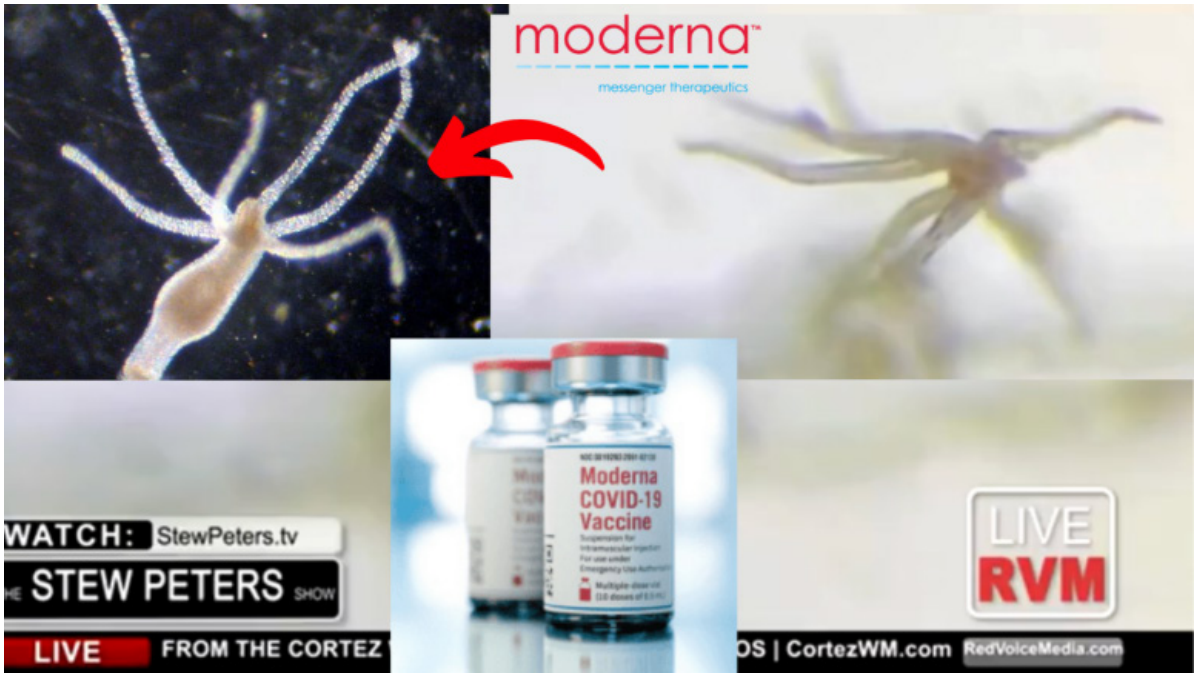
November 1, 2021 by Dr. Ariyana Love

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 **Hydra findings** on the Stew Peter's Show on September 29th, 2021, followed by Dr Zandre Botha's **stunning discovery** 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 **also identified** 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 Aluminium, Carbon, and Bromium. This means the Hydras 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 commentary (1) on Stew Peter's Show about Dr Zalewski's findings, emphasizing that the dormant Hydra "eggs" become active, grow and *multiply* when exposed to Graphite tape and *heat*.

(1) Jab: Scientist Discovers Hatching Eggs, Parasites Birthed After Injection

Earlier in August, *parasites* and other horrors were **identified by Dr Robert Young** in four Covid-19 vials.

Scanning & Transmission Electron Microscopy Reveals Graphene Oxide in CoV-19 Vaccines

Updated: Oct 19

2021, February 5th, Updated October 1st, 2021

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15th Revision

Dr Jane Ruby again **joined Stew Peters** to give crucial commentary on Dr Young's findings.

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

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

Pfizer whistle-blower 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 **patent filings** and was deliberately concealed [as a trade secret] 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 **declared ingredients** 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 vaccines 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 "***super spreaders***" and transmit deadly pathogens to healthy individuals.

There's an AI component to these vaccines 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 vaccine 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. Hydras 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 Hydras 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 **slice open the membrane** 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 **Workshop on DNA Nanotechnology** 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 vaccines that everyone is talking about is called a Lentivirus**.** 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 **Stanford study** 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

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

Morphogenesis and stem-cell control using the Hydras were developed to learn the neurobiological functions of humans and for **in vivo tracing of cells**. Transgenic Hydra allows **in vivo tracking** 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 transfection. 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 ***transfecting humans***. Once inside the human body, these transgenic Hydra polyps serve to *rewire* and *control* the ancestral circuitry of human beings.

BLAST Sequence **technology is being used** 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 **Gain-Of-Function research project** from Germany, demonstrates how RNA extraction and quantitative reverse transcription-polymerase chain reaction or *reverse genetics* is used to knockout and knockdown genes using Hydras and CRISPR/Cas9.

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

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 **bacteria being chimerically enhanced** to transfect Hydras for **implantation into humans** is **E. coli**, which causes about 36% of the infections in humans.

PARASITES

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

The human **Malaria Genome Project** developed at Stanford University, used CRISPR technology and bacterial plasmids which can **replicate rapidly inside parasites**. 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 Chloroquine-*resistant* transgenic parasite strain called Dd2.

LUCIFERASE

Hydra polyps are also being coded with the **overexpressed *chimeric protein*** 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 **transplanting Hydra into humans** 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 ***assimilated into the human host***. The transplanting of Hydras into humans is called Homoplastic transplantation using induced Hydranth as "implants".

Epithelial cells are stem cell lineages responsible for cell signalling. Transgenic Hydras reporter genes are cell-signalling with each other inside humans, much like *neurons* in a neural network. Transgenic Hydras cell signalling becomes synthesized with human cell signalling in a process called **catenin signalling**, 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 Hydras, parasites, and newly hybridized humans.

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

A **team of scientists** 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 behaviour. This technology was developed over the last decade through the **Human Brain Project**.

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

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 **initiative called 2045** 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) Hydras 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

Hydras 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.

Did you think the Starlink satellite network's *"Precision Tracking Space System"* had something to do with *defense?* 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 **CRISPR parasites** to be used as **gene vectors** for human transfection. These are not "vaccines" at all but a **WEAPONS SYSTEM** (*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).

ProSplign 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 *new neural network* in Hydras. This technology is being deployed in humans through the Covid-19 Quackccine [vaccine] program now.

Dr Carrie Madej also disclosed in her **latest interview on Stew Peter's Show** that the vaccine operating system is building an artificial neural network in humans.

ADDGene offers a **Tetracycline off system** 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 **Hydra genome assembly** offers a Nano DNA kit called Illumina. Illumina Inc figured out how to **reduce the cost of sequencing** 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 **Vector Biolabs** 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 **Vector Builder**. You can create artificial genome sequences and merge genomes of different species.

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

The **Genome Data Viewer (GDV)** 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 **shRNA cloning options** 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 catalogue of over **27,000 shRNA plasmid sets** targeting the human genome.

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

ADDGene's **lentiviral genome** 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 **Adenovirus (AAV) expressing shRNA** for the knockout (gene silencing) of Human SP5. When developing this technology during the **animal trials**, social recognition, spatial learning, and memory were impaired after 4 weeks.

In an **animal study** 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 behavioural impairment.

REPRODUCTION & FERTILITY

The transgenic Hydras 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**.

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

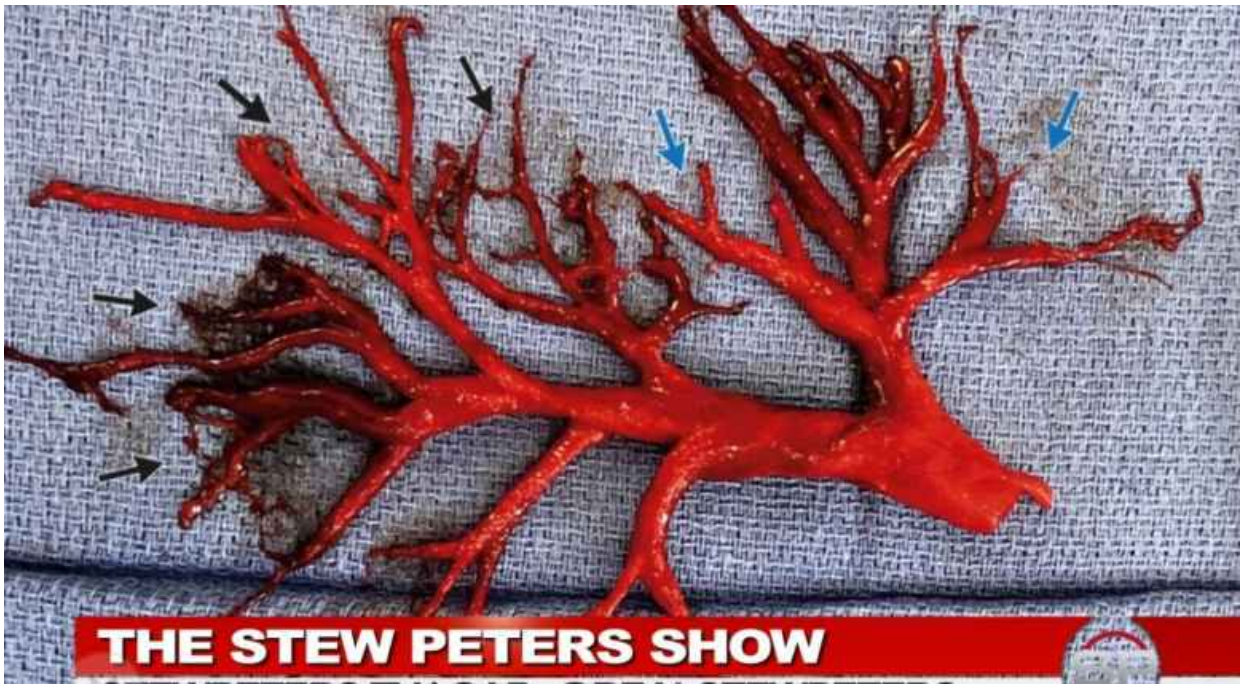
Proteins control gene expression. Transgenic **Hydra is instrumental** in encoding the human SP5 (shRNA **silencing AAV**) which is a gene on chromosome 2q31.1 that encodes a protein that binds to the GC-box promoter elements, thought to play a role in coordinating the intricate changes in transcription 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 Hydras, targeting human ***embryos*** and ***embryonic cells***, leading to developmental alterations from binding genes to the Wnt/β-catenin signalling pathway.

Do you understand what this means? 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) ***integrates randomly*** 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!



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 vax.

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 **78th Annual Meeting of Association of American Physicians and Surgeons**, 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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