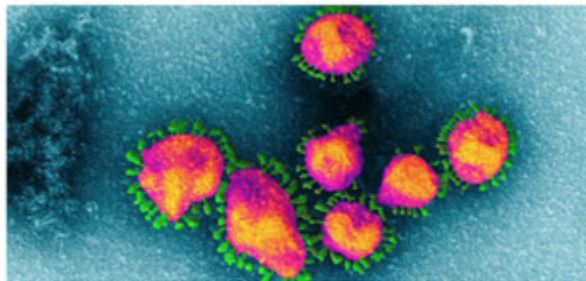


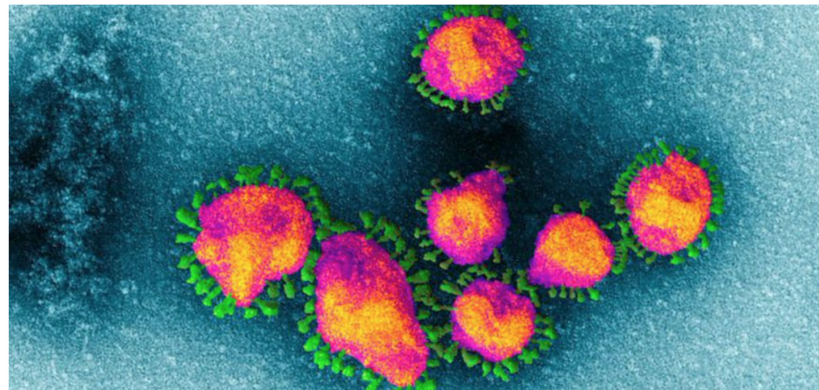
Dismantling The Viral Theory

59-74 minutes

HAS THE EXISTENCE OF POLIO, MEASLES, HIV, CMV, EBV, HEP C, EBOLA, THE FLU, ZIKA AND NOW CORONA VIRUSES BEEN DEMONSTRATED AND SCIENTIFICALLY PROVEN?



A Micrograph of an aggregation of infected red blood cells with the phantom Coronavirus. These so-called infected cells are nothing more than biological transforming red blood cells that are going through pleomorphic changes due to increased acidity and a declining pH - <7.2. The biological transformation of blood or body cells is a natural process that takes place in an acidic environment of the interstitial fluids of the Interstitium compartments and then spilling over into the blood plasma via hydrostatic pressure caused by the buildup of dietary and metabolic acidic waste which has not been properly eliminated by the lymphatic system via the four channels of elimination - urination, defecation, perspiration and/or respiration.. <https://www.drrobertyoung.com/post/dismantling-the-viral-theory>



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The first isolation of a virus was achieved in 1892 by Russian bacteria hunter Dimitri Iwanowski, who gathered fluid from diseased tobacco plants. He passed this liquid through a filter fine enough to retain bacteria; yet to Iwanowski's surprise, the bacteria-free filtrate

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easily made healthy plants sick. In 1898 a Dutch botanist, Martinus Willem Beijerinck, repeating the experiment, also recognized that there was an invisible cause and named the infectious agent “tobacco mosaic virus.” In the same year as Beijerinck’s report, two German scientists purified a liquid containing filterable viruses that caused foot-and-mouth disease in cattle (viruses were at one time called “filterable viruses,” but eventually the term “filterable” came to apply only to viruses, and was dropped). Walter Reed followed in 1901 with a filtrate responsible for yellow fever, and soon dozens of other disease-causing viruses were found.

History of Virology

- 1892 –Dmitri Iwanowski shows that extracts from diseased tobacco plants can transmit disease to other plants after passage through filters fine enough to retain the smallest known bacteria.
 - **This is generally recognized as the beginning of Virology!** But, nobody understood the significance until...
- 1898 – Beijernick made the same discovery, but suggested that the pathogen is a distinct agent, not just really small bacteria




Dmitri Iwanowski
(1864-1920)




Martinus Beijerinck
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In 1935 another American, Wendell M. Stanley, went back to the beginning and created pure crystals of tobacco mosaic virus from a filtered liquid solution. He affirmed that these crystals could easily infect plants, and concluded that a virus was not a living organism, since it could be crystallized like salt and yet remained infectious. Subsequently, bacteriologists all over the world began filtering for viruses, and a new area of biology was born-virology.

Dismantling The Viral Theory

Historically, medical science has vacillated on the question of whether a virus is alive. Originally it was described as nonliving, but is currently said to be an extremely complex molecule or an extremely simple microorganism, and is usually referred to as a parasite having a cycle of life. (The term “killed” is applied to certain viral vaccines, thus implying an official conviction that viruses live.) Commonly composed of either DNA or RNA cores with protein coverings, and having no inherent reproductive ability, viruses depend upon the host for replication. They must utilize the nucleic acids of living cells they infect to reproduce their proteins (i.e., trick the host into producing them), which are then assembled into new viruses like cars on an assembly line. Theoretically, this is their only means of surviving and infecting new cells or hosts.

The Replicating Virus Theory

Then it was discovered that, when bacteria slowly begin to die, bacteria create tiny, apparently lifeless forms of survival, the so-called spores. It was then suspected that these spores were toxic and that they were the so-called pathogenic poisons. This was then refuted, since the spores are rapidly developing into bacteria when their vital resources are being restored. When scientists in the laboratory observed that the weak, highly inbred bacteria perished very quickly while turning into much smaller structures than the spores, it was first believed that the bacteria were being killed by the alleged pathogenic poisons, called viruses, and that the viruses were thereby replicating.





The micrograph above was done using Dark Field Microscopy showing red blood cells and the evolution of bacterial phages and bacterial spores (the white spots) from red blood cell biological transformation

The Invention of Bacterial Viruses

Due to the belief that these – at the time of their discovery still invisible- structures were killing the bacteria, they were called phages/bacteriophages, “eaters of bacteria”. Only later it was determined that merely highly inbred and therefore almost non-viable bacteria can be made to turn into phages, or bacteria which are being destroyed so fast that they do not have time to form spores.

The introduction of the electron microscopy led to the discovery of the structures resulting from the biological transformation or pleomorphism of bacteria when these were suddenly dying or when the metabolism of the highly inbred germs was overwhelmed by processes triggered by the adding of “phages”. It was also discovered that there are hundreds of types of different-looking “phages”. The discovery of phages, the so-called bacterial “viruses”, reinforced the wrong assumption and the belief that there were human and animal viruses that looked the same and had the same structure. This is not and cannot be the case, for several different reasons.

After introducing chemical examination techniques in biology, it was discovered that there are thousands of types of phages and that phages of one type always have the same structure. They consist of a particular molecule, made of nucleic acid, which is covered in a

shell of proteins of a given number and composition. It was only later discovered that merely the bacteria which had been highly inbred in the test tube could turn into phages themselves, by contact with phages, but this never applied to natural bacteria or bacteria which had just been isolated from their natural environment. In this process, it was discovered that these “bacterial viruses” actually serve to provide other bacteria with important molecules and proteins, and that the bacteria themselves emerged from such structures.

Before it could be established that the “bacterial viruses” cannot kill natural bacteria, but they are instead helping them to live and that bacteria themselves emerge from such structures, these “phages” were already used as models for the alleged human and animal viruses. It was assumed that the human and animal viruses looked like the “phages”, were allegedly killing cells and thereby causing diseases, while at the same time producing new disease poisons and in this way transmitting the diseases. To date, many new or apparently new diseases have been attributed to viruses if their origin is unknown or not acknowledged.

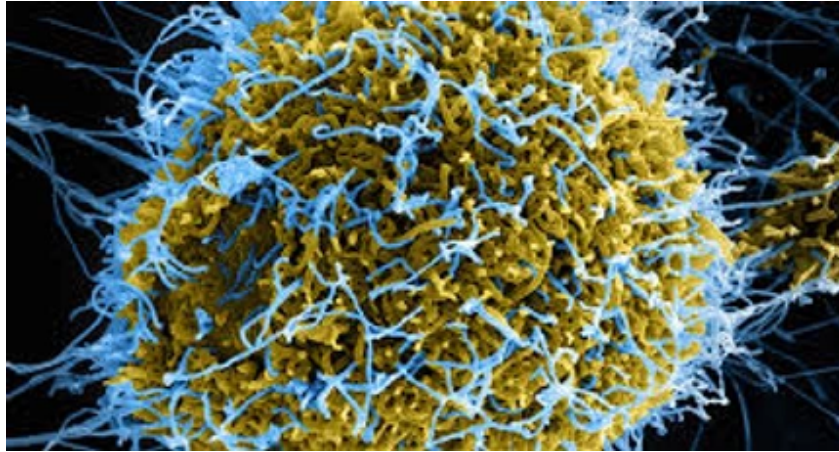
This reflex found an apparent confirmation in the discovery of the “bacterial viruses”.

It is important to note that the theories of fight and infection were accepted and highly praised by a majority of the specialists only if and when the countries or regions where they lived were also suffering from war and adversity. In times of peace, other concepts dominated the world of science.[272]

It is very important to note that the theory of infection – starting from Germany – has only been globalized through the third Reich, when the Jewish researchers, most of which had opposed and refuted the politically exploited theories of infection, were removed from their positions.[273]

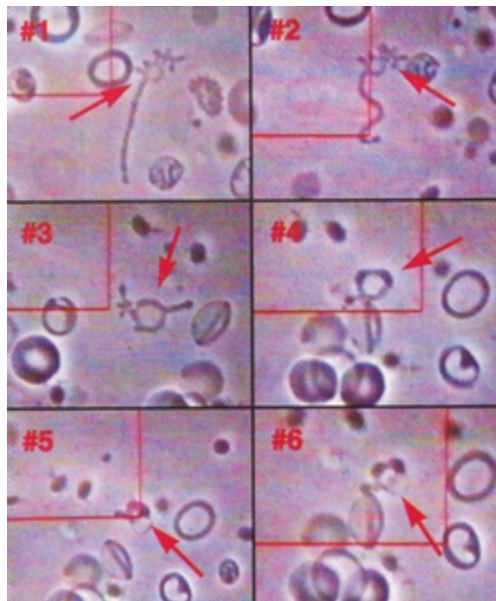
The Detection of Phages and Biological Transformation - The existence of phages can be proved rapidly

Dismantling The Viral Theory

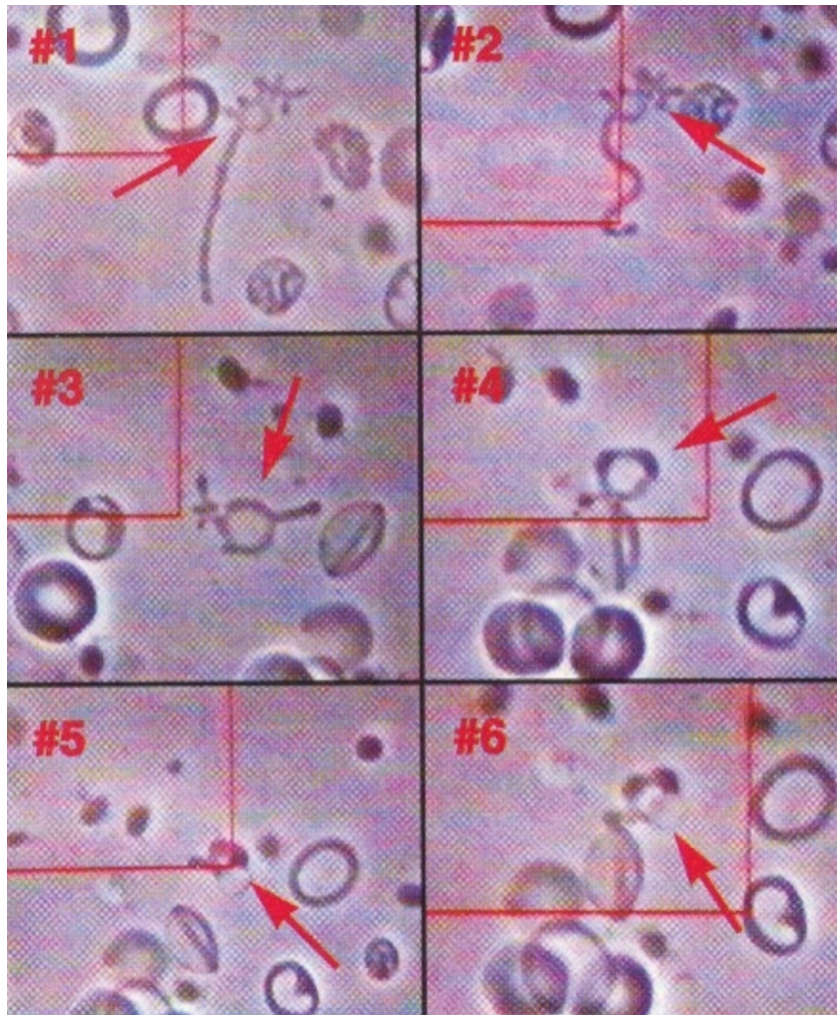


Bacterial pHage being born out of a blood and/or body cell. A biological process known as pleomorphism

First step: their presence is confirmed through an effect, namely the transformation of bacteria into phages, and also through an electron micrograph of those phages. The control experiments show that phages do not appear if bacteria do not change or if bacteria randomly start decomposing due to extrinsic sudden annihilation, without forming phages.



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The micrographs (micrographs #1 through #6] above show the cellular transformation of red blood cells, using phase contrast microscopy, into rod bacteria, cell-wall deficient bacteria, Y-form yeast and then bacteria phages

Second step: the liquid containing the phages is concentrated and applied on another liquid, which has a high concentration at the bottom of the test tube and a low concentration at the top of the test tube. The test tube with the phages is then powerfully spun (centrifuged) and all the particles gather according to their mass and weight to the place of their own density. The density is the ratio of weight (mass) per unit of volume, expressed as Kg/l or g/mg, respectively. That is why this concentration and purification step for particles with the same density is called density gradient centrifugation.

The layer where many particles of the same density gather becomes “cloudy”, which is called a “band.” This step is being documented, then the particles concentrated, purified and

sedimented in a “band” are removed with a syringe needle. The extracted concentrated amount of particles is called an isolate. A fast and simple electron micrograph will confirm the presence of phages in the isolate, which at the same time is an indication for the purity of the isolate, if the micrograph shows no other particles but the phages. The appearance and the diameter of the phages will also be established with the help of this micrograph.

The control experiment performed for this step consists in treating and centrifuging the liquid from bacteria which did not form any phages, where no phages appear at the end of the procedure.

After the step of successfully isolating the phages, the decisive biochemical characterization of the phages follows. The biochemical characterization of their composition is essential for identifying the specific type of phage, since different types of phages often appear to be similar. The isolate obtained through the density gradient centrifugation is now divided in two parts. One part is used to determine the size, type and composition of the nucleic acid; in a separate procedure, the other part is used to determine the amount, size and morphology of the proteins of the phages. Since the 1970s, these tests have been simple standard techniques that are learned by every biology student in their first semesters.

These tests represent the biochemical characterization of the phages. In almost every case, these results have been and are being published in only one publication, since a phage has a very simple structure which is very easy to analyze. The control experiments for these tests use liquid from bacteria which do not form phages and thus cannot present any biochemical proof. The existence of approximately two thousand different types of phages have been scientifically demonstrated this way

The So-Called Pathogenic Viruses

The “bacteriophages,” correctly defined as incomplete mini spores and building blocks of the bacteria, have been scientifically isolated, while the so-called pathogenic viruses have never been observed in humans or animals or in their body fluids and have never been isolated and subsequently biochemically analyzed. To date, none of the researchers involved in virology research seems to have realized this very important point.

The use of electron microscopy and the biochemistry were very

slowly returning to normal after 1945 and no one had realized that not one pathogenic virus had ever been isolated in humans or animals; thus, as of 1949 researchers started applying the same idea used for the (bacterio) phages, in order to replicate the human and animal “viruses.” John Franklin Enders, born in 1897 in the family of a rich financier, was active in various fraternities after having finished his studies, then he worked as a real estate agent and studied foreign languages for four years before turning to bacterial virology, which fascinated him. He then simply transferred the ideas and concepts that he learned in this area of research to the supposed pathogenic viruses in humans.

UnScientific Experiments and Interpretations Gave Birth to Virology

With his unscientific experiments and interpretations that he had never confirmed through negative controls, Enders brought the entire “viral” infectious medicine to a dead end. It is important to note at this point that Enders, like many infectious diseases specialists, worked for the U.S. military, which had always been and remains to date a huge victim of the fear of contagions. It was mainly the U.S. military which spread its erroneous belief that besides chemical weapons there were also biological weapons in the form of bacteria and viruses.

In 1949, Enders announced that he had managed to cultivate and grow the alleged polio virus in vitro on various tissues. The American expert opinion believed everything immediately. What Enders did was to add fluids from patients with poliomyelitis to tissue cultures which he claimed to have had sterilized, then he alleged that the cells were dying because of the virus, that the virus was replicating in this way and that a vaccine could be harvested from the respective culture. At that time, summer polio epidemics (polio = flaccid paralysis) were very frequent during summer and they were believed to be caused by the polio virus.

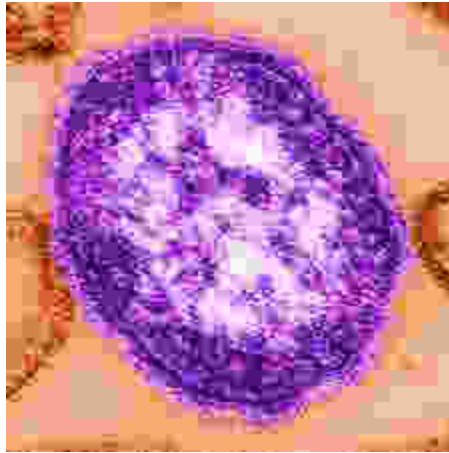
A vaccine was to help eradicate the alleged virus. After the polio vaccine was introduced, the symptoms were then re-diagnosed among other things as multiple sclerosis, flaccid acute paralysis, aseptic meningitis etc. and later polio was claimed to have been eradicated. During his experiments, Enders et al. sterilized the tissue cultures in order to exclude the possibility of bacteria killing the cells. What he didn't take into consideration was that the

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sterilization and the treatment of the cell culture when preparing it for the alleged infection was exactly what was destroying and killing the cells. Instead, he interpreted the cytopathic effects as the existence and the action of a so-called polio virus, without ever having isolated a single virus and describing its biochemistry.

The necessary negative control experiments, which would have shown that the sterilization and the treatment of the cells prior to the “infection” in the test tube was killing the cells, have never been performed. However, for this “performance” Enders received the Nobel prize in 1954.

The Invention of the Polio Virus and ‘YES’ the Measles Virus Too! NOW We Have The Invention of the Coronavirus!



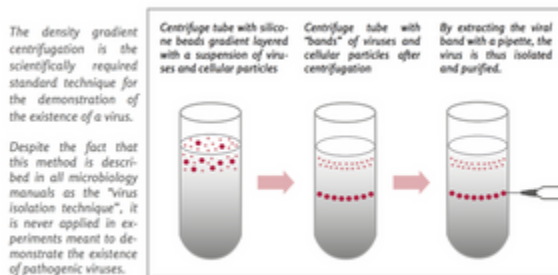
Measles virus or a bacterial pHage or Just Cellular Debris?

Dismantling The Viral Theory

1954 is also the year in which Enders applied and introduced the same technique in order to allegedly replicate the measles virus. As he had been awarded the Nobel prize for the alleged polio virus the same year, all researchers believed his technique to be scientifically valid. Thus, to date, the entire concept of polio and measles has been based upon this unscientific technique and fraud.

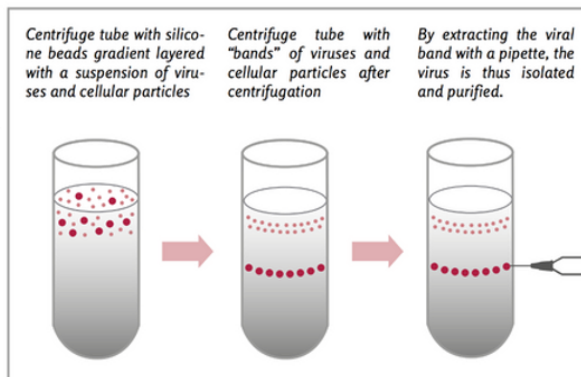
Thus, the polio and measles vaccines do not contain viruses, but particles of dead monkey kidney tissue or human cancerous body cells. To date, no negative control experiments have been done with respect to the so-called polio and measles viruses either, which would have shown that it was the laboratory procedures that lead to the cytopathic effects on the cells.

Additionally, all claims and experiments made by Enders et al. and subsequent researchers lead to the only objective conclusion, that in fact they were observing and analyzing the cellular particles or fragments and the activity thereof in the test tube, misinterpreting these as particles and characteristics of the alleged polio and/or measles viruses.



The density gradient centrifugation is the scientifically required standard technique for the demonstration of the existence of a virus.

Despite the fact that this method is described in all microbiology manuals as the "virus isolation technique", it is never applied in experiments meant to demonstrate the existence of pathogenic viruses.



Cellular Debris NOT So-Called Viruses

ALL Viruses from HIV, EBV, CMV, Hepatitis C, West Nile Virus, Ebola, Measles, Zika, and Now the Coronavirus, are ALL Phantom Viruses - Viral Existence Has NEVER Been Scientifically Demonstrated and Never Proven!

The following explanations applies to all the so-called (human or animal) “pathogenic viruses”. The six papers provided by Dr. Bardens in the course of the “measles trial” as proof for the existence of the measles virus described in a didactically ideal way the various steps of the chain of misinterpretations up to the belief in the existence of a measles virus.

The first paper was published in 1954 by Enders et al.:
“Propagation in tissue cultures of cytopathogenic agents from patients with measles” (Proc Soc Exp Biol Med. 1954 Jun; 86 (2): 277–286).

This publication can be found on the internet, like all the other publications presented at the measles trial. In that experiment, Enders et al. cut down dramatically on the nutrient solution and added cell-destroying antibiotics to the cell culture before introducing the allegedly infected fluid. The subsequent dying of the cells was then misinterpreted as presence and also isolation of the measles virus. No control experiments were performed to exclude the possibility that it was the deprivation of nutrients as well as the antibiotics which led to the cytopathic effects.

Enders’ and his colleagues’ blindness can be explained by the fact that he truly wanted to help people, while the ‘virus hysteria’ was intensifying after the war and during the cold war. It can also be explained by the fact that Enders and many of his colleagues had no idea about medicine or biochemistry and they were competing with the Soviet Union for the development of the first measles vaccine. Such a pressure for success can also explain why Enders and his colleagues ignored their own reservations and cautions expressed in 1954, when they had observed and noted that many cells also died after being treated normally (i.e. without being “infected”), which they thought to have been caused by unknown viruses and other factors. All these facts and cautions were subsequently disregarded.

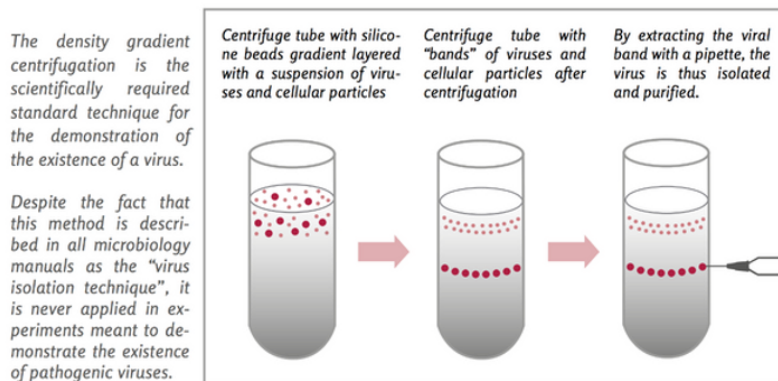
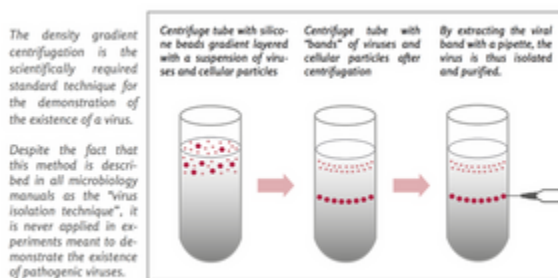
The second paper presented by the claimant in the ‘measles trial’

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was published in 1959[274] and, for the reasons presented above, the authors concluded that the technique introduced by Enders was not appropriate for the isolation of ANY virus. This rebuttal is not only NOT being discussed by ALL the other researchers, but it is being ignored completely!

The 'Viral Dogma' of Pathogenic Viruses is Still Being Promoted Today!

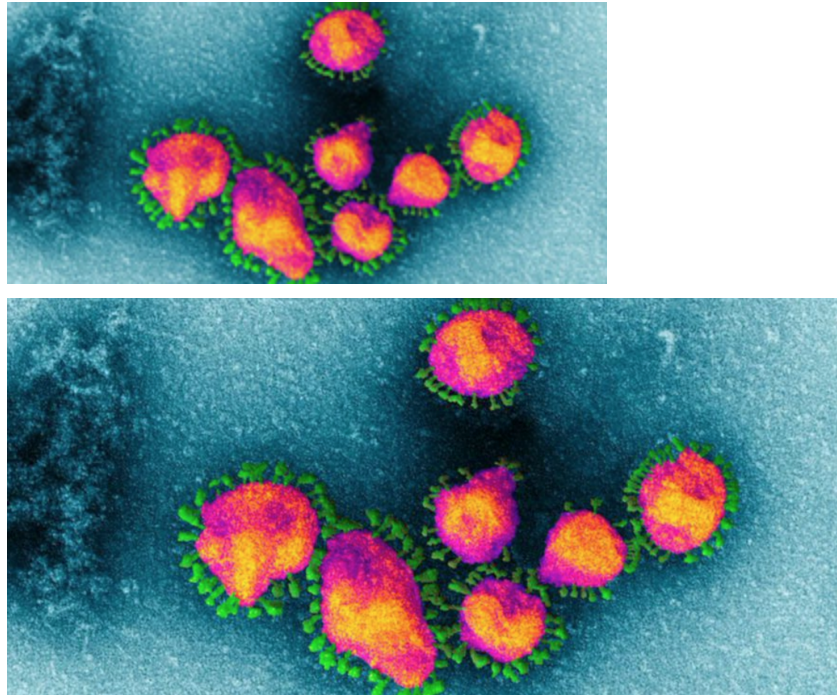
In a third paper[275], the authors photographed typical cellular particles inside the cells and misinterpreted these as measles virus. They did not isolate any virus. For unexplained reasons, they failed to determine and describe the biochemical structure of what they were presenting as a virus in a separate experiment. In the short description of the methods used, one can read that the authors did not apply the standard isolation technique for viruses, i.e. the density gradient centrifugation. They simply centrifuged fragments of dead cells at the bottom of a test tube and then, without describing their biochemical structure, they misinterpreted the cellular debris as viruses.



Cellular Debris NOT So-Called Viruses

From the way the experiments were performed, one can only conclude that cellular particles were misinterpreted as viruses. We find the same situation in the fourth[276] and the sixth[277] publication put forward by the claimant as proof of the existence of a measles virus. The fifth publication[278] is a review describing the consensus process as to which nucleic acid molecules from the dead cells would represent the so-called genome of the polio or measles virus. The result is that dozens of research teams work with short pieces of cell-specific molecules, after which -following a given model – they put all the pieces together on paper. However, this jigsaw puzzle made of so many pieces was never scientifically proven to exist as a whole and was never isolated from a virus, for a polio, measles, HIV or Hepatitis C, Ebola or Zika viruses have never been seen, neither in humans nor in a test tube. Referring to this publication, the court-appointed expert stated that it described the gold standard, i.e. the entire virus genome. It is obvious that the expert did not read this paper, whose authors stated that the exact molecular composition and functions of the measles virus genome will have to be the object of further research, which is why they had to rely on other virus models in order to achieve a consensus on the structure and functions of ANY virus genome. The easiest thing for anyone to notice is that in all of these publications, as well as in all other publications on the “measles virus” and other pathogenic viruses, including HIV, EBV, CMV, Ebola and Zika, no control experiments have ever been performed. No researchers used the density gradient centrifugation technique; instead, they only centrifuged cellular debris at the bottom of a test tube. This technique, used to collect all the particles from a fluid, is called pelletizing. From a logical and scientific perspective, it can be said that in all publications on the so-called “pathogenic viruses”, the researchers demonstrated in fact only particles and characteristics of cells. I would also like to point out that the so-called giant viruses[279] , i.e. an enwrapped nucleic acid can be found everywhere in the sea and in basic organisms. Like all bacterial phages, not only are they harmless, but they have beneficial functions. They can be also isolated by using the density gradient centrifugation, which proves their existence (see the graphic above).

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A Micrograph of an aggregation of infected red blood cells with the phantom Coronavirus. These so-called infected cells are nothing more than biological transforming red blood cells that are going through pleomorphic changes due to increased acidity and a declining pH - <7.2. The biological transformation of blood or body cells is a natural process that takes place in an acidic environment of the interstitial fluids of the Interstitium compartments and then spilling over into the blood plasma via hydrostatic pressure caused by the buildup of dietary and metabolic acidic waste which has not been properly eliminated by the lymphatic system via the four channels of elimination - urination, defecation, perspiration and/or respiration.. <https://www.drrobertyoung.com/post/dismantling-the-viral-theory>

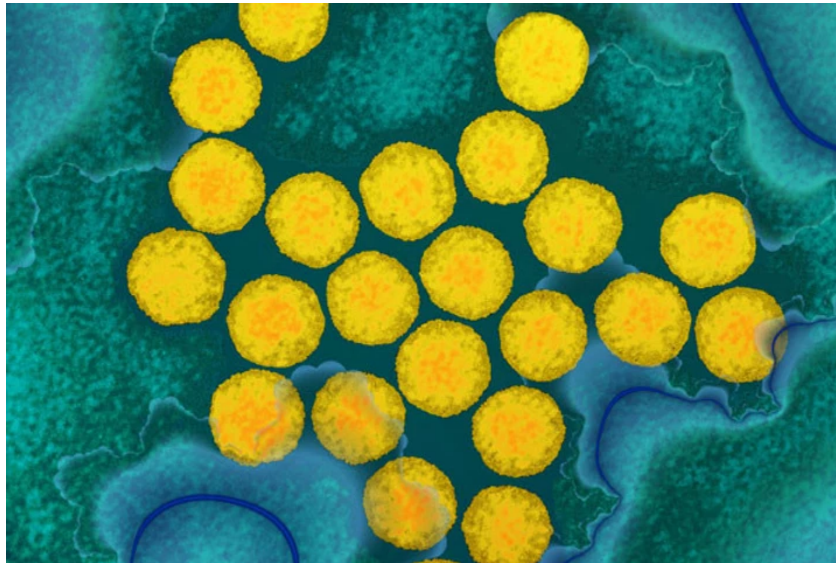
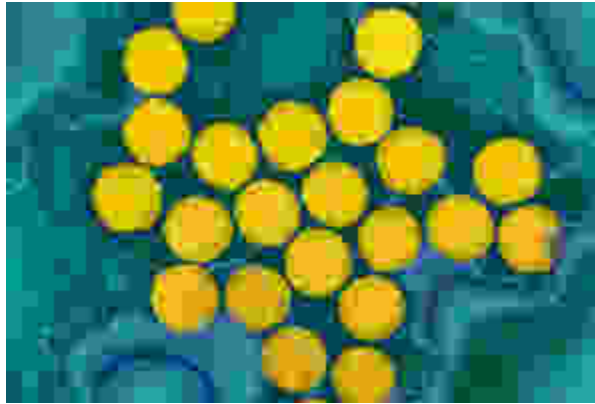
I also recommend Prof. Lüdtke's relevant review (1999).[280] He noted that at the early beginnings of virology, the majority of virologists always concluded that the structures they had mistaken for viruses turned out to be components of the cells and thus, they were only the result of the experiment and not the cause of the changes observed.

After the discovery and characterization of the phages and after introducing the dogma that the nucleic acid was the genome of all cells and viruses, the consensus was born, according to which such viruses must exist in humans and animals as well. In 1992, the dogma stating that the nucleic acid is the genotype of all cells was

Dismantling The Viral Theory

retracted in the scientific community. The 'viral dogma' of pathogenic viruses, however, is still being promoted today to the harm of billions of people. – for what?

The Bottom Line Concerning Phantom Viruses and the Polio and Measles Virus



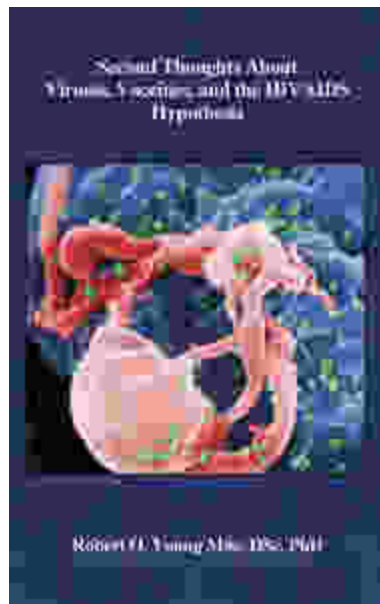
An Electron micrograph of the so-called Polio virus that has never been demonstrated scientifically to cause the symptoms of paralysis. Illustration has been colorized for effect

My bottom line still holds the truth that the terrain or internal environment is everything and the germ or so-called virus is NOTHING! The germ or so-called virus can only be a symptom of cellular breakdown due to an imbalance of the delicate alkaline pH balance of the body fluids and NOT the cause of that breakdown. That is why years ago I offered any scientist in the World a finders fee of 5 million US dollars if they could prove the existence of the HIV virus using Koch's postulates. It has now been over 20 years

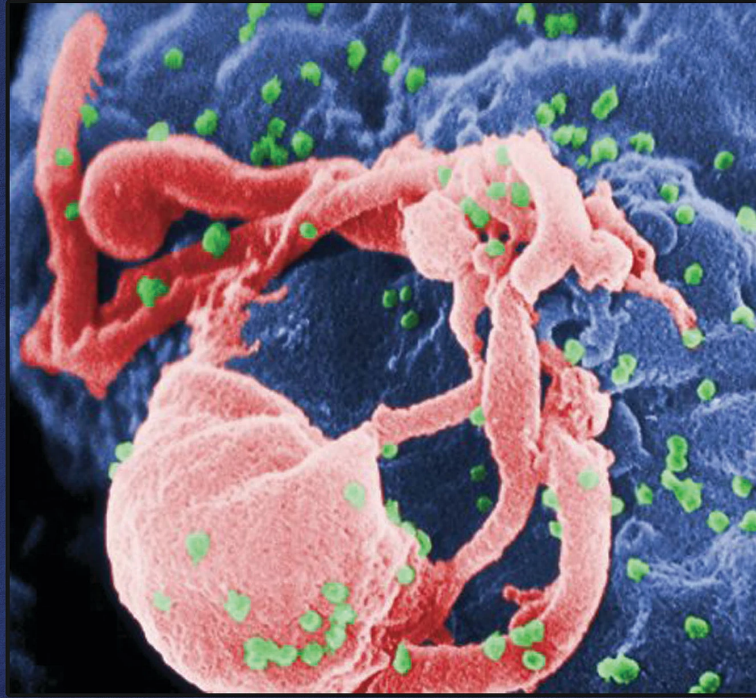
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and I am still waiting even though currently I no longer have the funds to pay the prize due to political assassination! It is unfortunate that a former 5 million US dollar prize offered 20 years ago was not enough money to change the current medical viral dogma that is currently paying out trillions of dollars to guess who?[281]

Dismantling The Viral Theory



**Second Thoughts About
Viruses, Vaccines, and the HIV/AIDS
Hypothesis**



Robert O. Young MSc, DSc, PhD

The Genesis of Severe Acute Respiratory (Syndrome) Disease or SARS (Coronavirus - COVID - 2 and COVID - 19) is Found in the Interstitial Fluids of Intestium

Abstract

Interstitial lung disease (IFLD), or diffuse parenchymal lung disease (DPLD),[3] is a group of lung diseases affecting the Interstitium (the interstitial fluids or space around the alveoli (air sacs of the lungs).[4] It concerns alveolar epithelium, pulmonary capillary endothelium, basement membrane, and perivascular and perilymphatic tissues. It occurs when metabolic, dietary, respiratory and environmental acids injure the lung tissues that triggers an abnormal healing response. Ordinarily, the body generates just the right amount of tissue to repair acid damage, but in interstitial lung disease, the repair process goes awry because of the acidic pH (ideal healing takes place at a pH of 7.365)[5]) of the interstitial fluids that effects the normal healing of the tissue around the air sacs (alveoli) and therefore becomes scarred and thickened. Micrograph of usual interstitial pneumonia (UIP). UIP is the most common pattern of interstitial pneumonia (a type of interstitial lung disease) and usually represents pulmonary fibrosis caused by decompensated acidosis of the interstitial fluids of the largest organ of the human body - the Interstitium. H&E stain. Autopsy specimen. This makes it more difficult for oxygen to pass into the bloodstream. The term ILD is used to distinguish these diseases from obstructive airways diseases but the cause of all lung disease is due to decompensated acidosis of the interstitial fluids (pH is below 7.2) that is systemic although affecting the weakest area of the lungs. These weaknesses can be attributed to lifestyle and dietary choices There are specific types of interstitial lung disease in children. The acronym chILD is used for this group of diseases and is derived from the English name, Children's Interstitial Lung Diseases – chILD.[6] Prolonged interstitial lung disease may result in

pulmonary fibrosis, but this is not always the case. Pulmonary fibrosis is specifically caused by the increase of acids in the interstitial fluids of the lung due to an inverted way of living, eating, drinking, breathing, thinking, feeling and believing. Interstitial lung disease of the Interstitium is associated with typical findings both radiographic (basal and pleural-based fibrosis with honeycombing) and pathologic (temporally and spatially heterogeneous fibrosis, histopathologic honeycombing, and fibroblastic foci). Our findings using advanced technologies with non-invasive, non-radioactive 3D Bio-Electro Scanning of the Interstitial fluids of the largest organ of the body, the Interstitium to determine a complete chemistry, including pH. This testing has shown in all cases of interstitial fluid lung disease, including viral diseases, and malignancies with 98 percent accuracy for decompensated metabolic acidosis, of the IFLD, high levels of lactic acid and high levels of calcium. Since the interstitial fluids pass through every organ, gland and tissues we now have a complete picture of the functionality and chemistry of every organ, every gland and every tissue, including all bones and muscles.

Title

What Do Viruses Like HIV & Corona Have In Common With Exosomes?

Abstract

There is only one sickness, one disease and one treatment. The one sickness and one disease is the over-acidification of the blood and then interstitial fluids due to an inverted way of living, eating, drinking, breathing, thinking, feeling and believing. There are six major contributing factors that lead to the declining acidic pH of the body fluids. As the pH of the body fluids become compensated by these six contributing factors and the body cell membranes and genetic material begin to degenerate the cells release exosomes as a defense to activate and support the lymphocytes to release oxygen species or antioxidants to reduce the acidic loads stored in the interstitial fluids of the Interstitium. The one treatment is to support the immune system with increased amounts of reduced oxygen (O-) and reduced hydrogen (H-) to restore the alkaline design of the body fluids, open up the channels of elimination in order to remove dietary, metabolic, respiratory and environmental

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toxic acidic waste held in the interstitial fluids of the Interstitium and thus restoring health, energy and vitality to the body.

Abstract

There is now over 100 years of documented history and research on the Polio virus and whether or not its treatment by inoculation has been successful in eradicating Polio. I am suggesting in this article and in my lecture that there are significant findings based on historical and past and current research, including my own that the viral theory of Polio and possibly other modern-day diseases, such as Post-Polio Syndrome, Polio Vaccine-Induced Paralysis, Legionnaires, CNS disease, Cancer, HIV/AIDS and now Zika may be caused by acidic chemical poisoning from DDT (dichloro-diphenyl-trichloroethane) and other related DDT pesticides, acidic vaccinations, and other factors including lifestyle and dietary factors rather than from a lone infectious virus. I will present ten historical graphs outlining the history of Polio, the production of DDT, BHC, Lead, Arsenic, Polio vaccinations and the author's theory that chemical poisoning, vaccination, and lifestyle and dietary choices are a more likely causes for the symptoms of Polio, neurological diseases, Cancer, HIV/AIDS and now Zika.

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