Transgenic Hydras & Parasites A Biological Weapons System For Rapid Human Cloning

By Dr. Ariyana Love

The transhumanist dystopian nightmare we find ourselves in is taking a new turn with the shocking discovery of *Hydra Vulgaris* and PARASITES in the so-called Covid-19 "vaccines".

Dr. Carrie Madej revealed her <u>Hydra findings</u> on the Stew Peter's Show on September 29th, 2021, followed by Dr. Zandre Botha's <u>stunning discovery</u> of microscopic, self-assembling medical devices in the blood of her vaxxed patients. The red blood cells are dangerously deformed and coagulated, things she says she's never seen before in her 15 years as a blood doctor.



Hydra Vulgaris identified in Pfizer & Moderna Covid-19 serums

About 10 days later, "That Thing" (Hydra Vulgaris) was <u>also identified</u> in Pfizer vials by Dr. Franc Zalewski. He took the science to a new level and did a chemical analysis of the Hydra, exposing that the chemical compound of the creature contains Aluminum, Carbon, and Bromium. This means the Hydra's are being genetically modified before they're injected into humans. The good doctor also identified *parasites* in the vials.

Dr. Jane Ruby, a pharmaceutical researcher, gave vital <u>commentary on Stew Peter's Show</u> about Dr. Zalewski's findings, emphasizing that the dormant Hydra "eggs" become active, grow and *multiply* when exposed to Graphite tape and *heat*.

Earlier in August, *parasites* and other horrors were <u>identified by Dr. Robert Young</u> in four Covid-19 vials. Dr. Jane Ruby again <u>joined Stew Peters</u> to give crucial commentary on Dr. Young's findings.

Investigative Journalist Ramola D. provided us with <u>further information</u> about the parasites discovered by Dr. Young and did an <u>expose</u> in October.

Back in July, La Quinta Columna <u>studied four "vaccines"</u>; Pfizer, Moderna, AstraZeneca, and Johnson & Johnson, and found toxic nano metallic particulates, particularly nano-graphene oxide, in significant amounts, as well as lipid nanoparticles and the parasite *Trypanosoma cruzii*, in the Pfizer-BioNTech serum.

Pfizer whistleblower Karen Kingston appeared on Stew Peter's Show in July and walked us through a presentation on how Graphene Oxide is in *all* Covid-19 serums. Graphene Oxide was not listed in the <u>patent filings</u> and was deliberately concealed under a trade secrete because it's known to be poisonous to humans.

As a result of these horrifying discoveries, I did my own research on Graphene Oxide Nanoparticles and Toxicity and wrote an article entitled "Graphene Oxide The Vector For Covid-19 Democide". I reveal how humanity is being saturated with Graphene Oxide Nanoparticles in a myriad of ways.

I also wrote an article on protocols for detoxifying Graphene Oxide from your body, **here**.

The openly <u>declared ingredients</u> in Covid-19 serums should be enough to dissuade anybody from taking them. Now it's clear there are additional poisonous and other horrors not being disclosed to the public by the Biotech pharma industry.

Karen Kingston has backed up all these terrifying discoveries with the patent filings and receipts, on Stew Peter's Show. Kingston explains that the vaxxines are a "gateway to an obedience platform and potentially an execution platform if you are not obedient to your score".

Informed Consent has been *waived* and therefore people didn't know they're being injected with smart devices and bioweapons. The patents also reveal that it was already *known* by **Pfizer** that the vaxxed would become "<u>super spreaders</u>" and transmit deadly pathogens to healthy individuals.

There's an Al component to these vaxxines Kingston explains, "they're committed to replacing the American people with Artificial Intelligence". She continues disclosing that "Hong Kong is ready to replace the American people with robots right now".

Due to the fact that patent filings do not reveal the components to Biotech's vaxxine ingredients, I began researching scientific peer-reviewed studies involving Hydra Vulgaris and parasites to see if I could identify why they're being injected into humans.

GAIN-OF-FUNCTION

Everything I'm writing here is based on evidence from open-source, peer-reviewed literature of scientific breakthroughs and technological developments that extend through the past *decades* and are linked in this article. As sci-fi thrilling as this information may sound, the technology has already been deployed and is being injected into the veins of our children as we speak. You can read the studies for yourself, as I have done.

DNA hybridization began in 1980 with Nadrian C. Seeman who started constructing self-assembled nanostructures. *Hydra Vulgaris* transgenesis technology was developed over the last 30 years. This is the process of transferring genes and organisms from one species to another which creates a new cloned species.

The Human Genome Project began in the year 2,000. Hydra's are used in the human genome assembly for *gene silencing* of humans. Messenger RNA (mRNA), SPIONS (Super Paramagnetic Iron Oxide Nanoparticles), DNA coated lipid-nanoparticles containing drugs and chemicals, transgenic Hydra's and parasites are all part of an "operating system" which is bypassing the human immune system. You can read more about Moderna's "operating system" from their own website, here.

Graphene Oxide sheets are used to <u>slice open the membrane</u> of your cells so that *programmable* Nanorobots can reach the cell nuclei to *turn off* undesired genes (gene silencing) and code artificial gene sequences. This process is called biohacking.

Graphene Oxide sheets are able to slice open every cell membrane of the human body within 15 minutes after inoculation, according to Dr. Robert Martin.

DARPA partly funded the development of protein-to-genomic sequence alignments for cross-species genomics. Gain-Of-Function and Loss-Of-Function studies using transgenic Hydra were funded by Fauci and the NIH and developed at the Wuhan Institute in China and in universities in the U.S. and China. Their scientific findings were published in 2013.

The Sixth International <u>Workshop on DNA Nanotechnology</u> was held August 26–28, in 2017, in Beijing, China where the forum showcased the applications of self-assembled DNA nanostructures.

CHIMERIC SPIKE PROTEIN

The "spike protein" in the Covid-19 vaxxines that everyone is talking about is called a <u>Lentivirus</u>. The Lentivirus contains a combination of **HIV types 1-3, SRV-1/AIDS, MERS,** and **SARS**. These are the *most* deadly Gain-Of-Function bioweapons ever developed, thanks to mass-murdering *Fauci*.

A <u>Stanford study</u> reveals that the Lentivirus is a "genus of retroviruses that cause chronic and deadly diseases characterized by long incubation periods, in humans". It enables long-term transgene expression. The best-known Lentivirus is the human immunodeficiency pathogen, which causes AIDS. This is why we're seeing an autoimmune and neurodegenerative decline after Covid-19 inoculation. This is an induced condition known as PRION.

The mRNA from the Lentivirus chimeric cocktail is inserted into the DNA of human cells through an invasive procedure that *permanently* changes the genome of that cell. Once inside the host cell's cytoplasm, lipid-coated nanobots take the reverse transcriptase enzyme in the Lentivirus to produce DNA from the mRNA genome, the reverse of the usual pattern, thus *retro*.

HYDRA 2.0 GENOME ASSEMBLY

Hydra's are used in <u>cross-species genomics</u>. They're being genetically modified in a lab at the University of Kiev to produce <u>transgenic clonal Hydra lines</u>. Since 2006, thousands of embryos have been microinjected and nearly 200 transgenic lines have been established in the <u>Hydra Transgenic Facility</u>.

Morphogenesis and stem-cell control using the Hydras were developed to learn the neurobiological functions of humans and for <u>in vivo tracing of cells</u>. Transgenic Hydra allows <u>in vivo tracking</u> of individual stem cells during morphogenesis (tissue and cell growth).

Transgenic Hydra lines are generated by embryo microinjection with plasmid DNA from self-replicating DNA found in *bacteria*. This is a permanent transmissible change of genetic material (DNA) resulting in the decreased production of a protein. The merging of the two species is a "cloning" process called <u>transfection</u>. A new generation of transgenic Hydra polyps continues reproducing the chimeric genetic expression in their offspring.

These GMO Hydra polyps are now genetically *coded vectors*, carrying a variety of programmed synthetic genomic sequences and mRNA (messenger RNA) for the purpose of <u>transfecting humans</u>. Once inside the human body, these transgenic Hydra polyps serve to *rewire* and *control* the ancestral circuitry of human beings.

BLAST Sequence <u>technology is being used</u> to create new DNA sequences and find similar genetic sequences between species, performing alignment functions for same-species and *cross-species genetic splicing* for the purpose of transcription.

Proteins regulate gene expression. This technology targets the cell organelles of the nuclei which store genetic information; mitochondria, which produce chemical energy; and ribosomes, which assemble proteins, using mRNA to make mitochondrial sequences.

A 2017 <u>Gain-Of-Function research project</u> from Germany, demonstrates how RNA extraction and quantitative reverse transcription-polymerase chain reaction or *reverse genetics* is used to knockout and knockdown genes using Hydra's and CRISPR/Cas9.

The genetically modified Hydra lines in the Covid-19 operating system is first <u>coded</u> <u>with chimeric gene sequences</u> (Lentivirus) which is then being <u>coded</u> into human cells using CRISPR-Cas9 technology and <u>electroporation</u>.

Electrodes attached to **gold programmable nanorobots** transfect human cells, *silencing* your innate God-given genetic sequences and coding your cells to reproduce the synthetic genetic sequence of the chimeric spike protein (Lentivirus), *indefinitely*. More simply stated, your cells will continue to replicate themselves over and over again with the new genetic sequence of the *chimeric pathogen* you were injected with. The same chimeric pathogen was funded by bioterrorist Anthony Fauci and developed in Wuhan, China.

One of the deadly <u>bacteria being chimerically enhanced</u> to transfect Hydra's for <u>implantation into humans</u> is <u>E. coli</u>, which causes about 36% of the infections in humans.

PARASITES

Parasites are also <u>transfected with bacteria</u> and used as <u>transfection vectors</u> for *DNA binding* and genetic sequencing in humans. Parasites can evade drugs, escape the immune system and regulate genes.

The human <u>Malaria Genome Project</u> developed at Stanford University, used CRISPR technology and bacterial plasmids which can <u>replicate rapidly inside parasites</u>. They transfected bacterial plasmids into parasites, disrupting a series of gene encoding molecules. In that study, scientists transfected Malaria parasites with Luciferase to use it for gene targeting and transgene expression in *humans*.

T. gondii and *P. falciparum* and other parasites were also used in transfection studies. It's important to be aware that from the P. falciparum they designed a <u>Chloroquine-resistant</u> transgenic parasite strain called Dd2.

LUCIFERASE

Hydra polyps are also being coded with the <u>overexpressed chimeric protein</u> called Luciferase, which is a Green Florescent Protein derived from the firefly. Transgenic Hydra also carries the Luciferase RNA trigger to code your cells with and silence genes in human cells.

Holstein lab investigated the repressing activity of HySp5 on the *HyWnt3* promoter, performing Luciferase reporter assays in human HEK293T cells for *DNA-binding* and <u>transplanting Hydra into humans</u> by invading human tissues.

The transgenic Hydra and parasites *replicate* and merge with humans during transfection. They are integrated with the transgenes (Luciferase and Lentivirus) into one of the epithelial cell lineages and <u>assimilated into the human host</u>. The transplanting of Hydra's into humans is called <u>Homoplastic transplantation</u> using induced Hydranth as "implants".

Epithelial cells are stem cell lineages responsible for cell signaling. Transgenic Hydra's reporter genes are cell-signaling with each other inside humans, much like *neurons* in a neural network. Transgenic Hydra's cell signaling becomes synthesized with human cell signaling in a process called <u>catenin signaling</u>, which is induced by mutations of genes in humans through upregulation (cell response) to the plasmids expressing activators in the Hydra (HySp5–2992:Luc); aka *transfection*.

Transgenic Hydra and parasites induce humans to generate a *new* electrochemical signal by organizing enzymes spatially to create a *programmable redox enzymatic cascade pathway*, changing the predictable generation of electrochemical signals in humans. The newly established synthetic gene sequences are now *shared* between the transgenic Hydra's, parasites, and newly hybridized humans.

In fact, Biotech's chimeric operating system establishes a <u>new neural network</u> in humans and an <u>artificial brain</u> by re-directing endogenous neural stem cells. Brain <u>implants can erase memories</u> and implant new, *artificial* memories while Graphene Oxide can <u>"hear your brain whisper"</u>.

A <u>team of scientists</u> from UC Davis and Rice University was boasting back in July about manipulating the nervous system of Hydra Vulgaris and humans to "build a new brain from the bottom up", in order to control neural pathways and human behavior. This technology was developed over the last decade through the **Human Brain Project**.

The European Union's 1.5 billion euro <u>Graphene Flagship</u> project developed <u>graphene-based implants</u> for <u>"future brain-computer interfaces"</u>. I have to wonder if the "implants" they're referring to contain transgenic Hydra's?

Graphene implants can record electrical activity in the brain at extremely low frequencies and over large areas, "unlocking the wealth of information found below 0.1 Hz".

A Russian <u>initiative called 2045</u> wants to use neural interfaces for an "improvement of man himself" because mankind is "standing at the edge of a total loss of the conceptual guidelines necessary for further evolution". This demonstrates the anti-human mindset of eugenicists who want to clone the entire human race.

The fluorescent (Luciferase) Hydra's were also tested with externally applied electrical fields to see how much voltage they could endure, to "facilitate the future use of electric fields as an experimental means to redistribute intracellular constituents in developing tissues". I presume this was to test Hydra's ability to survive 5G frequency?

THE OPERATING SYSTEM

Hydra's and parasites also serve as a *reporter system*. Luciferase exhibits bright green fluorescence when exposed to light in the blue to the ultraviolet range, enabling the vaxxed to be traced externally. Genes of interest can be turned off occasionally or turned on at will by your patent holders through what's called transregulation.

This means you're not only externally traced 24/7 but you'll also be externally controlled. Your patent holders will be able to upregulate and downregulate your genetic codes through an external database, through the **Eukaryotic Genome Annotation Pipeline** for transgenic humans.

<u>Did you think the Starlink satellite network's "Precision Tracking Space System" had something to do with defense?</u> Don't worry, you'll be "happy" owning nothing so long as they get your dopamine levels worked out.

ADDGene is selling a variety of <u>CRISPR parasites</u> to be used as <u>gene vectors</u> for human transfection. These are not "vaccines" at all but a <u>WEAPONS SYSTEM</u> (*my words*) for the **RAPID CLONING** (*their words*) of humans, through gene knockout (silencing), artificial gene sequencing (coding) and to monitor transfectants inside of humans (tracing).

<u>ProSplign</u> is a worldwide protein-to-genome alignment tool enabling Human DNA to be easily synthesized from a single-stranded RNA template and catalyzed by an enzyme for *reverse transcriptase*.

ADDGene also offers a **Lentiviral Envelope** and Packaging Plasmids for transfecting humans using transgenic Hydra. They offer "Non-overlapping **NEURAL NETWORKS**" (*their words*) using Hydra Vulgaris for building a <u>new neural network</u> in Hydra's. This technology is being deployed in humans through the Coivd-19 Quackccine program now.

Dr. Carrie Madej also disclosed in her <u>latest interview on Stew Peter's Show</u> that the vaxxine operating system is building an artificial neural network in humans.

ADDGene offers a <u>Tetracycline off system</u> for on/off gene expression, "fusing tetR with the C-terminal domain of VP16 (virion protein 16), an essential transcriptional activation domain from **HSV** (*herpes simplex virus*) which is being used for "reduced gene expression" in humans. This uses the chimeric E. coli bacteria and Lentivirus.

The <u>Hydra genome assembly</u> offers a Nano DNA kit called <u>Illumina</u>. Illumina Inc figured out how to <u>reduce the cost of sequencing</u> a human genome down from \$1 million to \$1,000 USD, back in 2007.

After Luciferase is infused and coded for targeted genes via a computer, it's then mapped onto the Human through the public *Galaxy* server to perform "differential expression analysis". Proteins can be targeted, upregulated, and downregulated.

Then there's <u>Vector Biolabs</u> whose selling Adenovirus' for human sp5 shRNA silencing. A Knockout vector system (adenovirus) for knocking down the expression of particular genes (gene silencing), is being marketed online and sold by <u>Vector Builder</u>. You can create artificial genome sequences and merge genomes of different species. [The God gene VMAT2 can be silenced this way]

You can preorder DNA sequences for humans on **HydraAtlas website**.

The <u>Genome Data Viewer (GDV)</u> will help you select genome assemblies (DNA sequences) for humans from primarily finished human clones, that were sequenced as part of the Human Genome Project.

VIGENE offers multiple <u>shRNA cloning options</u> for your gene silencing experiments. They're packaging transfer Plasmids, Adenovirus' (AAV) and Lentivirus' and they guarantee at least a 70% knockdown of your gene of interest. They have a catalog of over <u>27,000 shRNA plasmid sets</u> targeting the human genome.

This <u>table lists common</u> Lentiviral envelope and packaging plasmids that can be used with 2nd and 3rd generation lentivirus technologies.

ADDGene's <u>lentiviral genome</u> is delivered to a target cell upon infection using CRISPR gRNA. They explain how the Lentiviral genome encodes genetic material that the "researcher" (*or patent holders and Big Pharma*) wants to be delivered to specific target cells. The genome is encoded by plasmids called "transfer plasmids," which can be modified to encode a wide range of gene products.

ADDGene admits their DNA-targeting enzymes very often will *delete*, *insert* or otherwise *alter* the targeted RNA or DNA, so don't let the fake media fool you.

Lentiviral Plasmids can be ordered through ADDGene *here*.

BEHAVIOR CONTROL

Vector Biolabs offers an <u>Adenovirus (AAV) expressing shRNA</u> for the knockout (gene silencing) of Human SP5. When developing this technology during the <u>animal trials</u>, social recognition, spatial learning, and memory were impaired after 4 weeks.

In an <u>animal study</u> using reverse transcriptase-polymerase chain reaction (RT-PCR) with an Adenovirus vector and drugs, scientists were able to induce Huntington's Disease by targeting the Corpus striatum of the brain which resulted in 100 fold neurodegeneration and motor behavioral impairment.

REPRODUCTION & FERTILITY

The transgenic Hydra's are used to induce gene silencing predominantly targeting *embryonic* cells in the testes of men and the ovaries of women and also *nerve* cells. This is why we're seeing neurological degeneration (PRION) after inoculation. It also

explains why 82% of expectant mothers who take the "jab" are having **spontaneous abortions**.

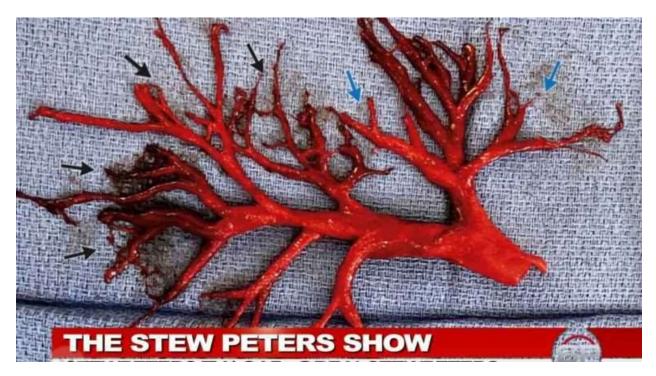
<u>Microinjection of foreign DNA</u> into the pro-nucleus of single-cell *embryos* of fertilized mice to control the genetic expression of future generations has been perfected, since 2008.

Proteins control gene expression. Transgenic <u>Hydra is instrumental</u> in encoding the human SP5 (shRNA <u>silencing AAV</u>) which is a gene on chromosome 2q31.1 that encodes a protein that binds to the GC-box promoter elements, thought to play a role in coordinating the intricate changes in transcription which occur in the developing *embryo*.

Wnt-3 is a protein that in humans is encoded by the WNT3 gene. These proteins have been implicated in oncogenesis and in several developmental processes, including regulation of cell fate and patterning during *embryogenesis*.

The point I'm making here is that the operating system is DNA-binding, downregulation, and upregulating genes using the transgenic Hydra's, targeting human *embryos* and *embryonic cells*, leading to developmental alterations from binding genes to the Wnt/β-catenin signaling pathway.

<u>Do you understand what this means?</u> It's not only the vaxxed who are being genetically modified, cloned, and hybridized, but SO ARE THEIR OFFSPRING! That is of course if you're still able to reproduce at all after the jab! Most people will just be *sterilized* and their babies aborted. This is a human cloning experiment as well as *extermination*.



Microinjection of Retrovirus transgenes (Lentivirus & Luciferase) <u>integrates randomly</u> into the genome which poses enormous risks for the vaxxed as well as their hybrid offspring. This can create strange and unpredictable mutations of DNA by the addition of one or more base pairs. This is precisely why we're seeing freaky mutations and why doctors are removing blood clots with Hydra-like tentacles from teenagers! Picture above on page 9.

Dr. Carrie Madej shares an image of a blood clot with Hydra-like mutational growth that was removed from the heart of an early teen who received a Covid vaxx.

If you still aren't convinced, please listen to Dr. Peter McCullough explain this biotechnology and how the chimeric spike proteins are being *coded into human cells*, at the <u>78th Annual Meeting of Association of American Physicians and Surgeons</u>, on October 2, 2021.

"It is a deadly protein" explains Dr. McCullough. "It is the first time in medicine that we are injecting vaccines and asking the human body to make a potentially lethal protein".

While the Covid-19 serums appear on the surface to be only a clear liquid, under microscopy you can visibly see all the many components of the computer-interface operating system, which is a sophisticated biological weapons system for the cloning and extermination of the human race.

FINAL NOTE

If you would like more information on detox protocols and disrupting the blood coagulation cascade which leads to blood clots from the jab or for protocols that will protect you from the adverse effects of transmission, please contact me directly at: metanutrients@mailfence.com

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If Dr. Ariyana Love's article does not concern you, then what I am going to share along with this report is further evidence that they want to harm or kill us, but an entirely different level. This report comes from the Liberty Beacon about Secret Military Experiments & Our Health. It goes without saying that I have been convinced "Depopulation is the World's #1 Issue" for half a century. The article and the interview that came with it between Mike Adams and Dr. Lee Merritt indicate there is a collective conspiracy world-wide if not nationally to kill us.

When Bill Gates tells you to prepare for a 'smallpox bioterror attack', you should believe him. Bill Gates is also now calling for the microneedle vaccine patch.

Bill Gates funded and hosted the dystopian Event 201 where he and his buddies imagined what a COVID global outbreak would look like, and amazingly, just a months

later a global COVID outbreak took place, killing millions. Now Bill Gates has warned governments to prepare for smallpox terror attacks and future pandemics by investing billions into research and development. Bill Gates is telling you the truth and you should believe him.

Secret Military Experiments & Our Health

November 5, 2021 This article was posted by TLB Staff



Secret Military Experiments and Our Health Analysis by Tessa Lena

This story is about secret experiments on people. It is also about the reality of living inside a Machine that plays games with us. Sometimes the Machine selectively shows its face — and sometimes it selectively hides it — and based on that, it plays its favorite game called, "Confuse, Divide and Conquer."

The story was specifically inspired by the fact that as I am typing this, a biosecurity experiment is taking place in my home town of New York. All safe, allegedly ...

Of course it's safe. It's always safe when the television says so! We in the West are used to being mostly spared by the Machine, and so we think that our bureaucracies are mostly under control. We believe that yes, there is corruption — and yes, there are bad apples here and there — but they cannot go very far because we have a Democracy. A Machine that is working for us. A working Machine.

And while our levels of trust do vary from person to person and from community to community — based on the fact that some people get to see the Machine more up close than others — overall, up until last year, we've been living in relative comfort, taught to generally trust our bureaucracies like sweet puppies generally trust their owners with their open-for-petting bellies (including those owners who, on a side note, sterilize their

pets at the beginning of their life journeys and then potentially euthanize the mat the end of it, in order to prevent greater suffering; such is our culture).

Relationships of love inside the Machine can be puzzling at times. And so, we've been living in relative comfort and relative trust of our bureaucracies, and feeling okay. And then last spring, we were asked to lockdown for two weeks to flatten the curve — and here we are.



1:50 AM - Oct 7, 2021 - SnapStream TV Search

The blood-chilling dehumanization is in vogue. It's 2021, and we are marching straight into the <u>Great Reset</u>. Everything is "fine."

And what about the Machine? The Machine is showing its face to most of us at last. It demands that we stop being human, stop touching each other, and bury our faces and our children's faces under The Mask, "to be good citizens." It demands that we don't think and comply. It proclaims that we have no bodily freedom and need to obey.

But another thing that is happening is that we the people are becoming awake. Many people are becoming "mechanics," looking under the hood — and discovering things we have not thought about before.

For example, how many of us just recently found out that during the Prohibition era, the U.S. government purposely <u>poisoned alcohol</u> to ween the unruly Americans off the forbidden habit?

That particular story went like this: At the time, there was a black market for re-distilled industrial alcohol — and, unhappy with the state of affairs, federal officials "ordered manufacturers to make their products far more deadly."

According to 'Slate', "In 1926, President Calvin Coolidge's government decided to turn to chemistry as an enforcement tool ... By mid-1927, the new denaturing formulas included some notable poisons ... The Treasury Department also demanded more methyl alcohol be added — up to 10 percent of total product. It was the last that proved most deadly."

"In 1926, in New York City, 1,200 were sickened by poisonous alcohol; 400 died. The following year, deaths climbed to 700. These numbers were repeated in cities around the country." (In 1926, the population of the United States was about 117 million people.) "Officially, the special denaturing program ended only once the 18th Amendment was repealed in December 1933."

And how many of us remember that in 1931, Dr. Cornelius Rhoads, a 1945 recipient of Legion of Merit, infected people with cancer cells while working with the Rockefeller Institute for Medical Investigations in Puerto Rico, "under the auspices of studying hookworm-caused anemia and tropical sprue anemia"?

How many of us know that fourteen years after the Tuskegee experiment started, the U.S. also <u>conducted syphilis experiments in Guatemala</u>, purposely infecting Guatemalan men and women with syphilis?

How many of us know that during the Cold War, <u>MIT researchers</u> and the Quaker Oats Company fed the mentally disabled children at the Walter E. Fernald School in Waltham, MA, doses of radiation in their cereal?

How many of us know that in 1950, in San Francisco, the Navy intentionally hosed a "<u>cloud of microbes</u> into the air and into the city's famous fog"? The microbe was Serratia marcescens. According to <u>Discover</u>, over a period of six days, "<u>members of the US Navy sprayed clouds of Serratia from giant hoses aboard a Navy minesweeper drifting two miles along the San Francisco coastline."</u>

As a result, "a week after the spraying, eleven patients were admitted to the now defunct Stanford University Hospital in San Francisco with severe urinary tract infections, resistant to the limited antibiotics available in that era. One gentlemen, recovering from prostate surgery, developed complications of heart infection as Serratia colonized his heart valves." Unfortunately, he passed away.

According to WJS, in 1950s, "Army researchers dispersed Serratia on Panama City, Fla., and Key West, Fla., with no known illnesses resulting. They also released fluorescent compounds over Minnesota and other Midwestern states to see how far they would spread in the atmosphere. The particles of zinc-cadmium-sulfide — now a known cancer-causing agent — were detected more than 1,000 miles away in New York state."

Another bacterium, Bacillus globigii, "was released in San Francisco, while still others were tested on unwitting residents in New York, Washington, D.C., and along the Pennsylvania Turnpike, among other places, according to Army reports released during the 1977 hearings."

How many of us know that in 1966, in New York, military researchers dispersed Bacillus subtilis variant Niger, believed to be harmless at the time and currently considered to be a pathogen, in the subway system, by dropping lightbulbs filled with the bacteria onto tracks in stations in midtown Manhattan?

WSJ <u>wrote</u> that the bacteria were carried for miles throughout the subway system. In a 1968 report, Army officials concluded: "Similar covert attacks with a pathogenic [disease-causing] agent during peak traffic periods could be expected to expose large numbers of people to infection and subsequent illness or death."

Also notable is the reaction from the public to being fumed over. At the time, in a <u>report</u> titled "A Study of the Vulnerability of Subway Passengers in New York City to Covert Attack with Biological Agents," the team noted: "When the cloud engulfed people, they brushed their clothing, looked up at the grating and walked on. People in the big city are moving too fast to see what is going on around them." Sounds familiar?

To further quote <u>WJS</u>, "army officials also found widespread dispersal of bacteria in a May 1965 secret release of Bacillus globigii at Washington's National Airport and its Greyhound bus terminal, according to military reports released a few years after the Senate hearings. More than 130 passengers who had been exposed to the bacteria traveling to 39 cities in seven states in the two weeks following the mock attack."

The Army kept the biological-warfare tests secret until word of them was leaked to the press in the 1970s. Between 1949 and 1969, when President Nixon ordered the Pentagon's biological weapons destroyed, open-air tests of biological agents were conducted 239 times, according to the Army's testimony in 1977 before the Senate's subcommittee on health.

In 80 of those experiments, the Army said it used live bacteria that its researchers at the time thought were harmless, such as the Serratia that was showered on San Francisco. In the others, it used inert chemicals to simulate bacteria."

In 1975, the 'New York Times' published an <u>article</u> detailing various military experiments that took place in 1960s. In particular, it talked about the 1966 subway experiment in New York and various other "exotic" experiments and weapons.

The 'New York Times' story focused on the testimony by Charles Senseney, "a project engineer who developed weapons such as an electric poison dart gun and a system to spread biological poison from a fluorescent bulb."

Charles Senseney told the Senate Select Committee on Intelligence that he "took part in the New York 'vulnerability study' as one of many such efforts aimed at testing the dangers of biological warfare ... Mr. Senseney said the studies [were] conducted by the staff of the Army laboratories at Fort Detrick, Md ... They included tests at the White House, the Pentagon, a Food and Drug Administration building in Washington and McGuire Air Force Base in New Jersey."

Additionally, Mr. Senseney confirmed that "the Fort Detrick labs had secretly placed a colored dye into the water system of a Washington building used by the Food and Drug Administration to test how fast the occupants could be killed or incapacitated by introducing a biological agent into the water system. Mr. Senseney said he had developed a special drill that allowed the dye to be punched into a water pipe without leakage or change in the water pressure."

A question: How many Americans at the time had any idea that all this was happening? I would bet that even today, this might come as a shock to many. Stay safe?

And how many of us are aware that from 1963 to 1973, under <u>Project SHAD</u>, numerous sets of tests were performed, "many involving multiple releases of biological or chemical agents in locations as far-ranging as the Deseret Test Center (80 miles from Dugway), Hawaii, the Panama Canal Zone, and off the coast of San Diego?"

Among the substances and organisms released were F. tularensis, Coxiella burnetti, Staphylococcal enterotoxin B), as well as the nerve agents sarin, tabun, soman, and VX. "One or more of these potential biological or chemical weapons were used in 26 SHAD tests."

"Biological simulants were also released: B. globigii in 24 tests, S. marcescens in 7 tests, and E. coli in 5 tests." This list of experiments is incomplete and doesn't even touch on Fauci or Daszac — but even an incomplete list should make us stop and ponder why we are living in a world like this.

Speaking of experiments, here is a thought-provoking interview with <u>Dr. Lee Merritt</u> in which she and Mike Adams discuss the possibility of spikes having been scattered over target cities. I could not locate the URL for that interview but I believe both Mike Adams and Dr. Lee Merritt have it posted on their sites.

And then my mind comes back to the experiment happening right now in the open in my city, where they are <u>releasing "harmless substances" on the New York subway and in various populous locations</u>. Among those substances are DNATrax, amorphous silica, sulfur hexafluoride, and perfluorocarboms.

Will we hear in ten or twenty or fifty years that it was a really bad idea, and that whatever they thought was harmless (wink, wink) was toxic? If history is any proof, we probably will. That is, if Mother Nature saves us again and we survive mad science and mad feudal owners for the millionth time.

If we do, I hope, we will wake up to our senses and stop accepting abuse. Just stop accepting abuse. Abuse is not cool.

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Blessings, Jesus is at the Door!

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