How Vaccines Dysregulate The Immune System and Impact Genetic Control Over Disease Expression

By Patricia Jordan, DVM, CVA, CTCVH, & Herbology
http://dr-jordan.com/

(Extracts): ***Vaccines lead to genetic mutations... Genetic impact on MHC (HLA) is what dictates genetic expression of disease susceptibility. Vaccines rob the individual of natural evolutionary selection pressures based on natural antigen risks. Vaccines are altering gene sequences, inserting genes, affecting genome and destroying the organic immunologically determined susceptibility that evolved with natural selection. Determined susceptibility is genetically impacted ahead of disease expression by the antigens presented to or encountered by the individual. The immune system has evolved naturally to promote life and what is happening with unnatural antigen environment delivered in unnatural route to dysregulate the immune system is resulting in unnatural selection, immune system corruption and species distortion.***

The genetic basis for susceptibility to disease is complex but well before man understood anything about the immune system and how it worked, he intruded on the evolved design with a hubris that is having collateral damage and unintended consequences of species de- evolution. There are many ways that vaccines dysregulate the immune system and de construct health. One effect of the vaccine is on the genetic expression of disease through the unnatural engagement of the MHC (Major Histocompatibility Complex), the major HLA (Human Leukocyte Antigens) then minor loci; cytokine genes, CD-encoding genes, T cell receptor genes, growth hormone and immunoglobulin genes any of the polygenes that cascade down to the intricacies of our many possible gene responses. The second effect of the vaccine is to skew the immune system and remove the balance of Th1 and Th2 between cell mediated immunity and humoral immunity.

Total dysregulation then comes, including a combination of mutations, gene expression, biochemical pathway alterations, enzyme disruptions, up regulation of the IgE expression and a down regulation of the IgA, and disrupts the cytokine profile and many, many other routes depending on the nature of the pathogen and toxins in the vaccine and the variable gene response of the individual. Expression of disease now, is a function of the unnatural exposure to unnatural pathogens and toxins and the expression of disease as varied as behavioural, Type I-IV hypersensitivity; allergies, asthma, anaphylaxis, atopy, eczema, cancer, autoimmune, bacterial, viral, yeast, fungal, internal and external parasites and genetic diseases.

The genetic expression of disease is predated by the link-up of the pathogen and the individual's gene. The unnatural selection pressure on the species by the use of vaccines is unnaturally evolving or de-evolving the species through genotoxicity and genetic disease increase. The genetic damage or "genetic susceptibility" is transferable to the next generation. The next generation when vaccinated, expresses easily the adverse events that vaccines are selecting for. The type of immune response that occurs after pathogen binding is determined by cytokine messengers that are triggered by certain elements of the pathogen. The vaccine contains a multiple number of ways to affect this: contamination with unknown viruses and microbial components, unnatural pathogens, chimeras and other genetically engineered products, unfiltered genetic pieces like virions, prions, viruses from other species, aluminum, mercury which can directly lead to abnormal cytokine messengers being produced via pathogen alterations/adulterations/mutations.

Modified live viruses (or "attenuated viruses") allow the vaccine viruses to migrate to and replicate onto the host's tissues. Another act of hubris has occurred because despite scientists' having played with viruses all of these last 300 years, it was only recently that science discovered that viruses are not dead, they are not alive; they are packets of genetic material that when in the presence of a susceptible and permissive living cell that has the necessary receptor can replicate and infect.

Sometimes, viral contamination in a vaccine can activate in the human body 30-40 years after inoculation. The presence of unknown viruses, the contamination of viruses, the recombination and reassortment of viral genes and the introduction of xenotropic viruses, infective DNA viruses have all again – due to the hubris of man – introduced disease and pathology into organisms receiving the jab. The process of injecting unnatural substances into the body started well before the identification of the fist virus! The contamination continues today with the filtering process not finding virions and prions and other smaller genetic impacting contaminants. [The rotavirus vaccines for children have been found to contain pig viruses]. The viral and even microbial antigens are all players in the genetic expression of disease and disease susceptibility to every genome via the MHC and other still unidentified pathways.

Adjuvants additionally adulterate the intelligence of the innate immune response. Adjuvants "add" inflammation and pathogen distortion and therefore cell signalling adulterations, thus impinging upon the evolutionarily perfected system and resulting in a loss of order. In 1988, Dr. Ron Schultz spoke out in a roundtable discussion over his concerns of the random addition to anything into vaccines without understanding in the least the impact that the addition of, for example, interleukins into vaccines. He framed the impact of the whole body or even just the immune system as a complete unknown yet the cavalier attitude from vaccine makers was that no caution was necessary.

Including interleukin in some vaccines in the 1980's has likely now produced children born to vaccinated populations with the genetic disease of missing interleukins! The "new" auto inflammatory syndrome DIRA deficiency of interleukin 1 receptor agonist where children display a constellation of serious and potentially fatal systemic symptoms from birth are inherited mutations in IL IRN – a gene that encodes a protein known as interleukin 1 receptor antagonist.

The irony that Dr. Ian Tizzard would recommend adding ingredients like alum to vaccines that have been used since 1926, without having any idea how they worked, is little comfort to the many parents of children suffering from brain cancer.

In 1999, the World Health Organization's International Agency for Research on Cancer (IARC) evaluated the carcinogeniticity of feline vaccines containing aluminium adjuvants, due to cats developing tumours at the site where they had been vaccinated. The IARC monograph stated: "There is limited evidence in cats for the carcinogenicity of certain feline vaccines

containing adjuvants" (Page 310)

http://monographs.iarc.fr/ENG/Monographs/vol74/mono74.pdf

It doesn't help either to understand now that aluminium will increase the permeability of the blood brain barrier and allow viruses (for example, viruses that have an affinity for the central nervous system like measles) into the brain along with the mercury and aluminium – both metals that can act synergistically to mutate. Seriously, they still don't see where the rise in childhood brain cancer is coming from?

Aluminium in the vaccines is also up-regulating IgE and compromising IgA, therefore the presence of aluminium in the vaccines is a much involved gene impactor which causes vaccines to result in allergies, atopy, anaphylaxis, asthma and eczema expression. The natural immune system has a variety of defence tools to use in the protection of the organism; however these systems are dysregulated when damaged pathways result from damaged pathogens or genetically engineered pathogens are artificially introduced. The effects of alum were never known even though the toxin has enjoyed a hierarchical rise in use and success.

The amount of aluminium adjuvant used in vaccines is not a "safe" amount, it is only the amount they found necessary to exert its inflammatory effect as an adjuvant! The lack of safety studies, lack of teratogenicity or carcinogenic studies or any long term effects signalling genetic defects from vaccines were never done. For any agency from the HHS, CDC, FDA, USDA,WHO, UN, UNICEF, and GAVI to endorse or project vaccines as "safe" is criminal, and investigations should be called for. Liability waivers put in force for the drug companies to escape prosecutorial litigation will not be upheld in the face of gross criminal action for failure to perform due diligence in the safety studies or even of the efficacy studies that are lacking for vaccine use in the first place.

The question as to what exactly was known as FDA-licensed products are unleashed upon the public gives rise to another question: why is it that the drug companies that makes vaccines and promotes their use, are the same drug companies that make the drug for the VACCINE DISEASE that follows the vaccine use? What exactly are the revelations that are bound behind "proprietary confidentiality clauses" and is this the way drug companies are pleading the Fifth Amendment for protection from self incrimination?

Would this be the reason the governments remove vaccine liability from the manufacturers of the experimental guise under which health care is purported?

The highly polymorphic HLA and DLA (Human Leukocyte Antigen and Dog Leukocyte Antigen) systems which are involved in antigen presentation clearly affect responses to vaccination and therefore this impact is unknown in any organism receiving the jab. This lack of knowing makes every vaccination "experimentation under the guise of health care delivery." Effects of vaccines on any individual are variable and therefore any expected result incalculable, the risk to any organism is therefore unknown.

Administering a jab is not synonymous with conveying immunity. Antibody production is not equated to immunity and vaccination does not mean immunisation. Damage from vaccines are cumulative; cell mediated immune suppression increases significantly with every jab. Multivalent vaccines are particularly damaging and immune-disrupting. Only vaccinated individuals were found to develop autoantibodies in a landmark study done at Purdue University.

Autoantibodies are made with the vaccines from the viruses, from the microbial antigens, from the aluminium and mercury and other ingredients that would mutate or disrupt the pathogen. The increase of molecular mimicry increases with vaccines and these examples of pathways to increase the number of autoantibodies formed the trigger necessary to promote genetic expression of autoimmune disease. Certainly, autoimmune disease expression is one step closer to genetic disease and that handicap will transfer vertically to the next generation in many instances.

The important understanding is that the adulteration of the genome came in via the injection of vaccines. Since not even a very heavy book could contain all the pathways to disease expression from genetic effects of the vaccine (the great immune adulterant), let us at least end this with the following understanding; vaccines have no environmental epidemiological studies to support the benefits over risks of vaccine administration, they are not safe nor innocuous and have not even been proven effective in conveying immunity which is the only reason one would consider their use in the first place. Vaccination use fits the definition of "a medical assumption" and according to Dr. Stephen Blake is certainly the biggest medical assumption ever made in the history of mankind and is directly responsible for more disease, death and disability than any other medical procedure or act. The precautionary principle is a moral and political principle which states that if an action or policy might cause severe or irreversible harm to the public or to the environment, in the absence of a scientific consensus that harm would not ensue, the burden of proof falls on those who would advocate taking the action. The principle aims to provide guidance for protecting public health and the environment in the face of uncertain risks, stating that the absence of full scientific certainty shall not be used as a reason to postpone measures where there is a risk of serious or irreversible harm to public health or the environment. It is obvious that the unnatural vaccine has unnaturally selected for genes that do not reflect a natural exposure from the real environment and thusly resulted in unnatural selection of genes that have dysregulated the immune system and disrupted the inflammatory pathway and distorted the populations genetically.

Unnatural gene selection is then leading to resistance and susceptibility to disease which is unnatural and not the real picture of the antigen state within our external environment for which an immune system is geared to provide survivability against encounter. Vaccination is resulting in abnormal disease expression and the making of disease previously not encountered. Although it is popular to blame our external environment, this is not the main environment our immune system is being pressured by.

In the madness, the species are being distorted and genomes are being corrupted. The rise of genetic susceptibility and genetic disease is a reflection of this distortion. Much of the illness that we see today is "Vaccine Disease", in that dysregulation of the immune system by vaccines has altered genetic susceptibility and expression of disease and is not evolving a better immune system and health. Instead, the effect of vaccination is to deconstruct the immune system and point the genome towards doom.

About Dr. Jordan

Patricia Jordan is a graduate of the North Carolina College of Veterinary Medicine. Having practiced conventional veterinary medicine for fifteen years and originated four different veterinary practices in North Carolina, Dr. Jordan found holistic medicine in 2000 at the AHVMA American Holistic Veterinary Medical Association Conference in Williamsburg, Va. Holistic medicine set her on a pathway towards many of the modalities that provided her with the inspiration to follow the path of healing with energy and intention. Working to complete a Master's Program in TCVM Traditional Chinese Veterinary Medicine with Dr. Xie of the Chi Institute and participating in Dr. Richard Pitcairn's Professional Course for Veterinarians has opened the way to naturopathic medicine for Dr. Jordan.

SOURCE: http://www.thedogplace.org/VACCINES/ImmuneSystem-Impact_Jordan-DVM-107.asp