen.

Watch the entire roundtable discussion here:

Public Record

Ban

ATRAZINE

Almost all of us go out of the way to avoid harmful toxins and poisons: We wouldn't leave bleach near a toddler, for instance, and most of know about the dangers of **Monsanto Roundup**. But what if the toxin is something that's quite hard to escape … and we're consuming it unwittingly, without knowing exactly what will happen to us?

Say hello to atrazine, the second-most widely used herbicide in the U.S. behind glyphosate (the active ingredient in Roundup) and likely just as dangerous, most infamously as an **endocrine disruptor**. While other countries have banned the herbicide, atrazine is still used in American crops — and often winds up in our water supply. In fact, it's the most common chemical contaminant in U.S. water supplies.

In June 2016, the Environmental Protection Agency released a preliminary risk assessment, its most damning criticism of the toxin to date. But with a public comments date that's been extended from the initial 60-day period and the federal wheels of bureaucracy moving ever so slowly, it's up to each of us to take steps to reduce our exposure to atrazine and avoid its toxic effects.

So what exactly *is* atrazine? And how is it that although it's quite prevalent in America and contributes to our **tap water toxicity**, most of us haven't ever heard of it? It's time to delve into the dirty nitty gritty of atrazine.

What Is Atrazine? A Look at the Toxin and Its History

Atrazine is an herbicide produced by Syngenta AG, a global company based in Switzerland. In the U.S., the product is used mainly to kill weeds. It was first registered by the United States Department of Agriculture (USDA) as an herbicide in 1958.

While 90 percent of atrazine, or 70 million pounds, is used annually in America is to prevent weeds in corn fields, atrazine is also used on sugarcane, sorghum, macadamia nuts, soybeans, schools, parks, playgrounds, guava, athletic fields and evergreen farms, where families buy their Christmas trees. (1, 2) In fact, 65 percent of sorghum and sugarcane fields are treated with atrazine. It's also used in other products for farming and landscaping purposes, about 200 in total.

When Monsanto's glyphosate came onto the scene, the idea was that atrazine use would be reduced. But because crops have become resistant to glyphosate, atrazine is still used as a weed killer, often in conjunction with glyphosate for a toxin double-whammy.

Having a toxin sprayed on **corn** and crops is bad enough but, like most pesticides, atrazine doesn't stay only where it's sprayed. It usually ends up in our surface water and ground water, which means it's in our nation's drinking water supply. (3, 4) Nearly 90 percent of the water tested by the USDA has atrazine residue in it. (5)

So if it atrazine has been approved for use, why is it so harmful — and why is the EPA finally taking notice?

Toxic Effects of Atrazine

Syngenta, the company behind atrazine, would have you believe that the herbicide is perfectly safe. According to them, "Atrazine is effective, safe, and integral to agriculture's success in the United States and worldwide." (6) But that couldn't be further from the truth.

Atrazine Is an Endocrine Disruptor

One of atrazine's scariest effects is that it is an endocrine disruptor. These are chemicals foreign to the human body that, after a certain level of exposure, disrupt our endocrine — also known as hormonal — systems. Endocrine disruptions can cause adverse developmental, reproductive, neurological and immune effects in people and wildlife.

This occurs because the endocrine system includes hormone-secreting glands and is in charge of regulating blood sugar, our reproductive systems, metabolism, brain function and the nervous system. Our bodies are kept in check with a delicate balance. When one hormone goes out of whack, it can have serious ripple effects throughout the body. (7)

When it comes to atrazine, its endocrine disruption abilities are frightening. A 2011 study published in *The Journal of Steroid Biochemistry and Molecular Biology* summarized a huge swath of research on atrazine, dating back to 1997. The study alone features 22 authors from around the globe. (8)

The study confirmed what researchers have been saying for years: atrazine "demasculinizes" and "feminizes" vertebrate male gonads. In other words, atrazine is a "decrease in male gonadal characteristics," because the herbicide shrinks testicles and reduces sperm counts. By "feminizing" male gonads, atrazine can lead to the growth of ovaries in males.

Frogs turning from males into females means they can now mate with male frogs. But since the female frogs are still genetically male, their offspring are all male. This leads to a major skewing of the sex ratios in a population, which leads to a decrease or even an elimination of the population. (9)

And while much of the media attention has been on how male frogs can turn into females, what this comprehensive study found is that the effects "do not occur merely across populations, species or even genera or orders, but across vertebrate classes." That means they occur across amphibian, fish, mammal and reptile species.

The researchers believe these scary changes occur because atrazine reduces production of male hormones, while increasing the effect of estrogen, a female hormone. The atrazine levels that frogs which change sex are exposed to is less than what's legally allowed in our water — it occurs at levels as low as 0.1 parts per billion, or

PPB. In comparison, the EPA allows atrazine at levels 30 times higher than this in our drinking water — 3 ppb.

Public Record

The Epoch Times

Unusual Toxic Components Found in COVID Vaccines, 'Without Exception': German Scientists

A group of independent German scientists found toxic components—mostly metallic—in all the COVID vaccine samples they analyzed, "without exception" using modern medical and physical measuring techniques. The Working Group for COVID Vaccine Analysis says that some of the toxic elements found inside the AstraZeneca, Pfizer, and Moderna vaccine vials were not listed in the ingredient [...]

AUGUST 22, 2022 BY ENRICO TRIGOSO

https://www.theepochtimes.com/unusual-toxic-components-found-in-covid-vaccines-without-exception-german-scientists_4673873.html?utm_source=ai&utm_medium=search

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Pfizer Has Stopped Its COVID Vaccine Clinical Trial in Pregnant Women: Internal Email

Pfizer and BioNTech's clinical trial to evaluate their COVID-19 vaccine in pregnant women has been discontinued, according to a newly disclosed internal Pfizer email filed as evidence in a recent court case. The case in part questions whether the vaccine has been adequately tested to assess for risks to women's reproductive health. A professor of [...]

SEPTEMBER 1, 2022 BY NOÉ CHARTIER

https://www.theepochtimes.com/pfizer-has-stopped-its-covid-vaccine-clinical-trial-in-pregnant-women-internal-email_4700143.html?utm_source=ai&utm_medium=search

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Leaked Video Suggests Israeli Health Officials Covered Up Serious Safety Problems With Pfizer COVID Vaccine

A leaked video recording reveals researchers in June shared data with the Israeli Ministry of Health showing serious and long-term side effects associated with Pfizer's COVID-19 vaccine, but Israeli health officials told the public in an August report that serious side effects were "rare" and short-term. This article was originally published by The Defender – [...]

SEPTEMBER 9, 2022 BY SUZANNE BURDICK, THE DEFENDER

https://www.theepochtimes.com/health/leaked-video-suggests-israeli-health-officials-covered-up-serious-safety-problems-with-pfizer-covid-vaccine_4720976.html?utm_source=ai&utm_medium=search

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Fibrous Clots, Foreign Matter in Blood After COVID Jabs: Is There a Way to Detox?

Recently unusual blood clots as well as metal-like foreign objects found in the vessels of COVID-19 jab recipients have been reported across the country. Both types of substances are unusual and are likely to be harmful to our bodies. What are the potential causes and ramifications of these substances, and is there any chance of [...]

SEPTEMBER 18, 2022 BY DR. YUHONG DONG, DR. ANN CORSON

https://www.theepochtimes.com/health/fibrous-clots-foreign-matter-in-blood-after-covid-jabs-is-there-a-way-to-detox_4738079.html?utm_source=ai&utm_medium=search

COVID Vaccines Contaminate Breastmilk With mRNA: Study

A new study published on the JAMA network concludes that mRNA from COVID vaccines can be transmitted in small amounts through breast milk. The authors of the study examined 11 "lactating individuals," after getting either the Pfizer or Moderna mRNA shots. Nine of them were white, one black, and one Asian. Five of the participants [...]

SEPTEMBER 27, 2022 BY ENRICO TRIGOSO

https://www.theepochtimes.com/covid-vaccines-contaminate-breastmilk-with-mrna-jama-study_4757809.html?utm_source=ai&utm_medium=search

COVID Vaccine Mandates Weakening US Military, Retired Vice Admiral Warns

A retired U.S. Coast Guard rear admiral warned that COVID-19 vaccine mandates are weakening the military and driving down recruitment efforts. In a letter published Monday, retired Coast Guard Vice Admiral William "Dean" Lee criticized the Biden administration's vaccine mandate. The rule, implemented in 2021 through the Department of Defense (DOD), has allowed for few *religious* [...]

OCTOBER 4, 2022 BY JACK PHILLIPS

https://www.theepochtimes.com/covid-vaccine-mandates-weakening-us-military-retired-real-admiral-warns_4774265.html?utm_source=ai&utm_medium=search

Unsettling Research Links COVID Vaccine to Parkinson's

The list of complications, conditions, and diseases resulting from the COVID shots is nearly endless and can affect any organ system in the body. Pfizer knew. Here's their document. Look at the last 8 pages, which lists more than 1100 serious side effects and life-threatening illnesses Pfizer knew would happen from the first shot. We [...]

OCTOBER 4, 2022 BY SHERRI TENPENNY

https://www.theepochtimes.com/unsettling-research-links-covid-vaccine-to-parkinsons_4765232.html?utm_source=ai&utm_medium=search

Free mRNA for Your Baby?

Scientists at New York University's Long Island School of Medicine have detected messenger RNA COVID-19 vaccines in human breast milk, according to a new study. This peer-reviewed research, published on September 26, 2022 in the journal JAMA Pediatrics, looked at the breastmilk of 11 healthy breastfeeding women, five of whom had received the Moderna vaccine [...]

OCTOBER 6, 2022 BY JENNIFER MARGULIS, JOE WANG

https://www.theepochtimes.com/health/free-mrna-for-your-infant-baby_4779439.html?utm_source=ai&utm_medium=search

The Science Behind Florida's Recent Recommendation Against mRNA COVID Vaccines for Men 18–39

IN-BRIEF Florida is the first state to recommend against mRNA COVID vaccination of children and men up to age 39, but it joins the UK, Sweden, and Denmark in some regards. The Florida Department of Health conducted a self-controlled case series (SCCS) with a 25-week observation period, similar to an analysis done in the UK. [...]

OCTOBER 17, 2022 BY DR. YUHONG DONG

https://www.theepochtimes.com/health/the-science-behind-floridas-recent-recommendation-against-mrna-covid-vaccines-for-men-18-39_4801126.html?utm_source=ai&utm_medium=search

Growing Number of Governors Reject COVID Vaccines for School Entry After CDC Vote

The GOP governors of several states indicated they will not implement mandates for children to receive a COVID-19 vaccine to enter school after a Centers for Disease Control and Prevention (CDC) advisory panel voted last week to recommend adding the vaccine to the childhood immunization schedule. Governors in Tennessee, South Carolina, Virginia, Montana, Alabama, Oklahoma, Florida, [...]

OCTOBER 23, 2022 BY JACK PHILLIPS

https://www.theepochtimes.com/growing-number-of-governors-reject-covid-vaccines-for-school-entry-after-cdc-vote_4814957.html?utm_source=ai&utm_medium=search

Coalition Presents Letter Signed by Canadian Scientists, Asking Health Officials to Affirm COVID Vaccine 'Safety Risks'

A coalition of over 20 public interest groups is calling on the federal government to publicly affirm the "material risks and scientific uncertainties" related to COVID-19 mRNA vaccines, presenting a letter signed by 19 doctors, scientists, and academics across the country. Coming together as the "Citizens' Group," the coalition wrote the letter to Jean-Yves Duclos and [...]

OCTOBER 25, 2022 BY ISAAC TEO

https://www.theepochtimes.com/coalition-presents-letter-signed-by-canadian-scientists-asking-health-officials-to-affirm-covid-vaccine-safety-risks_4818542.html?utm_source=ai&utm_medium=search

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Biden Calls on Americans as Young as 5 to Get 'COVID Vaccine Once a Year'

President Joe Biden called on Americans five years and up to get a dose of an "updated" COVID-19 booster, recommending the vaccine "once a year" as he received a dose of the newly authorized shot on Tuesday. "For Americans over five years of age who are fully vaccinated, our nation's health experts recommend that they get [...]

OCTOBER 26, 2022 BY LORENZ DUCHAMPS

https://www.theepochtimes.com/biden-calls-on-americans-as-young-as-5-to-get-covid-vaccine-once-a-year_4821219.html?utm_source=ai&utm_medium=search

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COVID Vaccine Mandates Expose Religious Discrimination Within the Military

Commentary The COVID-19 vaccination mandates impacted Americans across the country, forcing many to choose between their bodily autonomy and, in many cases, their livelihoods. But many Americans had to choose between not just their preferred health choices but also their sincerely held religious beliefs. Sadly, the very men and women who defend constitutional *rights, such* [...]

OCTOBER 27, 2022 BY EMILY ALLISON

https://www.theepochtimes.com/covid-vaccine-mandates-expose-religious-discrimination-within-the-military_4820763.html?utm_source=ai&utm_medium=search

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Hyperbaric Oxygen Therapy for Long COVID and Post COVID Vaccine Symptoms

Hyperbaric oxygen therapy (HBOT) is a treatment that increases blood oxygen levels to boost wound healing and clear bacterial infections. Recent studies and doctors' clinical experiences suggest that it may be useful for treating long COVID and post COVID vaccine symptoms. "When I first heard about it [HBOT] I thought, 'this is goofy,'" said Dr. [...]

OCTOBER 27, 2022 BY MARINA ZHANG

https://www.theepochtimes.com/health/hyperbaric-oxygen-therapy-for-long-covid-and-post-covid-vaccine-symptoms_4823119.html?utm_source=ai&utm_medium=search

More Adverse Events: It's Time to Halt COVID Vaccine Recommendations for Pregnant Women

According to the recommendations from the Centers for Disease Control and Prevention (CDC) and the Food and Drug Administration (FDA), pregnant women should be vaccinated against COVID-19. However, the main research work to back up this policy was the influential CDC-sponsored article by Shimabukuro et al. (2021), published in the New England Journal of Medicine, [...]

OCTOBER 28, 2022 BY DR. SEAN LIN

https://www.theepochtimes.com/health/more-adverse-events-its-time-to-halt-covid-vaccine-recommendations-for-pregnant-women_4824656.html?utm_source=ai&utm_medium=search

Toxic Components Found in Blood of Patients Who Took COVID Vaccines: German Scientists

In this episode of Frontline Health, we look at the blood of patients who took the COVID-19 mRNA vaccine. These blood samples were taken and analyzed by a group of 60 interdisciplinary scientists, including physicists, chemists, pharmacologists, microbiologists, and doctors. They found strange foreign objects floating among the blood cells. The team previously observed these [...]

AUGUST 25, 2022 BY DAN SKORBACH

https://www.theepochtimes.com/toxic-components-found-in-blood-of-patients-who-took-covid-vaccines-german-scientists_4689152.html?utm_source=ai&utm_medium=search

COVID Vaccine Injury Data: 143,233 Percent Increase in Cancer

When something goes wrong inside a cell's DNA, the cell fires a signal that tells other cells, "I've gone bad. Please kill me." This early warning helps the immune system stop cancer cells from multiplying. It stops a virus from replicating. So it's a very important early response tool that works across the whole body. [...]

AUGUST 30, 2022 BY DAN SKORBACH

https://www.theepochtimes.com/covid-vaccine-injury-data-143233-percent-increase-in-cancer_4699328.html?utm_source=ai&utm_medium=search

Posted by ClarkCountyToday.com
Date: Tuesday, October 25, 2022

Study finds Dr. Fauci's 'fingerprint' on origin of COVID virus

All of it says, my God, there was really a big, very risky research agenda underway'

Art Moore
WND News Center

A new pre-print study has concluded the virus that causes COVID-19 has a unique "fingerprint" indicating it originated in a laboratory rather than in nature.

Dr. Alex Washburne, a mathematical biologist, worked with researchers in the U.S. and Germany who studied the SARS-CoV-2 genome sequence and compared it to previously discovered coronaviruses.

They detected "peculiar patterns" they concluded were the hallmark of a manufactured virus, describing it having a "synthetic fingerprint."

Meanwhile, Jeffrey Sachs, chairman of The Lancet COVID-19 Commission, a task force that investigated the origins of COVID-19, has concluded after 22 months of study that SARS-CoV-2 probably was laboratory-generated and that the technology likely came from gain-of-function research funded by the U.S. National Institutes of Health.

The "synthetic fingerprint" discovered in the new study led by Washburne, says Sachs in an article published by Children's Health Defense, points to the work of Dr. Ralph Baric, a virologist at the University of North Carolina known for his NIH-funded gain-of-function research in cooperation with the Wuhan Institute of Virology.

Baric has developed a controversial "seamless ligation" technique designed to conceal evidence of human tampering in laboratory-created viruses. Baric nicknamed his invention the "no-see'm" method.

"It's the artist that doesn't sign his name to the painting; the virologist that doesn't put his signature into the virus to let us know whether or not it is emerging naturally or whether it is produced in a laboratory," said Sachs.

"All of it says, my God, there was really a big, very risky research agenda underway."

Evolutionary biologist Bret Weinstein called Baric's technique is the "exact opposite of what you would do if your interest was public health." Public health scientists would be marking their enhancements with red flags – not devising ways to hide them. The only reason you would want a concealer is to advance a sinister purpose – such as illegal bioweapons development – some mischief that the scientist didn't want traceable back to his lab."

Baric taught his technique to the Wuhan Institute of Virology's leading coronavirus researcher, Shi Zhengli, in 2016, CHD reported. In return, Baric received Chinese coronaviruses collected by Shi from bats in China's Yunnan province.

'If I am proven wrong I will change my mind'

In a post on Substack, Washburne reacted to criticism that his study was "very poorly controlled" and "cherry-picked."

"The topic is personally relevant to every person capable of being infected by a virus or impacted by pandemic policies," he wrote. "I invite people to prove us wrong and, if they do so, even if there are flaws in their work, I will not call them names or attack their credentials.

"I will celebrate their ingenuity and commitment to the Truth, and if I am proven wrong I will change my mind," he promised.

Washburne said science "can save lives and revolutionize our civilization, but only if scientists and our broader society remain honest, curious, and open-minded."

Professor Francois Balloux, a professor of computational biology at University College London, called the study "an important piece of work."

"To me, it looks solid both conceptually and methodologically," he wrote on Twitter. "I was given advance warning and was able to replicate the key findings. To the best of my knowledge, I confirm the reported patterns are genuine."

Balloux said the "distribution of restriction sites in SARS-CoV-2 is highly atypical when compared to related viruses in circulation, and far more in line with previous lab-engineered coronaviruses."

A critic of the study, Texas A&M University virologist Dr. Benjamin Neuman, has called the lab theory of the origin of COVID-19 "discredited." The new study, he said, is "very poorly controlled, cherry-picked and making a big deal out of lumps and bumps that are of no significance to the virus."

Critic was part of COVID-origin teleconference with Fauci

Another critic, Kristian Andersen, a virologist at Scripps Research in California, famously joined with three other virologists in a January 2020 email to Fauci stating they saw strong evidence the virus that causes COVID-19 was engineered in a lab, as WND reported. But after a teleconference the next day with Fauci to discuss the virologists' conclusion, Andersen began dismissing the lab-leak possibility as among "crackpot theories" that "relate to this virus being somehow engineered with intent and that is demonstrably not the case."

In April 2020, Fauci was asked by a reporter during a White House briefing if the research at the Wuhan lab might be responsible for the pandemic. Fauci insisted a "group of highly qualified evolutionary virologists" had concluded the virus was "totally consistent with a jump of a species from an animal to a human."

The next day, Peter Daszak – the EcoHealth Alliance founder who received funding from Fauci's agency to conduct research engineering coronaviruses – sent a thank you email to Fauci. Daszak thanked the National Institutes of Health and Infectious Disease director for "publicly standing up and stating that the scientific evidence supports a natural origin for COVID-19 from a bat-to-human spillover, not a lab release from the Wuhan Institute of Virology."

"From my perspective, your comments are brave, and coming from your trusted voice, will help dispel the myths being spun around the virus's origins," Daszak wrote to Fauci on April 18, 2020.

The new study by Washburne, Andersen said, is "so deeply flawed that it wouldn't pass kindergarten molecular biology."

"The study is a clear example of motivated reasoning with a heavy dose of technobabble to make it sound legitimate – but it's nothing more than poppycock dressed up as science," said Andersen.

"In plain language — this is uninformed nonsense and it's simply not worth engaging with."

Among the studies cited by scientists who believe the virus had a natural origin is one published in Scientific Reports that showed a total of about 47,000 wild animals from 38 species were sold in four markets in Wuhan between May 2017 and November 2019. The researchers claimed they had evidence the the necessary conditions were in place for animal-to-human transmission.

However, they admitted that had no evidence that any of the animals had SARS-CoV-2.

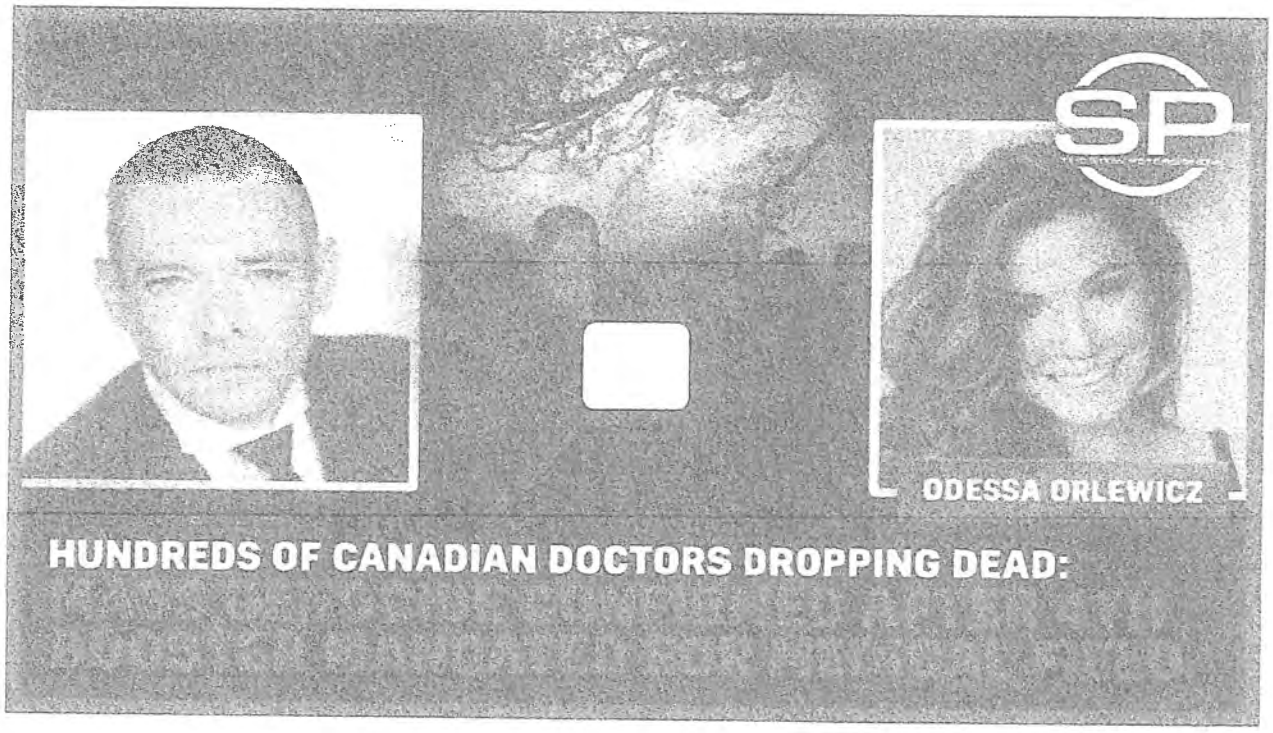
In the new study, Washburne and his colleagues concluded it's "extremely unlikely" that the "synthetic fingerprint" appeared "by random evolution," finding similarities to many engineered coronavirus genomes.

Significantly, it differs from the closest relatives found in nature.

"Our findings strongly suggest a synthetic origin of SARS-CoV2."

Hundreds Of Canadian Doctors Dead: Genocide Confirmed After 4th Booster Mandated For Medical Field

Stew Peters Network · Published August 22, 2022 · 136,378 Views SUBSCRIBE 1.0K SHARE



1.29K rumbles EMBED

Doctors are dropping like flies in Canada, and the culprit is the deadly vaccine forced upon them by the tyrannical government.
Canada is raising the requirements to be considered fully jabbed, and it is killing people!

Odessa Orlewicz joins us today to discuss this ongoing problem.
Prepare your family for famine and shortages by purchasing food through: https://heavensharvest.com/.

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Prepare your family for famine and shortages by purchasing food through: https://heavensharvest.com/

Righteously stand against vaccine tyranny by supporting Christopher Key:
https://www.vaccine-police.com/

Fauci Says COVID-19 and Influenza Vaccines Don't Work Well

Dr. Anthony Fauci is among the growing number of officials who are acknowledging that the COVID-19 vaccines don't work well against infection.

Vaccines against both COVID-19 and influenza have "deficiencies," including that they "elicit incomplete and short-lived protection against evolving virus variants that escape population immunity," Fauci, until recently President Joe Biden's chief medical adviser, and other top National Institutes of Health officials wrote in a recent paper.

"With the imperfections of these vaccines, it seems a public health imperative to aggressively pursue better vaccines and vaccination strategies," they wrote.

CDC Adds COVID-19 Vaccines to Child Immunization Schedule

The U.S. Centers for Disease Control and Prevention (CDC) has added COVID-19 vaccines to its routine immunization schedule for children and adults.

According to the CDC's 2023 immunization schedule for children and adolescents, two or three doses of COVID-19 vaccines have been recommended beginning with infants who are just 6 months old. Children in the age group of 6 months to 4 years, and 5 years to 11 years are recommended COVID-19 vaccines from Moderna or Pfizer. Among children aged 12 to 18, Novavax vaccines also are recommended in addition to Pfizer and Moderna.

In the list for adults, two or three doses of COVID-19 vaccines have been recommended from the age of 19 years. The 2023 COVID-19 vaccine recommendation for kids and adults is included among other typically recommended vaccines for measles, flu, rubella, and so on.

Advisers to the CDC in 2022 recommended adding the vaccines to the schedule.

Messenger RNA Sequences Found in Blood 28 Days After COVID-19 Vaccination

Messenger RNA sequences from the Pfizer and Moderna COVID-19 vaccines were found in the blood of multiple individuals weeks after vaccination, according to a study.

Researchers in Denmark analyzed samples from the vaccinated and detected partial or even full sequences of the messenger RNA (mRNA) following vaccination. The sequences were found as late as 28 days after vaccination, or the longest time period the study analyzed.

The findings mean that the mRNA, which is situated in lipid nanoparticles for delivery into the body, lingers for much longer than officials in the United States and elsewhere acknowledge.

Top Officials Discussed Vaccine 'Adverse Event Issue' and Pregnancy: Emails

CONTINUED FROM A1

Dr. Amanda Cohn, chief medical officer of the CDC's National Center for Immunizations and Respiratory Diseases, replied.

"We have a meeting with Rochelle at 3:30 about if we should say anything or wait until we have more definitive information ... I will let you know where we land," Cohn wrote. Dr. Rochelle Walensky is the head of the CDC.

"I'm not sure there is a right answer," Cohn added.

The emails were obtained by the nonprofit Judicial Watch, which sued the U.S. government for failing to appropriately respond to a Freedom of Information Act request for messages regarding adverse events, deaths, or injuries caused by the COVID-19 vaccines.

Adverse events include health issues such as arthritis or heart inflammation.

No other emails about the "adverse event issue" were included in the latest tranche obtained by the nonprofit.

"I respectfully decline to comment," Marks told The Epoch Times in an emailed statement when asked what the issue was.

Cohn and the CDC didn't respond to requests for comment.

The CDC currently recommends virtually all Americans aged 6 months and older, including pregnant women, get a COVID-19 vaccine and multiple booster shots.

The original trials didn't include enough information "to make conclusions about the safety of the" vaccines from Pfizer, Moderna, and Johnson & Johnson, according to FDA documents.

Authorities have relied on observational data, including a study from the CDC, a corrected version of which was published in October 2021.

Pfizer conducted a post-authorization trial of its vaccine in pregnant women that was labeled completed in mid-2022 but results haven't been reported publicly as of yet.

"We think that part of the reason is because the results are so bad," Linda Wastila, a professor at the University of Maryland whose expertise is in pharmacotherapy and drug policy, told The Epoch Times.

Pfizer didn't respond to a request for comment.

Confidentiality Agreement

The FDA vaccine advisory committee met on Dec. 10, 2020, to consider whether to advise the FDA to authorize Pfizer's vaccine.

During the meeting, officials revealed that two postvaccination cases of severe allergic shock (anaphylaxis) had been recorded in the United Kingdom out of 6,000 doses administered on the day the events occurred.

"At this point, we are seeking further information from the MHRA under our confidentiality agreement to learn more about this and to really tease that out," Marion Gruber, an FDA official at the time, told the panel.

CHRISTOF STACHE/AFP VIA GETTY IMAGES

A health worker fills a syringe with a COVID-19 vaccine in Freising, Germany, on Feb. 2, 2021.

The Medicines and Healthcare products Regulatory Agency (MHRA) is the British equivalent of the FDA.

The newly obtained set of emails showed U.S. and UK officials discussing the cases of anaphylaxis throughout December, including as early as Dec. 9, 2020.

Marks wrote that it would "be very helpful if our Office of Vaccines could receive additional details" from MHRA "under the terms of our mutual confidentiality agreement."

Jonathan Mogford, a UK official, sent back data, stating, "If I can just remind—information shared under our confidentiality agreement."

The MHRA declined to provide a copy of the confidentiality agreement, and the FDA didn't respond to emailed questions about the pact.

"It again took a lawsuit for the Biden administration to hand over, albeit heavily redacted, information regarding the safety of the COVID vaccines that the public has every right to know," Judicial Watch President Tom Fitton said in a statement. "This disturbing batch of new documents have uncovered a secret confidentiality agreement tied to COVID vaccine safety issues and emails that raise new questions about the vaccines and pregnancy."

The Epoch Times has submitted Freedom of Information Act requests to try to uncover more information about the adverse event issue and the confidentiality agreement. Previous requests have unearthed secret data on COVID-19 vaccines, including showing how the CDC identified hundreds of potential safety issues with the Pfizer and Moderna vaccines.

People with a history of severe allergic reactions to any vaccine were excluded from Pfizer's trial and people with a history of allergic reactions to any components of the Moderna vaccine were excluded from Moderna's trial. No such exclusions were reported for the Johnson & Johnson trial.

The fact sheets for all three vaccines in the United States were updated to include warnings about severe allergic shock, including saying the vaccine shouldn't be administered to people with a known history of severe allergic reaction to any components of the shot.

COVID VACCINES

States That Have Banned COVID-19 Vaccine Mandates in School, Bucking CDC

JACK PHILLIPS

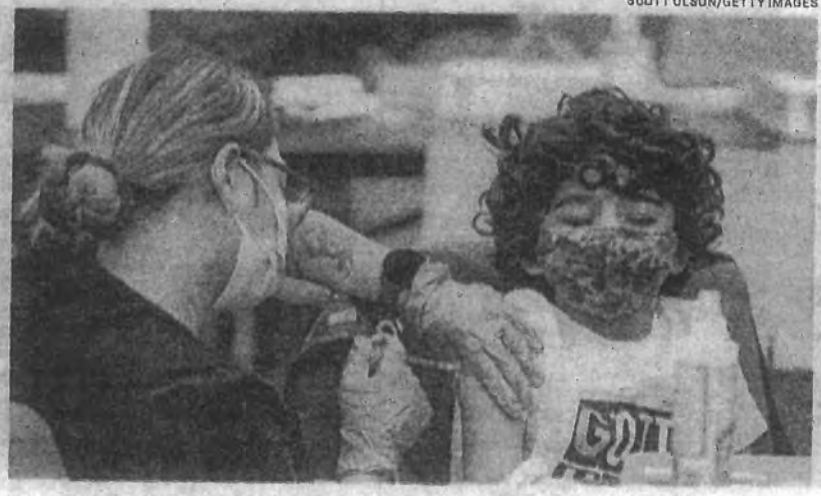
As the Centers for Disease Control and Prevention (CDC) adds COVID-19 vaccines to its routine immunization schedule for children and adults, at least 20 states have passed legislation or issued rules barring school systems from mandating the vaccines.

The National Academy for State Health Policy, in a Feb. 6 update, shows there are about two dozen states that have barred COVID-19 vaccines from being included in mandates for schools and students: Alabama, Arizona, Arkansas, Florida, Georgia, Idaho, Indiana, Iowa, Kansas, Michigan, Mississippi, Montana, New Hampshire, Ohio, Oklahoma, South Carolina, South Dakota, Tennessee, Texas, Utah, and West Virginia.

Officials in other states, including California, have announced they will not require students to get the shots. On Feb. 3, California's Department of Public Health said "that the state will not require the COVID-19 vaccine for school attendance," the website noted.

To date, not a single state has passed legislation or implemented an order to mandate COVID-19 vaccines to attend class, according to the academy. But the District of Columbia has a student COVID-19 vaccine mandate, although it has been delayed until the start of the 2023–2024 school year, after the D.C. City Council voted last November to do so.

SCOTT OLSON/GETTY IMAGES



A child receives a COVID-19 vaccine in Chicago on Nov. 12, 2021.

As long as I'm around and as long as I'm kicking and screaming, there will be no COVID shot mandates for your kids. That is your decision to make as a parent.

Florida Gov. Ron DeSantis

The academy has issued updates on COVID-19 vaccine mandates and bans on them since the CDC signaled that it could add those vaccines to its recommended child immunization schedule. While the CDC's guidelines are only recommendations, many states and municipalities rely on them to set policies.

Laws in at least 31 states and the District of Columbia require the vaccines on the CDC schedules to be taken by children for school attendance, said the Policy, Practice, and Prevention Research Center at the University of Illinois Chicago's School of Public Health last year. Other states sometimes impose requirements that mostly align with the schedules.

Response

Last year, when the CDC advisory panel voted to recommend adding the COVID-19 vaccines to the childhood immunization schedule, a number of governors had indicated they would not make them mandatory.

"Under my watch, there will be no COVID vaccine mandates for kids—period," said Republican Gov. Kim Reynolds of Iowa. "In fact, we signed a law that prevents it. It's the parent's decision, not the government's."

Florida Gov. Ron DeSantis noted that some parents are concerned about the CDC's guidelines, telling reporters there is a "fear" that schools would force shots on children. "So I just want to let everyone be clear. You know, as long as I'm around and as long as I'm kicking and screaming, there will be no COVID shot mandates for your kids. That is your decision to make as a parent," he said.

And at the time, Montana Gov. Greg Gianforte, a Republican, wrote: "I trust parents to raise their kids and do what's best for their kids' health. On my watch, the State of Montana will not mandate the COVID-19 vaccine."

A spokesperson for California Gov. Gavin Newsom last year said that the "main impact" of the CDC's recommendation would be that "health insurance companies will be required to cover the cost of the immunization and that the federal government can continue to provide it for free to low-income families."

Schedule Update

The CDC's 2023 immunization schedule for children and adolescents, released on Feb. 9, includes shots for flu, MMR (measles, mumps, and rubella), polio, and other inoculations—and now COVID-19. The CDC doesn't have the authority to issue mandates for vaccines.

Last year, all members of the CDC's Advisory Committee on Immunization Practices voted to add the Moderna, Pfizer, and Novavax vaccines to the 2023 schedules, asserting that the vaccines can prevent severe disease—despite studies showing waning effectiveness of the shots.

"We view this as COVID is here to stay," Dr. Matthew Daley, one of the advisers, said in a meeting. "When I think about the routine immunization schedule as a pediatrician, I think of it as an opportunity to prevent serious disease and death. And if something is added to the schedule, it's because I feel like the benefits continue to strongly outweigh the risks."

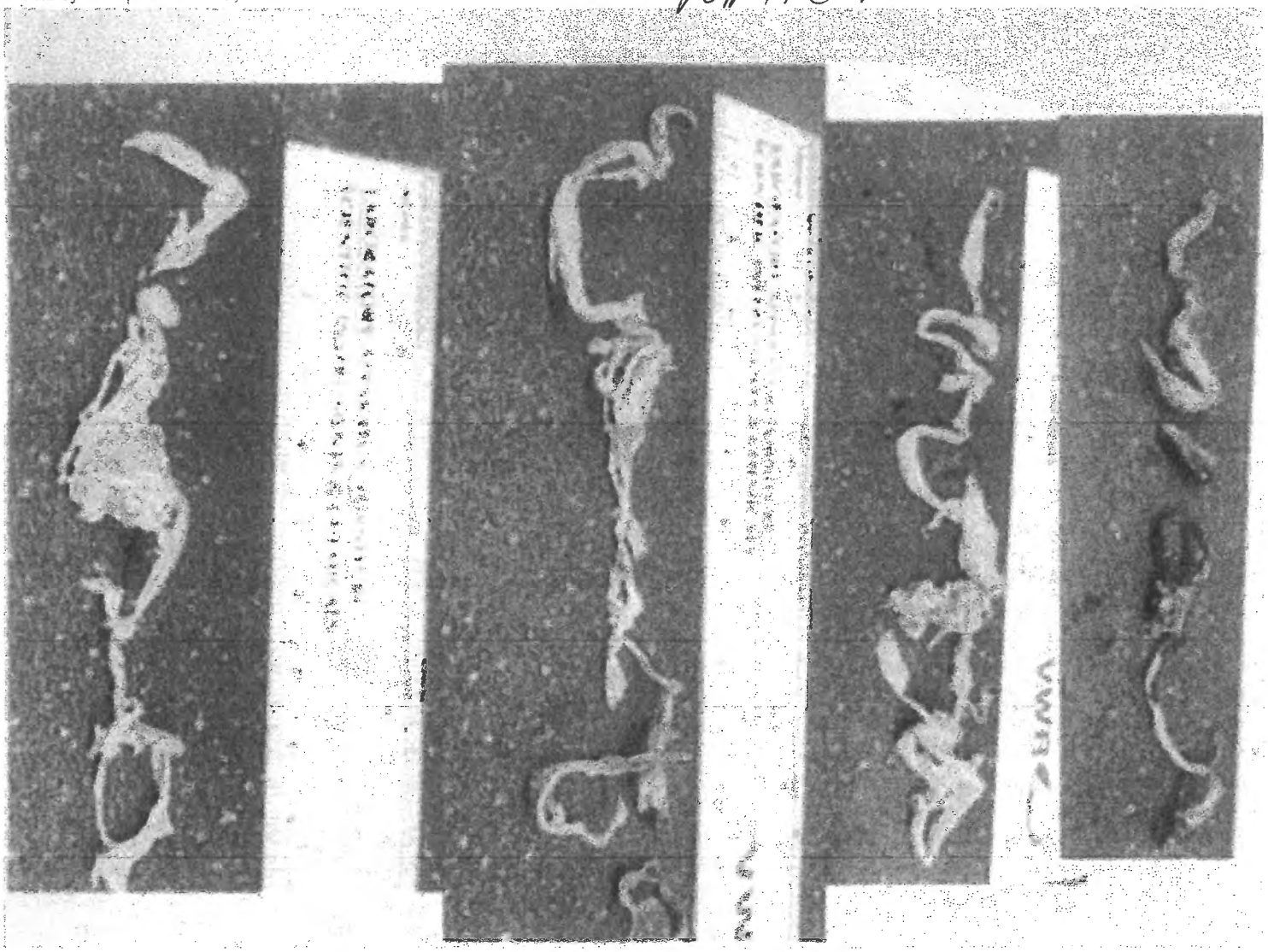
Zachary Stieber contributed to this report.

1

Experienced Pathologist Explains Blood Clots, Nano Tech and Parasites in COVID Vaccines

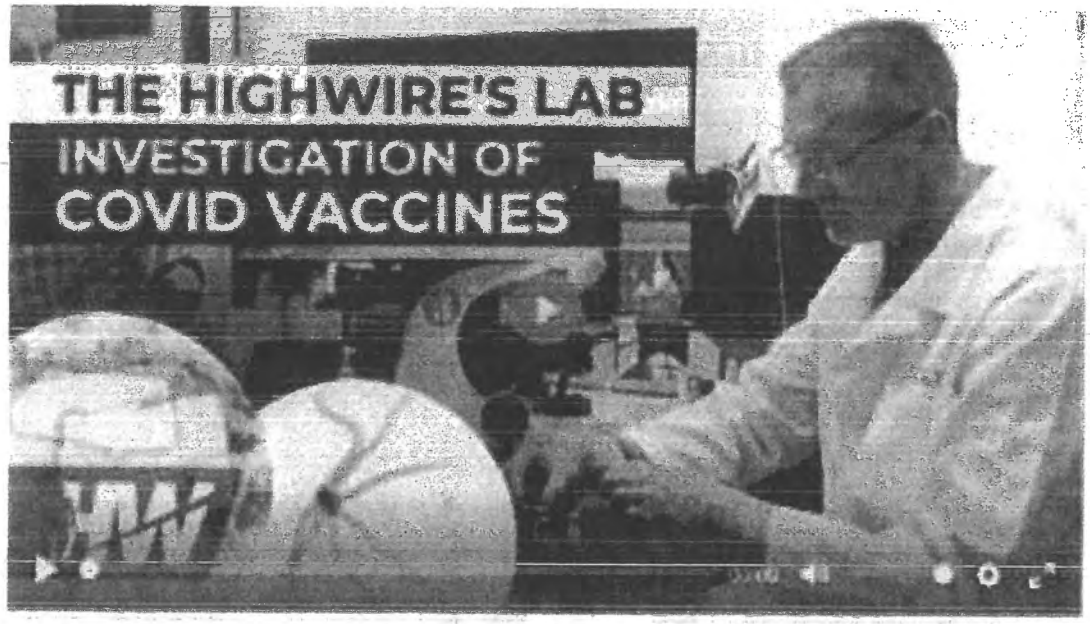
Public Record

written by GEG | December 19, 2022



Dr. Ryan Cole, a pathologist with 26 years of experience, explains that the blood clots that have been found by morticians, and also in living patients, are congealed protein made from an amyloid-like material. Dr. Cole said the microscopic rod and circuit-like structures others have identified as graphene oxide are actually cholesterol crystals, stacked-layered cholesterol, salt flakes and sugar crystals. He identified the the structures identified by others as parasites as being a leaf that came from environmental contamination.

Rumble: Dr. Jane Ruby on Blood clots



Link for video: **https://www.bitchute.com/video/BrVlJTsYZ6x4/**

Dr. Ryan Cole, who has been a pathologist for 26 years, identified the rubbery fibrous structures or clots that embalmers have been finding in deceased after the rollout of Covid injections as congealed protein, an amyloid-like material. he said the spike protein can induce inflammation in blood vessels that can cause blood clots. The spike protein, covered by a lipid nano-layer, was designed to cross the blood-brain barrier.

Dr. Cole examined the Johnson & Johnson, Moderna and Pfizer ingredients. He found a microscopic glass shard and said that there are problems with contamination from debris in the 'vaccines'. According to Johnson & Johnson's application to the FDA, their product contains trace human DNA and human proteins from aborted fetal tissue. Proteins from other people trigger a immune response in the injected person.

Dr. Cole said the microscopic rod and circuit-like structures others have identified as graphene oxide are actually cholesterol crystals, stacked-layered cholesterol, salt flakes and sugar crystals. He said that there are metallic and mineral contaminants in COVID shots. He affirmed that the injections do contain polyethylene glycol ("PEG"), which can trigger allergies. He said that the samples that looked like parasites were simply part of a leaf that was a contaminant from the environment. Dr. Cole said that COVID shots that were not properly frozen and preserved lost their potency. Some of the samples had broken RNA that produce shortened proteins, which are known carcinogens.

Lipid nanopartical and a gene sequence that makes vaccine recipients make foreign proteins are the cause of harm in vaccinated people. The lipid nanopartical is hyper-inflammatory and can be toxic to cells. Lipid nanopartical injections were designed to only be given once. There may be a cumulative toxicity. He said that the more the gene that triggers production of the spike protein is injected into a person's body, the more spike protein is produced, which has numerous side effects including blood clotting, potential increase in cancer, and more.

Vascular and organ damage induced by mRNA vaccines: irrefutable proof of causality

Michael Palmer, MD and Sucharit Bhakdi, MD

doctors4covidethics.org

Thursday 18th August, 2022

Abstract

This article summarizes evidence from experimental studies and from autopsies of patients deceased after vaccination. The collective findings demonstrate that

1. mRNA vaccines don't stay at the injection site by instead travel throughout the body and accumulate in various organs,
2. mRNA-based COVID vaccines induce long-lasting expression of the SARS-CoV-2 spike protein in many organs,
3. vaccine-induced expression of the spike protein induces autoimmune-like inflammation,
4. vaccine-induced inflammation can cause grave organ damage, especially in vessels, sometimes with deadly outcome.

We note that the damage mechanism is which emerges from the autopsy studies is not limited to COVID-19 vaccines only but is completely general—it must be expected to occur similarly with mRNA vaccines against any and all infectious pathogens. This technology has failed and must be abandoned.

While clinical case reports (e.g. [1, 2]) and statistical analyses of accumulated adverse event reports (e.g. [3, 4]) provide valuable evidence of damage induced by mRNA-based COVID-19 vaccines, it is important to establish a causal relationship in individual cases. Pathology remains the gold standard for proof of disease causation. This short paper will discuss some key findings on autopsy materials from patients who died within days to several months after vaccination. For context, some experimental studies are briefly discussed as well.

1 Most of the evidence presented here is from the work of pathologist Prof. Arne Burkhardt, MD

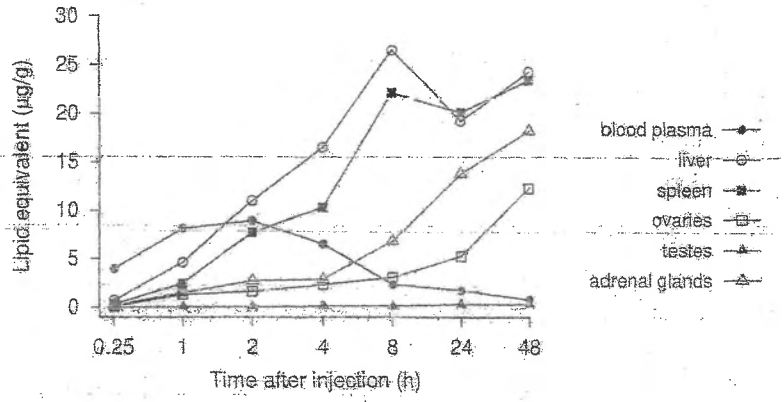
Prof. Burkhardt is a very experienced pathologist from Reutlingen, Germany. With the help of his colleague Prof. Walter Lang, he has studied numerous cases of death which occurred within days to several months after vaccination. In each of these cases, the cause of death had been certified as "natural" or "unknown." Burkhardt became involved only because the bereaved families doubted these verdicts and sought a second opinion.

It is remarkable, therefore, that Burkhardt found not just a few but the majority of these deaths to be due to vaccination.

- Dr. Burkhardt was approached by the families of patients deceased after "vaccination"
- Autopsy materials were examined by standard histopathology and immunohistochemistry
- Based on the findings, most deaths were attributed to "vaccination" with a high to very high degree of likelihood

While all four major manufacturers of gene-based vaccines were represented in the sample of patients studied by Burkhardt and Lang, most patients had received an mRNA vaccine from either Pfizer or Moderna. Some of the deceased patients had received both mRNA- and viral vector-based vaccines on separate occasions.

2 Pfizer's own animal experiments show that the vaccine quickly distributes throughout the body



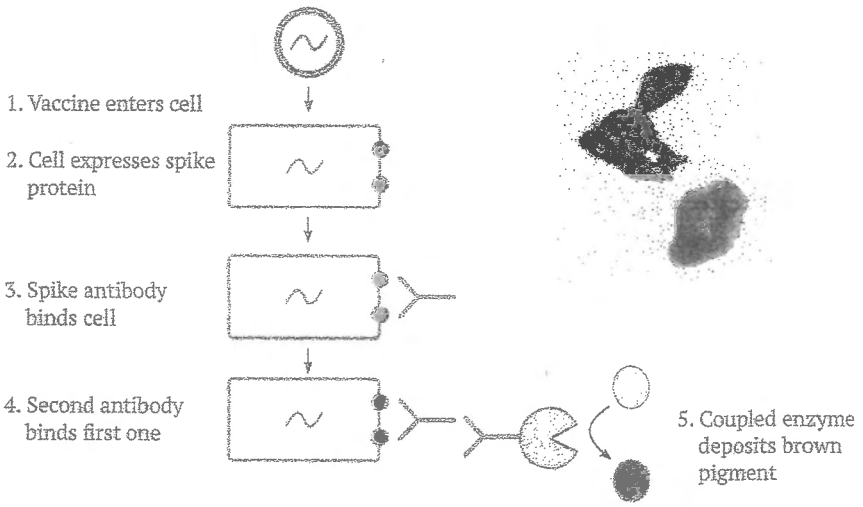
In order to cause potentially lethal damage, the mRNA vaccines must first distribute from the injection site to other organs. That such distribution occurs is apparent from animal experiments reported by Pfizer to Japanese authorities with its application for vaccine approval in that country [5]. Rats were injected intramuscularly with a radioactively labelled model mRNA vaccine, and the movement of the radiolabel first into the bloodstream and subsequently into various organs was followed for up to 48 hours.

The first thing to note is that the labelled vaccine shows up in the blood plasma after a very short time—within only a quarter of an hour. The plasma level peaks two hours after the injection. As it drops off, the model vaccine accumulates in several other organs. The fastest and highest rise is observed in the liver and the spleen. Very high uptake is also observed with the ovaries and the adrenal glands. Other organs (including the testes) take up significantly lower levels of the model vaccine. We note, however,

that at least the blood vessels will be exposed and affected in every organ and in every tissue.

The rapid and widespread distribution of the model vaccine implies that we must expect expression of the spike protein throughout the body. For a more in-depth discussion of this biodistribution study, see Palmer and Bhakdi [6].

3 Expression of viral proteins can be detected with immunohistochemistry



While the distribution of the model vaccine leads us to expect widespread expression of the spike protein, we are here after solid proof. Such proof can be obtained using *immunohistochemistry*, which method is illustrated in this slide for the vaccine-encoded spike protein.

If a vaccine particle—composed of the spike-encoding mRNA, coated with lipids—enters a body cell, this will cause the spike protein to be synthesized within the cell and then taken to the cell surface. There, it can be recognized by a spike-specific antibody. After washing the tissue specimen to remove unbound antibody molecules, the bound ones can be detected with a secondary antibody that is coupled with some enzyme, often horseradish peroxidase. After another washing step, the specimen is incubated with a water-soluble precursor dye that is converted by the enzyme to an insoluble brown pigment. Each enzyme molecule can rapidly convert a large number of dye molecules, which greatly amplifies the signal.

At the top right of the image, you can see two cells which were exposed to the Pfizer vaccine and then subjected to the protocol outlined above. The intense brown stain indicates that the cells were indeed producing the spike protein.

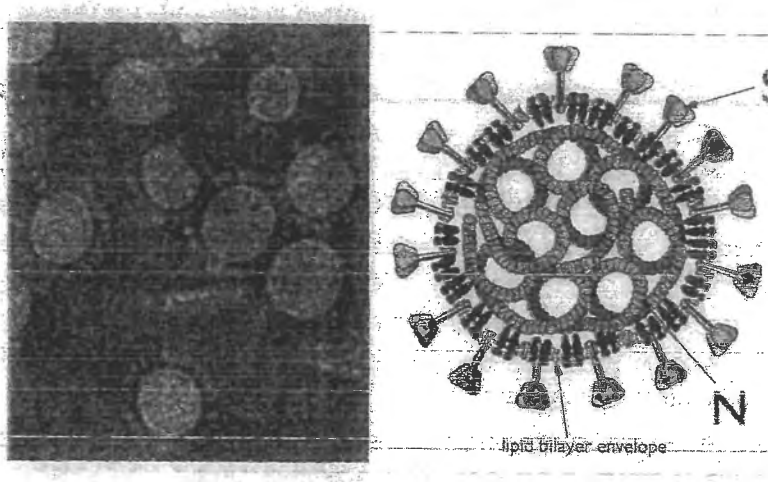
In short, wherever the brown pigment is deposited, the original antigen—in this example, the spike protein—must have been present. Immunohistochemistry is widely used not only in clinical pathology but also in research; it could readily have been used to detect widespread expression of spike protein in animal trials during preclinical development. However, it appears that the FDA and other regulators never received or demanded such experimental data [7].

4 Expression of spike protein in shoulder muscle after vaccine injection

This slide (by Dr. Burkhardt) shows deltoid muscle fibres in cross section. Several (but not all) of the fibres show strong brown pigmentation, again indicating spike protein expression.

While the expression of spike protein near the injection site is of course expected and highly suggestive, we would like to make certain that such expression is indeed caused by the vaccine and not by a concomitant infection with the SARS-CoV-2 virus. This is particularly important with respect to other tissues and organs which are located far away from the injection site.

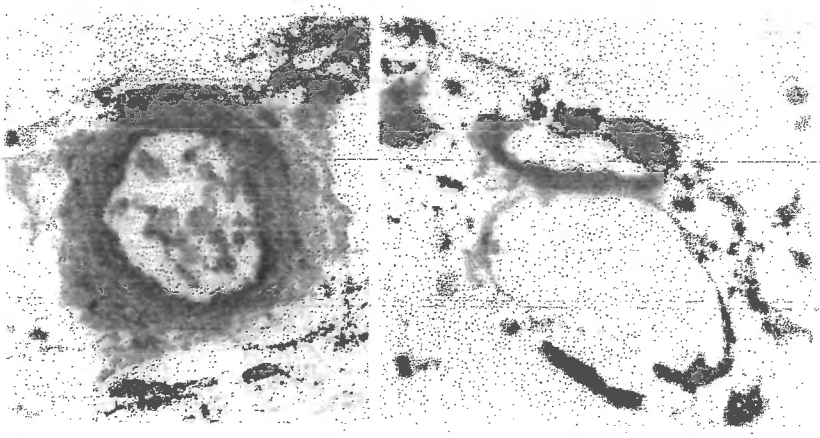
5 Coronavirus particles contain two prominent proteins: spike (S) and nucleocapsid (N)



To distinguish between infection and injection, we can again use immunohistochemistry, but this time apply it to another SARS-CoV-2 protein—namely, the nucleocapsid, which is found inside the virus particle, where it enwraps and protects the RNA genome. The rationale of this experiment is simple: cells infected with the virus will express all viral proteins, including the spike and the nucleocapsid. In contrast, the mRNA-based COVID vaccines (as well as the adenovirus vector-based ones produced by AstraZeneca and Janssen) will induce expression only of spike.

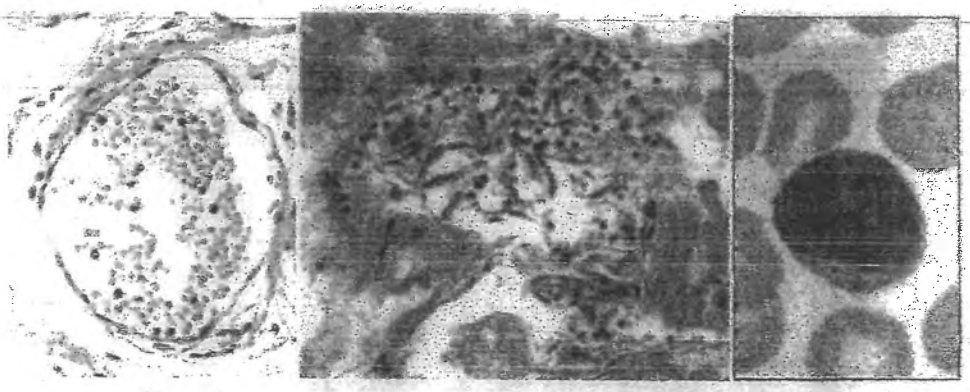
We will see shortly that the strong expression of spike protein in heart muscle after vaccination correlates with significant inflammation and tissue destruction.

8 Expression of spike protein within the walls of small blood vessels



We see spike protein expression in arterioles (small arteries; left) as well as in venules (small veins) and capillaries (right). Expression is most prominent in the innermost cell layer, the *endothelium*. This makes the endothelial cells "sitting ducks" for an attack by the immune system.

9 Endothelial stripping and destruction of a small blood vessel after vaccination



We now turn to the evidence of immune attack on the endothelial cells which produce the spike protein. On the left, a normal venule, delimited by an intact endothelium and containing some red blood cells and few white blood cells (stained blue) inside.

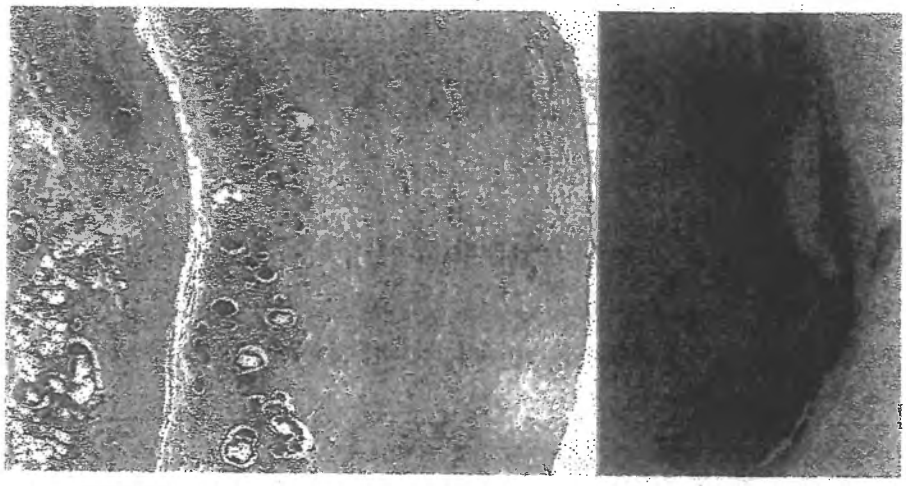
The image on at the centre shows a venule that is being attacked and destroyed by the immune system. The outline is already dissolving, and the spindle-shaped (and swollen) endothelial cells have peeled off from the vessel wall. Furthermore, we see lymphocytes—the small cells with dark, round nuclei and with very little cytoplasm around them; a single lymphocyte (at much higher magnification) is shown on the right.

Lymphocytes are the backbone of the specific immune system—whenever antigens are recognized and antibodies are produced, this is done by lymphocytes. Also among

the lymphocytes we find cytotoxic T cells and natural killer cells, which serve to kill virus-infected cells—or ones that look to them as if infected, because they have been forced to produce a viral protein by a so-called vaccine.

A crucial function of the endothelium is to prevent blood clotting. Thus, if the endothelium is damaged, as it is in this picture, and the tissues beyond it make contact with the blood, this will automatically set off blood clotting.

10 A crack in the wall of the aorta, lined by clusters of lymphocytes, leading to aortic rupture

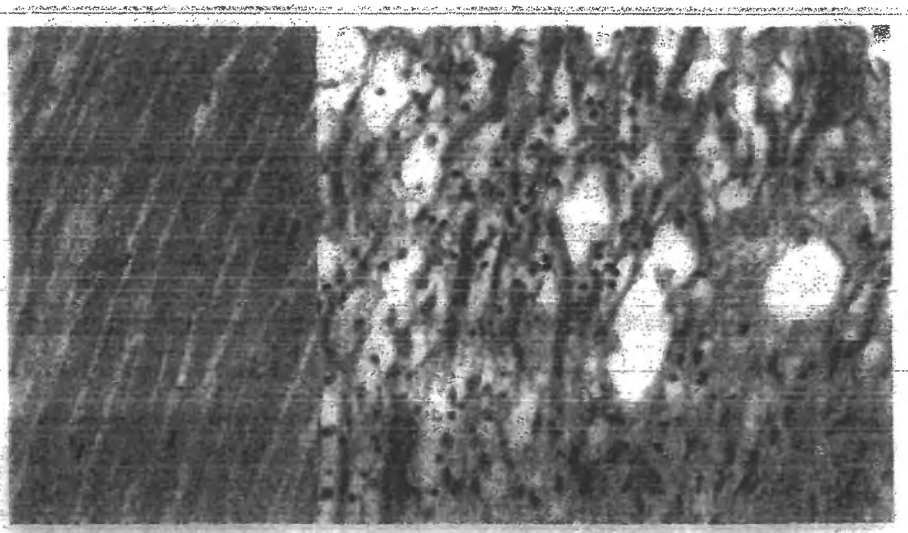


On the left, a section through the wall of an aorta. This picture is taken at an even lower magnification than the one before; the lymphocytes now appear as just a cloud of tiny blue specks. To the left of this blue cloud, we see a vertical crack running through the tissue. Such a crack is also visible macroscopically in the excised specimen of an aorta shown on the right.

The aorta is the largest blood vessel of the body. It receives the highly pressurized blood ejected by the left ventricle of the heart, and it is thus exposed to intense mechanical stress. If the wall of the aorta is weakened by inflammation, as it is here, then it may crack and rupture. Aortic rupture is normally quite rare, but Prof. Burkhardt found multiple cases in his limited number of autopsies. Some of the affected aortas were also shown to have expressed the spike protein.

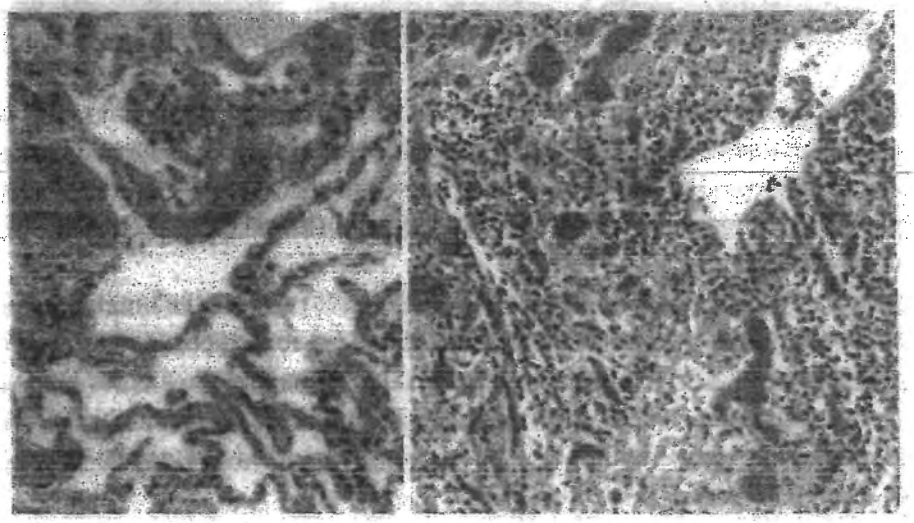
11 Healthy heart muscle tissue, and lymphocytic myocarditis

In Slide 7, we saw that heart muscle cells strongly expressed the spike protein after vaccine injection. Here, we see the consequences. The picture on the shows a sample of healthy heart muscle tissue, with regularly oriented and aligned heart muscle fibres. On the right, we see a heart muscle sample from one of the autopsies. The muscle fibres are disjointed and disintegrating, and they are surrounded by invading lymphocytes. Burkhardt found myocarditis in multiple of his deceased patients.



12 **Lymphocytic infiltration and proliferative inflammation in lung tissue**

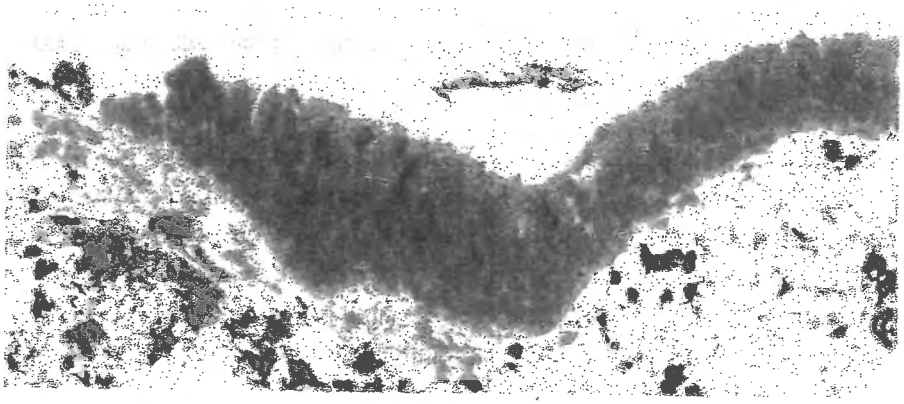
On the left, we see healthy lung tissue, with air-filled spaces (the alveoli), delimited by delicate alveolar septa with embedded, blood-filled capillaries. We also see some larger blood vessels.



On the right hand side, we see lung tissue overrun by lymphocytes. The air-filled spaces have largely disappeared and been filled with scar (connective) tissue. This vaccine-injected patient would obviously have had very great trouble breathing.

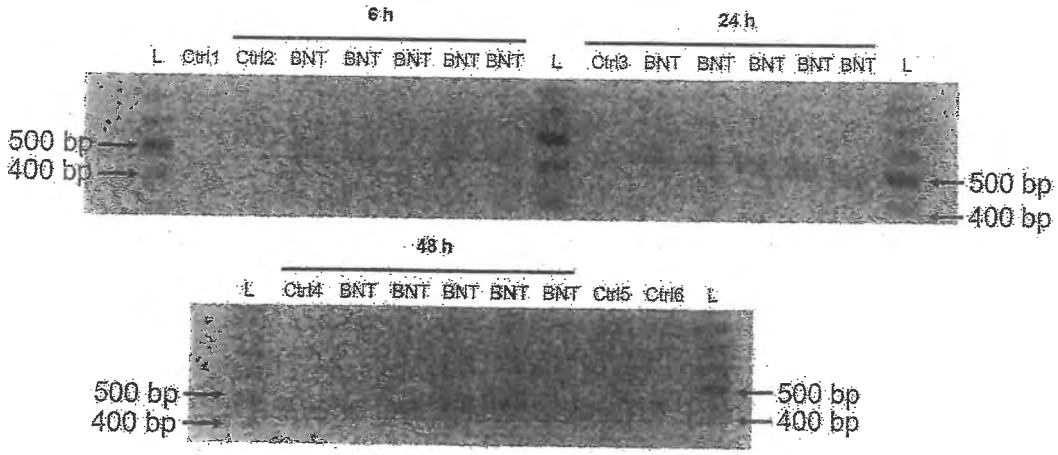
Lymphocytic infiltration, inflammation and destruction were also observed in many other organs, including the brain, the liver, the spleen, and multiple glands. However, instead of illustrating them all, we will conclude the pathological evidence with another immunohistochemistry result, which strikingly shows the long duration of spike protein expression.

13 Vaccine-induced expression of spike protein in a bronchial biopsy nine months after vaccination



The slide shows a sample of bronchial mucous membrane, from a patient who is alive but has suffered respiratory symptoms ever since being vaccinated. We see several cells in the uppermost cell layer that strongly express spike protein—and this even nine months after his most recent vaccine injection! While this is indeed the most extreme case of long-lasting expression, there is evidence both from Burkhardt's autopsies and from published studies on blood samples [8] or lymph node biopsies [9] to indicate that expression does last several months.

14 The Pfizer vaccine mRNA gets copied ("reverse-transcribed") into DNA and inserted into the cellular genome



The official mRNA vaccine narrative maintains that the modified mRNA contained in the vaccine will not be replicated in vivo; expression of the spike protein should therefore cease once the injected RNA molecules have been degraded.

The limited experimental studies available [10, 11] suggest that the injected modified mRNA should be degraded within days to a few weeks of the injection. This is obviously difficult to square with the observed long-lasting expression; in some form or other, the genetic information appears to be perpetuated in vivo.

A recent experimental study from Sweden [12] has shown that human-derived cells can copy the Pfizer mRNA vaccine into DNA and then insert it into their own chromosomal DNA. The image shows the key evidence from this study. The cells were exposed to the vaccine for the lengths of time indicated. Cellular DNA was then isolated, and inserted DNA copies of the vaccine mRNA detected by PCR amplification of a fragment 444 base pairs (bp) in length.

All samples labelled with "BNT" had been treated with the vaccine, and they all show a PCR product of the expected length, as is evident from comparison to a DNA fragment length standard ("L"). Samples labelled with "Ctrl n" were controls: Ctrl 1–4 contained DNA from cells not incubated with vaccine, Ctrl 5 contained RNA (not DNA) from vaccine-treated cells; Ctrl 6 contained the same but was additionally treated with RNAse, which step was also performed in the purification of DNA samples. As expected, none of the control samples contain the PCR product.

Considering Aldén's observation of DNA insertion in every single experimental sample, it seems highly likely that this will also occur in vivo. Beyond providing a plausible mechanism for perpetuating the expression of spike protein, DNA insertion also poses risks of genetic damage, leading to cancers and leukemias.

15 Summary

The evidence presented here clearly demonstrates a chain of causation from vaccine injection to

- rapid distribution of the vaccine through the bloodstream,
- widespread spike protein expression, prominently in blood vessels, and
- autoimmune-like inflammation and organ damage.

Vaccine-induced vascular damage will promote blood clotting, and clotting-related diseases such as heart attack, stroke, lung embolism are very common in the adverse events databases [4, 13].

In addition to autoimmune-like inflammation, other disease mechanisms, including prion-mediated CNS degeneration [14], aberrant vascular protein deposition (amyloidosis) [15, 16], and lipid nanoparticle toxicity [6], are plausible but require further study and corroboration. Overall, these vaccines can no longer be considered experimental—the "experiment" has resulted in the disaster that many medical doctors and scientists predicted from the outset [17]. The vaccination must be stopped, and all approvals and authorizations of their use must be revoked.

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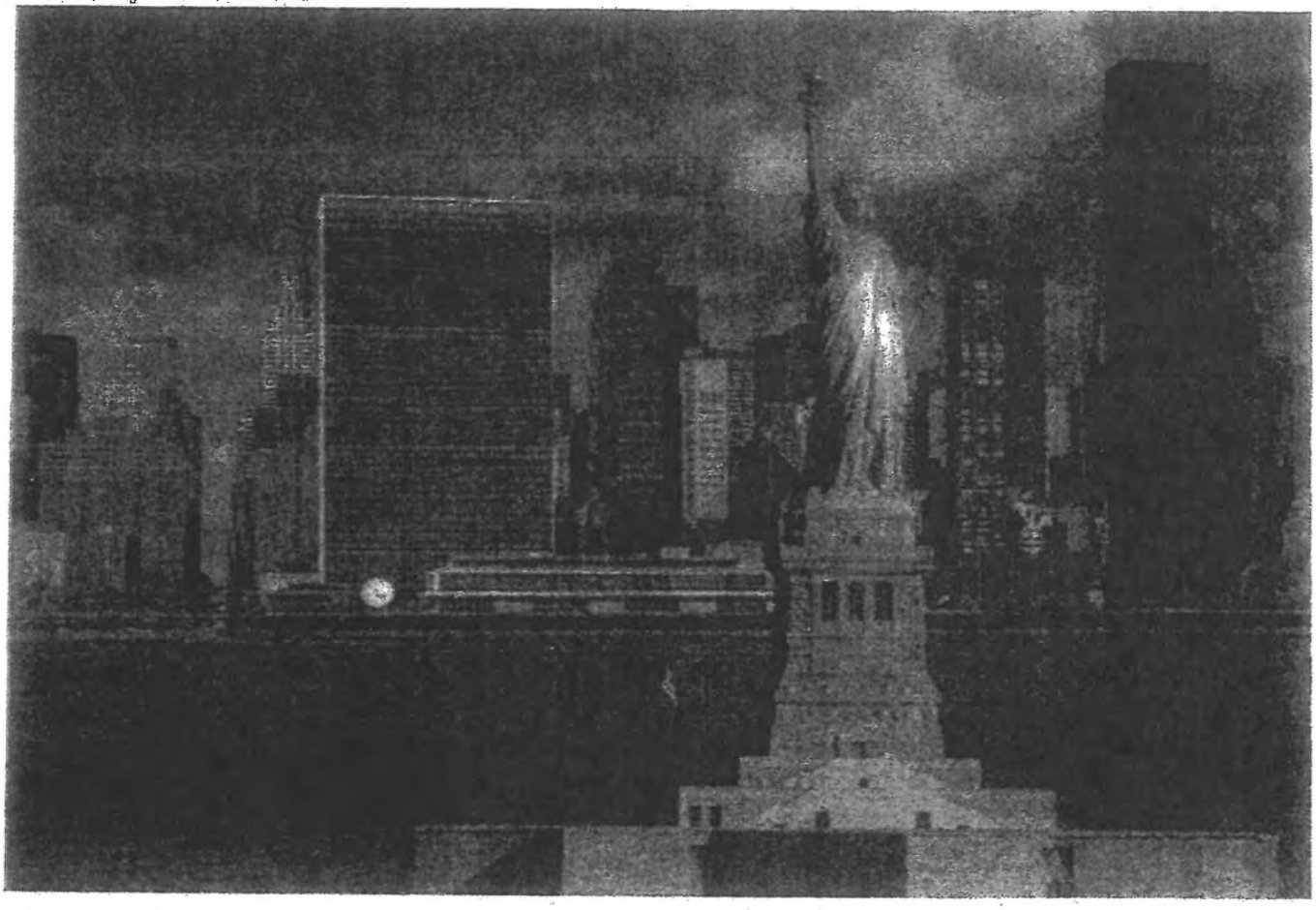
Blasting CCP Influence, Lawmakers File Bill to Get US Out of UN & WHO

written by Alex Newman



Outraged by the growing influence of the mass-murdering Chinese Communist Party and other perceived problems, U.S. lawmakers recently re-introduced legislation that would end U.S. membership in the United Nations and its agencies, such as the UN World Health Organization (WHO).

In addition to ending U.S. government involvement with the UN, the American Sovereignty Restoration Act (H.R. 7806) would remove the UN's controversial headquarters from U.S. soil and protect American troops from having to serve under UN command.

U.S. Representative Mike Rogers, a conservative Republican representing eastern Alabama, has been the lead sponsor of the bill in several congresses so far. He has raised numerous concerns over the years, including corruption, waste, hostility to Israel, opposition to fundamental American principles, the UN's hatred of the Second Amendment, and more.

"The United Nations has repeatedly proven itself to be an utterly useless organization," explained Rogers in a statement announcing the re-introduction of the bill last month, doubling down on previous comments referring to the UN as a "disaster."

Some of the congressman's major concerns are the UN's growing hostility to genuine human rights and its increasing subservience to the dictatorship in Beijing and others hostile to individual liberty and the United States.

"The UN's founding charter states the UN's mission 'to reaffirm faith in fundamental human rights, in the dignity and worth of the human person, in the equal rights of men and women and of nations large and small,'" added Rogers in the statement. "However, the UN High Commissioner for Human Rights Michelle Bachelet has proven herself to be nothing more than a puppet for the Chinese Communist Party — aiding the CCP in playing down the very real and horrifying genocide being carried out against Uyghurs."

Indeed, *The New American* has been exposing the "Socialist" Bachelet for years. From her close ties to the communist movement in Latin America and Beijing to her ongoing anti-American diatribes calling for restricting rights in America, Bachelet has become extremely controversial. Concerns about the UN official's abuse of diplomatic immunity to shield her and others from criminal probes are also growing.

Rogers blasted the UN's cozy relationship with the Chinese Communist Party. "It's unconscionable that China continues to sit on the UN Human Rights Council even as it carries out this disturbing genocide on top of its numerous and daily violations of basic human rights," the Republican congressman said.

"It's clear the UN has abandoned the ideals set in its founding charter and that's why, among many other reasons, I've reintroduced legislation to withdraw the United States from the UN," he added.

When introducing the bill in 2019, Rogers blasted the UN as an "inefficient bureaucracy" and a "complete waste of American tax dollars." Saying the legislation was one of his top priorities, Rogers noted that the global organization "works against America's interests around the world" and continues to "attack our rights as U.S. citizens."

Another key element of the bill would end U.S. involvement in the disgraced World Health Organization. Among other scandals, the UN agency is led by a former communist terror leader backed by Beijing, and was repeatedly exposed parroting the CCP's talking points.

"The WHO lost all credibility when they chose to put public health second to the Chinese Communist Party by helping the CCP cover up the origins of COVID-19," continued Rogers, blasting the UN WHO as "corrupt."

Reacting to similar concerns, President Donald Trump started the process to remove the U.S. from the WHO, drawing widespread applause from conservatives and Republicans across America.

Joe Biden promptly re-engaged with the UN agency after taking power, though numerous congressional efforts to stop funding for and end U.S. involvement in WHO continue. (Trump did get the United States out of UNESCO, the UN's "education" agency, and so far Biden has not been able to reverse that.)

Co-sponsors of the latest iteration of the American Sovereignty Restoration Act include Rep. Thomas Massie (R-Ky.), Rep. Diana Harshbarger (R-Tenn.), Rep. Paul Gosar (R-Ariz.), and Rep. Ronny Jackson (R-Texas).

Massie, a longtime champion of the #Amexit movement to get the United States out of the UN, previously told *The New American* that there are many reasons why the U.S. government should cut all ties with the controversial global organization.

Congress Redefines Marriage

AP Images

"Today, we stand up for the values the vast majority of Americans hold dear: a belief in the dignity, beauty and divinity — spark of divinity — in every person, and abiding respect for love so powerful that it binds two people together."

House Speaker **Nancy Pelosi** *made this statement just prior to the actual House vote to codify on the federal level what is commonly called same-sex marriage. Her inclusion of the term "divinity" in her remarks means she either doesn't know or doesn't care about the biblical boundary that identifies marriage as the bond between one man and one woman. To find out how all members of the House and Senate voted on this legislation, see the Freedom Index, House vote no. 39 and Senate vote no. 39, in this issue.*

AP Images

Who Is the World's Most Dangerous Person?

"I get asked 'Who's the most dangerous person in the world? Is it Chairman Kim, is it Xi Jinping?' The most dangerous person in the world is Randi Weingarten. It's not a close call."

Former secretary of state under President Donald Trump and possible 2024 GOP presidential candidate **Mike Pompeo** *made this incredible assertion during an interview by the global news platform known as Semafor. Explaining his rationale, Pompeo added, "If you ask, 'Who's the most likely to take this republic down?' It would be the teachers unions, and the filth that they're teaching our kids, and the fact that they don't know math and reading or writing." Weingarten is president of the American Federation of Teachers.*

White House Press Secretary Claims President Biden Has Been to the U.S.-Mexico Border

"White House press secretary Karine Jean-Pierre insisted that President Biden has been to the U.S.-Mexico border despite no record of him having been there — even as vice president."

As reported by **Fox News**, *when reporter Peter Doocy asked White House Press Secretary Karine Jean-Pierre to indicate when the president had actually visited the border, she refused to provide any date and then amazingly insisted that the border was in a shambles because the GOP has refused to work with Biden on fixing the problem.*

Prominent Doctor and Former Trump White House Coronavirus Task Force Member Blasts FDA

"In my 30 years as a doctor, the unprecedented approach taken by the FDA where ivermectin was somehow forbidden if you used it for an off-label purpose was a shocking interference of the ability of a doctor to do his job. [In the past] ivermectin was approved by the FDA and found so safe that billions of doses have been given without the need for a doctor's prescription until the FDA prohibited its use to treat Covid-19."

A senior fellow in health care policy at Stanford University, Dr. **Scott Atlas** *has termed the FDA's position on ivermectin eminently deserving of the suit recently filed against the agency by three prominent doctors challenging that position.*

White House Press Secretary Dismisses Twitter's Recent Disclosures of Its 2020 Coverup of Hunter Biden's Activities

AP Images

"[This] is full of old news, if you think about it."

The damaging information on Hunter Biden's laptop was not big news during the 2020 presidential campaign because the story was either ignored or dismissed as Russian disinformation when it was reported. But now that it is uncontested that the laptop story is legitimate, and Twitter under new CEO Elon Musk has begun providing details about the coverup in a series of releases, White House Press Secretary **Karine Jean-Pierre** *cavalierly refused to comment when a reporter asked if the censorship was appropriate, dismissing the story as "old news."* ■

— Compiled by John F. McManus

Fauci Says COVID-19 and Influenza Vaccines Don't Work Well

Dr. Anthony Fauci is among the growing number of officials who are acknowledging that the COVID-19 vaccines don't work well against infection.

Vaccines against both COVID-19 and influenza have "deficiencies," including that they "elicit incomplete and short-lived protection against evolving virus variants that escape population immunity," Fauci, until recently President Joe Biden's chief medical adviser, and other top National Institutes of Health officials wrote in a recent paper.

"With the imperfections of these vaccines, it seems a public health imperative to aggressively pursue better vaccines and vaccination strategies," they wrote.

CDC Adds COVID-19 Vaccines to Child Immunization Schedule

The U.S. Centers for Disease Control and Prevention (CDC) has added COVID-19 vaccines to its routine immunization schedule for children and adults.

According to the CDC's 2023 immunization schedule for children and adolescents, two or three doses of COVID-19 vaccines have been recommended beginning with infants who are just 6 months old. Children in the age group of 6 months to 4 years, and 5 years to 11 years are recommended COVID-19 vaccines from Moderna or Pfizer. Among children aged 12 to 18, Novavax vaccines also are recommended in addition to Pfizer and Moderna.

In the list for adults, two or three doses of COVID-19 vaccines have been recommended from the age of 19 years. The 2023 COVID-19 vaccine recommendation for kids and adults is included among other typically recommended vaccines for measles, flu, rubella, and so on.

Advisers to the CDC in 2022 recommended adding the vaccines to the schedule.

Messenger RNA Sequences Found in Blood 28 Days After COVID-19 Vaccination

Messenger RNA sequences from the Pfizer and Moderna COVID-19 vaccines were found in the blood of multiple individuals weeks after vaccination, according to a study.

Researchers in Denmark analyzed samples from the vaccinated and detected partial or even full sequences of the messenger RNA (mRNA) following vaccination. The sequences were found as late as 28 days after vaccination, or the longest time period the study analyzed.

The findings mean that the mRNA, which is situated in lipid nanoparticles for delivery into the body, lingers for much longer than officials in the United States and elsewhere acknowledge.

Top Officials Discussed Vaccine 'Adverse Event Issue' and Pregnancy: Emails

CONTINUED FROM A1

Dr. Amanda Cohn, chief medical officer of the CDC's National Center for Immunizations and Respiratory Diseases, replied.

"We have a meeting with Rochelle at 3:30 about if we should say anything or wait until we have more definitive information ... I will let you know where we land," Cohn wrote. Dr. Rochelle Walensky is the head of the CDC.

"I'm not sure there is a right answer," Cohn added.

The emails were obtained by the nonprofit Judicial Watch, which sued the U.S. government for failing to appropriately respond to a Freedom of Information Act request for messages regarding adverse events, deaths, or injuries caused by the COVID-19 vaccines.

Adverse events include health issues such as arthritis or heart inflammation.

No other emails about the "adverse event issue" were included in the latest tranche obtained by the nonprofit.

"I respectfully decline to comment," Marks told The Epoch Times in an emailed statement when asked what the issue was.

Cohn and the CDC didn't respond to requests for comment.

The CDC currently recommends virtually all Americans aged 6 months and older, including pregnant women, get a COVID-19 vaccine and multiple booster shots.

The original trials didn't include enough information "to make conclusions about the safety of the" vaccines from Pfizer, Moderna, and Johnson & Johnson, according to FDA documents.

Authorities have relied on observational data, including a study from the CDC, a corrected version of which was published in October 2021.

Pfizer conducted a post-authorization trial of its vaccine in pregnant women that was labeled completed in mid-2022 but results haven't been reported publicly as of yet.

"We think that part of the reason is because the results are so bad," Linda Wastila, a professor at the University of Maryland whose expertise is in pharmacotherapy and drug policy, told The Epoch Times.

Pfizer didn't respond to a request for comment.

Confidentiality Agreement

The FDA vaccine advisory committee met on Dec. 10, 2020, to consider whether to advise the FDA to authorize Pfizer's vaccine.

During the meeting, officials revealed that two postvaccination cases of severe allergic shock (anaphylaxis) had been recorded in the United Kingdom out of 6,000 doses administered on the day the events occurred.

"At this point, we are seeking further information from the MHRA under our confidentiality agreement to learn more about this and to really tease that out," Marion Gruber, an FDA official at the time, told the panel.

CHRISTOF STACHE/AFP VIA GETTY IMAGES

A health worker fills a syringe with a COVID-19 vaccine in Freising, Germany, on Feb. 2, 2021.

The Medicines and Healthcare products Regulatory Agency (MHRA) is the British equivalent of the FDA.

The newly obtained set of emails showed U.S. and UK officials discussing the cases of anaphylaxis throughout December, including as early as Dec. 9, 2020.

Marks wrote that it would "be very helpful if our Office of Vaccines could receive additional details" from MHRA "under the terms of our mutual confidentiality agreement."

Jonathan Mogford, a UK official, sent back data, stating, "If I can just remind—information shared under our confidentiality agreement."

The MHRA declined to provide a copy of the confidentiality agreement, and the FDA didn't respond to emailed questions about the pact.

"It again took a lawsuit for the Biden administration to hand over, albeit heavily redacted, information regarding the safety of the COVID vaccines that the public has every right to know," Judicial Watch President Tom Fitton said in a statement. "This disturbing batch of new documents have uncovered a secret confidentiality agreement tied to COVID vaccine safety issues and emails that raise new questions about the vaccines and pregnancy."

The Epoch Times has submitted Freedom of Information Act requests to try to uncover more information about the adverse event issue and the confidentiality agreement. Previous requests have unearthed secret data on COVID-19 vaccines, including showing how the CDC identified hundreds of potential safety issues with the Pfizer and Moderna vaccines.

People with a history of severe allergic reactions to any vaccine were excluded from Pfizer's trial and people with a history of allergic reactions to any components of the Moderna vaccine were excluded from Moderna's trial. No such exclusions were reported for the Johnson & Johnson trial.

The fact sheets for all three vaccines in the United States were updated to include warnings about severe allergic shock, including saying the vaccine shouldn't be administered to people with a known history of severe allergic reaction to any components of the shot.

COVID VACCINES

States That Have Banned COVID-19 Vaccine Mandates in School, Bucking CDC

JACK PHILLIPS

As the Centers for Disease Control and Prevention (CDC) adds COVID-19 vaccines to its routine immunization schedule for children and adults, at least 20 states have passed legislation or issued rules barring school systems from mandating the vaccines.

The National Academy for State Health Policy, in a Feb. 6 update, shows there are about two dozen states that have barred COVID-19 vaccines from being included in mandates for schools and students: Alabama, Arizona, Arkansas, Florida, Georgia, Idaho, Indiana, Iowa, Kansas, Michigan, Mississippi, Montana, New Hampshire, Ohio, Oklahoma, South Carolina, South Dakota, Tennessee, Texas, Utah, and West Virginia.

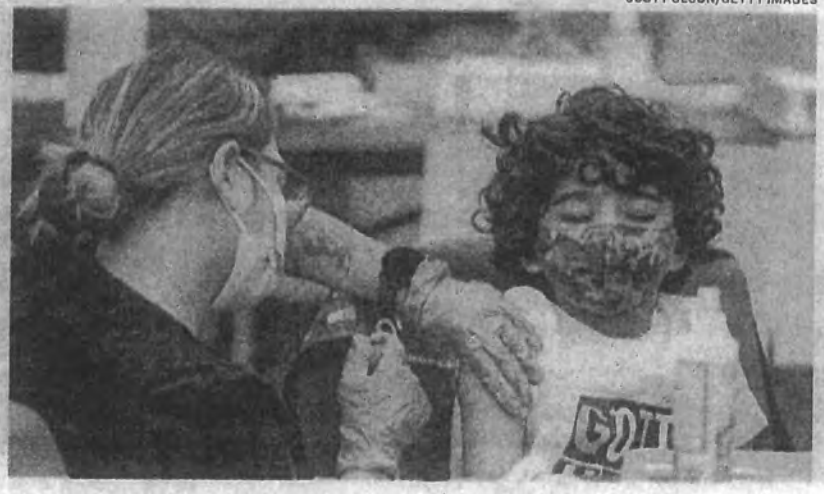
Officials in other states, including California, have announced they will not require students to get the shots. On Feb. 3, California's Department of Public Health said "that the state will not require the COVID-19 vaccine for school attendance," the website noted.

To date, not a single state has passed legislation or implemented an order to mandate COVID-19 vaccines to attend class, according to the academy. But the District of Columbia has a student COVID-19 vaccine mandate, although it has been delayed until the start of the 2023–2024 school year, after the D.C. City Council voted last November to do so.

The academy has issued updates on COVID-19 vaccine mandates and bans on them since the CDC signaled that it could add those vaccines to its recommended child immunization schedule. While the CDC's guidelines are only recommendations, many states and municipalities rely on them to set policies.

Laws in at least 31 states and the District of Columbia require the vaccines on the CDC schedules to be taken by children for school attendance, said the Policy, Practice, and Prevention Research Center at the University of Illinois Chicago's School of Public Health last year. Other states sometimes impose requirements that mostly align with the schedules.

SCOTT OLSON/GETTY IMAGES



A child receives a COVID-19 vaccine in Chicago on Nov. 12, 2021.

As long as I'm around and as long as I'm kicking and screaming, there will be no COVID shot mandates for your kids. That is your decision to make as a parent.

Florida Gov. Ron DeSantis

Response

Last year, when the CDC advisory panel voted to recommend adding the COVID-19 vaccines to the childhood immunization schedule, a number of governors had indicated they would not make them mandatory.

"Under my watch, there will be no COVID vaccine mandates for kids—period," said Republican Gov. Kim Reynolds of Iowa. "In fact, we signed a law that prevents it. It's the parent's decision, not the government's."

Florida Gov. Ron DeSantis noted that some parents are concerned about the CDC's guidelines, telling reporters there is a "fear" that schools would force shots on children. "So I just want to let everyone be clear. You know, as long as I'm around and as long as I'm kicking and screaming, there will be no COVID shot mandates for your kids. That is your decision to make as a parent," he said.

And at the time, Montana Gov. Greg Gianforte, a Republican, wrote: "I trust parents to raise their kids and do what's best for their kids' health. On my watch, the State of Montana will not mandate the COVID-19 vaccine."

A spokesperson for California Gov. Gavin Newsom last year said that the "main impact" of the CDC's recommendation would be that "health insurance companies will be required to cover the cost of the immunization and that the federal government can continue to provide it for free to low-income families."

Schedule Update

The CDC's 2023 immunization schedule for children and adolescents, released on Feb. 9, includes shots for flu, MMR (measles, mumps, and rubella), polio, and other inoculations—and now COVID-19. The CDC doesn't have the authority to issue mandates for vaccines.

Last year, all members of the CDC's Advisory Committee on Immunization Practices voted to add the Moderna, Pfizer, and Novavax vaccines to the 2023 schedules, asserting that the vaccines can prevent severe disease—despite studies showing waning effectiveness of the shots.

"We view this as COVID is here to stay," Dr. Matthew Daley, one of the advisers, said in a meeting. "When I think about the routine immunization schedule as a pediatrician, I think of it as an opportunity to prevent serious disease and death. And if something is added to the schedule, it's because I feel like the benefits continue to strongly outweigh the risks."

Zachary Stieber contributed to this report.

On March 21, 2023, the City received a written public comment for the City Council's regular meeting of March 22, 2023, in the form of photocopied pages from the book *It's Perfectly Normal: Changing Bodies, Growing Up, Sex, Gender, and Sexual Health*. Because the City does not have permission from the copyright owners of the book, the photocopied pages are not being distributed on the City's website or through its agenda system.

Members of the public interested in viewing the photocopies may do so in the Office of the City Clerk during normal business hours. The City Clerk's Office is located in City Hall at 400 Grand Avenue.