Evidence that the "H1N1 Flu" vaccine is toxic, especially the adjuvant Squalene

The willful use of toxic substances known to cause injury and death in a vaccine or injection is prohibited. The prohibition of such use is a universally accepted as a part of national and international law.

There is scientific evidence that the swine flu vaccine, especially the adjuvants, specifically squalene, cause serious injury and death.

Adjuvants have not been approved by the FDA for use in the USA .

According to Dr. Meryl Nass, there are only three vaccines in existence using an approved squalene adjuvant. None of the three are approved for use in the U. S. Novartis's proprietary squalene adjuvant for their H1N1 vaccine is MF59. Glaxo's is ASO3. MF59 has yet to be approved by the FDA for use in any U. S. vaccine, despite its history of use in other countries.

A 2000 study published in the American Journal of Pathology demonstrated a single injection of the adjuvant squalene into rats triggered "chronic, immune-mediated joint-specific inflammation," also known as rheumatoid arthritis.

Specific evidence of the dangers of the swine flu vaccine with adjuvants, especially squalene, include

- The UK Health Protection Agency warnied that the "swine flu" vaccine could cause neurological damage in a letter leaked to The Mail on Sunday, a UK newspaper, which was not made public.

Dr Mercola explains: "Gulf War veterans with Gulf War Syndrome (GWS) received anthrax vaccines which contained squalene. [ix] MF59 (the Novartis squalene adjuvant) was an unapproved ingredient in experimental anthrax vaccines and has since been linked to the devastating autoimmune diseases suffered by countless Gulf War vets. [x]

The Department of Defense made every attempt to deny that squalene was indeed an added contaminant in the anthrax vaccine administered to Persian Gulf war military personnel – deployed and non-deployed – as well as participants in the more recent Anthrax Vaccine Immunization Program (AVIP).

However, the FDA discovered the presence of squalene in certain lots of AVIP product. A test was developed to detect anti-squalene antibodies in GWS patients, and a clear link was established between the contaminated product and all the GWS sufferers who had been injected with the vaccine containing squalene.

A study conducted at Tulane Medical School and published in the February 2000 issue of Experimental Molecular Pathology included these stunning statistics:

" … the substantial majority (95%) of overtly ill deployed GWS patients had antibodies to squalene. All (100%) GWS patients immunized for service in Desert Shield/Desert Storm who did not deploy, but had the same signs and symptoms as those who did deploy, had antibodies to squalene.

In contrast, none (0%) of the deployed Persian Gulf veterans not showing signs and symptoms of GWS have antibodies to squalene. Neither patients with idiopathic autoimmune disease nor healthy controls had detectable serum antibodies to squalene. The majority of symptomatic GWS patients had serum antibodies to squalene."[xi]

- Adjuvants are used in vaccines designed to impair fertility in animals according to patents.

WO/1999/034825) FERTILITY IMPAIRING VACCINE AND METHOD OF USE This application claims the benefit of U. S. Provisional Application No. 60/070,375, filed January 2,1998, U. S. Provisional Application No. 60/071,406, filed January 15,1998 “The vaccine of the invention preferably additionally includes an immunological adjuvant to enhance the immunological response of the subject to the glycoprotein antigen. Examples of adjuvants include Freund’s Complete Adjuvant, Freund’s Incomplete Adjuvant, and an adjuvant comprising an immunostimulant such as synthetic trehalose dicorynomycolate (STDCM) and an oil such as squalene oil (see P. Willis et al., J. Equine Vet. Sci., 14,364-370 (1994)). An adjuvant comprising synthetic trehalose dicorynemycolate, squalene oil, and a surfactant such as lecithin is preferred. Lecithin typically includes phosphatidyl choline. In a preferred embodiment the vaccine comprises oil, preferably a biodegradable oil such as squalene oil. Typically, the vaccine is prepared using an adjuvant concentrate which contains lecithin in squalene oil. The aqueous solution glycoprotein is typically a phosphate-buffered saline (PBS) solution, and additionally preferably contains Tween 80.”

- There is clear evidence all vaccinations are bad and that vaccine companies have been involved in concealing the dangers from us by systematically manipulating data. Dr Rebecca carley, court qualified expert on vaccines, has called innoculations, the true weapons of mass destruction.

In spite of the scientific evidence that vaccines, especially those with adjuvants, particularly squalene, are dangerous, the WHO prescribed by virtue of the authority vested in it by its Constitution and the International Health Regulations 2005 that vaccinations against the swine flu must contain adjuvants.

Scientific evidence offers proof that adjuvants such as squalene are dangerous just as there is substantial proof a loaded gun will be dangerous when fired at you without getting hold of the specific bullets and exminaing them in a lab - and yet the WHO choses to include that dangerous substance even though vaccine company's own published clinical trials show that adjuvants are deletrious.

Reports in full length

- The report in the Mail on Sunday: Swine flu jab link to killer nerve disease: Leaked letter reveals concern of neurologists over 25 deaths in America

By Jo Macfarlane Last updated at 11:05 PM on 15th August 2009

Prevention: Is the swine flu jab safe?

A warning that the new swine flu jab is linked to a deadly nerve disease has been sent by the Government to senior neurologists in a confidential letter.

The letter from the Health Protection Agency, the official body that oversees public health, has been leaked to The Mail on Sunday, leading to demands to know why the information has not been given to the public before the vaccination of millions of people, including children, begins.

It tells the neurologists that they must be alert for an increase in a brain disorder called Guillain-Barre Syndrome (GBS), which could be triggered by the vaccine.

GBS attacks the lining of the nerves, causing paralysis and inability to breathe, and can be fatal.

The letter, sent to about 600 neurologists on July 29, is the first sign that there is concern at the highest levels that the vaccine itself could cause serious complications.

It refers to the use of a similar swine flu vaccine in the United States in 1976 when:

* More people died from the vaccination than from swine flu. * 500 cases of GBS were detected. * The vaccine may have increased the risk of contracting GBS by eight times. * The vaccine was withdrawn after just ten weeks when the link with GBS became clear. * The US Government was forced to pay out millions of dollars to those affected.

Concerns have already been raised that the new vaccine has not been sufficiently tested and that the effects, especially on children, are unknown.

It is being developed by pharmaceutical companies and will be given to about 13million people during the first wave of immunisation, expected to start in October.

Top priority will be given to everyone aged six months to 65 with an underlying health problem, pregnant women and health professionals.

The British Neurological Surveillance Unit (BNSU), part of the British Association of Neurologists, has been asked to monitor closely any cases of GBS as the vaccine is rolled out.

One senior neurologist said last night: 'I would not have the swine flu jab because of the GBS risk.'

There are concerns that there could be a repeat of what became known as the '1976 debacle' in the US, where a swine flu vaccine killed 25 people – more than the virus itself.

A mass vaccination was given the go-ahead by President Gerald Ford because scientists believed that the swine flu strain was similar to the one responsible for the 1918-19 pandemic, which killed half a million Americans and 20million people worldwide.

The swine flu vaccine being offered to children has not been tested on infants

Within days, symptoms of GBS were reported among those who had been immunised and 25 people died from respiratory failure after severe paralysis. One in 80,000 people came down with the condition. In contrast, just one person died of swine flu.

More than 40million Americans had received the vaccine by the time the programme was stopped after ten weeks. The US Government paid out millions of dollars in compensation to those affected.

The swine flu virus in the new vaccine is a slightly different strain from the 1976 virus, but the possibility of an increased incidence of GBS remains a concern.

Shadow health spokesman Mike Penning said last night: 'The last thing we want is secret letters handed around experts within the NHS. We need a vaccine but we also need to know about potential risks.

'Our job is to make sure that the public knows what's going on. Why is the Government not being open about this? It's also very worrying if GPs, who will be administering the vaccine, aren't being warned.'

Two letters were posted together to neurologists advising them of the concerns. The first, dated July 29, was written by Professor Elizabeth Miller, head of the HPA's Immunisation Department.

It says: 'The vaccines used to combat an expected swine influenza pandemic in 1976 were shown to be associated with GBS and were withdrawn from use.

'GBS has been identified as a condition needing enhanced surveillance when the swine flu vaccines are rolled out.

'Reporting every case of GBS irrespective of vaccination or disease history is essential for conducting robust epidemiological analyses capable of identifying whether there is an increased risk of GBS in defined time periods after vaccination, or after influenza itself, compared with the background risk.'

The second letter, dated July 27, is from the Association of British Neurologists and is written by Dr Rustam Al-Shahi Salman, chair of its surveillance unit, and Professor Patrick Chinnery, chair of its clinical research committee.

Halted: The 1976 US swine flu campaign

It says: 'Traditionally, the BNSU has monitored rare diseases for long periods of time. However, the swine influenza (H1N1) pandemic has overtaken us and we need every member's involvement with a new BNSU survey of Guillain-Barre Syndrome that will start on August 1 and run for approximately nine months.

'Following the 1976 programme of vaccination against swine influenza in the US, a retrospective study found a possible eight-fold increase in the incidence of GBS.

'Active prospective ascertainment of every case of GBS in the UK is required. Please tell BNSU about every case.

'You will have seen Press coverage describing the Government's concern about releasing a vaccine of unknown safety.'

If there are signs of a rise in GBS after the vaccination programme begins, the Government could decide to halt it.

GBS attacks the lining of the nerves, leaving them unable to transmit signals to muscles effectively.

It can cause partial paralysis and mostly affects the hands and feet. In serious cases, patients need to be kept on a ventilator, but it can be fatal.

Death is caused by paralysis of the respiratory system, causing the victim to suffocate. It is not known exactly what causes GBS and research on the subject has been inconclusive.

However, it is thought that one in a million people who have a seasonal flu vaccination could be at risk and it has also been linked to people recovering from a bout of flu of any sort.

The HPA said it was part of the Government's pandemic plan to monitor GBS cases in the event of a mass vaccination campaign, regardless of the strain of flu involved. But vaccine experts warned that the letters proved the programme was a 'guinea-pig trial'.

Dr Tom Jefferson, co-ordinator of the vaccines section of the influential Cochrane Collaboration, an independent group that reviews research, said: 'New vaccines never behave in the way you expect them to. It may be that there is a link to GBS, which is certainly not something I would wish on anybody.

'But it could end up being anything because one of the additives in one of the vaccines is a substance called squalene, and none of the studies we've extracted have any research on it at all.'

He said squalene, a naturally occurring enzyme, could potentially cause so-far-undiscovered side effects.

Jackie Fletcher, founder of vaccine support group Jabs, said: 'The Government would not be anticipating this if they didn't think there was a connection. What we've got is a massive guinea-pig trial.'

Professor Chinnery said: 'During the last swine flu pandemic, it was observed that there was an increased frequency of cases of GBS. No one knows whether it was the virus or the vaccine that caused this.

'The purpose of the survey is for us to assess rapidly whether there is an increase in the frequency of GBS when the vaccine is released in the UK. It also increases consultants' awareness of the condition.

Panic over? The number of swine flu cases has fallen sharply in the past few weeks

'This is a belt-and-braces approach to safety and is not something people should be substantially worried about as it's a rare condition.'

If neurologists do identify a case of GBS, it will be logged on a central database.

Details about patients, including blood samples, will be collected and monitored by the HPA.

It is hoped this will help scientists establish why some people develop the condition and whether it is directly related to the vaccine.

But some question why there needs to be a vaccine, given the risks. Dr Richard Halvorsen, author of The Truth About Vaccines, said: 'For people with serious underlying health problems, the risk of dying from swine flu is probably greater than the risk of side effects from the vaccine.

'But it would be tragic if we repeated the US example and ended up with more casualties from the jabs.

'I applaud the Government for recognising the risk but in most cases this is a mild virus which needs a few days in bed. I'd question why we need a vaccine at all.'

Professor Miller at the HPA said: 'This monitoring system activates pandemic plans that have been in place for a number of years. We'll be able to get information on whether a patient has had a prior influenza illness and will look at whether influenza itself is linked to GBS.

'We are not expecting a link to the vaccine but a link to disease, which would make having the vaccine even more important.'

The UK's medicines watchdog, the Medicines and Healthcare Products Regulatory Agency, is already monitoring reported side effects from Tamiflu and Relenza and it is set to extend that surveillance to the vaccine.

A Department of Health spokesperson said: 'The European Medicines Agency has strict processes in place for licensing pandemic vaccines.

'In preparing for a pandemic, appropriate trials to assess safety and the immune responses have been carried out on vaccines very similar to the swine flu vaccine. The vaccines have been shown to have a good safety profile.

'It is extremely irresponsible to suggest that the UK would use a vaccine without careful consideration of safety issues. The UK has one of the most successful immunisation programmes in the world.'

I COULDN''T EAT OR SPEAK... IT WAS HORRENDOUS

Victim: Hilary Wilkinson spent three months in hospital after she was diagnosed with Guillain-Barre Syndrome When Hilary Wilkinson woke up with muscle weakness in her left arm and difficulty breathing, doctors initially put it down to a stroke.

But within hours, she was on a ventilator in intensive care after being diagnosed with Guillain-Barre Syndrome.

She spent three months in hospital and had to learn how to talk and walk again. But at times, when she was being fed through a drip and needed a tracheotomy just to breathe, she doubted whether she would survive.

The mother of two, 57, from Maryport, Cumbria, had been in good health until she developed a chest infection in March 2006. She gradually became so weak she could not walk downstairs.

Doctors did not diagnose Guillain-Barre until her condition worsened in hospital and tests showed her reflexes slowing down. It is impossible for doctors to know how she contracted the disorder, although it is thought to be linked to some infections.

Mrs Wilkinson said: 'It was very scary. I couldn't eat and I couldn't speak. My arms and feet had no strength and breathing was hard.

I was treated with immunoglobulin, which are proteins found in blood, to stop damage to my nerves. After ten days, I still couldn't speak and had to mime to nurses or my family.

‘It was absolutely horrendous and I had no idea whether I would get through it. You reach very dark moments at such times and wonder how long it can last.

But I’m a very determined person and I had lots of support.’

After three weeks, she was transferred to a neurological ward, where she had an MRI scan and nerve tests to assess the extent of the damage.

Still unable to speak and in a wheelchair, Mrs Wilkinson eventually began gruelling physiotherapy to improve her muscle strength and movement but it was exhausting and painful.

Three years later, she is almost fully recovered. She can now walk for several miles at a time, has been abroad and carries out voluntary work for a GBS Support Group helpline.

She said: ‘It makes me feel wary that the Government is rolling out this vaccine without any clear idea of the GBS risk, if any. I wouldn’t wish it on anyone and it certainly changed my life.

‘I’m frightened to have the swine flu vaccine if this might happen again – it’s a frightening illness and I think more research needs to be done on the effect of the vaccine.’

Hotline staff given access to confidential records

Confidential NHS staff records and disciplinary complaints could be accessed by hundreds of workers manning the Government’s special swine flu hotline.

They were able to browse through a database of emails containing doctors’ and nurses’ National Insurance numbers, home addresses, dates of birth, mobile phone numbers and scanned passport pages – all details that could be used fraudulently.

And private and confidential complaints sent by hospitals about temporary medical staff – some of whom were named – were also made available to the call-centre workers, who were given a special password to log in to an internal NHS website.

It could be a breach of the Data Protection Act.

The hotline staff work for NHS Professionals, which was set up using taxpayers’ money to employ temporary medical and administrative staff for the health service.

The not-for-profit company runs two of the Government’s swine flu call centres – with 300 staff in Farnborough, Hampshire, and 900 in Watford, Hertfordshire.

Shadow Health Secretary Andrew Lansley described the revelations as ‘disturbing’.

Anne Mitchell, a spokeswoman for Unison, said: ‘There’s no excuse for such a fundamental breach of personal security. Action needs to be taken as soon as possible to make sure this does not happen again.’

A spokeswoman for NHS Professionals would not confirm whether access to the confidential files had been granted.

- This is a report by Dr Mercola on squalene:

“Novartis’s proprietary squalene adjuvant for their H1N1 vaccine is MF59. Glaxo’s is ASO3. MF59 has yet to be approved by the FDA for use in any U. S. vaccine, despite its history of use in other countries.

Per Dr. Nass, there are only three vaccines in existence using an approved squalene adjuvant. None of the three are approved for use in the U. S.

What Squalene Does to Rats

Oil-based vaccination adjuvants like squalene have been proved to generate concentrated, unremitting immune responses over long periods of time. [vi]

A 2000 study published in the American Journal of Pathology demonstrated a single injection of the adjuvant squalene into rats triggered “chronic,

immune-mediated joint-specific inflammation," also known as rheumatoid arthritis. [vii]

The researchers concluded the study raised questions about the role of adjuvants in chronic inflammatory diseases.

What Squalene Does to Humans

Your immune system recognizes squalene as an oil molecule native to your body. It is found throughout your nervous system and brain. In fact, you can consume squalene in olive oil and not only will your immune system recognize it, you will also reap the benefits of its antioxidant properties.

The difference between "good" and "bad" squalene is the route by which it enters your body. Injection is an abnormal route of entry which incites your immune system to attack all the squalene in your body, not just the vaccine adjuvant.

Your immune system will attempt to destroy the molecule wherever it finds it, including in places where it occurs naturally, and where it is vital to the health of your nervous system. [viii]

Gulf War veterans with Gulf War Syndrome (GWS) received anthrax vaccines which contained squalene. [ix] MF59 (the Novartis squalene adjuvant) was an unapproved ingredient in experimental anthrax vaccines and has since been linked to the devastating autoimmune diseases suffered by countless Gulf War vets. [x]

The Department of Defense made every attempt to deny that squalene was indeed an added contaminant in the anthrax vaccine administered to Persian Gulf war military personnel – deployed and non-deployed – as well as participants in the more recent Anthrax Vaccine Immunization Program (AVIP).

However, the FDA discovered the presence of squalene in certain lots of AVIP product. A test was developed to detect anti-squalene antibodies in GWS patients, and a clear link was established between the contaminated product and all the GWS sufferers who had been injected with the vaccine containing squalene.

A study conducted at Tulane Medical School and published in the February 2000 issue of Experimental Molecular Pathology included these stunning statistics:

" … the substantial majority (95%) of overtly ill deployed GWS patients had antibodies to squalene. All (100%) GWS patients immunized for service in Desert Shield/Desert Storm who did not deploy, but had the same signs and symptoms as those who did deploy, had antibodies to squalene.

In contrast, none (0%) of the deployed Persian Gulf veterans not showing signs and symptoms of GWS have antibodies to squalene. Neither patients with idiopathic autoimmune disease nor healthy controls had detectable serum antibodies to squalene. The majority of symptomatic GWS patients had serum antibodies to squalene."[xi]

According to Dr. Viera Scheibner, Ph. D., a former principle research scientist for the government of Australia:

"… this adjuvant [squalene] contributed to the cascade of reactions called "Gulf War Syndrome," documented in the soldiers involved in the Gulf War.

The symptoms they developed included arthritis, fibromyalgia, lymphadenopathy, rashes, photosensitive rashes, malar rashes, chronic fatigue, chronic headaches, abnormal body hair loss, non-healing skin lesions, aphthous ulcers, dizziness, weakness, memory loss, seizures, mood changes, neuropsychiatric problems, anti-thyroid effects, anaemia, elevated ESR (erythrocyte sedimentation rate), systemic lupus erythematosus, multiple sclerosis, ALS (amyotrophic lateral sclerosis), Raynaud's phenomenon, Sjorgren's syndrome, chronic diarrhoea, night sweats and low-grade fevers."[xii]

Post Vaccination Follow-Up Might as Well Be Non-Existent

There is virtually no science to support the safety of vaccine injections on your long-term health or the health of your children. Follow-up studies last on average about two weeks, and look only for glaring injuries and illnesses.

Autoimmune disorders like those seen in Gulf War Syndrome frequently take years to diagnose due to the vagueness of early symptoms. Complaints like headaches, fatigue and chronic aches and pains are symptoms of many different illnesses and diseases.

- This is a patent for a vaccine to impair fertility in animals contains squalene.

The plan to use this fertility-impairing adjuvant in the "swine flu" vaccine underscores concerns

Adjuvants made of monophosforyl lipid A (MPL) MF59TM (containing a polysorpate TweenTM 80) or AS03, AS04 also known as squalene in the proposed vaccines, which are immunosterilant or an immunocontraceptive, says Daniel Solis.

The patent of the veterinary FERTILITY IMPAIRING VACCINE can be found on-line. It mentions both, the lipoid adjuvans squalene and the polysorbate TM80.

Below are the quotes from the patent and further some clinical studies about the toxicity of both.

(WO/1999/034825) FERTILITY IMPAIRING VACCINE AND METHOD OF USE This application claims the benefit of U. S. Provisional Application No. 60/070,375, filed January 2,1998, U. S. Provisional Application No. 60/071,406, filed January 15,1998 "The vaccine of the invention preferably additionally includes an immunological adjuvant to enhance the immunological response of the subject to the glycoprotein antigen. Examples of adjuvants include Freund's Complete Adjuvant, Freund's Incomplete Adjuvant, and an adjuvant comprising an immunostimulant such as synthetic trehalose dicorynomycolate (STDCM) and an oil such as squalene oil (see P. Willis et al., J. Equine Vet. Sci., 14,364-370 (1994)). An adjuvant comprising synthetic trehalose dicorynemycolate, squalene oil, and a surfactant such as lecithin is preferred. Lecithin typically includes phosphatidyl choline. In a preferred embodiment the vaccine comprises oil, preferably a biodegradable oil such as squalene oil. Typically, the vaccine is prepared using an adjuvant concentrate which contains lecithin in squalene oil. The aqueous solution glycoprotein is typically a phosphate-buffered saline (PBS) solution, and additionally preferably contains Tween 80."

Abstract: A vaccine comprising an antigen derived from a zona pellucida glycoprotein is effective to impair fertility in animals, preferably carnivores. The vaccine can be used as an immunosterilant or an immunocontraceptive. http://www.wipo.int/pctdb/en/wo.jsp?wo=1999034825 Description: http://www.wipo.int/pctdb/en/wo.jsp?IA=US1998027658&wo=1999034825&DISPLAY=DESC

_____ Annals of Allergy, Asthma and Immunology, Volume 95, Number 6, December 2005, pp. 593-599(7) "Polysorbate 80 was identified as the causative agent for the anaphylactoid reaction of nonimmunologic origin in the patient. Conclusions: Polysorbate 80 is a ubiquitously used solubilizing agent that can cause severe nonimmunologic anaphylactoid reactions."

Gajdova M, Jakubovsky J, Valky J. Institute of Preventive and Clinical Medicine, Limbová, Bratislava. Delayed effects of neonatal exposure to Tween 80 on female reproductive organs in rats. Food Chem Toxicol. 1993 Mar; 31(3): 183-90. PMID: 8473002. "Baby female rats were injected with polysorbate 80at days 4-7 after birth. It accelerated the maturing of the rats and caused changes to the vagina and womb lining, hormonal changes, ovary deformities and degenerative

follicles."
http://www.ncbi.nlm.nih.gov/pubmed/8473002

The Endogenous Adjuvant Squalene Can Induce a Chronic T-Cell-Mediated Arthritis in Rats Barbro C. Carlson*, Åsa M. Jansson*, Anders Larsson, Anders Bucht and Johnny C. Lorentzen* http://ajp.amjpathol.org/cgi/content/abstract/156/6/2057

_____ Now, how can WHO claim the adjuvans is harmless: http://www.who.int/vaccine_safety/topics/adjuvants/squalene/questions_and_answers/en/index.html

when there is clear evidence of its effects provoking AI deseases: ANTI-SQUALENE ANTIBODIES LINK GULF WAR SYNDROME TO ANTHRAX VACCINE http://www.autoimmune.com/GWSGen.html

or

"Dr. Jules Freund creator of this oil-based adjuvant warned in 1956 that animals injected with his formulation developed terrible, incurable conditions: allergic aspermatogenesis (stoppage of sperm production), experimental allergic encephalomyelitis (the animal version of MS), allergic neuritis (inflammation of the nerves that can lead to paralysis) and other severe autoimmune disorders. Source: : Gary Matsumoto, Vaccine A-The Covert Government Experiment That's Killing our Soldiers and Why GI's are Only the First Victims, Kapitola 3. "The Greatest Story Never Told" http://www.vaccine-a.com/excerpt.html"

- **Innoculations:** The True Weapons Of Mass Destruction Rebecca Carley, M.D. Infowars June 28, 2009

"One basic truth can be used as a foundation for a mountain of lies, and if we dig down deep enough in the mountain of lies, and bring out that truth, to set it on top of the mountain of lies; the entire mountain of lies will crumble under the weight of that one truth. And there is nothing more devastating to a structure of lies than the revelation of the truth upon which the structure of lies was built, because the shock waves of the revelation of the truth reverberate, and continue to reverberate throughout the Earth for generations to follow, awakening even those people who had no desire to be awakened to the truth." (by Delamar Duvaris as written in the preface of "Behold the Pale Horse" by William Cooper). featured stories Innoculations: The True Weapons Of Mass Destruction

The basic truth that served as the foundation for the mountain of lies known as vaccinations was the observation that mammals who recover from infection with microorganisms acquire natural immunity from further infections. Whenever T cells (the little packman cells which kill viruses, bacteria, and cancer cells, thus conferring cellular immunity) and B cells (antibody producing cells which confer humoral immunity) are activated by various substances foreign to the body called antigens, some of the T and B cells become memory cells. Thus, the next time the individual meets up with that same antigen, the immune system can be quickly triggered to demolish it. This is the process known as immunity.

This truth gave birth to a beLIEf that if a foreign antigen was injected into an individual, that individual would then become immune to a future infection. This beLIEf, (you see the lie in the middle), was given the name, "vaccinations". What the promoters of vaccination failed to realize is that the respiratory tract of ALL mammals (since animals are just as devastated by these inoculations with disease as are humans) contain secretory IgA (an antibody which initiates the natural God given immune response) within the respiratory tract mucosa. Bypassing this mucosal aspect of the immune system by directly injecting organisms into the bloodstream leads to a corruption in the immune system itself. As a result, the pathogenic viruses or bacteria cannot be eliminated by the immune system and remain in the body, where they will further grow and/or mutate as the individual is exposed to ever more antigens and

toxins in the environment which continue to assault the immune system.

The mechanism by which the immune system is corrupted can best be realized when you understand that the two poles of the immune system (the cellular and humoral mechanisms) have a reciprocal relationship. Thus, when one is stimulated, the other is inhibited. Since vaccines activate the B cells to secrete antibody, the T cells are subsequently suppressed. This suppression of the cell mediated response is a key factor in the development of cancer and life threatening infections. In fact, the "prevention" of a disease via vaccination is, in reality, an inability to expel organisms due to the suppression of the cell-mediated response. Thus, rather than preventing disease; they actually prevent the disease from ever being resolved. The organisms continue circulating through the body, mutating and transforming into other organisms (as demonstrated by the work of Professor Antoine Bechamp), depending on the acidity and toxicity of the internal terrain of the body. Note that Bechamp proved that Louis Pasteur's "germ theory" of disease was incorrect due to this ability of organisms to transform and mutate based on the body's internal terrain (as Pasteur admitted on his deathbed). Thus, treatment of infection with antibiotics as well as "prevention" of disease with vaccines are both just examples of cutting off the branches of dis-ease, when the root of the cause is a toxic internal environment. However, since Pasteur's germ theory was conducive to the profits of the burgeoning pharmaceutical companies who only manage dis-ease, no mention of the work of Professor Bechamp has been made in medical school curricula.

To make matters worse than the suppression of cellular immunity which occurs when vaccines are injected, adjuvants (which are substances added to vaccines to enhance the antibody response) can actually lead to serious side effects themselves. Adjuvants include oil emulsions, mineral compounds (which may contain the heavy metal aluminum), bacterial products, liposomes (which allow delayed release of substances), and squalene. The side effects of adjuvants themselves include hyperactivity of B cells leading to pathologic levels of antibody production, as well as allergic reaction to the adjuvants themselves (as demonstrated in Gulf War I soldiers injected with vaccines containing the adjuvant squalene, to which antibodies were found in many soldiers). Note that the pathologically elevated hyperactivity of antibody production caused by adjuvants also results in a distraction from the other antigens that the immune system encounters "naturally", which must be addressed to maintain health.

This hyperactivity of the humoral (antibody producing) pole of the immune system is, in this author's opinion, the sole cause of all autoimmune diseases. The only thing which determines which autoimmune disease you develop is which tissues in your body are attacked by auto-antibodies. If the inside lining of the gastrointestinal tract (the mucosa) is attacked by auto-antibodies you develop leaky gut syndrome (which leads to food allergies when partially digested food particles are released into the bloodstream, are recognized as antigens foreign to the body, and elicit an antibody response against those food particles that becomes heightened every time that same food is eaten and released into the bloodstream partially digested again). Crohn's disease and colitis are also caused by auto-antibody attack on the mucosa of the GI tract itself. If the islet (insulin producing) cells of the pancreas are attacked by auto-antibodies, you develop insulin dependent (juvenile) diabetes. If the respiratory mucosa is attacked by auto-antibodies, you develop "leaky lung" syndrome where, just as with leaky gut, antigens recognized as foreign to the body which are inhaled are able to traverse the lining of the respiratory tract, causing the creation of antibodies against those antigens (usually dust, mold, pet or pollen antigens). When these substances are inhaled again, the allergic response producing constriction of the bronchioles is called asthma. If the components of the articular surface of the joints are attacked by auto-antibodies, you develop rheumatoid (or juvenile) arthritis. If the skin is attacked, you develop "leaky

skin" syndrome, where contact antigens which could not otherwise traverse the skin are allowed in, leading to skin allergies to contact antigens. Additionally, depending on which level of the skin is attacked by auto-antibodies, (i. e., the epidermis or dermis), you develop eczema, psoriasis or scleroderma. If the kidney tissue is attacked by auto-antibodies, you develop one of the many types of nephritis, depending on which component of renal tissue is attacked (for example, with glomerulonephritis, the basement membrane of the glomerular apparatus within the kidney (which filters blood to form urine) is attacked by auto-antibodies, thus allowing protein to escape from the serum into the urine). If you develop auto-antibodies against thyroid gland tissue, you develop Grave's disease. If you develop auto-antibodies against the tissue of the thymus gland (which is crucial in T cell production and function), you develop myasthenia gravis. If you develop auto-antibodies against the very DNA in the nucleus of all cells, you develop systemic Lupus (thus, the potential autoimmune potential of DNA vaccines being developed now is self evident; worse yet, DNA components from these vaccines can be incorporated into your DNA, leading to actual genetic changes which could cause extinction of all (vaccinated) life on the Earth). And on, and on, and on.

The brain and spinal cord can also be attacked with auto-antibodies (which this author refers to as vaccine induced encephalitis), leading to a variety of neurological diseases. The most severe of these, leading to death, are sudden infant death syndrome (SIDS) and most cases of "shaken baby syndrome". If components of the myelin sheath (the insulating covering of nerve fibers which allows proper nerve conduction) or the actual neurofilaments themselves are attacked by auto-antibodies, the resultant condition is determined solely by the location of the damage done. Such neurological conditions include but are not limited to minimal brain dysfunction, ADD/ADHD, learning disabilities, mental retardation, criminal behavior, the spectrum of pervasive developmental disorders (including autism), multiple sclerosis, Parkinson's, Lou Gehrig's disease, Guillen Barre,, seizure disorders, etc., etc. etc. (Please note that other toxins are also sometimes involved, such as: aspartame, Lymes and mercury in cases of MS; aspartame in seizures; or pesticides in cases of Parkinson's). Thus, when detoxing to reverse these diseases, these other substances must also be detoxed to obtain a full recovery. However, the corruption of the immune system caused by the injection of vaccines is a key component in these disease states leading to immune malfunction, and is the reason why an autistic child may also have

leaky gut or eczema, etc. Note that myelin production, for the most part, does not begin until after birth. Most myelin is apparently laid down by age 5 years and usually completed by age 10 years, judging by the level of success at various ages in reversing autistic and other neurological VIDS symptoms that this author has observed in hundreds of children by detoxing the viruses with homeopathic nosodes, and repairing the immune corruption by simultaneous administration of bovine colostrum (i. e., after 10 years of age, the ability to stop and repair auto-antibody induced damage in the myelin sheath and neurofilaments themselves is dramatically decreased).

Thus, the hyperactivity of the humoral arm of the immune system in autoimmune disease is caused by adjuvants added just for that purpose. However, the autoimmunity itself (i. e., antibody against self) has several mechanisms, including the following:

1. The antigens present in the culture media itself cannot be completely filtered and separated from the organisms cultured thereon. Thus, any antibodies formed against antigens from the culture cells themselves (for example myelin basic protein from chick embryos or the 13 vaccines which now contain aborted fetal cells) can cross-react to form an autoimmune reaction against the myelin basic protein in your myelin sheath, etc.

2. Molecular mimicry is due to similarity of proteins contained in organisms and mammals. (For example, the measles virus is made up of

proteins similar to myelin basic protein; thus, antibodies formed against the measles virus antigens subsequently also cause an auto-antibody attack against myelin basic protein in the myelin sheath due to cross reactivity of these antibodies).

3. intentional inclusion of antigens in vaccines to cause formation of antibodies that attack specific hormones or races (for example, experiments done on women of childbearing age in the Philippines and probably other locations where HCG (human chorionic gonadotropin, the hormone produced when women first become pregnant) placed into vaccines given these women resulted in antibodies against the HCG hormone, and subsequent spontaneous abortion thus occurred when the women became pregnant. It is also this author's hypothesis that the epidemic of vitiligo in people of color (hypo pigmentation of skin caused by auto-antibody attack on melanocytes (melanin producing cells) in skin) is also occurring due to intentional inclusion of melanin in vaccines given to people of color.

In addition to the above phenomena which lead to simultaneous depression of cellular immune function and hyperactivity of humoral immune function, vaccines also contain other toxic substances which can cause serious side effects themselves. The following ingredients are actually listed on the CDC website with this introductory statement: "Many things in today's world, including food and medicines, have chemicals added to them to prevent the growth of germs and reduce spoilage." Translation: you, re already toxic, so what's the big deal with adding more poison? This author's answer to that question is that any immunotoxin can end up being the "straw that breaks the immune system's back" in that individual, leading to dis-ease. This is where genetics is key; i. e., not that what disease you develop is actually caused by some "gene" in most cases; but rather that your genes determine the strength of your immune system (i. e., how many assaults your immune system can take before it reaches critical mass, and you develop a dis-ease).Some additional ingredients in vaccines (as listed by the CDC on their website) include antibiotics, aluminum gels, formaldehyde, monosodium glutamate (MSG), egg protein, and sulfites. Thus, we have antibiotics (which you could be allergic to) ; aluminum (highly involved in causing Alzheimer's disease) ; formaldehyde (a toxic carcinogenic substance used to pickle dead people) ; MSG (a potent excitotoxin which, like aspartame, can cause seizures, brain tumors, etc.) ; egg protein (to which you could have a life threatening anaphylactic reaction) ; and sulfites (another toxin which we are advised not to consume much of orally, but in vaccines, it is injected directly into the blood stream). Is this not a veritable witch's brew of chemicals, organisms, and animal or human (aborted fetus) body parts? Note in this list that they do not mention the ethyl-mercury containing preservative thimerosol, which has been the only dangerous substance in vaccines to receive mainstream media attention (albeit most of that being disinformation) after the explosion in the rate of occurrence of autism in the last generation became self-evident proof that vaccines are the causative factor. For, although the scientists working for the medical mafia continue to use statistics to twist and spin their data to make us beLIEve that vaccines are not the cause, too many thousands of parents have watched their children enter the downward spiral into autism after their children received the vaccine which was the straw that broke the back of their child's immune system. No matter what the "white coats" tell these parents; they know the truth!

Mercury (also in dental amalgam fillings) is a highly toxic heavy metal, has been documented to cause cancer, and can be absorbed through the digestive track, skin, and respiratory track. Mercury is 1,000 times more toxic than lead, and is second only to uranium as the most toxic metal. If children receive all recommended vaccines, they will receive 2,370 times the "allowable safe limit" for mercury in the first two years of life (as if there is such a thing as a "safe" amount of a toxic poison). Yet, even after Congressional hearings instigated by Congressman Dan Burton (whose own grandchild became autistic after receiving

vaccines) resulted in the FDA requesting (not ordering) vaccine manufacturers to remove this toxic heavy metal from their products, mercury is still present in many vaccines.

Although the symptoms of mercury poisoning are identical to the symptoms of autism, it should be noted that most children who descend into the hellish state known as autism do so after the MMR vaccine. The MMR vaccine is one of the few vaccines that do not contain mercury. Thus, it is self-evident that the removal of mercury will not make vaccines "safe". (This is why the mercury is the only thing being addressed at all; because when the people reading this paper realize that the very mechanism by which vaccines corrupt the immune system means that NO vaccine is safe and effective; there will be an evolution of consciousness where the structure of lies telling us vaccines are safe and effective disintegrates.) In the autistic community, this will lead to an exodus from the multiple autism groups saying it is all about the mercury or worse yet, that autism genes are "inherited", to the only group which has their focus on the actual problem. This group is named TAAP (the Autism Autoimmunity Project at www.taap.info/), and is led by April Oakes. In this author's opinion, it will be TAAP in alliance with the vaccine damaged soldiers and vets of the American Gulf War Veterans Association at www.agwva.org led by Peter Kawaja which, working together, will stop this holocaust on humanity called VIDS. The good news is that these VIDS can be reversed using natural remedies contained in the Hippocrates Protocol (www.drcarley.com). This "surgical strike" detoxification approach which has the potential to reverse ALL of the aforementioned conditions under the VIDS umbrella as long as detoxification is started early enough (before age 10 for neurological VIDS) will be the one truth put on top of the mountain of lies (that vaccines are safe and effective) that will cause the entire mountain of vaccine lies to crumble. Combined with a massive outpouring of public support to these two organizations, the root of the tree of vaccine evil will be exposed. Thus, the holocaust on humanity (where instead of people being put in concentration camps, the concentration camps are being put into the people) will finally be put to an end.

Unfortunately, we can no longer pretend that this epidemic of VIDS is merely a "mistake" made by good intentioned, albeit misguided mad scientists. Because it's even worse than the above, folkswe are talking TREASON and CRIMES AGAINST HUMANITY, PETS, and even PLANTS, (which are also being genetically modified to create vaccines). The evidence for this is as follows:

As concern for population growth started to grow and the final plans to bring in the New World Order were put in place, this lie called vaccines was transformed into pure evil, as it was realized that such delivery systems could be used to intentionally cause disease, which is now being done under the US Code, Title 50, Chapter 32, § 1520 and 1524. You can read it for yourself at your local library, or go to

http://www.drcarley.com/Title%2050%20US%20Code. htm.

law has been in place since the 1960's, and it was last modified in April of 2000. The only stipulation made for experimentation on human subjects is that local civilian officials be notified 30 days before the experiment is started. Section 1524 adds that the Secretary of Defense may enter into agreements with the Secretary of Health and Human Services to provide support for vaccination programs through use of excess peacetime biological weapons (i. e., weapons of mass destruction). In April 2000, § 1520 (a) was passed to put alleged restrictions on the use of human subjects for testing of chemical or biological agents after a caller on C Span mentioned this law in 1999, which revealed this treasonous law to a huge audience of listeners (including this author, who has been including it in lectures and written materials since that call came into "Washington Journal"). However, the exceptions written to Title 50, chapter 32 under § 1520 subsection (b) in the

2000 law passed by our aiders and abettors of treason in Congress not only loophole back in a test carried out for "any peaceful purpose that is related to a medical, therapeutic, pharmaceutical, agricultural, industrial, or research activity"; but add that such biological and chemical warfare agents can now be also used for any law enforcement purpose, including "any purpose related to riot control" (just in case those C Span listeners should actually get off the couch at the horror of what the traitors in Washington, D.C. are doing to God's people). Subsection (c) of this law now mandates that "informed consent" be required. In reality, not a single vaccine has ever been tested for its long term side effects (including carcinogenic potential). Additionally, the intentional introduction of stealth viruses, (including man-made viruses that cause cancer, mycoplasma and the HIV virus), antigens which target certain races, (and surely a microchip in the future) into vaccines makes it self evident that informed consent is impossible, as it would initiate impeachment proceedings and war crimes trials against every "public servant" involved in perpetrating these crimes against the American people, in violation of the Nuremberg Code (which was written after the end of WW II to prevent the barbaric experiments that occurred in the Nazi concentration camps) .

What most people don't know is that the top level mad scientists from Nazi Germany were actually brought to the United States through "Operation Paperclip", and have been continuing their work to this day in places like Brookhaven labs, Cold Spring Harbor and Plum Island in this author's backyard on Long Island. (To see the document proving that American scientists created the HIV virus, this author refers you to p. 442 of Death in the Air by Dr. Leonard G. Horowitz, where you can read the 1969 document in which the U. S. military/CIA and Rockefeller directed National Academy of Sciences-National Research Council (NAS-NRC) announced that a research program to explore the feasibility of "creating a new infective microorganismwhich would be refractory to the immunological and therapeutic processes upon which we depend to maintain our relative freedom from infectious disease" could be completed at a total cost of $10 million.) Yes, this is what your tax dollars are going towards, folks. But hang on to your hat, because it only gets worse.

The most heinous, bone chilling and evil piece of this puzzle has been revealed to the world by an American hero named Peter Kawaja, who worked in the late 1980's as a security and counter terrorism expert for the United States government (a service for which he has been rewarded with the murder of his wife, torching of his home, issuance of a war crimes subpoena to (they thought) confiscate all his evidence, illegal IRS liens on all subsequent income, and multiple attempts on his own life, all funded by YOUR tax dollars). Please go to www.agwva.org/mission.htm and read some of the 34 counts that Mr. Kawaja brought against the domestic traitors to America (in both their individual and governmental capacities) in a federal lawsuit in which the perpetrators, again, used your tax dollars to hire themselves attorneys from the Department of "Justice" whose defense of their war criminal clients was that they are "immune, under color of law". (You can listen to Mr. Kawaja on one of his multiple internet radio shows, including "What's Ailing America?" which he co-hosts with this author at www.highway2health.net every Wednesday at 10 PM, EST.

Dr. James R. Shannon, former director of the National Institute of Health reported in December, 2003 that "the only safe vaccine is one that is never used". However, the reverberating truth, "the shot heard round the world" which will lead to the evolution of consciousness necessary to stop the holocaust against humanity known as vaccinations, will be that not only are vaccinations not safe or effective, but that they are actually weapons of mass destruction being perpetrated upon humanity in the name of health, for the purpose of genocide and to bring in the New World Order. Part 2 of the genocidal plan could drop anytime with activation

of the Model State Health Emergency Powers Act whenever the next fabricated terrorist attack using biological agents occurs. Worse yet, the Congressional traitors in Washington posing as public "servants" are doing all they can to pass "Codex" legislation which will make the natural remedies and supplements used in the Hippocrates Protocol developed by this author to reverse all dis-eases only available by prescription. So, you didn't hear about that on your local news station either? Please go to the site of another American hero, John Hamill of the International Alliance for Health Freedom (who reversed his schizophrenia symptoms with these natural supplements and has dedicated his life to stop Codex from passing) at www.iahf.com .

Wake up, GUYS-it's getting very late. It is time for the mountain of lies to crumble. Please spread the world to everyone you know we can make it happen. The time to stop chopping at branches and get to the root of this evil is now.

Jane Burgermeister