Coronavirus: Proof 5G Exposure Causes Oxygen Deprivation

Part 2

5G Network and Corona Virus

In Part One of this series, I cited two known examples of where people received in their vaccinations, pathogens that should never have been in their vaccines or they were "murdered" with "injected" poison pathogens. The most famous example is that of the SV40 virus in which 65 million children during the 1950s were vaccinated with a cancer agent known as SV40, inserted in their Salk/Sabin polio vaccines. Millions died from cancer prematurely cutting them down in the prime of life. The second incident appears to have been given to medical doctors and scientists because they were natural-pathic physicians and researchers in the area of natural cures and non-traditional medicine. Their mysterious deaths, i.e., murders were linked to the pathogen Nagalase.

The article that follows will enrage you, if you think your country would not use **5G** to kill every one of us, and then use plausible deniability to refute it had anything to do with the virus outbreak that first occurred in Wuhan, China, and then Milan, Italy, then you are just another vulnerable sheep waiting to be sheared. The author of the below article has proven that we are mere guinea pigs or lab rats in its effort to find pathogens that it can develop "gain of function", a term used to describe intensified virility and lethality. The objective has always been to maximize the "bang for the buck" when it comes to waging war on armies or as Dr. Henry Kissinger would say, "useless eaters"!

The following article below is by Donald W. Scott MA MSc President The Common Cause Medical Research Foundation. It first appeared in the August/September 2001, issue of *'Nexus'* magazine. Donald W. Scott's credentials are at the close of this most important article on Mycoplasma. It is foundational reading for those interested in the pathogen known as Mycoplasma. It is one of many papers and research findings on

weaponized Mycoplasma. My interest in Mycoplasma came about in my research three years about the dangers of **5G** EMF wireless communications. I once subscribed to *'Nexus'* magazine and this article was one of many great articles that I have never forgotten. I have Lymphedema and Neuropathy, in which the evidence shows Mycoplasma can be activated through wireless EMF communications. My problems began six years ago in October, 2014. It began two weeks after moving to where we live now. I'm in a direct line of sight 2,500 feet from a cell tower. Since I am a serious health observant person using vitamin and nutrient supplements, and Vitamin D^3 has kept me alive!

As the article states, as a result of the weaponization of the pathogen Mycoplasma, nearly all of us have a hidden time-bomb, that when activated, which wireless EMF energy can and will attack each individual with Mycoplasma at their genetic weakest area of the human biome, we all have a weak point, a predisposed weak point if you will, that can attack and kill us. It is the central element of decomposition of all life forms. I cannot say this any plainer, 5G EMF will through its millimeter-wave beam spectrum activate a molecular storm in your body that will kill you. From this article, we know the U.S. Military Biowarefare Labs are knowledgeable of the technology, as far back as fifty+ years ago, if not longer.

Mycoplasma: Linking Pathogen

The Linking Pathogen in Neurosystemic Diseases

Several strains of Mycoplasma have been "engineered" to become more dangerous. They are now being blamed for AIDS, cancer, CFS, MS, CJD and other neurosystemic diseases.

I – PATHOGENIC MYCOPLASMA

A Common Disease Agent Weaponized

There are 200 species of Mycoplasma. Most are innocuous and do no harm; only four or five are pathogenic. Mycoplasma fermentans (incognitus strain) probably comes from the nucleus of the Brucella bacterium. This disease agent is not a bacterium and not a virus; it is a mutated form of the Brucella bacterium, combined with a visna virus, from which the mycoplasma is extracted. The pathogenic Mycoplasma used to be very innocuous, but biological warfare research conducted between 1942 and the present time has resulted in the creation of more deadly and infectious forms of Mycoplasma. Researchers extracted this Mycoplasma from the Brucella bacterium and actually reduced the disease to a crystalline form. They "weaponized" it and tested it on an unsuspecting public in North America.

Dr. Maurice Hilleman, chief virologist for the pharmaceutical company Merck Sharp & Dohme, stated that this disease agent is now carried by everybody in North America and possibly most people throughout the world. Despite reporting flaws, there has clearly been an increased incidence of all the neuro/systemic degenerative diseases

since World War II and especially since the 1970s with the arrival of previously unheard-of diseases like chronic fatigue syndrome and AIDS.

According to Dr. Shyh-Ching Lo, senior researcher at The Armed Forces Institute of Pathology and one of America's top mycoplasma researchers, this disease agent causes many illnesses including AIDS, cancer, chronic fatigue syndrome, Crohn's colitis, Type I diabetes, multiple sclerosis, Parkinson's disease, Wegener's disease and collagen-vascular diseases such as rheumatoid arthritis and Alzheimer's. Dr. Charles Engel, who is with the U.S. National Institutes of Health, Bethesda, Maryland, stated the following at an NIH meeting on February 7, 2000: *"I am now of the view that the probable cause of chronic fatigue syndrome and fibromyalgia is the mycoplasma…"*

I have all the official documents to prove that Mycoplasma is the disease agent in chronic fatigue syndrome/fibromyalgia as well as in AIDS, multiple sclerosis and many other illnesses. Of these, 80% are U.S. or Canadian official government documents, and 20% are articles from peer-reviewed journals such as the *'Journal of the American Medical Association', 'New England Journal of Medicine'* and the *'Canadian Medical Association Journal.'* The journal articles and government documents complement each other.

How the Mycoplasma Works

The Mycoplasma acts by entering into the individual cells of the body, depending upon your genetic predisposition. You may develop neurological diseases if the pathogen destroys certain cells in your brain, or you may develop Crohn's colitis if the pathogen invades and destroys cells in the lower bowel. Once the Mycoplasma gets into the cell, it can lie there doing nothing sometimes for 10, 20 or 30 years, but if a trauma occurs like an accident or a vaccination that doesn't take, the Mycoplasma can become triggered. Because it is only the DNA particle of the bacterium, it doesn't have any organelles to process its own nutrients, so it grows by up taking pre-formed sterols from its host cell and it literally kills the cell; the cell ruptures and what is left gets dumped into the bloodstream. [It lives on the host's cells.]

II – CREATION OF THE MYCOPLASMA

A Laboratory-Made Disease Agent

Many doctors don't know about this Mycoplasma disease agent because it was developed by the U.S. military in biological warfare experimentation and it was not made public. This pathogen was patented by the United States military and Dr. Shyh-Ching Lo. I have a copy of the documented patent from the US Patent Office.[1]

All the countries at war were experimenting with biological weapons. In 1942, the governments of the United States, Canada and Britain entered into a secret agreement to create two types of biological weapons (one that would kill, and one that was disabling) for use in the war against Germany and Japan, who were also developing biological weapons. While they researched a number of disease pathogens, they primarily focused on the Brucella bacterium and began to weaponize it. From its

inception, the biowarfare program was characterized by continuing in-depth review and participation by the most eminent scientists, medical consultants, industrial experts and government officials, and it was classified Top Secret.

The U.S. Public Health Service also closely followed the progress of biological warfare research and development from the very start of the program, and the Centers for Disease Control (CDC) and the National Institutes of Health (NIH) in the United States were working with the military in weaponizing these diseases. These are diseases that have existed for thousands of years, but they have been weaponized-which means they've been made more contagious and more effective. And they are spreading.

The Special Virus Cancer Program, created by the CIA and NIH to develop a deadly pathogen for which humanity had no natural immunity (AIDS), was disguised as a war on cancer but was actually part of MKNAOMI.[2] Many members of the Senate and House of Representatives do not know what has been going on. For example, the U.S. Senate Committee on Government Reform had searched the archives in Washington and other places for the document titled *"The Special Virus Cancer Program: Progress Report No. 8",* and couldn't find it. Somehow they heard I had it, called me and asked me to mail it to them. Imagine: a retired schoolteacher being called by the United States Senate and asked for one of their secret documents! The U.S. Senate, through the Government Reform Committee, is trying to stop this type of government research.

Crystalline Brucella

The title page of a genuine U.S. Senate Study, declassified on February 24, 1977, shows that George Merck, of the pharmaceutical company, Merck Sharp & Dohme (which now makes cures for diseases that at one time it created), reported in 1946 to the US Secretary of War that his researchers had managed "for the first time" to "isolate the disease agent in crystalline form."[3] They had produced a crystalline bacterial toxin extracted from the Brucella bacterium. The bacterial toxin could be removed in crystalline form and stored, transported and deployed without deteriorating. It could be delivered by other vectors such as insects, aerosol or the food chain (in nature it is delivered within the bacterium). But the factor that is working in the Brucella is the Mycoplasma.

Brucella is a disease agent that doesn't kill people; it disables them. But, according to Dr. Donald MacArthur of the Pentagon, appearing before a congressional committee in 1969,[4] researchers found that if they had mycoplasma at certain strength-actually, 10 to the 10th power (10^{10}) - it would develop into AIDS, and the person would die from it within a reasonable period of time because it could bypass the natural human defenses. If the strength was 10^8, the person would manifest with chronic fatigue syndrome or fibromyalgia. If it was 10^7, they would present as wasting; they wouldn't die and they wouldn't be disabled, but they would not be very interested in life; they would waste away.

Most of us have never heard of the disease brucellosis because it largely disappeared when they began pasteurizing milk, which was the carrier. One salt shaker of the pure

disease agent in a crystalline form could sicken the entire population of Canada. It is absolutely deadly, not so much in terms of killing the body but disabling it. Because the crystalline disease agent goes into solution in the blood, ordinary blood and tissue tests will not reveal its presence. The Mycoplasma will only crystallize at 8.1 pH, and the blood has a pH of 7.4 pH. So the doctor thinks your complaint is "all in your head".

Crystalline Brucella and Multiple Sclerosis

In 1998 in Rochester, New York, I met a former military man, PFC Donald Bentley, who gave me a document and told me: *"I was in the U.S. Army, and I was trained in bacteriological warfare. We were handling a bomb filled with brucellosis, only it wasn't brucellosis; it was a Brucella toxin in crystalline form. We were spraying it on the Chinese and North Koreans."*

He showed me his certificate listing his training in chemical, biological and radiological warfare. Then he showed me 16-pages of documents given to him by the U.S. military when he was discharged from the service. They linked brucellosis with multiple sclerosis, and stated in one section: *"Veterans with multiple sclerosis, a kind of creeping paralysis developing to a degree of 10% or more disability within two years after separation from active service, may be presumed to be service-connected for disability compensation. Compensation is payable to eligible veterans whose disabilities are due to service."* In other words: *"If you become ill with multiple sclerosis, it is because you were handling this Brucella, and we will give you a pension. Don't go raising any fuss about it."* In these documents, the government of the United States revealed evidence of the cause of multiple sclerosis, but they didn't make it known to the public-or to your doctor.

In a 1949 report, Drs. Kyger and Haden suggested *"the possibility that multiple sclerosis might be a central nervous system manifestation of chronic brucellosis."* Testing approximately 113 MS patients, they found that almost 95% also tested positive for Brucella.[5] We have a document from a medical journal, which concludes that one out of 500 people who had brucellosis would develop what they call neurobrucellosis; in other words, brucellosis in the brain, where the Brucella settles in the lateral ventricles-where the disease multiple sclerosis is basically located.[6]

Contamination of Camp Detrick Lab Workers

A 1948 *'New England Journal of Medicine'* report titled "Acute Brucellosis Among Laboratory Workers" shows us how actively dangerous this agent is.[7] The laboratory workers were from Camp Detrick, Frederick, Maryland, where they were developing biological weapons. Even though these workers had been vaccinated, wore rubberized suits and masks and worked through holes in the compartment, many of them came down with this awful disease because it is so absolutely and terrifyingly infectious.

The article was written by Lt. Calderone Howell, Marine Corps, Captain Edward Miller, Marine Corps, Lt. Emily Kelly, United States Naval Reserve, and Captain Henry Bookman. They were all military personnel engaged in making the disease agent Brucella into a more effective biological weapon.

III – COVERT TESTING OF MYCOPLASMA

Testing the Dispersal Methods

Documented evidence proves that the biological weapons they were developing were tested on the public in various communities without their knowledge or consent. The government knew that crystalline Brucella would cause disease in humans. Now they needed to determine how it would spread and the best way to disperse it. They tested dispersal methods for Brucella suis and Brucella melitensis at Dugway Proving Ground, Utah, in June and September 1952. Probably, 100% of us now are infected with Brucella suis and Brucella melitensis.[8]

Another government document recommended the genesis of open-air vulnerability tests and covert research and development programs to be conducted by the Army and supported by the Central Intelligence Agency. At that time, the Government of Canada was asked by the U.S. Government to cooperate in testing weaponized Brucella, and Canada cooperated fully with the United States. The U.S. Government wanted to determine whether mosquitoes would carry the disease and also if the air would carry it. A government report stated that "open-air testing of infectious biological agents is considered essential to an ultimate understanding of biological warfare potentialities because of the many unknown factors affecting the degradation of micro-organisms in the atmosphere."[9]

Testing via Mosquito Vector in Punta Gorda, Florida

A report from *'The New England Journal of Medicine'* reveals that one of the first outbreaks of chronic fatigue syndrome was in Punta Gorda, Florida, back in 1957.[10] It was a strange coincidence that a week before these people came down with chronic fatigue syndrome, there was a huge influx of mosquitoes. The National Institutes of Health claimed that the mosquitoes came from a forest fire 30 miles away. The truth is that those mosquitoes were infected in Canada by Dr. Guilford B. Reed at Queen's University. They were bred in Belleville, Ontario, and taken down to Punta Gorda and released there. Within a week, the first five cases ever of chronic fatigue syndrome were reported to the local clinic in Punta Gorda. The cases kept coming until finally 450 people were ill with the disease.

Testing via Mosquito Vector in Ontario

The Government of Canada had established the Dominion Parasite Laboratory in Belleville, Ontario, where it raised 100 million mosquitoes a month. These were shipped to Queen's University and certain other facilities to be infected with this crystalline disease agent. The mosquitoes were then let loose in certain communities in the middle of the night, so that the researchers could determine how many people would become ill with chronic fatigue syndrome or fibromyalgia, which was the first disease to show. One of the communities they tested it on was the St. Lawrence Seaway valley, all the way from Kingston to Cornwall, in 1984. They let out hundreds of millions of infected mosquitoes. Over 700 people in the next four or five weeks developed myalgic encephalomyelitis, or chronic fatigue syndrome.

IV – COVERT TESTING OF OTHER DISEASE AGENTS

Mad Cow Disease/Kuru/CJD in the Fore Tribe

Before and during World War II, at the infamous Camp 731 in Manchuria, the Japanese military contaminated prisoners of war with certain disease agents. They also established a research camp in New Guinea in 1942. There they experimented upon the Fore Indian tribe and inoculated them with a minced-up version of the brains of diseased sheep containing the visna virus which causes "mad cow disease" or Creutzfeldt-Jakob disease. About five or six years later, after the Japanese had been driven out, the poor people of the Fore tribe developed what they called Kuru, which was their word for "wasting," and they began to shake, lose their appetites and die. The autopsies revealed that their brains had literally turned to mush. They had contracted "mad cow disease" from the Japanese experiments.

When World War II ended, Dr. Ishii Shiro-the medical doctor who was commissioned as a General in the Japanese Army so he could take command of Japan's biological warfare development, testing and deployment-was captured. He was given the choice of a job with the United States Army or execution as a war criminal. Not surprisingly, Dr. Ishii Shiro chose to work with the U.S. military to demonstrate how the Japanese had created mad cow disease in the Fore Indian tribe. In 1957, when the disease was beginning to blossom in full among the Fore people, Dr. Carleton Gajdusek of the U.S. National Institutes of Health headed to New Guinea to determine how the minced-up brains of the visna-infected sheep affected them. He spent a couple of years there, studying the Fore people, and wrote an extensive report. He won the Nobel Prize for "discovering" kuru disease in the Fore tribe.

Testing Carcinogens over Winnipeg, Manitoba

In 1953, the U.S. Government asked the Canadian Government if it could test a chemical over the city of Winnipeg. It was a big city with 500,000 people, miles from anywhere. The American military sprayed this carcinogenic chemical in a 1,000%-attenuated form, which they said would be so watered down that nobody would get very sick; however, if people came to clinics with a sniffle, a sore throat or ringing in their ears, the researchers would be able to determine what percentage would have developed cancer if the chemical had been used at full strength. We located evidence that the Americans had indeed tested this carcinogenic chemical-zinc cadmium sulphide-over Winnipeg in 1953. We wrote to the Government of Canada, explaining that we had solid evidence of the spraying and asking that we be informed as to how high up in the government the request for permission to spray had gone. We did not receive a reply.

Shortly after, the Pentagon held a press conference on May 14, 1997, where they admitted what they had done. Robert Russo, writing for the *Toronto Star*[11] from Washington, DC, reported the Pentagon's admission that in 1953 it had obtained permission from the Canadian Government to fly over the city of Winnipeg and spray out this chemical-which sifted down on kids going to school, housewives hanging out

their laundry and people going to work. U.S. Army planes and trucks released the chemical 36 times between July and August 1953. The Pentagon got its statistics, which indicated that if the chemical released had been full strength, approximately a third of the population of Winnipeg would have developed cancers over the next five years.

One professor, Dr. Hugh Fudenberg, MD, twice nominated for the Nobel Prize, wrote a magazine article stating that the Pentagon came clean on this because two researchers in Sudbury, Ontario-Don Scott and his son, Bill Scott-had been revealing this to the public. However, the legwork was done by other researchers! The U.S. Army actually conducted a series of simulated germ warfare tests over Winnipeg. The Pentagon lied about the tests to the mayor, saying that they were testing a chemical fog over the city, which would protect Winnipeg in the event of a nuclear attack. A report commissioned by U.S. Congress, chaired by Dr. Rogene Henderson, lists 32 American towns and cities used as test sites as well.

V – BRUCELLA MYCOPLASMA AND DISEASE

AIDS

The AIDS pathogen was created out of a Brucella bacterium mutated with a visna virus; then the toxin was removed as a DNA particle called a Mycoplasma. They used the same Mycoplasma to develop disabling diseases like MS, Crohn's colitis, Lyme disease, etc. In the previously mentioned U.S. congressional document of a meeting held on June 9, 1969,[12] the Pentagon delivered a report to Congress about biological weapons. The Pentagon stated: *"We are continuing to develop disabling weapons."* Dr. MacArthur, who was in charge of the research, said: *"We are developing a new lethal weapon, a synthetic biological agent that does not naturally exist, and for which no natural immunity could have been acquired."*

Think about it. If you have a deficiency of acquired immunity, you have an acquired immunity deficiency. Plain as that, AIDS. In laboratories throughout the United States and in a certain number in Canada including at the University of Alberta, the U.S. Government provided the leadership for the development of AIDS for the purpose of population control. After the scientists had perfected it, the government sent medical teams from the Centers for Disease Control-under the direction of Dr. Donald A. Henderson, their investigator into the 1957 chronic fatigue epidemic in Punta Gorda-during 1969 to 1971 to Africa and some countries such as India, Nepal and Pakistan where they thought the population was becoming too large.[13] They gave them all a free vaccination against smallpox; but five years after receiving this vaccination, 60% of those inoculated were suffering from AIDS. They tried to blame it on a monkey, which is nonsense.

A professor at the University of Arkansas made the claim that while studying the tissues of a dead chimpanzee she found traces of HIV. The chimpanzee that she had tested was born in the United States 23 years earlier. It had lived its entire life in a U.S. military laboratory where it was used as an experimental animal in the development of these diseases. When it died, its body was shipped to a storage place where it was deep-frozen and stored in case they wanted to analyze it later. Then they decided that they

didn't have enough space for it, so they said, *"Anybody want this dead chimpanzee?"* and this researcher from Arkansas said: *"Yes. Send it down to the University of Arkansas. We are happy to get anything that we can get."* They shipped it down and she found HIV in it. That virus was acquired by that chimpanzee in the laboratories where it was tested.[14]

Chronic Fatigue Syndrome/ Myalgic Encephalomyelitis

Chronic fatigue syndrome is more accurately called myalgic encephalomyelitis. The chronic fatigue syndrome nomenclature was given by the U.S. National Institutes of Health because it wanted to downgrade and belittle the disease. An MRI scan of the brain of a teenage girl with chronic fatigue syndrome displayed a great many scars or punctate lesions in the left frontal lobe area where portions of the brain had literally dissolved and been replaced by scar tissue. This caused cognitive impairment, memory impairment, etc. And what was the cause of the scarring? The Mycoplasma. So there is very concrete physical evidence of these tragic diseases, even though doctors continue to say they don't know where it comes from or what they can do about it.

Many people with chronic fatigue syndrome, myalgic encephalo-myelitis and fibromyalgia who apply to the Canada Pensions Plan Review Tribunal will be turned down because they cannot prove that they are ill. During 1999, I conducted several appeals to Canada Pensions and the Workers Compensation Board (WCB, now the Workplace Safety and Insurance Board) on behalf of people who have been turned down. I provided documented evidence of these illnesses, and these people were all granted their pensions on the basis of the evidence that I provided.

In March 1999, for example, I appealed to the WCB on behalf of a lady with fibromyalgia who had been denied her pension back in 1993. The vice-chairman of the board came to Sudbury to hear the appeal, and I showed him a number of documents which proved that this lady was physically ill with fibromyalgia. It was a disease that caused physical damage, and the disease agent was a mycoplasma. The guy listened for three hours, and then he said to me: *"Mr. Scott, how is it I have never heard of any of this before?* I said: *"We brought a top authority in this area into Sudbury to speak on this subject and not a single solitary doctor came to that presentation."*

VI – TESTING FOR MYCOPLASMA IN YOUR BODY

Polymerase Chain Reaction Test

Information is not generally available about this agent because, first of all, the Mycoplasma is such a minutely small disease agent. A hundred years ago, certain medical theoreticians conceived that there must be a form of disease agent smaller than bacteria and viruses. This pathogenic organism, the Mycoplasma, is so minute that normal blood and tissue tests will not reveal its presence as the source of the disease.

Your doctor may diagnose you with Alzheimer's disease, and he will say: "Golly, we don't know where Alzheimer's comes from. All we know is that your brain begins to deteriorate, cells rupture, the myelin sheath around the nerves dissolves, and so on." Or

if you have chronic fatigue syndrome, the doctor will not be able to find any cause for your illness with ordinary blood and tissue tests.

This Mycoplasma couldn't be detected until about 30 years ago when the polymerase chain reaction (PCR) test was developed, in which a sample of your blood is examined and damaged particles are removed and subjected to a polymerase chain reaction. This causes the DNA in the particles to break down. The particles are then placed in a nutrient, which causes the DNA to grow back into its original form. If enough of the substance is produced, the form can be recognized, so it can be determined whether Brucella or another kind of agent is behind that particular Mycoplasma.

Blood Test

If you or anybody in your family has myalgic encephalomyelitis, fibromyalgia, multiple sclerosis or Alzheimer's, you can send a blood sample to Dr. Les Simpson in New Zealand for testing.

If you are ill with these diseases, your red blood cells will not be normal doughnut-shaped blood cells capable of being compressed and squeezed through the capillaries, but will swell up like cherry-filled doughnuts which cannot be compressed. The blood cells become enlarged and distended because the only way the Mycoplasma can exist is by up taking pre-formed sterols from the host cell. One of the best sources of pre-formed sterols is cholesterol, and cholesterol is what gives your blood cells flexibility. If the cholesterol is taken out by the Mycoplasma, the red blood cell swells up and doesn't go through, and the person begins to feel all the aches and pains and all the damage it causes to the brain, the heart, the stomach, the feet and the whole body because blood and oxygen are cut off.

And that is why people with fibromyalgia and chronic fatigue syndrome have such a terrible time. When the blood is cut off from the brain, punctate lesions appear because those parts of the brain die. The Mycoplasma will get into portions of the heart muscle, especially the left ventricle, and those cells will die. Certain people have cells in the lateral ventricles of the brain that have a genetic predisposition to admit the mycoplasma, and this causes the lateral ventricles to deteriorate and die. This leads to multiple sclerosis, which will progress until these people are totally disabled; frequently, they die prematurely. The Mycoplasma will get into the lower bowel, parts of which will die, thus causing colitis. All of these diseases are caused by the degenerating properties of the Mycoplasma.

In early 2000, a gentleman in Sudbury phoned me and told me he had fibromyalgia. He applied for a pension and was turned down because his doctor said it was all in his head and there was no external evidence. I gave him the proper form and a vial, and he sent his blood to Dr. Simpson to be tested. He did this with his family doctor's approval, and the results from Dr. Simpson showed that only 4% of his red blood cells were functioning normally and carrying the appropriate amount of oxygen to his poor body, whereas 83% were distended, enlarged and hardened, and wouldn't go through the

capillaries without an awful lot of pressure and trouble. This is the physical evidence of the damage that is done.

ECG Test

You can also ask your doctor to give you a 24-hour Holter ECG. You know, of course, that an electrocardiogram is a measure of your heartbeat and shows what is going on in the right ventricle, the left ventricle and so on. Tests show that 100% of patients with chronic fatigue syndrome and fibromyalgia have an irregular heartbeat. At various periods during the 24 hours, the heart, instead of working happily away going "bump-BUMP, bump-BUMP", every now and again goes "buhbuhbuhbuhbuhbuhbuhbuhbuh". The T-wave (the waves are called P, Q, R, S and T) is normally a peak, and then the wave levels off and starts with the P-wave again. In chronic fatigue and fibromyalgia patients, the T-wave flattens off, or actually inverts. That means the blood in the left ventricle is not being squeezed up through the aorta and around through the body.

My client from Sudbury had this test done and, lo and behold, the results stated: "The shape of T and S-T suggests left ventricle strain pattern, although voltage and so on is normal." The doctor had no clue as to why the T-wave was not working properly. I analyzed the report of this patient who had been turned down by Canada Pensions and sent it back to them. They wrote back, saying: *"It looks like we may have made a mistake. We are going to give you a hearing and you can explain this to us in more detail."*

So it is not all in your imagination. There is actual physical damage to the heart. The left ventricle muscles do show scarring. That is why many people are diagnosed with a heart condition when they first develop fibromyalgia, but it's only one of several problems because the mycoplasma can do all kinds of damage.

Blood Volume Test

You can also ask your doctor for a blood volume test. Every human being requires a certain amount of blood per pound of body weight, and it has been observed that people with fibromyalgia, chronic fatigue syndrome, multiple sclerosis and other illnesses do not have the normal blood volume their body needs to function properly. Doctors aren't normally aware of this. This test measures the amount of blood in the human body by taking out 5 cc, putting a tracer in it and then putting it back into the body. One hour later, take out 5 cc again and look for the tracer. The thicker the blood and the lower the blood volume, the more tracer you will find.

The analysis of one of my clients stated: *"This patient was referred for red cell mass study. The red cell volume is 16.9 ml per kg of body weight. The normal range is 25 to 35 ml per kg. This guy has 36% less blood in his body than the body needs to function."* And the doctor hadn't even known the test existed. If you lost 36% of your blood in an accident, do you think your doctor would tell you that you are alright and should just take up line dancing and get over it? They would rush you to the nearest hospital and start transfusing you with blood. These tragic people with these awful diseases are functioning with anywhere from 7% to 50% less blood than their body needs to function.

VII – UNDOING THE DAMAGE

The body undoes the damage itself. The scarring in the brain of people with chronic fatigue and fibromyalgia will be repaired. There is cellular repair going on all the time. But the Mycoplasma has moved on to the next cell. In the early stages of a disease, doxycycline may reverse that disease process. It is one of the tetracycline antibiotics, but it is not bactericidal; it is bacteriostatic-it stops the growth of the Mycoplasma. And if the Mycoplasma growth can be stopped for long enough, then the immune system takes over.

Doxycycline treatment is discussed in a paper by Mycoplasma expert Professor Garth Nicholson, PhD, of the Institute for Molecular Medicine.[15] Dr. Nicholson is involved in a U.S. $8-million Mycoplasma research program funded by the U.S. military and headed by Dr. Charles Engel of the NIH. The program is studying Gulf War veterans, 450 of them, because there is evidence to suggest that Gulf War syndrome is another illness (or set of illnesses) caused by Mycoplasma.

Endnotes:

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4. Dr. Donald MacArthur, Pentagon, Department of Defense Appropriations for 1970, Hearings before Subcommittee of the Committee on Appropriations, House of Representatives, Ninety-First Congress, First Session, Monday June 9, 1969, pp 105-144, esp. pp. 114, 129.
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- Scott Donald W. and William L. C. Scott, The Extremely Unfortunate Skull Valley Incident The Chelmsford Publishers Canada 1996 (revised extended edition available from mid-September 2001; US$16.00 pre-pub. price + US$3 s&h in US).
- The Journal of Degenerative Diseases (Donald W. Scott Editor) The Common Cause Medical Research Foundation (Box 133 Stat B. Sudbury Ontario P3E 4N5) Canada (quarterly journal; annual subscription: US$25.00 in USA $30 foreign).

Additional Contacts:

- Ms Jennie Burke Australian Biologics Level 6 383 Pitt Street Sydney NSW 2000 Australia tel +61 (0)2 9283 0807 fax +61 (0)2 9283 0910. Australian Biologics does tests for mycoplasma.
- Consumer Health Organization of Canada 1220 Sheppard Avenue East #412 Toronto Ontario Canada M2K 2S5 tel +1 (416) 490 0986 website www.consumerhealth.org/.
- Professor Garth Nicholson PhD Institute for Molecular Medicine 15162 Triton Lane Huntington Beach CA 92649-1401 USA tel +1 (714) 903 2900.
- Dr. Les Simpson Red Blood Cell Research Ltd 31 Bath Street Dunedin 9001 New Zealand tel +64 (0)3 471 8540 email rbc.research.limited@xtra.co.nz. (Note: Dr. Simpson directs his study to red cell shape analysis not the mycoplasma hypothesis.)
- The Mycoplasma Registry for Gulf War Illness S. & L. Dudley 303 47th St J-10 San Diego CA 92102-5961 tel/fax +1 (619) 266 1116 fax (619) 266 1116 email mycoreg@juno.com.

About the Author:

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Donald Scott MA MSc is a retired high school teacher and university professor. He is also a veteran of WWII and was awarded the North Atlantic Star, the Burma Star with Clasp, the 1939-1945 Volunteer Service Medal, and the Victory Medal. He is currently President of The Common Cause Medical Research Foundation a not-for-profit organization devoted to research into neurosystemic degenerative diseases. He is also Adjunct Professor with the Institute for Molecular Medicine and he produces and edits the Journal of Degenerative Diseases. He has extensively researched neurosystemic degenerative diseases over the past five years and has authored many documents on the relationship between degenerative diseases and a pathogenic mycoplasma called Mycoplasma fermentans. His research is based upon solid government evidence.

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The simple primal act of getting LOTS of the high noon Sun on your bare "unprotected" skin in large amounts just may be the most important single thing you can do for your health. If the Sun casts a shadow longer than you are tall you cannot make adequate vitamin D from the Sun. You MUST take vitamin D3 at the daily rate of about 1000 IU/20 pounds of body weight.

I consider this the single best thing that I could have done to be still alive. I don't take quite the recommendation above, but I do take 4,000-6,000 IU of D3 daily. There are other nutrients and vitamins that I take that have proven successful in Lymphedema. Diet is also a major factor and going gluten-free has been a positive as well.

An important, very important thing to keep aware in protection against Mycoplasma is to know that Mycoplasma is monitoring three things, or is drawn to a weak area of the human biome. These are:

1. **Your pH level (ideal is 7 in balance) between alkaline or acidic**
2. **Your oxygen level in your blood**
3. **Your immunity system**

The pH range is 0 to 14, with seven being neutral. Acidic is 0 to 7, Alkaline is 7 to 14. There are abundant food category charts available that show whether the various foods are acidic or alkaline. The ideal is pH of 7 or neutral, meaning the body is in balance.

As the author in the above study reveals, Mycoplasma was weaponized during WWII, and through lab mistakes, failed precautions, and errors in experiments, the world's population has been infected with Mycoplasma. The "cover-up" in this case has become worse than the crime itself. Government and military sources have chosen to withhold from the public what is known concerning the harm of this pathogen.

Mycoplasma is the basic element (pathogen) of composting and decomposition. When you see a dead bird or animal on the road or a mouse on your sidewalk, you will note that sick odor as the smell of death. Do not touch, pick-up, or attempt to remove without EPA protective clothing such as gloves, plastic garbage bags, or a shovel to dispose of the dead animal.

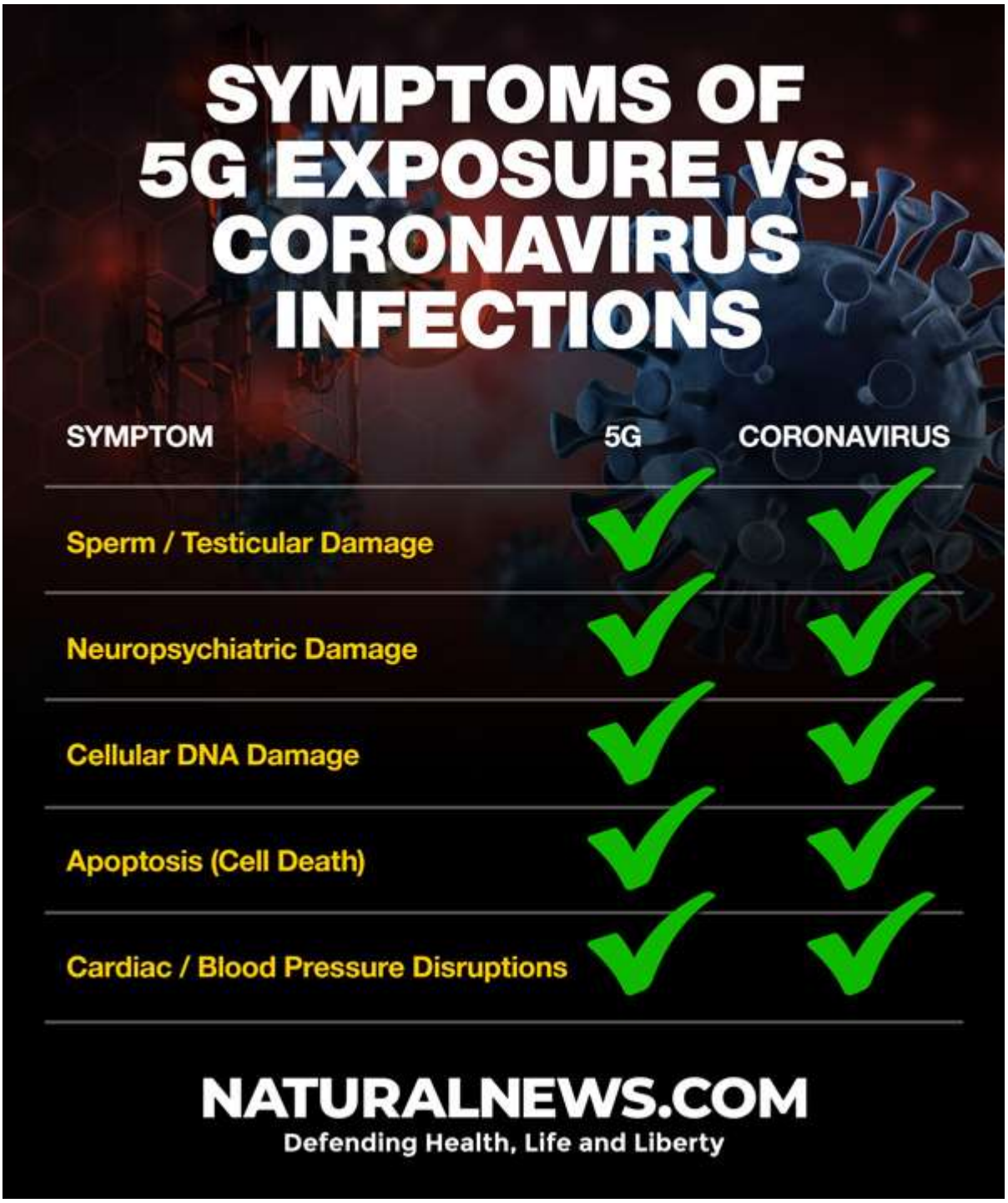
Now as to why these two elements are so critical: Mycoplasma and **5G**, are such a dangerous combination, is because they are across two fields of science. One is biological (Mycoplasma) and the second is electrical (**5G**). Dr. Martin Pall is a retired scientist from the University of Washington, cross-trained in two specialties of science, the academic fields of Electro-Biology, and medicine, one of the few leading experts in how **5G** impacts the human biome. Dr. Pall is perhaps the major nemesis of the telecommunications industry precisely because of his expertise in biology and electricity.

From the evidence we now know that **5G** can be electro-ported [electroporation]. This put into laymen terms means: the action or process of introducing DNA or chromosomes into bacteria or other cells using a pulse of electricity to briefly open the pores in the cell membranes. We have abundant evidence now that proves wireless energy, from 1G to 4G EMF radiation causes many issues to all life forms. The latency [delayed effects] of cell phone use has taken two decades to reveal its harm.

From a peak speed perspective, **5G** is 20X times faster than 4G. This means that during the time it took to download just one piece of data with 4G (like a movie); the same could have been downloaded 20 times over a **5G** network. Looking at it another way: you could download close to 10 movies before 4G could deliver even the first half of one!

5G has a minimum peak download speed of 20 Gbps while 4G sits at just 1 Gbps. These numbers refer to devices that aren't moving, like in a fixed wireless access (FWA) setup where there's a direct wireless connection between the cell tower and the user's device. Speeds vary once you start moving, like in a car or train. However, these aren't usually referred to as the "normal" speeds that devices experience, since there are often many factors that affect bandwidth. Instead, it's more important to look at the realistic speeds, or the average measured bandwidth.

There are many things that set **5G** apart from anything we have ever experienced in the way of communications, data transfer, but for our purpose here, the danger is in the millimeter wave band spectrum. The beam-wave signal is in millimeters, and not the typical 120-degree spectrum span of 4G. The standard cell-tower signal range is 120-degrees for up to 25 miles. If you have observed a teacher using a laser pen pointing that red laser dot on a blackboard, it is highly focused on a single point. With **5G** the EMF signal is 1,000 times more powerful than what we are using today, and will be able to send infinitely more signal as well. The intensity enables electroporation of the **5G** energy EMF into the human biome. The 5G beam can penetrate through the sweat gland pores and hair follicles and do great damage to the internal organs of the body.



In microbiology, the process of electroporation is often used to transform bacteria, yeast, or plant protoplasts by introducing new coding DNA. If bacteria and plasmids are mixed together, the plasmids can be transferred into the bacteria after electroporation, though depending on what are being transferred cell-penetrating peptides or CellSqueeze could also be used. Electroporation works by passing thousands of volts across a distance of one to two millimeters of suspended cells in an electroporation

cuvette (1.0 – 1.5 kV, 250 – 750 V/cm). Afterwards, the cells have to be handled carefully until they have had a chance to divide, producing new cells that contain reproduced plasmids. This process is approximately ten times more effective than chemical transformation.

Electroporation is also highly efficient for the introduction of foreign genes into tissue culture cells, especially mammalian cells. For example, it is used in the process of producing knockout mice, as well as in tumor treatment, gene therapy, and cell-based therapy. The process of introducing foreign DNA into eukaryotic cells is known as transfection. Electroporation is highly effective for transfecting cells in suspension using electroporation cuvettes. Electroporation has proven efficient for use on tissues in vivo, for in utero applications as well as in ovo transfection. Adherent cells can also be transfected using electroporation, providing researchers with an alternative to trypsinizing their cells prior to transfection. One downside to electroporation, however, is that after the process the gene expression of over 7,000 genes can be affected. This can cause problems in studies where gene expression has to be controlled to ensure accurate and precise results.

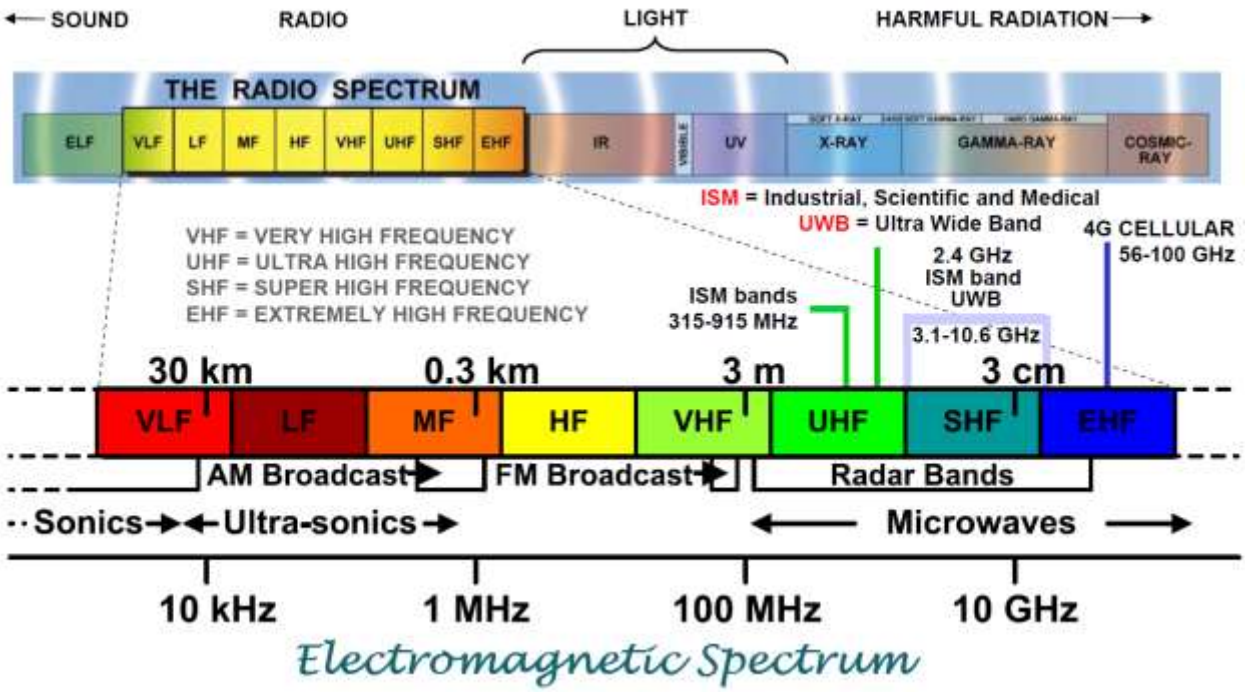
Since **5G** can introduce electroporation, it can do serious damage, invisible so that the damage done by the spectrum beaming, can and will activate the Mycoplasma to attack the cells of s human body, or any living object. Examples of known damage are noted on the chart above on page 16 &17.

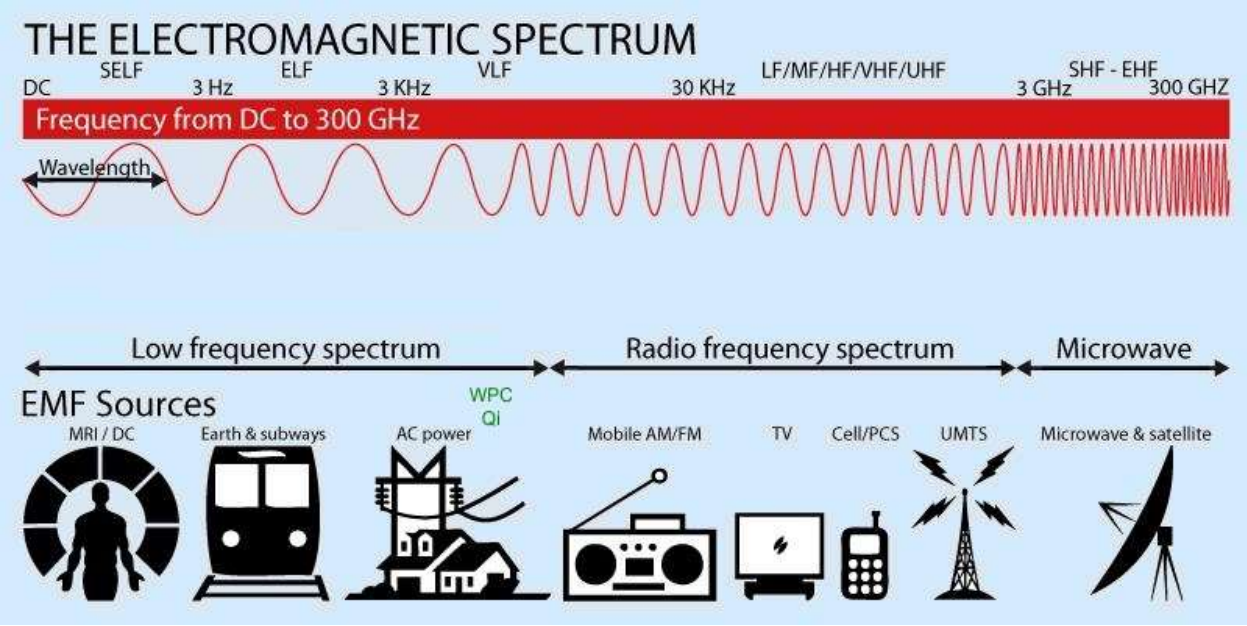
On the next two pages are three different light spectrum charts. No single light spectrum chart is capable of including all the different detail features and where various electronic devices fall in the larger scale of electronics, communications, devices, etc. The chart on page 17 notes ionizing and non-ionizing energy. Until recently, non-ionizing energy was considered in the safe range. However, recent studies have raised serious doubt about non-ionizing energy being safe. The regulations as to what qualifies was written in 1996, and with the expansion of what we call Broad-band, Wireless, Cellular, has presented serious issues that have yet to be fully studied and measured for damage or harm to the human body.

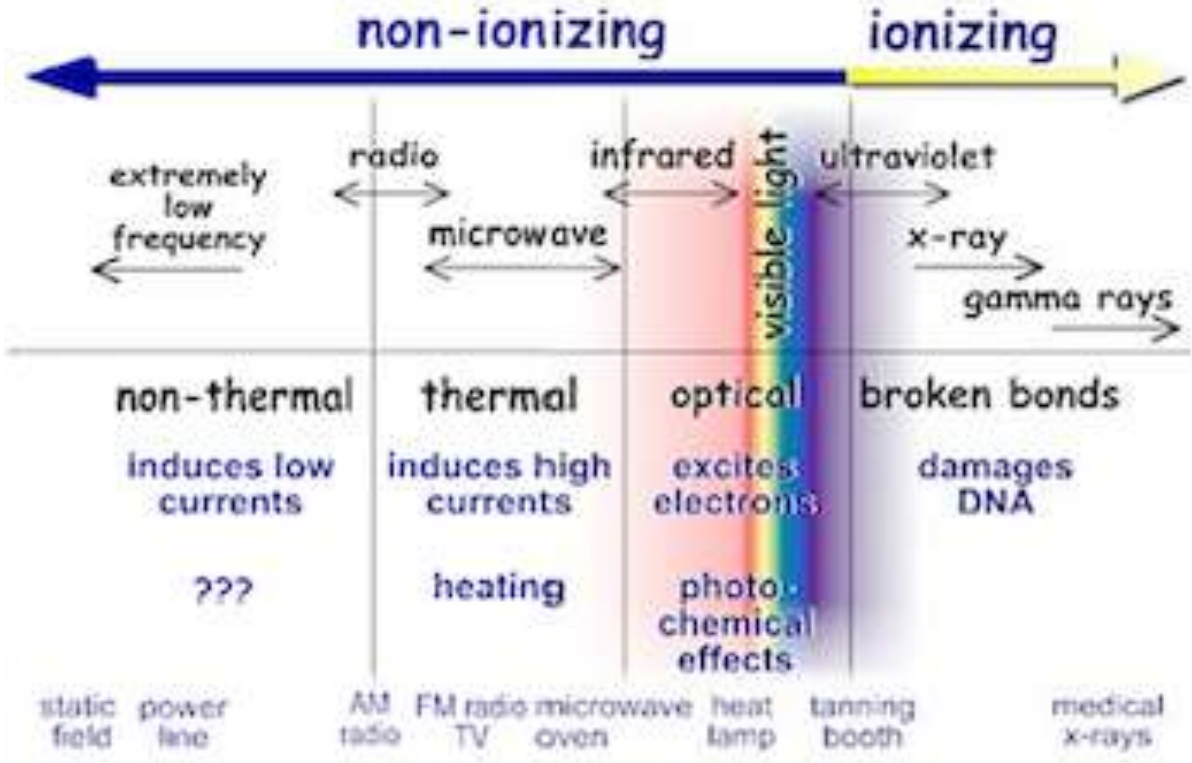
5G has certainly pushed the window limit for safe or harmless use. The very fact that the U.S. government and military have kept secret knowledge of Mycoplasma is troubling for this reason. They have the potential of using **5G** both as a “soft kill” or a “hard kill” weapon.

To this extent **5G** can be modulated in terms of energy level, an through electroporation, the Mycoplasma can be activated with as little as 10Ghz, that could lead to individuals unaware they have been infected by the existing Mycoplasma by **5G** remote electroporation. That individual may die, depending on the area where Mycoplasma attacks the body, in three days, three weeks, three months, or three years; and more important the time elapse from electroporation to death is likely to vary. A “soft kill” approach leads to deniability by the perpetrator with success in hiding the truth. The

"hard kill" approach would be to turn up the **5G** to 40GHz and higher that brings relatively quick death.






non-ionizing
ionizing
extremely low frequency
radio
microwave
infrared
visible light
ultraviolet
x-ray
gamma rays
non-thermal
thermal
optical
broken bonds
induces low currents
induces high currents
excites electrons
damages DNA
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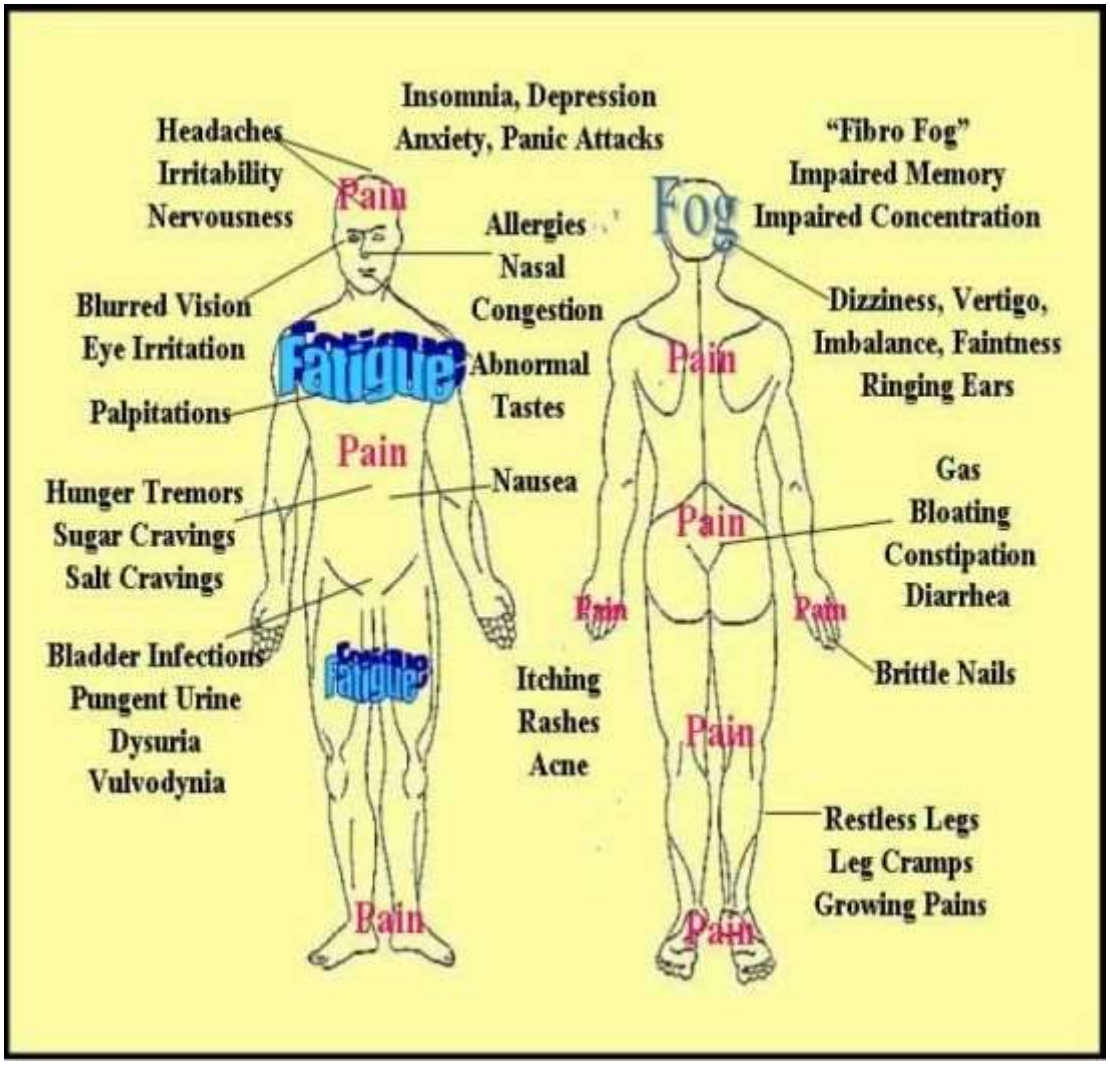
heating
photo-chemical effects
static field
power line
AM radio
FM radio TV
microwave oven
heat lamp
tanning booth
medical x-rays



The chart above lists many of the known physical issues that have been related to cellular Wi-Fi EMF radiation sickness. The hurdle for most people is over the issue of invisibility. What you can't see can hurt you whether you know it or not!

Mycoplasma Protection and Treatment Protocol

The simple primal act of getting LOTS of the high noon Sun on your bare "unprotected" skin in large amounts just may be the most important single thing you can do for your health. If the Sun casts a shadow longer than you are tall you cannot make adequate vitamin D from the Sun. You MUST take vitamin D^3 at the daily rate of about 1000 IU/20 pounds of body weight.

Very high body levels of Vitamin D^3 and sunshine seem in my opinion to be very protective against mycoplasma. Those pushing a vaccine for Covid-19 never promoted Vitamin C, $D^{3,}$ Elderberry, and Zinc as a natural treatment for Covi-19. Why not?

Simply put, it did not fit the narrative of their agenda. You never heard Dr. Fauci or his buddy Bill Gates discussing the answer to Covid-19 is Vitamin C, D^3, Elderberry, or Zinc. Their one-track minds have an objective that wants you dead!

"Mycoplasma is the co-factor that alters the human immune system and opens the door for the autoimmune degenerative diseases such as AIDS, Alzheimer's disease, Bi-Polar Disease, Creutzfeldt-Jakob disease, Chronic Fatigue/ Myalgic Encephalomyelitis, Diabetes Type One, Fibromyalgia, Huntington's disease, Multiple Sclerosis, Parkinson's disease." -(Doctor Harold Clark)

"According to Dr. Shyh-Ching Lo, senior researcher at The Armed Forces Institute of Pathology and one of America's top mycoplasma researchers, this disease agent causes many illnesses including AIDS, cancer, chronic fatigue syndrome, Crohn's colitis, Type I diabetes, multiple sclerosis, Parkinson's disease, Wegener's disease and collagen-vascular diseases such as rheumatoid arthritis and Alzheimer's. I have all the official documents to prove that mycoplasma is the disease agent in chronic fatigue syndrome/fibromyalgia as well as in AIDS, multiple sclerosis and many other illnesses." -(Donald W. Scott MA)

"Due to the misdirection of medical science compartmentalizing the human body into 10 separate specialty fields (dermatology, endocrinology, urology, neurology, psychology, oncology, gastric specialty, general practice etc.), like an auto mechanic would segregate engine parts, none of the mainstream physicians understand how all 10 body systems work synergistically as a whole like a flowing river. This has led medical science to perpetuate trash can labels to terms for symptoms of Mycoplasma to hide their ignorance." –(Gary Tunsky)

"Three years ago, several mishaps occurred with vials of viruses that could have been potential bioweapons in the wrong hands. A moratorium on deadly virus creation was installed, but the ban was just reversed on Tuesday allowing scientists to begin manipulating benign pathogens into deadly, epidemic-worthy human disease. Not only do the scientists want permission to create with abandon, they'd like to keep the results secret – i.e., keep them unpublished so that the work doesn't get into "the wrong hands" and cause an act of Bioterror … Now that the government has lifted the ban, scientists essentially have carte blanche to breed laboratory horrors." Scientists Get Approval to Create Air Transmissible Ebola and Other Lethal Viruses

If ever there was any doubt about how this DEADLY GAME is being played this should REMOVE IT. WE CANNOT TRUST THE GOVERNMENT AGENCIES FOR PUBLIC HEALTH SINCE THEY HAVE KEPT SECRET THE INFORMATION SHARED IN THIS ARTICLE. The kept it Top Secret for near 50 years, the public, the government employees that contracted it, not to mention the unknown number they treated as lab rats in a cage.

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Below the reader will find an accumulation of professional research, studies, opinions, and therapy known to help those with health issues that track back to Mycoplasma. If you know of anyone with serious health issues, they need to learn more about how this pathogen has eluded many in the medical community.

“Because certain species of the Mycoplasma have an absolute growth requirement for the up-take of pre-formed sterols, including cholesterol they can cause the ‘spontaneous degeneration’ of the cells that they invade. If they do not cause sufficient damage to kill the cell, they at least compromise its capacity to defend itself from other disease agents, such as those which present as Kaposi’s sarcoma, pneumoniae carinii pneumonia, lymphadenopathy, and so on.” Donald W. Scott and William L.C. Scott, Note here it is the key to understanding AIDS

“From its inception, the biowarfare program was characterized by continuing in-depth review and participation by the most eminent scientists, medical consultants, industrial experts and government officials, and it was classified Top Secret. The US Public Health Service also closely followed the progress of biological warfare research and development from the very start of the program, and the Centers for Disease Control (CDC) and the National Institutes of Health (NIH) in the United States were working with the military in weaponizing these diseases. These are diseases that have existed for thousands of years, but they have been weaponized—which means they’ve been made more contagious and more effective. And they are spreading.” -(Donald W. Scott)

“Doctor David Williams cites a recent study in the USA in which 28 people suffering from severe Lyme Disease were treated for 6 months. 14 of the patients received the standard doses of antibiotics. The other 14 took doses of [the Samento form of] Cat's Claw, an herb from the Brazilian rainforest with a legendary reputation. The result? Three of the fourteen who took antibiotics showed slight improvement. But ALL of the 14 who took Cat's Claw showed "dramatic improvement." Indeed, 12 of the 14 were tested at the end of six months, and NO trace of LD remained!” Note Lyme disease is often a symptom of a mycoplasma infection

"A study to examine the effect of undenatured bioactive whey on increasing cellular glutathione levels in CFS patients was run in the clinic of Doctor Paul Cheney. The clinic conducted a six-month study of the first patented bioactive whey product (Immunocal), and discovered it significantly improved glutathione function. Though it was a small study (eight patients), the results were consistent with the feedback from Cheney's patient population as a whole. Each patient who tested positive for mycoplasma and chlamydia pneumoniae at the beginning of the study tested negative six months after treatment. Doctor Cheney found undenatured whey protein was the best way he explored to increase glutathione levels and function." Note here we see high glutathione appears to be a strong preventative measure against all the mycoplasma driven diseases including AIDS.

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conducted a six-month study of the first patented bioactive whey product (Immunocal), and discovered it significantly improved glutathione function. Though it was a small study (eight patients), the results were consistent with the feedback from Cheney's patient population as a whole. Each patient who tested positive for mycoplasma and chlamydia pneumoniae at the beginning of the study tested negative six months after treatment. Doctor Cheney found undenatured whey protein was the best way he explored to increase glutathione levels and function.

"The MRFIT study included more than 300,000 young and middle-aged men.16 years after the first cholesterol analysis the number of men whose cholesterol was lower than 160 and who had died from AIDS was four times higher than the number of men who had died from AIDS with a cholesterol above 240." –(Neaton et al, AIDS 11, 929–930, 1997)

"Mycoplasma can cause a respiratory flu-like illness that can progress to systemic chronic fatigue syndrome-like or fibromyalgia syndrome-like illness, sometimes advancing to multiple sclerosis-like amyotrophic lateral sclerosis and arthritic-like symptoms." –(Life Extension)

"Thousands of soldiers with Gulf War Syndrome are being helped when their mycoplasma infections are identified and killed." -(Nicolson et al 1998, Note it has been established GWS was caused by the "vaccinations" the soldiers received which apparently contained mycoplasma. Support our troops)

"Auto-immune doctors are finding organisms in patients that appear to be part virus, part bacteria and part fungus." -(William Thomas Medical Researcher, Note these are obviously bio-lab creations)

"It is my contention that the disease we now call Lyme Disease, originated in a lab at Plum Island. The funding for research would be covered in the Mycoplasma grant. The very first case of Lyme Disease was isolated in a youth, in 1975, within a few miles of the dock where the Plum Island Government Ferry boat lands on the Connecticut side of Long Island Sound." -(Patricia Doyle)

"One unique feature of M. fermentans incognitus is its ability to catabolize glucose both aerobically and anaerobically and also to hydrolyze arginine. M. fermentans incognitus cannot hydrolyze urea in a biochemical assay. When grown in culture, M. fermentans incognitus produces a prominent alkaline shift in pH after an initial brief acidic shift. The only other human mycoplasma which is known to metabolize both glucose and arginine is the rarely isolated M. fermentans." -(U.S. Army Mycoplasma Fermentans Incognitus Patent)

IMO Pathogenic Mycoplasma is a Bio-Lab Created Problem

"Sometime over the past 30 years, the organism has been altered to become more lethal. The Mycoplasmas found by the Nicolson's, in their lab, contain unusual gene

sequences that were probably inserted into the Mycoplasma by a specific laboratory procedure. This discovery has led them to conclude that the new forms of mycoplasma were specifically engineered for germ warfare. (9) In its laboratory evolution, the Mycoplasmas have become more invasive, more difficult to find, and capable of causing severe diseases in humans. Diseases, like Gulf War Illness, CFS, FMS, MCS, Rheumatoid Arthritis, and AIDS, for instance." -(Mycoplasma made in a lab)

"Dr. Lo has been credited with discovering the new pathogenic form of Mycoplasmas, and he currently holds several patents on methods for special handling of the organisms for study and development. (10) In one of his patents (in 1991), Dr. Lo lists the following diseases that are caused by Mycoplasma: HIV infection, AIDS, Aids Related Complex (ARC), Chronic Fatigue Syndrome, Wegener's Disease, Sarcoidosis, Respiratory Distress Syndrome, Kibuchi's Disease, Alzheimer's Disease, and Lupus. (10) In addition, Baseman and Tully have reviewed the literature on the role of Mycoplasmal infections in human disease and have concluded that they are important factors or co-factors in a variety of chronic illnesses." -(Mycoplasma made in a lab)

Mycoplasma is often described as a virus without its outer shell or a bacterium without a cell wall. Mycoplasma in many forms has existed in nature for centuries. It was once a fairly rare problem in humans because our bodies have many ways to defeat small natural amounts of Mycoplasma.

Pathogenic man-made Mycoplasma is the guts of a perniciousness designed to kill. Mycoplasma appears to be a bio-weapon created as part of the AIDS project. It is able to burrow into almost every recess of our bodies including, brain, and bones within individual cells. I believe many of us are exposed to low levels of this often weaponized disease vector more often than we might think. IMO "vaccination" may be the chief transfer vector for weaponized mycoplasma.

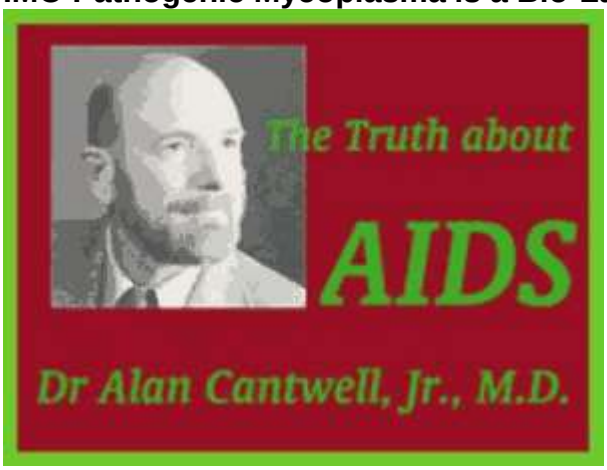
Mycoplasma Symptoms

- Chronic Fatigue **note CFS sufferers**
- Low Body Temperature
- Recurring Flu like illness **one after the other without end**

If you are having any mycoplasma associated disease symptoms and taking any "Cholesterol Lowering Substances" IMO stop them as soon as possible. If your doctor refuses to help you get off the drugs IMO tell him you are going off the drugs with or without his help. In any case make sure you are eating plenty of cholesterol food sources.

Luc Montagnier head of the Department of Virology at the Pasteur Institute and civilian discoverer of the HIV virus believes mycoplasmas are a necessary "co-factor" that allows progression to full-blown AIDS (See AIDS Protection Protocol). Mycoplasma can be communicable.

IMO Pathogenic Mycoplasma is a Bio-Lab Created Problem



"Researchers found that if they had mycoplasma at certain strength—actually, 10 to the 10th power—it would develop into AIDS, and the person would die from it within a reasonable period of time because it could bypass the natural human defenses. If the strength was 10 to 8, the person would manifest with chronic fatigue syndrome or fibromyalgia. If it was l0 to 7, they would present as wasting; they wouldn't die and they wouldn't be disabled, but they would not be very interested in life; they would waste away." **http://www.whale.to/m/scott7.html**

"A high prevalence of M. fermentans incognitus infection has been found in patients with AIDS by using the polymerase chain reaction. The genetic material specific for M. fermentans incognitus has been isolated from spleens, Kaposi's sarcoma, livers, lymph nodes, peripheral blood mononuclear cells and brains of patients with AIDS." **US Army Mycoplasma Fermentans Incognitus Patent**

"Mycoplasma fermentans (incognitus) has been tested on the Texas Department of Corrections prisoners in the late 1980s prior to the Gulf War. It was tested on death row inmates as well as other inmates in Huntsville, Texas. The guards then contracted it from the inmates, and the guards then gave it to their families and community. This mycoplasma vaccine testing was funded by the U.S. Army, and today there is an outbreak of 350 people in the Huntsville area with a strange disease resembling GWS." **Doctor Nicolson**

Clearly weaponized mycoplasma is a KEY bio-weapon of our time. It is involved in MANY "diseases" that have popped up like ripe mushrooms since 1970 or so. Please research this key topic until you understand what has been created.
http://www.whale.to/m/mycoplasma5.html

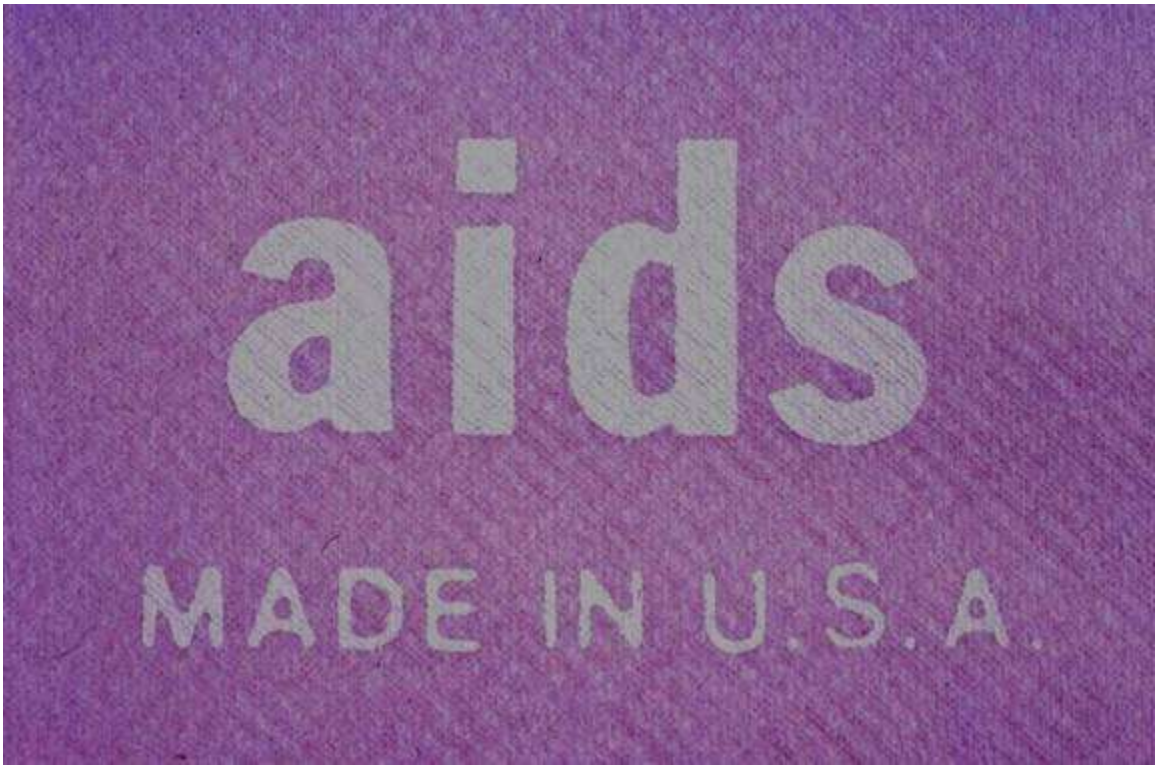
Diseases Associated with Weaponized Mycoplasma

- AIDS
- Alzheimer's disease
- Bi-Polar Disease
- Cancer
- Collagen-vascular Diseases

Creutzfeldt-Jakob disease
Chronic Fatigue/ Myalgic Encephalomyelitis
Crohn's Colitis
Diabetes Type One
Fibromyalgia,
Huntington's disease
Lime Disease
Multiple Sclerosis
Parkinson's disease
Rheumatoid Arthritis
Wegener's disease

What is Disease?

Weaponized Mycoplasma born in a lab
"Some call it AIDS – I call it MURDER!" **Doctor Eva Snead MD**



"Researchers Dr. Garth Nicolson and his wife Nancy have found a tiny bacterial microbe (a "mycoplasma") in the blood of nearly half the ill vets with GWI. Amazingly, this infectious agent has a piece of HIV (the AIDS virus) attached to it. This microbe could never have occurred naturally. On the contrary, the composition of the microbe suggests a man-made and genetically-engineered biological warfare agent." **Doctor Cantwell MD** **Mycoplasma made in a lab**

"Before AIDS, Kaposi's sarcoma was never seen in healthy young men. Identified a decade after HIV, in 1994, this KS virus is closely related to a primate cancer-causing herpes virus extensively studied and transferred in animal laboratories in the decade before AIDS. Also downplayed to the public is a new microbe (Mycoplasma penetrans), also of unknown origin, that was introduced into homosexuals, along with HIV and the new herpes virus. Thus, not one but three new infectious agents were inexplicably transferred into the gay population at the start of the epidemic (HIV, the herpes KS virus, and M.penetrans)." **--Alan Cantwell MD (Mycoplasma made in a lab)**

"Mycoplasmas are the smallest free-living bacteria, that can assume multiple shapes including round, pear shaped and even filamentous because they lack a cell wall. This makes it possible for them to pass through some filters used to remove bacteria; thus, unaffected by many common antibiotics. Because mycoplasmas have lost most of their genetic material; a strict dependence on the host for nutrients and refuge determines its ability to survive and grow." **Mycoplasma and AIDS, Degenerative Disease**

Once very uncommon in humans one or more strains Mycoplasma was successfully weaponized by one of our bio-weapons programs **The Special Virus Cancer Program** and now seems to be in everyone. As a biological weapon mycoplasma may be delivered as an aerosol airborne agent, via insects and in other ways. Weaponized mycoplasma may be concentrated billions of times over that found in nature. The taxpayers of the U.S. have inadvertently and indirectly at least in part given the world this scourge.

In 1970, just one year after President Nixon officially ended our offensive biowar research, Plum Island was granted 10 million dollars by the Federal Govt. The purpose of the grant was to research the use of mycoplasma for use in germ warfare. The original $10 million dollar grant was for a 5 year period. In 1975, the funding was continued as the Mycoplasma research was deemed very successful. IMO this was not research as we have seen most of the research into Mycoplasma was done in the 1950s and 1960s. This grant probably was involved with the AIDS project.

US Army Mycoplasma Fermentans Incognitus Patent

The **United States Armed Forces Institute of Pathology** holds a patent on weaponized crystallized Mycoplasma. In fairness to them they claim to have started work on this project in the late 1980s after the AIDS outbreak.

Mycoplasma is associated with MANY Diseases

"Mycoplasma is the co-factor that alters the human immune system and opens the door for the autoimmune degenerative diseases such as AIDS, Alzheimer's disease, Bi-Polar Disease, Creutzfeldt-Jakob disease, Chronic Fatigue/ Myalgic Encephalomyelitis, Diabetes Type One, Fibromyalgia, Huntington's disease, Multiple Sclerosis, Parkinson's disease." **Doctor Harold Clark**

Take a look at ALL these diseases which are associated with mycoplasma. You can use the Treatments here, especially Vitamin D, to Treat all these diseases. Please do it.

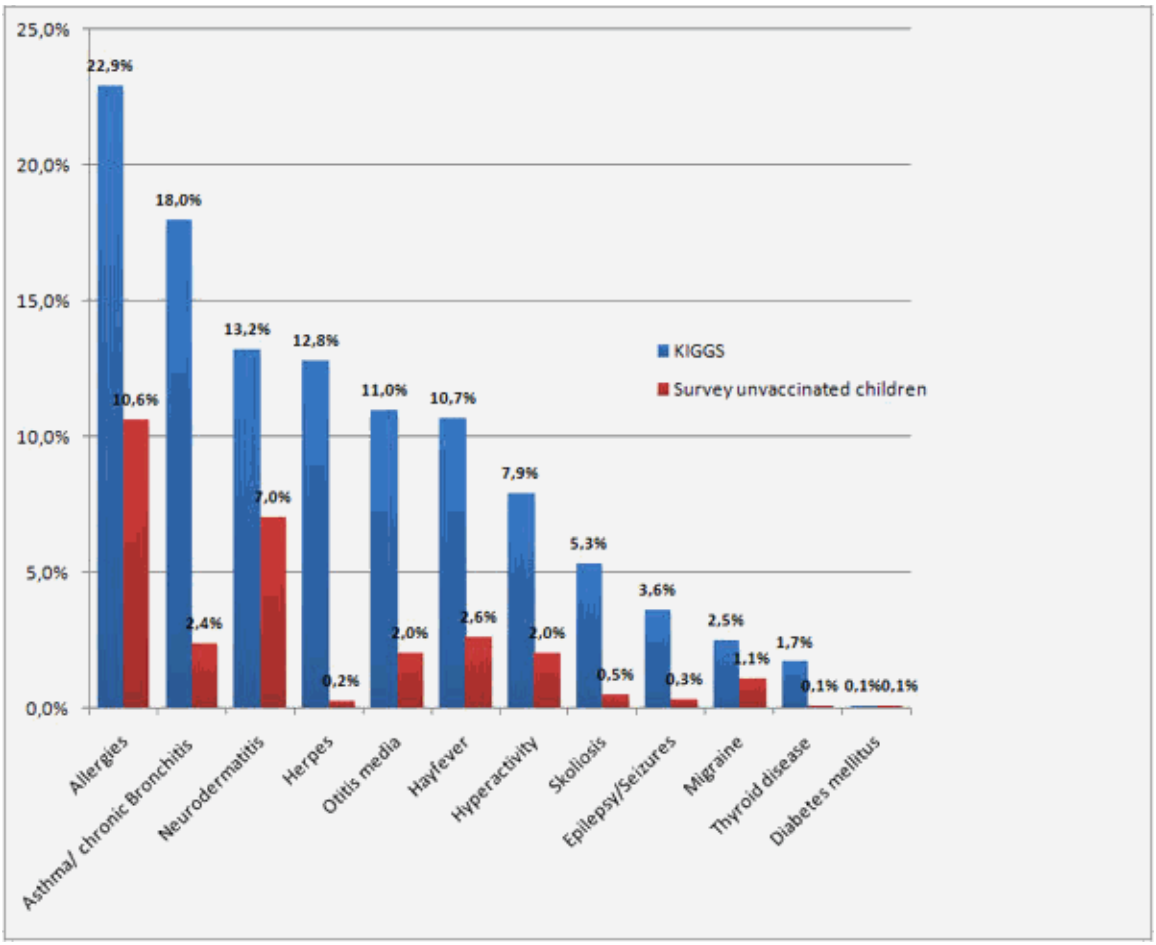
Vaccination and Autoimmune Disease

"Circulating Immune Complexes are the antigen (bad guy) plus antibody (policeman) pairs that clog up the bloodstream, can plug the kidneys and even implant in tissue so that the body recognizes itself as being contaminated and attacks itself. That last one is the very definition of autoimmunity. They call it something different in medical texts to cover up the fact that they caused it with vaccines and don't want anyone to know about it." **Patrick Jordan, Note allergies usually proceed autoimmune disease, Jordan shows us the cascade of disease initiated by vaccination and ending in autoimmune disease or death.**

If you have an autoimmune disease or do not wish to have an autoimmune disease Please don't consider the an autoimmune disease producing practice of vaccination. There is little question that vaccination is a cause of many allergies and autoimmune diseases including asthma. Patrick Jordan explains how autoimmune diseases including asthma often follow the barbaric practice of "vaccination".

Listen to Doctor Nicolson tell us why We Dare not "Vaccinate"

Doctor Garth Nicholson PhD "Vaccination" A Cause of MANY Chronic Degenerative Diseases. EXPLOSIVE! Doctor Nicholson gives us a TREATMENT for these diseases.



Please REJECT "Vaccination"
Please investigate the practice of "vaccination" and IMO reject it as a deadly scam. IMO this is possibly the best gift you can give your immune system; or more importantly the immune systems of your children and grandchildren. There is little question that vaccination is a cause of many allergies and autoimmune diseases.

Doctor Moulden, Patrick Jordan and others explain how ischemia, chronic inflammation and allergies often follow the barbaric practice of "vaccination" in a deadly cascade. After "vaccination" ischemia, chronic inflammation, leaky gut and allergy often is induced often followed in turn by delayed hypersensitivity, serum sickness and finally autoimmune disease including asthma. Of course autoimmune disease as bad as it is often not the WORST effect of "vaccination", there is autism and death.

"Vitamin D deficiency is now recognized as a pandemic. The major cause of vitamin D deficiency is the lack of appreciation that sun exposure in moderation is the major source of vitamin D for most humans. Very few foods naturally contain vitamin D, and foods that are fortified with vitamin D are often inadequate to satisfy either a child's or an adult's vitamin D requirement. Vitamin D deficiency

causes rickets in children and will precipitate and exacerbate Osteopenia, osteoporosis, and fractures in adults. Vitamin D deficiency has been associated with increased risk of common cancers, autoimmune diseases, hypertension, and infectious diseases." **Vitamin D deficiency: a worldwide problem with health consequences**

"When I studied Lyme Disease in 1997 I found the same antibiotic treatment was used to treat Lyme as is used to treat mycoplasmal infections. My son was diagnosed with parvovirus B19 not Lyme Disease, but he responded to the antibiotic. This didn't make sense until I found that ticks carry mycoplasmas." **Candace Brown, Mycoplasma made in a lab**

"Those of us with chronic Lyme disease are quite familiar with the names of the better known Lyme coinfections. Babesia, Bartonella, and Ehrlichia have become everyday words. As much as we would like to rid ourselves of these illness producing pathogens, they have become a part of our daily struggle to regain a sense of health and wellness. Unfortunately, these are not the only co-infections seen in chronic Lyme disease. For some reason, Mycoplasma infections are not only lesser known by patients, but seemingly often overlooked by doctors as well. It is important for us, as patients, to educate ourselves on the topic of Mycoplasma and to ask our practitioners how we are being evaluated and treated for these infections." **Scott Forsgren**

Some 70% or so of the strange recent diseases like AIDS, MS, Fibromyalgia, Chronic Fatigue Syndrome, Lyme Disease and other strange maladies like Gulf War Syndrome and even oldies such as Rheumatoid Arthritis test positive for Mycoplasma. Mycoplasma weakens your immune system allowing simple usually harmless infections as well as AIDS retroviruses to take hold. The method Mycoplasma uses to weaken your immune system is to exhaust your cholesterol stores leading to endocrine disruption. Mycoplasma has a very serious adverse effect upon your liver due to its voracious up-take of liver-produced cholesterol, In the end your liver fails to produce enough cholesterol. The cells themselves will then produce their own cholesterol. which may satisfy the cell's membrane cholesterol needs but not your endocrine needs?
Mycoplasma and vitamin D in RA

MANY Major Diseases are Associated with Mycoplasma
1) AIDS
2) Lyme Disease
3) Cancer

To see the evidence linking RA and MANY other diseases with Weaponized Mycoplasma go here

Please don't forget both mycoplasma and the Disease can be delivered in "vaccines" and Vitamin D appears to be HIGHLY protective against mycoplasma and many other diseases.

"Scientists have discovered that vitamin D effectively blocks development of MS in animals. When the biologically active, hormone form of vitamin D was administered to animals in a laboratory, the disorder was prevented. Conversely, a deficiency of vitamin D tended to increase the animals' susceptibility to the induced disease. When animals were given vitamin D after developing the disease, progression of symptoms was blocked. When vitamin D supplementation was withdrawn, the disease resumed." **Cantorna et al 1996**

Vitamin D3 and curcumin work together to clear deadly brain plaques
Vitamin D deficiency has been mistaken for fibromyalgia, chronic fatigue or peripheral neuropathy.

"Activated vitamin D in the adrenal gland regulates tyrosine hydroxylase, the rate limiting enzyme necessary for the production of dopamine, epinephrine and norepinephrine. Low vitamin D may contribute to chronic fatigue and depression." **-(Puchacz et al)**

"We now know that multiple sclerosis is almost certainly caused by insufficient sunlight and/or insufficient vitamin D, probably combined with a brief virus infection." **Good Vitamin D Treatise, Note sunlight and vitamin D seem to be highly protective of mycoplasma**
Mycoplasma and vitamin D in RA

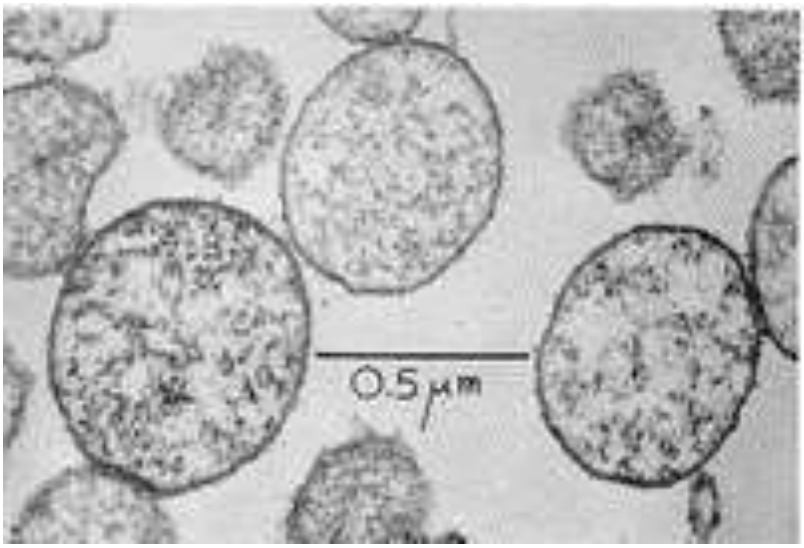
Autism and Mycoplasma
"It is most commonly claimed by both patients and doctors that the ASD is indeed associated with multiple **minor infections in young children. They notice an increase in the amount of antibiotics that are being used and the number of times that they have to take the child to the doctor for some investigation." Mycoplasma Bovis as a Mycoplasma Model for study**

"Usually Mycoplasma bovis is a secondary infection in that it typically needs another bacterial infection to lower the immune system of the calf to have effective growth (Cree 2002)." **Mycoplasma Bovis as a Mycoplasma Model for study, Note I found this paper on Mycoplasma Bovis VERY instructive**

Infections associated with Autism
As we see in the paper above there is plenty of evidence that an interplay between mycoplasma the standard medical mis-treatment of our, vaccines, antibiotics, our children and ASD.

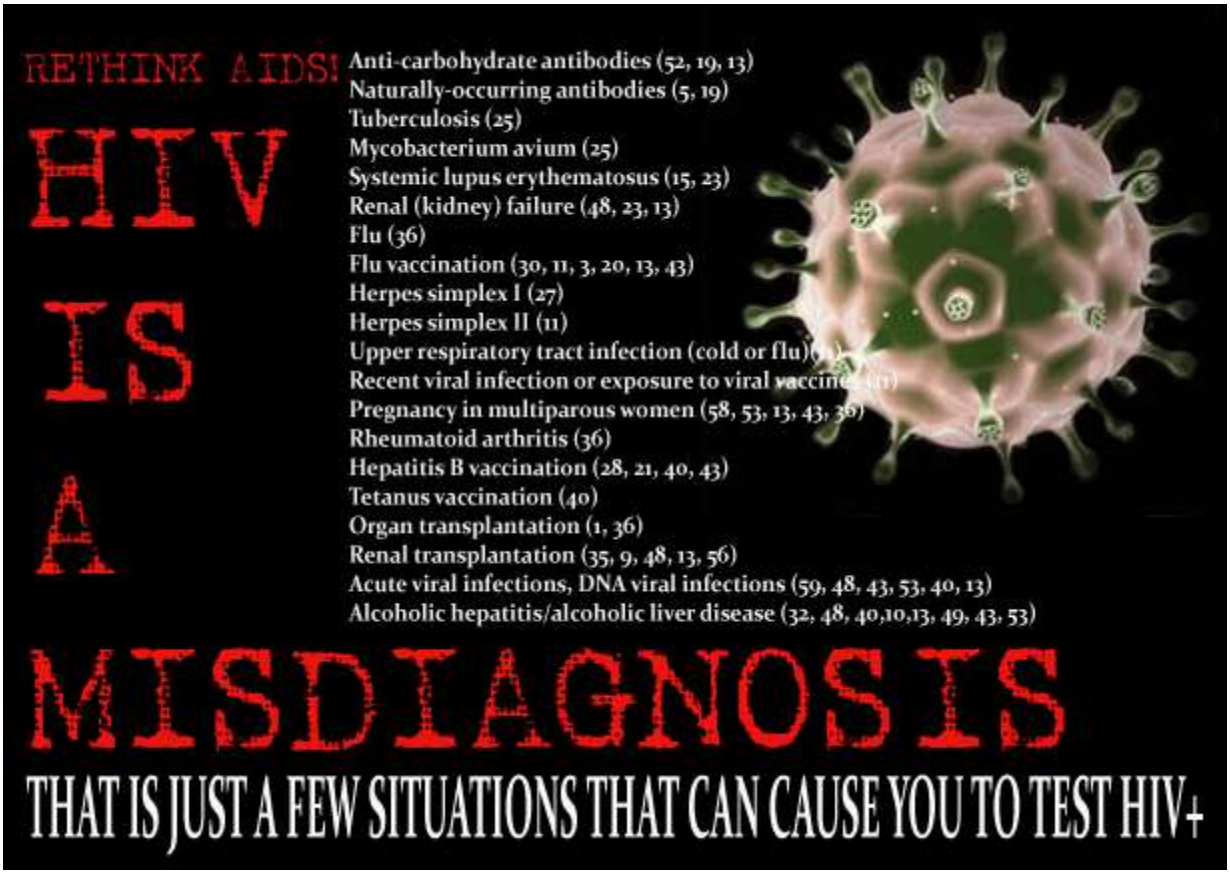
So now it appears that something besides mercury, aluminum and all the other toxins injected into our children may be in those "vaccination" needles. Of

course I don't not KNOW this but considering that "vaccination" confers FAR more risk than benefit it is certainly yet another reason to STOP VACCINATING. With a case of autism exhibiting the mycoplasma symptoms you may want to be tested for mycoplasma. This is not easy.



There is now a simple blood test for Mycoplasma
If you have any of the autoimmune like diseases that seem intractable to solution please get the test for mycoplasma.
TESTING FOR MYCOPLASMA IN YOUR BODY

IMO AIDS is largely a Man Made problem using Mycoplasma, it is NOT a natural disease



http://www.hivmisdiagnosis.com/

One of the first successful alternative treatments for mycoplasmas

Continuation of Mycoplasma Protection and Treatment Protocol (See Mycoplasma Protection and Treatment Protocol Continuation)

Five years ago when I began researching Lymphedema and Neuropathy, I searched the Internet day and night, for weeks and months. The Mayo Clinic, along with many others said, there is no known cause and there is no cure. That kind of answer was not an answer that I could or would accept. My research into **5G** technologies has been ongoing since 1996, by reading and study of bee colony collapse about bees dying from an unknown cause [really EMF radiation poisoning] and so I continued exploring, asking questions of anyone who would answer me back.

I am grateful for the work of Dr. Arthur Firstenberg, and Dr. Tom Cowen. I had read Dr. Leonard Horowitz's book *'Emerging Viruses: Aids and Ebola'* in 1997. I have read Dr. Nancy Mikovits recent books on vaccines and pandemic fraud in the vaccine industry. Dr. Andy Kaufman and others have taught me so much about biology, immunology, molecular biology, biochemistry. The experts in Radar, Electrophysics, and Electrical Engineering like Mark Stein, and Dr. Barry Trower, along with Dr. Martin Pall, emeritus scholar from the University of Washington and expert in the rare field of Electro-biology. The most recent reading which has enabled me to fine-tune my self-medication being

'Lymphedema and Lipedema Nutrition Guide: foods, vitamins, minerals, and supplements' by Chuck Ehrlich and team. Their work applied to my diet in a single week showed significant relief. Into my third week I am experiencing improvements that confirm diet and nutrition should not be overlooked in diagnosing health issues.

I am not at all reticent about saying that in all of this I give God the Glory for His direction in this work to get truth. ***"Thy word is a lamp unto my feet, and a light unto my path"*** –(Psalm 119:105. I have sensed being closer to my Lord, in ways that I could not fully explain to folks. Many a night when I went to bed in extreme pain, I felt an assurance that the Lord would direct me to important clues in my research and provide answers that at times were not always clear. I never faltered for a moment, always sensing the prodding and direction He wanted me to explore or follow. Almost daily, I would awake with such sharp clarity and intuitive insights to explore. For those of you who do not understand this, God frequently spoke to me intuitively in blazing fashion at times focusing my attention on something so profound and insightful. This may not mean anything, and yet it has seemed to me that Mycoplasma has a relationship with what the Bible calls sin. The first time we find the word "sin" in the Bible is Genesis 4:7

Cain still had an opportunity to do what was good and right. If he did God would accept him. But if he did not do what was good and right, sin was crouching at the door like a wild beast ready to spring. It was "at the door" because Cain's attitude brought him very close to sin. Sin was desiring or striving to get at Cain. But he could still reject the desire of sin and rule or take control over it. God was thus calling on Cain to repent of his wrong attitude and exercise self-discipline. But Cain would not worship God when He did not favor him. Cain put a higher priority on himself than on God. I am humbled by the fact that God has taught me a lot in this work that magnifies His Glory!

Blessings,

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