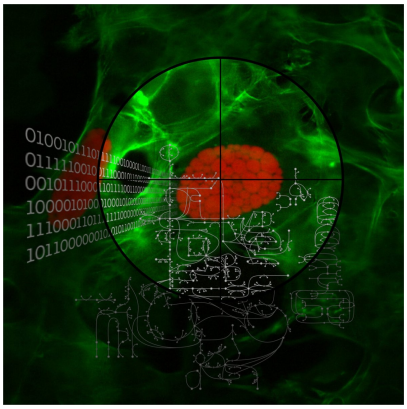

EPIGENETICS: Vaccines Are Deleting Human Genes & Transfecting Cells With Ebola/Marburg

Description

Stockholm University just released a ***scientific horror***. The “spike” protein in the Covid-19 “vaccines” are penetrating the cells of the vaccinated, reaching the cell nuclei, and impairing your cell’s ability to repair damaged DNA.

Pharmaceutical “vaccines” are silencing the genes responsible for DNA repair and deleting them *forever* in humans.

I joined Stew Peter’s Show once again, to ***break this shocking discovery*** on Red Voice Media.

Johnson & Johnson ***uses Adenovirus 26*** (Ad26) in its vaxxine. J&J openly admits that their Ad26 vector “codes your cells to produce a spike protein” but they don’t tell you they’re also deleting your genes.

The **U.S. patent #20140017278** for Adenovirus 26 and 35 Filovirus, openly states that it codes your cells with the Ebolaand Marburg chimeric proteins.

> “The filovirus antigenic protein is usually a glycoprotein from an Ebola virus or a Marburg virus.”

The J&J Adenovirus 26 vector ***deletes your E1 gene***. The patent also states that it deletes the E1 gene in Humans. This is known as the ***X Chromosome***. The E1 gene is required for accurate and instant repair of damaged DNA. Deletion of this gene is lethal.

E1 gene deletion causes ***embryonic lethality*** which means permanent sterility for men and women. It causes Lactic Acidosis in children which is the lack of oxygen in the blood. E1 gene deletion ***causes rapid cancer*** growth, ***Thrombosis***, and the *coagulation of the blood* which leads to clotting. Blood clotting is the main reason people are dying from the Covid vaxxines.

E1 gene deletion causes Mitochondrial DNA-Associated Syndrome which is a process of glucose metabolism deficiency that exists in various diseases such as Alzheimer's, epilepsy, diabetes-associated cognitive decline, and severe neurological disorders such as ***Leigh's Syndrome***. There's a progressive loss of mental cognition and typically results in death within *two* or three years, usually due to *respiratory failure*.

Without your E1 gene, your cells will literally *self-suicide*.

U.S. patent #10695417 is the Human Adenovirus 5 vector that contains E1 and E2B gene deletions. The Adenovirus 5 vector is ***used in Sinovax*** and it ***encodes the cells*** with Ebola Glycoprotein. Glycoprotein -41 and glycoprotein -120 are ***HIVviral coat Proteins*** and Glycoprotein plays a ***key role in reproduction***.

Sinovac is used in China, Chile, Brazil, Turkey, and the Philippines.

The Adenovirus 5 vector was developed through ***Gain-of-Function research***.

Knocking out the E2B gene caused ***sterilization in male mice***.

You can order the chimeric messenger RNA of the Lentivirus and the Adenovirus 5 vectors or Baculoviridae online from ***Thermo Fischer***, for recombinant cross-species genomics (cloning).

Thermo Fischer explains how the Adenovirus 5 targets and***entersthe bronchial epithelial cells***(lungs) and *deletes* the E3 and E4 genes, intentionally inducing ***Sjögren's syndrome*** which is long-term autoimmunity (AIDS).

Loss of your E4 gene deletes your cognitive function. Deletion of your E3 gene degenerates your brain, causes dementia, gradual loss of memory, judgment, and the overall ability to function.

Here's a study describing how you ***knockout the E3 gene***.

The knockout of these genes affects the moisture-producing glands of your body. It's not the "spike protein" that's causing the blood of the vaxxed to coagulate, it's the gene silencing (deletion). Without moisture, your blood coagulates and clots. The deletion of these genes also causes gastrointestinal disorders.

Adenovirus 5 also alters the cell signaling pathways and leads to Lymphoma due to destruction to the immune system. This causes cancers to grow and the blood to coagulate.

This is proof positive they're creating the next "pandemic" with lethal injections that will gradually induce AIDS in the inoculated masses through gene deletion.

The pharmaceutical cartel has not only injected Ebola and Marburg into people but they're also ***transfecting people's cells*** with these catastrophic chimeric pathogens. The vaxxed will battle chronic infections and lifelong disabilities while the cells of the vaxxed continuously replicate with the synthetic genetic sequences of Ebola and Marburg until it kills them unless they ***detoxify continuously***.

The immune system of the vaxxed is ***depreciating 5% each week*** according to a recent UK Government study. Everyone who vaxxed age 30 and above, will have no immunity left by Christmas. But that's not all.

As the cells of the vaxxed replicate Ebola and Marburg "spike proteins" and their cells decay and die, they will shed the chimeric disease throughout the population via ***transmission***. Therefore it's crucial for the unvaxxed to ***continuously detoxify*** as well.

According to a ***UK government declaration*** from the NIH, we are presently in a Phase III clinical trial on *Humans* using the *Adenovirus 5* vector to "fight Covid-19" which began on January 22, 2021.

> "This is a global phase III clinical trial to evaluate efficacy, safety, immuogenicity of Ad5-nCoV manufactured by Cansino and Beijing Institute of Biotechnology in health adults aged 18 years old and above."

So the UK Government is partners with the Chinese Communist Party (CCP) to exterminate Humans.

The World Health Organization (WHO) also published on their website that we are in an Ebola Vaccine Stage III Clinical Trial.

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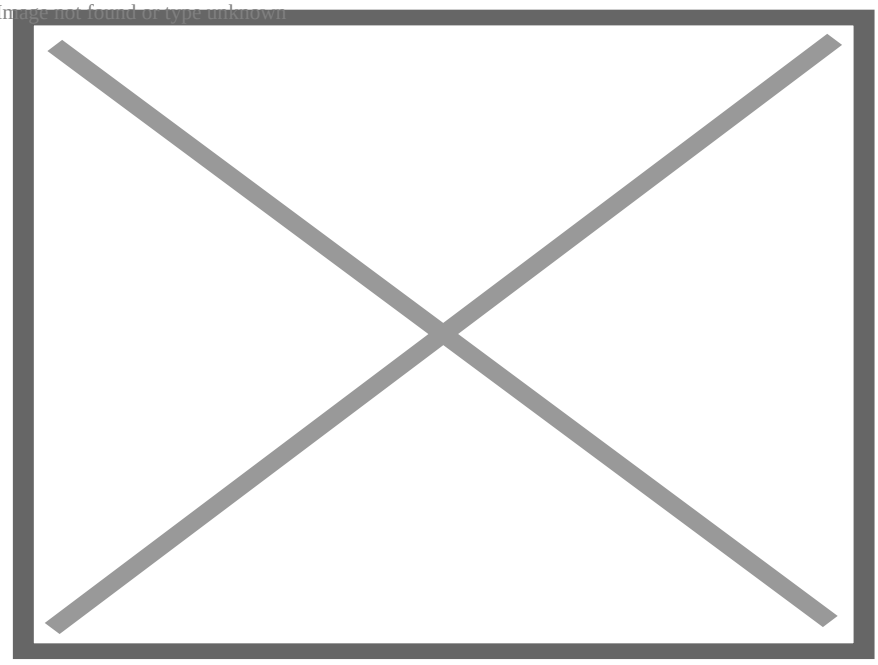

WHO published that we are in an Ebola Vaccine Stage III Clinical Trial

Later the WHO ***scrubbed it*** from their website.

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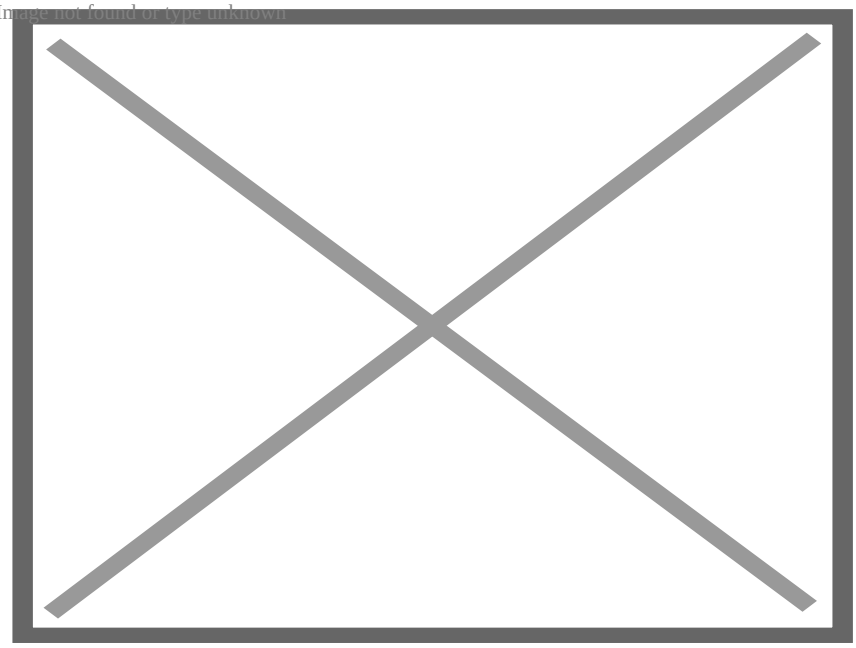

Viruses are still an***unproven theory***. Given the advances in scientific lab equipment and considering that we're able to observe nanotechnology under microscopy and spectroscopy, would somebody please explain to me why we still can't identify a virus? Could Germ Theory be the big pharmacopeia lie in modern medicine and ***Terrain Theory***be the more relevant truth?

Governments ***enhanced the airborne transmissibility*** in mammals (Humans) of highly virulent avian influenza strains. The history of making pathogens transmissible goes back at least to the synthesis of viable influenza H1N1 *from 1918*. Incidentally, the ***2009 swine flu Pandemic***was also induced by inoculation using the H1N1 vaxxine. So this is nothing new under the sun.

Sinister shadow governments have been weaponizing nature, producing diseases through vaxxine injection and genetically manipulating Humans for at least the past 100 years. But where did they get this technology to do it?

The U.S. National Library of Medicine revealed something rather interesting. The USSR's 'invisible anthrax' is a Gain-of-Function bioweapon created by introducing an ***"alien gene"*** into Bacillus anthracis (bacteria). That's how they made Anthrax. They used an *alien gene* and genetically altered bacterial immunological properties to produce a deadly pathogen to Humans. Where did they get an alien gene from? A UFO crash perhaps? Negotiations with *other beings*? Your guess is as good as mine.

The U.S. Government has been testing this germ warfare technology on its own military troops since the 1950s, using Adenovirus 4, Ad5, and Ad7 vectors with ***HIV encoding Envelope*** (clade C.1086). The Ad7 vector delivered in enteric capsules has been used to***"vaccinate" U.S. military personnel*** "against respiratory and gastrointestinal illness", since the 1970s.

Monkeypox is also made from the same bacterial pathogens as Ebola and Marburg. Ebola and Marburg can kill nine out of ten people it infects. Although this is not really an infection, it's *transfection*

(human cloning).

Ebola and Marburg ***cause hemorrhagic fevers*** (VHF). They simultaneously affect multiple organ systems in the body and may be accompanied by hemorrhage, or bleeding. These pathogens cause high fever, chills, muscle aches, and vomiting. The patients worsen rapidly until they bleed from every orifice in their body, including needle puncture wounds. They usually die within 1-3 days.

Bill and Melinda Gates say on their Gavi website to ***expect the next pandemic*** to be a Marburg outbreak. These two psychics or *psychos* also claim that Ebola and Marburg are carried by African Green Monkeys.

The***lying CDC claims***that people can get “Ebola Virus disease” through direct contact with an infected animal (bat or nonhuman primate). They’re taking the absolute piss out of us! Never in the history of medicine has an animal disease infected Humans! The only possible way to make a Human diseased with an animal disease is through cross-species genomics using Adenvirus or mRNA and Nanotechnology.

Polio vaccines used in the late 1950s and early 1960s were intentionally contaminated with a *bacterial* pathogen called the “Simian Virus” 40 (SV40) present in monkey kidney cells. The Simian Virus is used for *infecting Humans*. It was ***“accidentally administered to Humans”*** through Polio vaxxines.

The “Vaccinia virus” is similar to the “smallpox virus” but it’s not naturally occurring after all. ***Scientific Direct reported*** that “vaccinee-to-cattle and cattle-to-human Transmissions occurred on Farms”, proving the transmission of pathogens between animal to human species is being done by genetic engineering and administered through vaxxines.

Knocking out the E1 and E3 genes is necessary when ***transfecting cross-species*** in order to make the pathogen replicate. The Vaccinia pathogen has been used on a wide scale to produce many different kinds of chimeric proteins, including HIV-1 and it encodes approximately 250 genes.

Ebola and Marburg are GAIN-and-Loss-of-Function bioweapons and both are created using the “Green Monkey disease“, a chimeric pathogen that you can ***purchase online****!*

Adenovirus’ are from human/monkey clone origin and are used with chimeric Lentiviruses, as well as the Filoviruses. Of course, none of these are actual viruses! All the Adenoviruses and messenger RNA (mRNA) are *chimeric weapons* used to code your cells to reproduce deadly proteins used to both silence genes and program artificial genetic sequences. They are inducing diseases and making up disease names and syndromes to hide the fact that it’s coming from vaxxines!

Ebola, Marburg, and Monkeypox are ***Gain-of-Function bioweapons*** created using the “Green Monkey disease”. It’s an Adenovirus made from E. coli bacteria from the decaying flesh of a human/monkey hybrid’s kidney tissue culture. Adenovirus vectors transfect Humans with monkey DNA, Ebola, and HIV. Adenovirus vectors are ***transfecting humans*** with monkey DNA and Human DNA from a Chimpanzee/Human clone to be exact.

There are 49 immunologically distinct types of adenovirus that can cause infection for long-term gene expression. They’re made with ***Sialic acid*** which is a group of derivatives of Neuraminic Acid found in *animal tissues*

. Sialic acid is the primary entry receptor used in Adenovirus.

Sialic Acid is an animal DNA from “Species D” Adenovirus which is used in both Adenovirus 5 and Adenovirus 26 (Sinovac and J&J) to transfect Ebola and HIV into Human cells using Sialic acid-bearing glycans (animal DNA) as a primary cell entry receptor. Adenovirus 5 is of***“Chimpanzee origin”***. There’s that Green Monkey clone again!

Sialic acid ***proliferates tumor growth*** and metastases. N-acetylneuraminic acid (Neu5Ac) is made from E. coli bacteria. See the study’s ***here*** and ***here***.

Polio vaxxines used in the late 1950s and early 1960s were ***“contaminated with a virus”*** or rather a bacterial pathogen called the Simian Virus 40 (SV40) which is present in *monkey kidney cells* used to grow the vaxxine.

The Pharma death cult is inducing diseases with their vaxxine racket and making up names and syndromes that they can then profit from by “treating” later when people become diseased. What would life be like without the Pharma cartel disabling our children and killing healthy people in the name of science and medicine?

There are no viruses involved in the making of any of the mRNA or Adenovirus vaxxines, only GAIN- and Loss-of-Function chimeric pathogens made from bacteria and other plasmids which Pharma keeps naming “viruses”.

There’s no “spike” from a virus particle being used in any of these vaxxine induced diseases. We should change our language from “spike protein” which is a half-truth with a half-lie and replace it with *“chimeric protein”* to be medically accurate because that’s what we’re dealing with.

E. coli bacteria are used as the base for all these chimeric diseases because bacteria DNA *replicates*. They’re also using other plasmids and mixing fungus, yeasts, and “several mammal-based systems” (Human/Chimpanzee clone), then genetically enhancing them to increase lethality.

They’re also using ***baculovirus-mediated insect cell expression***. This means the Pharma cult is transfecting human cells with insect DNA. This could explain the strange mutations and Morgellons.

Marburg is simply Ebola with ***Ricin*** added to make it more lethal. Both cause hemorrhagic fevers (VHF) and attack multiple organ systems in the body, accompanied by bleeding.

The Pharma death cartel and the Eugenicists already have a **PCR kit** for “testing” for Marburg disease and a vaxxine to immunize against, called RiVax. The main component of RiVax is “a genetically altered version of a Ricin Toxin.” Ricin is more toxic than Graphene Oxide, by the way.

I think it’s high time people stop trusting our governments, stop relying on government and take our children out of public schools, as Dr. Zev Zelenko said to Alex Jones on Info Wars. Don’t sacrifice your kids to Satan.

My detox protocol works for the vaxxed and the unvaxxed to kill the Micro-plasmids (parasites, transgenic Hydra’s and bacteria) and reverse the coagulation cascade which leads to blood clots.

By Dr. Ariyana Love

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