Bethany Schoeff
Vaccine File:

I post a lot on the Facebook page Vaccine Freedom for Washington State. Here's one great article.

https://www.facebook.com/VFFWS/posts/1322458837776282

Great resource:

http://www.learntherisk.org/studies/

Studies - May need google account

https://docs.google.com/document/d/1ic6fA21vxHUY5-kGAiTrgwHUcVXyVhP_ArGYYDkjCmk/edit

https://docs.google.com/document/d/1ic6fA21vxHUY5-kGAiTrgwHUcVXyVhP_ArGYYDkjCmk/mobilebasic

Shedding

For anyone that may be reading this who doesn't believe in vaccines shedding:

The Emerging risks of live virus & virus vectored vaccines
http://web.archive.org/web/20161208182006/http://www.nvic.org/CMSTemplates/NVIC/pdf/Live-Virus-Vaccines-and-Vaccine-Shedding.pdf

What is shedding?
http://www.westonaprice.org/press/studies-show-that-vaccinated-individuals-spread-disease/

Flu (FluMist Intranasal) - Shedding (Section 5.4)
https://web.archive.org/web/20160530024811/http://www.fda.gov/downloads/BiologicalBloodVaccines/Vaccines/ApprovedProducts/UCM123743.pdf

Flu vaccine shedding
http://www.ncbi.nlm.nih.gov/m/pubmed/21513761/

Chicken Pox (Varivax) - Shedding (Section 5.4)
http://www.merck.com/product/usa/pi_circulars/v/varivax/varivax_pi.pdf

Shingles (Zostavax} - Shedding (Section 5.2)
http://www.merck.com/product/usa/pi_circulars/z/zostavax/zostavax_pi2.pdf

MMR Shedding (Page 5, Under Precaution)

http://www.merck.com/product/usa/pi_circulars/m/mmr_ii/mmr_ii_pi.pdf

Rotavirus (Rotarix)- Shedding (Section 5.4)
https://www.gsksource.com/gskprm/htdocs/documents/ROTARIX-PI-PIL.PDF

Smallpox (ACAM2000) - Shedding (Section 5.4)
http://www.fda.gov/downloads/biologicalBloodVaccines/vaccines/approvedProducts/UCM142572.pdf

Detection of Measles Virus RNA in Urine Specimen from Vaccine Recipients

http://jcm.asm.org/content/33/9/2485.long

Did you know that there are such things as "live virus vaccines"? These obviously contain live viruses. What does that really mean for us? They may be weakened, but they still come with inherent risks. Since they are alive they can still replicate given the right environment and infect the vaccinated individual and/or "shed" to people around them for weeks after the vaccine is given.

Live virus vaccines and the links to the package insert position that discusses shedding: http://www.immunize.org/packageinserts/

But does that REALLY happen? Yes. Yes it can and does as these studies illustrate:

Varicella transfer after vaccine to pregnant mom:
http://www.ncbi.nlm.nih.gov/pubmed/9255208

Pub Med article on Rotavirus shedding:
http://www.ncbi.nlm.nih.gov/pubmed/18922486

Small Pox shed to toddler via father (Military)
http://mobile.reuters.com/article/idUSN1744524120070518

http://mobile.nytimes.com/2007/05/18/health/18smallpox.html

"The five children with chickenpox, also called varicella, attend Poplin and Hemby Bridge elementary schools and all had been vaccinated, said Union County Health Director Phillip Tarte."

http://www.charlotteobserver.com/living/health-family/article64604577.html

Mumps Vaccine sheds:
http://www.ncbi.nlm.nih.gov/pubmed/24772647

Mumps vaccine sheds:
http://www.ncbi.nlm.nih.gov/pubmed/16266774

Mumps outbreak in Netherlands linked to those vaccinated twice with MMR: (open in safari - Facebook app malfunctions)
http://wwwnc.cdc.gov/eid/article/20/4/13-1681_article

(Vaccinated) Student Diagnosed With Mumps at California State University San Marcos

http://www.nbcsandiego.com/on-air/as-seen-on/Cal-State-San-Marcos-Student-Diagnosed-With-Mumps-395189031.html

Harvard Mumps Outbreak - all students vaccinated

http://m.huffpost.com/us/entry/us_57276bc7e4b0b49df6abc402

Mumps outbreak Whitworth University. All 3 students fully vaccinated.

http://www.khq.com/story/33328845/three-confirmed-mumps-cases-in-whitworth-university-students

Mumps outbreak - 31 students all vaccinated

http://www.kansascity.com/news/state/missouri/article115467353.html

Measles virus sheds for 1-13 days after vaccination:
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC228449/

Measles outbreak in a fully immunized secondary-school population
http://www.ncbi.nlm.nih.gov/pubmed/3821823

Catching measles in an appropriately vaccinated group: a well-circumscribed outbreak in the South East of Ireland, September-November 2013
http://www.ncbi.nlm.nih.gov/m/pubmed/27431259/

Measles vaccinated child responsible for outbreak in British Columbia:
http://www.eurosurveillance.org/images/dynamic/EE/V18N49/art20649.pdf

New York Measles outbreak linked to vaccinated:
http://cid.oxfordjournals.org/content/early/2014/02/27/cid.ciu105

Measles outbreak among the vaccinated:
http://www.ncbi.nlm.nih.gov/pubmed/8053748

Polio shedding:

**Shedding of virulent poliovirus revertants during immunization with oral poliovirus vaccine after prior immunization with inactivated polio vaccine.**

"Fecal shedding of revertant poliovirus after OPV challenge was observed in 50%-100% of subjects previously immunized with > or = 3 doses of the EIPV. These findings suggest that prior immunization with EIPV does not prevent fecal shedding of revertant polioviruses after subsequent reexposure to OPV."

https://www.ncbi.nlm.nih.gov/m/pubmed/8228342/

We don't know for certain how long shedding occurs because we don't test for it long term or regularly but in rare instances, it has gone on for years: http://www.westernmorningnews.co.uk/Vaccinated-man-spread-polio-30-years/story-27693988-detail/story.html

**Rotavirus vaccines: viral shedding and risk of transmission.**

https://www.ncbi.nlm.nih.gov/m/pubmed/18922486/

Detection of fecal shedding of rotavirus vaccine in infants following their first dose of pentavalent rotavirus vaccine

http://www.sciencedirect.com/science/article/pii/S0264410X11004567

"N. meningitidis inhabits the mucosal membrane of the nose and throat, where it usually causes no harm. Up to 5-10% of a population may be asymptomatic carriers. These carriers are crucial to the spread of the disease as most cases are acquired through exposure to asymptomatic carriers."

"The disease mainly affects young children, but is also common in older children and young adults."

http://www.who.int/csr/disease/meningococcal/en/

Additionally, the Dtap/Tdap is NOT a live virus vaccine, and has been shown by the FDA to cause the vaccinated to become asymptomatic carriers whenever exposed to the bacteria (not virus) for both pertussis and diphtheria, thus spreading the illness without knowing (carrying the bacteria in their throat).

Pertussis carrier:
http://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm376937.htm

"Study: Is the whooping cough resurgence due to vaccinated people not knowing they're infectious?"

https://web.archive.org/web/20160123024723/http://www.santafe.edu/news/item/althouse-scarpino-whooping-cough-asymptomatic/

Diphtheria carrier: http://www.cdc.gov/diphtheria/clinicians.html

You can also find that most medical facilities are aware of this. Johns Hopkins and St. Jude hospitals are just a few of many who post precautions for recently vaccinated visitors.

Why does this matter?

Because society has become so deathly afraid of these illnesses that they rush out and load up on vaccines and want to even pass laws forcing others to do the same, but the science shows that it's the vaccinated that are at a higher risk of infections and spreading the illnesses. Be aware of the infection and carrier risk each vaccine has when making your choice, and take care when around the immune compromised if you have been recently vaccinated or around someone with the illnesses even if you don't show symptoms.

http://www.vaccinationcouncil.org/2011/11/17/smoke-mirrors-and-the-disappearance-of-polio/

https://therefusers.com/studies-show-measles-is-spread-by-vaccinated-individuals/

https://web-beta.archive.org/web/20160324212629/http://www.cnbc.com/2015/03/03/globe-newswire-public-health-officials-know-recently-vaccinated-individuals-spread-disease.html

Shingles (Zostavax} - Shedding (Section 5.2)
http://www.merck.com/product/usa/pi_circulars/z/zostavax/zostavax_pi2.pdf

https://www.facebook.com/permalink.php?story_fbid=233005433804554&id=232581190513645

http://www.news-medical.net/news/2005/09/01/12896.aspx

https://www.sciencedaily.com/releases/2015/08/150811103555.htm

This is gross:

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3096786/

HIB

Via Elizabeth Aven:

"Hib

The chances of a child under 5 getting bacterial meningitis is pretty slim...according to the CDC, fewer than 55 cases per year are in children under 5 in the U.S. (And fewer than 5 deaths). So that's over a 1 in 5 million chance. And they are estimating.

http://www.cdc.gov/vaccines/vpd-vac/hib/downloads/dis-hib-color-office.pdf

According to WHO however, in 2010 there were only 8 reported cases and in 2011 there were only 2 reported cases in the U.S. http://apps.who.int/immunization_monitoring/globalsummary/timeseries/tsincidencehib_mening.html

2 deaths in 2006 and 2007 combined

http://www.health.gov.au/internet/publications/publishing.nsf/Content/cda-cdi34suppl.htm~cda-cdi34suppl-3-vpd.htm~cda-cdi34suppl-3-vpd2.htm

Most older children and adults who develop Hib meningitis have underlying medical conditions that interfere with immune function.

http://emedicine.medscape.com/article/1164916-overview#a4

The vaccine only "protects"against a few strains...there are many other strains out there. This vaccine also has a lot of reported side effects on VAERS...and on average about 38 people die yearly after getting this vaccine according to VAERS (38 from the vaccine, just a few from the disease itself)...and remember, VAERS is severely under reported!

VAERS received 29,747 reports after Hib vaccines; 5179 (17%) were serious, including 896 reports of deaths. http://www.ncbi.nlm.nih.gov/pubmed/25598306

Hib is not as contagious as the cold or the flu...my friend had it right after she gave birth and her breastfed baby never got it. Many healthy people carry this bacteria in their nose and throat and they never get sick.

http://www.cdc.gov/meningitis/bacterial.html

So if you get the bacteria, that doesn't always mean you will get meningitis. If your immune system is strong, it will fight the bacteria off before it reaches the blood, and then the brain. Things like garlic, colloidal silver, olive leaf extract, chlorella, and ginseng are all great things to take to prevent serious complications if someone does come down with bacterial meningitis (in addition to medical treatment). These

things help prevent the bacteria from spreading and damaging cells and they help fight it off.

Read about some natural remedies here:

http://www.healthnfairness.com/2014/09/herbal-remedies-for-meningitis.html?m=1

A vaccine bypasses the bodies first line of defense, putting that disease straight into the bloodstream. Once in the bloodstream it has easy access to vital organs and can cause meningitis. It also depletes your bones and organs of vital vitamins. Many vaccine inserts list meningitis as reported reactions to the vaccine. The best defense is a strong immune system!

"The precise level of antibody required for protection against invasive disease is not clearly established. "

"Secondary Hib disease is defined as illness occurring 1-60 days following contact with an ill child, and accounts for less than 5% of all invasive Hib disease. Among household contacts, six studies have found a secondary attack rate of 0.3% in the month following onset of the index case, which is about 600-fold higher than the risk for the general population. Attack rates varied substantially with age, from 3.7% among children 2 years of age and younger to 0% among contacts 6 years of age and older. In these household contacts, 64% of secondary cases occurred within the first week (excluding the first 24 hours) of disease onset in the index patient, 20% during the second week, and 16% during the third and fourth weeks."
http://www.cdc.gov/vaccines/pubs/pinkbook/hib.html

Pentacel (HIB combo vaccine they wanted to give my son) had a 1 in 25 serious reaction rate according to one of the studies. The top reported reactions were pneumonia, asthma, bronchiolitis, gastroenteritis, and dehydration. 1 in 1,196 died (they aren't sure if its related or not, but that's the facts and the potential risks).

Menhibrix had a 1 in 54 serious adverse reaction rate within a 31 day period following vaccination. 1 in 20 had a serious reaction after a 6 month period.

A total of 443 cases of Hib infection occurred in children eligible for vaccination; 363 (82%) were fully vaccinated. The incidence of Hib infection has been increasing predominantly in vaccinated children
http://jid.oxfordjournals.org/content/188/4/481.full

Hib in two vaccinated children
http://www.ncbi.nlm.nih.gov/m/pubmed/14743044/

Clustering of cases of insulin dependent diabetes (IDDM) occurring three years after hemophilus influenza B (HiB) immunization support causal relationship between

immunization and IDDM. http://www.greenmedinfo.com/article/exposure-hib-immunization-associated-increased-risk-insulin-dependent-diabetes

To date, no one has examined the global change in Hib disease epidemiology and whether Hib diseases are decreasing due to vaccination.
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC100154/

Vaccinated still get it http://www.greenmedinfo.com/article/between-may-1985-and-september-1987-228-reports-disease-due-haemophilus

"among Hib case-patients aged <5 years with age-appropriate vaccine status reported during 2002–2012 in the United States, 16% had completed the primary Hib series, and 43% had completed the full Hib series"
http://www.cdc.gov/mmwr/preview/mmwrhtml/rr6301a1.htm"

Information via Elizabeth Aven:

Meningococcal Meningitis:

http://theviennareport.us/meningitis/

http://vaxtruth.org/2011/10/meningococcal/

The manufacturer product inserts for meningococcal vaccine list adverse events reported during clinical trials or post licensure, including irritability, abnormal crying, fever, drowsiness, fatigue, injection site pain and swelling, sudden loss of consciousness (syncope), diarrhea, headache, joint pain, Guillain Barre Syndrome, brain inflammation, convulsions, and facial palsy.
As of September 1, 2015, there had been 47 claims filed in the federal Vaccine Injury Compensation Program (VICP) for injuries and deaths following meningococcal vaccination, including 2 deaths and 45 serious injuries.
Using the MedAlerts search engine, as of September 30, 2015, the federal Vaccine Adverse Events Reporting System (VAERS), which includes only a small fraction of the health problems that occur after vaccination in the U.S., had recorded more than 1,846 serious health problems, hospitalizations and injuries following meningococcal shots, including 99 deaths with about 34% of the deaths occurring in children under age six.

http://www.nvic.org/Vaccines-and-Diseases/Meningitis.aspx

The vaccine insert for it is listed here. That alone is startling.

http://www.immunize.org/fda/

Meningococcal Meningitis

550 total cases of meningococcal DISEASE in 2013 in the US (not all turned into meningitis).

Meningitis is observed in approximately 50% of invasive cases.

http://www.cdc.gov/vaccines/pubs/surv-manual/chpt08-mening.html

So that's a 1 in 575,000 chance of contracting...and over a 1 in 1,000,000 chance of it developing into meningitis.

Anyone can get meningococcal disease, but rates of disease are highest in children younger than 1 year, followed by a second peak in adolescence.

About 10-15 out of 100 will die according to the CDC.

http://www.cdc.gov/meningococcal/clinical-info.html

But according to WHO, 5-10 out of 100 will die:

"Even when the disease is diagnosed early and adequate treatment is started, 5% to 10% of patients die..."

http://www.who.int/mediacentre/factsheets/fs141/en/

Cases: 450 (0.14/100,000)
Deaths: 65 (0.02/100,000)

¥ In 2014, 564 cases of meningococcal disease (confirmed and probable cases) were reported to the National Notifiable Disease Surveillance System (incidence 0.18/100,000 population)

So that's a 1 in 565,000 (or 1 in 629,000 if you go by confirmed cases) chance of contraction and over a 1 in 1,000,000 chance of it developing into meningitis. Over 1 in 4,900,000 died.

http://www.cdc.gov/abcs/reports-findings/survreports/mening14.html

VACCINES

Menactra (meningococcal vaccine for groups A, C, Y, W-135) has a serious adverse event rate of: 1 in 40-50 (children 9-12 months), 1 in 167 (children 2-10), and 1 in 100 (11-18). Guillain-Barre syndrome (autoimmune disease where the immune system attacks the nerves, can result in paralysis) has been reported in temporal relationship following this vaccine.

Menomune (vaccine for groups A, C, Y, W-135) had a serious adverse event rate of: 1 in 152 (2-10 years old), 1 in 168 (11-18 years old), 1 in 59 (18-55 years old). Some reported reactions are Guillain-Barré syndrome, paresthesia (tingling or numbness usually caused by neurological disease or traumatic nerve damage), and dyspnea (difficulty breathing).

Menveo (vaccine for groups A, C, Y, W-135) had a serious adverse event rate of: 1 in 37 (during infant series, whereas 1 in 45 had a SAE during routine vaccinations without Menveo). The most common SAE's reported were wheezing, pneumonia, gastroenteritis, and convulsions. At 12 months of age the rate of SAE's were 1 in 26. 1 in 4,000 were diagnosed with Kawasaki disease (autoimmune disease dealing with inflammation of blood vessels), 1 out of 12,049 was diagnosed with acute disseminated encephalomyelitis (intense attack of inflammation in the brain and spinal cord), and 1 in around 6,000 died (sudden death and sepsis reported, none were assessed as related but there were no deaths in the control group). In children 2-10 years, the SAE rate was 1 in 137. There were several cases of pneumonia and appendicitis. Staph infection and convulsions were also reported. For subjects 11-55 years old, the SAE rate was 1 in 155. Several cases of appendicitis were reported, as well as other syndromes and diseases.

Bexsero (meningococcal group B vaccine) had a SAE rate of 1 in 46.

Trumenba (meningococcal group B vaccine) had a SAE rate of 1 in 49.

Vaccine inserts found here:
http://www.immunize.org/packageinserts/

"N. meningitidis inhabits the mucosal membrane of the nose and throat, where it usually causes no harm. Up to 5-10% of a population may be asymptomatic carriers. These carriers are crucial to the spread of the disease as most cases are acquired through exposure to asymptomatic carriers."

"The disease mainly affects young children, but is also common in older children and young adults."

http://www.who.int/csr/disease/meningococcal/en/

Countries at risk are in Africa
http://gamapserver.who.int/mapLibrary/Files/Maps/Global_MeningitisRisk_ITHRiskMap.png?ua=1&ua=1

While all the risk factors for meningococcal outbreaks in Africa are not understood, several conditions have been associated with the development of epidemics in the meningococcal belt. They include:

Medical conditions: immunological susceptibility of the population

Demographic conditions: travel and large population displacements
Socioeconomic conditions: poor living conditions and overcrowded housing
Climatic conditions: drought and dust storms
http://www.cdc.gov/meningococcal/global.html

Five to ten percent of adults are asymptomatic nasopharyngeal carriers of N. meningitidis. The frequency of carriage, like that of invasive disease, also varies by age. Adolescents and young adults have the highest rates of meningococcal carriage. Although asymptomatic carriage of both pathogenic and nonpathogenic strains is common, few carriers develop invasive disease. For the majority of people, carriage is an immunizing process that results in a systemic, serogroup-specific protective antibody response.

http://www.cdc.gov/meningococcal/clinical-info.html

The bacteria can be carried in the throat and sometimes, for reasons not fully understood, can overwhelm the body's defenses allowing infection to spread through the bloodstream to the brain. It is believed that 10% to 20% of the population carries Neisseria meningitidis in their throat at any given time. However, the carriage rate may be higher in epidemic situations.

The largest burden of meningococcal disease occurs in an area of sub-Saharan Africa known as the meningitis belt, which stretches from Senegal in the west to Ethiopia in the east

http://www.who.int/mediacentre/factsheets/fs141/en/

There is a natural cyclical pattern of meningococcal disease with peaks of disease occurring every 7-10 years (CDC, unpublished data). Current rates of meningococcal disease are at historic lows; the disease pattern of the past 10 years is outside the previously observed periodicity of disease.

Risk factors for meningococcal disease include organism, host, and environmental factors. Persons with persistent complement component deficiencies (e.g., C5-C9, properidin, factor H, or factor D) or functional or anatomic asplenia are at increased risk for invasive meningococcal disease.

Meningococcal disease rates in children younger than 1 year peak at 0-6 months. More than 50% of meningococcal disease in children 0-6 months is caused by serogroup B; serogroup Y is also more prevalent in this age group. In time, children gradually become exposed to meningococci and develop bactericidal antibodies. By the time they reach adulthood, 65%-85% of persons possess bactericidal antibody against meningococcal disease.

The use of antibiotics has dramatically reduced mortality due to meningococcal disease.

http://www.cdc.gov/vaccines/pubs/surv-manual/chpt08-mening.html

About 1 out of 10 people have this type of bacteria in the back of their nose and throat with no signs or symptoms of disease; this is called being 'a carrier'. But sometimes Neisseria meningitidis bacteria can invade the body causing certain illnesses, which are known as meningococcal disease.

Fortunately, these bacteria are not as contagious as germs that cause the common cold or the flu. The bacteria are not spread by casual contact or by simply breathing the air where a person with meningococcal disease has been.

http://www.cdc.gov/meningococcal/about/causes-transmission.html

Several different bacteria can cause meningitis. Neisseria meningitidis is the one with the potential to cause large epidemics. There are 12 serogroups of N. meningitidisthat have been identified, 6 of which (A, B, C, W, X and Y) can cause epidemics. Geographic distribution and epidemic potential differ according to serogroup.

http://www.who.int/mediacentre/factsheets/fs141/en/

Polio History:

THE HEALTH FINDER 1956
J.I. Rodale
Polio and Inoculations
Page 594. “A few days ago we received our first copy of the British medical journal. The Lancet, (issue of April 29, 1950) on our new subscription, and I was amazed to find in it an article called "Inoculation and Poliomyelitis," which draws attention to the association between various inoculations given for immunization against whooping cough and diphtheria, and resulting in polio.

"Professor Burnet contended that McCloskey had clearly shown in the 1949 epidemic in Victoria that paralytic poliomyelitis in the injected limb occurred within two months of inoculation with pertussis vaccine, either alone or in combination with diphtheria."

" Much evidence was brought forward at this meeting that the Australian experience was not unique but that there were many polio cases in England which had been tracked down to inoculations, of various kinds, in one instance to a series of measles cases."

Page 599. "Poliomyelitis and other diseases are caused by wrong personal habits of living and unfavorable environment. DISEASE GERMS CANNOT TAKE HOLD IN A HEALTHY BODY. That is certainly true for polio and all the common contagions

(measles, scarlet fever, whooping cough, diphtheria, syphilis, tuberculosis, etc. When I eat the correct foods in the minimum amounts to supply my essential requirements and drink enough water, I fear no infection whatever."

“In 1908 Dr. Karl Landsteiner, Viennese pathologist, published his theory, still held today, that it is a "virus" that launches epidemic polio. Landsteiner based his theory on the fact that he had been able to induce polio in monkeys by injecting them with nerve tissues from fatal human cases. But, believing that "the poliomyelitis produced in experimental animals is a pseudo-poliomyelitis produced by a pseudo-virus," Dr. Scobey reasoned that "the disease they produced in the monkey was an artificial disease and not a natural illness as it occurred in man." After exhausting medical literature in the form of case-histories of the subject in all lands and at all known times of its appearance, he concluded: It never has been proved that the "virus" of polio can cause that disease in human beings."

"It has never been explained how persons with the disease who have no contact with no one else having the disease got it."
Page 658.
" In talking about various theories as to the cause of polio the author says, “I am very glad to note that our ideas are being upheld by these men in Europe because the VIRUS THEORY HAS BEEN SO DEEPLY EMBEDDED IN THE MINDS OF THE AMERICAN PEOPLE BY THE NATIONAL FOUNDATION FOR INFANTILE PARALSIS. (This was 1956)You will recall that the Australian and British physicians forced the recognition in this country of a relationship between inoculations and polio."

http://vaccinationcouncil.org/2011/11/17/smoke-mirrors-and-the-disappearance-of-polio/

Book - "Dissolving Illusions" by Dr. Suzanne Humphries MD.

FDA Alum adjuvants:

How about some math along with some science.
Perhaps asking why it's magically ok to give an unsafe amount of alum adjuvant to to a baby as well. The FDA determined 5 mcg per pound of alum adjuvant to be the safe daily injectable (IV) amount (http://www.drdahlman.com/aluminum-in-vaccines/). The hep b vaccine is 250 mcg. So a 7 lb baby would have a max of 35 mcgs. How does 7x the safe amount magically end up being okay? vaccines go up to 850 mcg's also. Doctors are giving multiples at once with zero safety tests ever having been done in that regard. Another question is, how and why the FDA could approve vaccines having this knowledge of the math? Here are a few of the thousands (yes thousands) of peer reviewed scientific studies explaining the many dangers that vaccines pose to humans and animals alike:

Links in regard to above statement in regard to FDA determined 5 mcg per pound injected (IV) alum adjuvants. No studies outside of IV injection of alum adjuvants have been done:

http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/CFRSearch.cfm?fr=201.323

http://www.ashp.org/menu/News/PharmacyNews/NewsArticle.aspx?id=1537

http://www.nejm.org/doi/pdf/10.1056/NEJM199705293362203

INGESTION VS INJECTION

"In healthy subjects, only 0.3% of orally administered aluminum is absorbed via the GI tract, and the kidneys effectively eliminate aluminum from the human body. Only when the GI barrier is bypassed, such as by intravenous infusion or in the presence of advanced renal dysfunction, does aluminum have the potential to accumulate. As an example, with intravenously infused aluminum, 40% is retained in adults and up to 75% is retained in neonates."
"If a significant aluminum load exceeds the body's excretory capacity, the excess is deposited in various tissues, including bone, brain, liver, heart, spleen, and muscle. This accumulation causes morbidity and mortality through various mechanisms."

https://www.ncbi.nlm.nih.gov/pubmed/26948677

Aluminum vaccine adjuvants: are they safe?

"Despite almost 90 years of widespread use of aluminum adjuvants, medical science's understanding about their mechanisms of action is still remarkably poor. There is also a concerning scarcity of data on toxicology and pharmacokinetics of these compounds. In spite of this, the notion that aluminum in vaccines is safe appears to be widely accepted. Experimental research, however, clearly shows that aluminum adjuvants have a potential to induce serious immunological disorders in humans."
http://www.ncbi.nlm.nih.gov/m/pubmed/21568886/

**A possible central mechanism in autism spectrum disorders, part 1**
http://www.ncbi.nlm.nih.gov/pubmed/19043938

**The role of mercury in the pathogenesis of autism.**
http://www.ncbi.nlm.nih.gov/pubmed/12142947

**Transcriptomic analyses of neurotoxic effects in mouse brain after intermittent neonatal administration of thimerosal**
http://www.ncbi.nlm.nih.gov/pubmed/24675092

**A dose-response relationship between organic mercury exposure from thimerosal-containing vaccines and neurodevelopmental disorders**
http://www.ncbi.nlm.nih.gov/pubmed/25198681

**Immunological findings in autism.**
http://www.ncbi.nlm.nih.gov/m/pubmed/16512356/

**A two-phase study evaluating the relationship between Thimerosal-containing vaccine administration and the risk for an autism spectrum disorder diagnosis in the United States**
http://www.ncbi.nlm.nih.gov/m/pubmed/24354891/

**Influence of pediatric vaccines on amygdala growth and opioid ligand binding in rhesus macaque infants: A pilot study**
http://www.ncbi.nlm.nih.gov/m/pubmed/20628439/

**Gender-selective toxicity of thimerosal**
http://www.ncbi.nlm.nih.gov/m/pubmed/18771903/

**The Role of the Immune System in Autism Spectrum Disorder.**
http://www.ncbi.nlm.nih.gov/m/pubmed/27534269/

**Comparison of VAERS fetal-loss reports during three consecutive influenza seasons: was there a synergistic fetal toxicity associated with the two-vaccine 2009/2010 season?**
http://www.ncbi.nlm.nih.gov/m/pubmed/23023030/

**Adverse event reports after tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccines in pregnant women.**
http://www.ncbi.nlm.nih.gov/m/pubmed/22727350/

**Comparison of VAERS fetal-loss reports during three consecutive influenza seasons**
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3888271/

**Administration of aluminium to neonatal mice in vaccine-relevant amounts is associated with adverse long term neurological outcomes.**
http://www.ncbi.nlm.nih.gov/pubmed/23932735

**Prenatal environmental exposures, epigenetics, and disease**
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3171169/

**Efficacy of Quadrivalent HPV Vaccine against HPV Infection and Disease in Males.**
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3495065/

**B-Lymphocytes from a Population of Children with Autism Spectrum Disorder and Their Unaffected Siblings Exhibit Hypersensitivity to Thimerosal**
http://www.hindawi.com/journals/jt/2013/801517/

Genetic markers and toxins in vaccines link to autism
http://www.ncbi.nlm.nih.gov/pubmed/23576057

Heavy Metals in Hair Samples from Severely Autistic Children
http://www.mdpi.com/1660-4601/9/12/4486

Conjugate Vaccines may predispose children to autism disorders
http://www.ncbi.nlm.nih.gov/pubmed/23576057

Abnormal measles-mumps-rubella antibodies and CNS autoimmunity in children with autism.
http://www.ncbi.nlm.nih.gov/pubmed/12145534

A positive association found between autism prevalence and childhood vaccination uptake across the U.S.
http://www.ncbi.nlm.nih.gov/pubmed/21623535

Causal relationship between vaccine induced immunity and autism
http://www.ncbi.nlm.nih.gov/pubmed/12849883

Subtle DNA changes and the overuse of vaccines in autism
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3364648/

Vaccine and Autism- a New Scientific Review
http://www.cbsnews.com/…/vaccines-and-autism-a-new-scienti…/

Summary of previous Journal of Immunology
http://danmurphydc.com/…/01/AR-10-12-rata-AUTISM-VACCINE.pdf

Autism and Resulting Medical Conditions:
http://www.tacanow.org/…/2…/09/autism-studies-april-2008.pdf

Mercury toxic encephalopathy manifesting with clinical symptoms of regressive autistic disorders. http://www.ncbi.nlm.nih.gov/pubmed/17454560

Relation of mercury to high autism rates in boys
http://www.ncbi.nlm.nih.gov/pubmed/16264412

Elevated levels of measles in children with Autism
http://www.ncbi.nlm.nih.gov/pubmed/12849883

Abnormal MMR antibodies in children with autism
http://www.ncbi.nlm.nih.gov/pubmed/12145534

Tylenol, MMR and Autism - A parent survey study
http://www.ncbi.nlm.nih.gov/pubmed/18445737

A Positive Association found between Autism Prevalence and Childhood Vaccination
http://www.ingentaconnect.com/…/2011/000…/00000014/art00002…

Peer reviewed study on fetal cell contamination with retro virus associated with autism and cancer
http://www.globalresearch.ca/new-study-in-journal-o…/5402912

Study documentation- Dr Deisher
http://www.ms.academicjournals.org/…/article1409245960_Deis…

Autism and mercury poisoning
http://www.ncbi.nlm.nih.gov/pubmed/11339848

Hypothesis: conjugate vaccines may predispose children to autism spectrum disorders
http://www.ncbi.nlm.nih.gov/pubmed/21993250

Rise in autism coincides with rise in vaccines
http://www.ncbi.nlm.nih.gov/pubmed/21623535

A two-phase study evaluating the relationship between Thimerosal-containing vaccine administration and the risk for an autism spectrum disorder diagnosis in the United States
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3878266/

A positive association found between autism prevalence and childhood vaccination uptake across the U.S. population.
http://www.ncbi.nlm.nih.gov/pubmed/21623535

Commentary--Controversies surrounding mercury in vaccines: autism denial as impediment to universal immunisation.
http://www.ncbi.nlm.nih.gov/pubmed/25377033

Methodological issues and evidence of malfeasance in research purporting to show thimerosal in vaccines is safe.
http://www.ncbi.nlm.nih.gov/pubmed/24995277

Abnormal measles-mumps-rubella antibodies and CNS autoimmunity in children with autism.
http://www.ncbi.nlm.nih.gov/pubmed/12145534

Hepatitis B vaccination of male neonates and autism diagnosis, NHIS 1997-2002.
http://www.ncbi.nlm.nih.gov/pubmed/21058170

Do aluminum vaccine adjuvants contribute to the rising prevalence of autism?
http://www.ncbi.nlm.nih.gov/pubmed/22099159

What is regressive autism and why does it occur? Is it the consequence of multi-systemic dysfunction affecting the elimination of heavy metals and the ability to regulate neural temperature?
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3364648/

A case series of children with apparent mercury toxic encephalopathies manifesting with clinical symptoms of regressive autistic disorders.
http://www.ncbi.nlm.nih.gov/pubmed/17454560

A comprehensive review of mercury provoked autism.
http://www.ncbi.nlm.nih.gov/pubmed/19106436

Thimerosal Exposure and the Role of Sulfation Chemistry and Thiol Availability in Autism
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3774468/

B-Lymphocytes from a Population of Children with Autism Spectrum Disorder and Their Unaffected Siblings Exhibit Hypersensitivity to Thimerosal
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3697751/

Theoretical aspects of autism: causes--a review.
http://www.ncbi.nlm.nih.gov/pubmed/21299355

Conjugate vaccines and autism.
http://www.ncbi.nlm.nih.gov/pubmed/21907498

Autism: a novel form of mercury poisoning
http://www.ncbi.nlm.nih.gov/pubmed/11339848

A prospective study of thimerosal-containing Rho(D)-immune globulin administration as a risk factor for autistic disorders.
http://www.ncbi.nlm.nih.gov/pubmed/17674242

The potential importance of steroids in the treatment of autistic spectrum disorders and other disorders involving mercury toxicity.
http://www.ncbi.nlm.nih.gov/pubmed/15780490

126 Research Papers Supporting the vaccine/autism link:

https://www.scribd.com/book/220807175

Identical Twin SIDS
http://www.ncbi.nlm.nih.gov/m/pubmed/17654772/

Allergies vax
http://www.ncbi.nlm.nih.gov/m/pubmed/10714532/

Interesting research studies
http://www.sciencedirect.com/science?_ob=ArticleListURL&_method=list&_ArticleListID=-1023520050&_sort=v&_st=17&view=c&_origin=related_art&panel=citeRelatedArt&_mlktType=Journal&md5=1c6103c0140faf460ee9fba5daae373d&searchtype=a

Diabetes and vax
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC1116914/

" vaccines have a modest effect in reducing influenza symptoms and working days lost. There is no evidence that they affect complications, such as pneumonia, or transmission. WARNING: This review includes 15 out of 36 trials funded by industry (four had no funding declaration). An earlier systematic review of 274 influenza vaccine studies published up to 2007 found industry funded studies were published in more prestigious journals and cited more than other studies independently from methodological quality and size. Studies funded from public sources were significantly less likely to report conclusions favorable to the vaccines. The review showed that reliable evidence on influenza vaccines is thin but there is evidence of widespread manipulation of conclusions and spurious notoriety of the studies. The content and conclusions of this review should be interpreted in light of this finding."

http://www.ncbi.nlm.nih.gov/m/pubmed/20614424/

"Increased risk of noninfluenza respiratory virus infections associated with receipt of inactivated influenza vaccine."
http://www.ncbi.nlm.nih.gov/m/pubmed/22423139/

"Increased Risk of Noninfluenza Respiratory Virus Infections Associated With Receipt of Inactivated Influenza Vaccine"

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3404712/

**Association between the 2008–09 Seasonal Influenza Vaccine and Pandemic H1N1 Illness during Spring–Summer 2009: Four Observational Studies from Canada**

http://journals.plos.org/plosmedicine/article?id=10.1371/journal.pmed.1000258

http://www.nvic.org/NVIC-Vaccine-News/April-2016/cdc-admits-flu-shots-fail-half-the-time.aspx

https://liveto110.com/flu-vaccines-are-toxic/

The flumist live virus vaccine was causing a mutation that was causing severe flu in mice, that's why the CDC doesnt recommend it any longer, not to mention it was shedding the flu virus to others.." we found that the FluMist vaccine backbone could regain virulence to cause severe disease in mice. The revertant also regained virulence and caused significant disease in mice, with severity comparable to that caused by a wild type 2009 H1N1 pandemic virus."

https://www.ncbi.nlm.nih.gov/pubmed/27440882

**Insight: Evidence grows for narcolepsy link to GSK swine flu shot**

http://mobile.reuters.com/article/idUSBRE90L07H20130122

**AS03 Adjuvanted AH1N1 Vaccine Associated with an Abrupt Increase in the Incidence of Childhood Narcolepsy in Finland**
http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0033536#close

**Increased Incidence and Clinical Picture of Childhood Narcolepsy following the 2009 H1N1 Pandemic Vaccination Campaign in Finland**
http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0033723

Thimerisol/Mercury brain damage:

http://personalhealthdiary.co/dementia-now-striking-people-in-their-40s-as-mercury-from-vaccines-causes-slow-degenerative-brain-damage/

http://www.royalrife.com/flu_shots.html

Pregnancy:

Flu Vaccine and miscarriage increase.

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3888271/

Maternal Transfer of Mercury on Developing Fetus

http://www.tandfonline.com/doi/full/10.1080/02772248.2012.724574#.VHEJAIvF_pc

2 vaccines cause increase in fetal deaths in 2009/2010

http://www.ncbi.nlm.nih.gov/pubmed/23023030

Pertussis epidemic despite high levels of vaccination coverage with acellular pertussis vaccine.

http://www.ncbi.nlm.nih.gov/m/pubmed/24216286/

"Further, we show that aP vaccination impedes host immunity against B. parapertussis-measured as reduced lung inflammatory and neutrophil responses. Thus, we conclude that aP vaccination interferes with the optimal clearance of B. parapertussis and enhances the performance of this pathogen. Our data raise the possibility that widespread aP vaccination can create hosts more susceptible to B. parapertussis infection."

http://www.ncbi.nlm.nih.gov/m/pubmed/20200027/

**Acellular pertussis vaccines protect against disease but fail to prevent infection and transmission in a nonhuman primate model**

"The observation that aP, which induces an immune response mismatched to that induced by natural infection, fails to prevent colonization or transmission provides a plausible explanation for the resurgence of pertussis and suggests that optimal control of pertussis will require the development of improved vaccines."

http://m.pnas.org/content/111/2/787.abstract

“Our unvaccinated and under-vaccinated population did not appear to contribute significantly to the increased rate of clinical pertussis. Surprisingly, the highest incidence of disease was among previously vaccinated children in the eight to twelve year age group.”

http://www.ncbi.nlm.nih.gov/pubmed/22423127

Vaccinations create more powerful and virulent strains of bacteria and viruses. The reason for the current whooping cough outbreak. Read more here from the CDC"Vaccination against 2 avian viruses, the Marek disease virus, and the infectious bursal disease virus, were associated with the emergence of more virulent strains (33). An important role of host immunity in selecting for virulence is also suggested by the co-evolution of the myxomatosis virus and rabbits (34). Furthermore, immune pressure was shown to select for more virulent Plasmodium chabaudi parasites in mice (35). Based on mathematical modeling, vaccines designed to reduce pathogen growth rate and/or toxicity may result in the evolution of pathogens with higher levels of virulence.

(36)."http://wwwnc.cdc.gov/eid/article/15/8/08-1511_article.htm

"After the fifth dose of DTaP, the odds of acquiring pertussis increased by an average of 42% per year."

http://www.nejm.org/doi/full/10.1056/NEJMoa1200850#discussion

This study shows that efficacy of the DTaP falls rapidly. At 2 years post-vaccination, it's just 75%. By 5 years, it's down to 11.9%.

http://www.ncbi.nlm.nih.gov/pubmed/24903664

Pertussis 53 - 64% effective in adolescents and adults:

http://www.ncbi.nlm.nih.gov/pubmed/23873919
-the pertussis vaccine doesn't work, and any immunity from it wanes

Importantly, we demonstrate that acellular vaccine antigen-encoding genes are evolving at higher rates than other surface protein-encoding genes. This was true even prior to the introduction of pertussis vaccines but has become more pronounced since the introduction of the current acellular vaccines. The fast evolution of vaccine antigen-encoding genes has serious consequences for the ability of current vaccines to continue to control pertussis.

http://www.ncbi.nlm.nih.gov/m/pubmed/25489002/

Tetanus is anaerobic, so it dies when it hits oxygen. Bleeding from an injury is a good thing. Tetanus is more common where farm animal feces is present. If tetanus is suspected, an immune globulin shot can be administered. Getting a tetanus shot at the time of injury is a moot point. It would take several weeks to create antibodies, and if tetanus were present, death could occur within a week. Tetanus is extremely rare.

**Severe tetanus in immunized patients with high anti-tetanus titers**

https://www.ncbi.nlm.nih.gov/m/pubmed/1565228/

**Adverse event reports after tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccines in pregnant women.**

https://www.ncbi.nlm.nih.gov/m/pubmed/22727350/

https://www.sciencedaily.com/releases/2015/06/150624071018.htm

Is pertussis actually reemerging? Insights from an individual-based model

http://www.scielo.br/scielo.php?script=sci_arttext&pid=S0102-311X2001000300005&lng=en&nrm=iso&tlng=en

**Human papillomavirus vaccine and systemic lupus erythematosus.**

http://www.ncbi.nlm.nih.gov/pubmed/23624585

"We retrospectively described a case series including 18 girls (aged 12–24 years) referred to our "Second Opinion Medical Network" for the evaluation of "neuropathy with autonomic dysfunction" after HPV vaccination. All girls complained of long-lasting and invalidating somatoform symptoms (including asthenia, headache, cognitive dysfunctions, myalgia, sinus tachycardia and skin rashes) that have developed 1–5 days (*n* = 11), 5–15 days (*n* = 5) and 15–20 days (*n* = 2) after the vaccination. These cases can be included in the recently described immune dysfunction named autoimmune/inflammatory syndrome induced by adjuvants (ASIA)."
**https://link.springer.com/article/10.1007/s12026-016-8820-z**

"958 hospitalizations
+ 19,351 ED (emergency room visits)
= 20,309 serious adverse events"

So 20,000 of 195,000 girls who got HPV vaccines
---- OR 10% ----
had a SERIOUS enough reaction that it warranted a trip to the ER within 42 days of vaccination.

"Rates of AEFI [adverse reactions] after HPV immunization in Alberta are low and consistent with types of events seen elsewhere."

http://www.sciencedirect.com/science/article/pii/S0264410X16002036

**Behavioral abnormalities in female mice following administration of aluminum adjuvants and the human papillomavirus (HPV) vaccine Gardasil.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27421722/

**Efficacy of Quadrivalent HPV Vaccine against HPV Infection and Disease in Males.**

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3495065/

**Association between MTHFR gene polymorphisms and the risk of autism spectrum disorders: a meta-analysis.**

https://www.ncbi.nlm.nih.gov/m/pubmed/23653228/

Via Elizabeth Aven

https://www.facebook.com/permalink.php?story_fbid=238789313226166&id=232581190513645

https://www.facebook.com/permalink.php?story_fbid=238203156618115&id=232581190513645

Click "see more." I'm pinning this post and adding a list of common topics that come up for your easy reference. I have many more that I'm still adding! This will all go up on my site too.

Vaccines and autism -

https://www.facebook.com/permalink.php?story_fbid=236135150158249&id=232581190513645

Polio - the story is not what we all thought it was.

https://www.facebook.com/permalink.php?story_fbid=249477342157363&id=232581190513645&substory_index=0

Vaccines in pregnancy should be a felony.

https://www.facebook.com/permalink.php?story_fbid=240558636382567&id=232581190513645

How I went about getting a medical exemption.

https://www.facebook.com/permalink.php?story_fbid=241080596330371&id=232581190513645

2,500 scientific studies from the medical literature on the dangers of vaccines -

https://www.facebook.com/permalink.php?story_fbid=238789313226166&id=232581190513645

The myth of herd immunity

https://www.facebook.com/permalink.php?story_fbid=243340292771068&id=232581190513645

Very long list of doctors that have spoken out along the Vaxxed tour. FB will oddly only allow it to be shared from this link.

https://www.facebook.com/groups/1660084387554758/permalink/1897579363805258/

Question Everything Now's list of doctors that speak out on the Vaxxed tour.

https://www.facebook.com/permalink.php?story_fbid=238730049898759&id=232581190513645

Flu shot

https://www.facebook.com/permalink.php?story_fbid=244226056015825&id=232581190513645&substory_index=0

Autism in the non-vaccinated

https://www.facebook.com/permalink.php?story_fbid=249526025485828&id=232581190513645

"Vitamin" K shot controversy

https://www.facebook.com/permalink.php?story_fbid=241884052916692&id=232581190513645

MTHFR gene mutation

https://www.facebook.com/permalink.php?story_fbid=241072942997803&id=232581190513645

Vaccines didn't save us, sanitation did.

https://www.facebook.com/permalink.php?story_fbid=239785466459884&id=232581190513645

Gardasil, many scientific studies and whistleblowers

https://www.facebook.com/permalink.php?story_fbid=237720059999758&id=232581190513645

Peanut allergies and vaccines

https://www.facebook.com/permalink.php?story_fbid=235264986911932&id=232581190513645

Del Bigtree candidly speaks on the "Andrew Wakefield fraud" rumor

https://www.facebook.com/permalink.php?story_fbid=233539483751149&id=232581190513645

Lyme disease and autism

https://www.facebook.com/permalink.php?story_fbid=241180906320340&id=232581190513645

Food allergies and vaccines

https://www.facebook.com/permalink.php?story_fbid=233113110460453&id=232581190513645

Vaccines and autoimmunity

https://www.facebook.com/permalink.php?story_fbid=233049810466783&id=232581190513645

Mumps vaccine fraud

https://www.facebook.com/permalink.php?story_fbid=250032208768543&id=232581190513645&substory_index=0

How to get out of pet vaccines

https://www.facebook.com/permalink.php?story_fbid=233010990470665&id=232581190513645

Fluoride

https://www.facebook.com/permalink.php?story_fbid=239827059789058&id=232581190513645

Shingles, lawfirm

https://www.facebook.com/permalink.php?story_fbid=233005433804554&id=232581190513645

Meningitis and the meningococcal vaccine

https://www.facebook.com/permalink.php?story_fbid=239805066457924&id=232581190513645&substory_index=0

Babies commonly having strokes after vaccines. Dr. Moulden.

https://www.facebook.com/permalink.php?story_fbid=233006530471111&id=232581190513645&substory_index=0

Whooping cough, science

https://www.facebook.com/permalink.php?story_fbid=238203156618115&id=232581190513645

Do not sign the refusal to vaccinate form.

https://www.facebook.com/permalink.php?story_fbid=238779579893806&id=232581190513645

Hormone disruptors affect on hormones, gender, and orientation

https://www.facebook.com/permalink.php?story_fbid=243421799429584&id=232581190513645

Ways to address anxiety naturally

https://www.facebook.com/permalink.php?story_fbid=243303899441374&id=232581190513645&substory_index=0

Colonoscopies

https://www.facebook.com/permalink.php?story_fbid=239912783113819&id=232581190513645

Urinary tract infections

https://www.facebook.com/permalink.php?story_fbid=236538406784590&id=232581190513645&substory_index=0

Dr. Daniel Neides at the Cleveland Clinic.

https://www.facebook.com/permalink.php?story_fbid=249186625519768&id=232581190513645

Are vaccines safe? Dr. Suzanne Humphries. This is a good place to send people if they are just starting to research.

https://www.facebook.com/permalink.php?story_fbid=233649867073444&id=232581190513645

Genetics:

Genetic Risk Factors Associated with Vaccine Injury. These are not as rare as people might think.

The bulk of studies that have found associations between specific genetic variants and vaccine injury have identified genes related to immunological responses, as well as those associated with autoimmunity, and methylation. This makes sense, given that vaccines induce immunological reactions from both humoral (antibody-based immune reactions), as well as cell-mediated immune activation (immune responses involving cell-mediated inflammatory activity).

A study conducted in 2015 used existing literature to amass data related to genetic-based vaccine injury susceptibility. A landmark book published in 2015 "Vaccines & Autoimmunity" (Shoenfeld, Agmon-Levin & Tomljenovic) cites dozens of studies identifying various genetic variants of the HLA (human leukocyte antigen) gene family, as being strongly associated with vaccine-induced autoimmune activation. Some of these HLA variants include:

HLA DRB1
HLA DRB2
HLA DR4
HLA DRQ8

It has been well known that certain autoimmune processes can be triggered by vaccinations. This may be especially true of lupus, rheumatoid arthritis, macrophagic myofascitis, Guillain Barre syndrome, among others.

Modern genetics research has elucidated some of these heritable risks through the identification of the above-mentioned class of HLA gene variants. The HLA class of genes are strongly implicated in autoimmune processes. HLA's are involved in the detection and removal of antigens during infection. They signal various immunological responses to T-cells. Imbalances between the TH1 and TH2 immune branches are hallmark features of autoimmune processes.

An important study conducted in 2008 found strong statistical correlations between 6 genetic variants and vaccine injury following smallpox vaccine. These variants include:

3 variants of IL4 (interleukin 4)
2 variants of IRF1 (interferon regulatory factor)
1 variant of MTHFR, the C677T allele (methylene tetra hydrofolate reductase)
IL4 is a cytokine (immunologic signaling protein) involved in the cell-mediated inflammatory immune response. IRF1, is a transcription factor, involved in the release of cytotoxic interferon, and cell apoptosis (cell-programmed death). The MTHFR 677 variant identified in this study is central to the methylation cycle, folate metabolism, DNA repair, cell proliferation, as well as phase 2 detoxification reactions.

Other studies have identified additional cytokine-mediated genes, such as IL-1 and IL-18 (18) as being involved in adverse vaccine reactions.

Vaccines and Autoimmunity
https://www.amazon.com/dp/1118663438/ref=cm_sw_r_cp_api_0fT7xb0T6K26E

"The first genetic risk factor we studied is a mutation in GABAA, a receptor on both neurons and immune cells. Immune cells use this receptor to control the release of cytokines. When we mutate the GABAA gene in the mom and fetus, her pups do not behave differently. But *****when we trigger an immune response,*****(vaccines, anyone?) the mutation causes her immune cells to release too many cytokines, resulting in dramatic damage to the placenta, and extreme autistic-like behaviors in the pups."

http://scopeblog.stanford.edu/2016/10/11/on-genetics-immunology-and-autism-a-qa-with-stanfords-theo-palmer/

**Genetic Basis for Adverse Events Following Smallpox Vaccination.**

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2746083/

MTHFR Genetic Variant/Mutation

http://www.mommypotamus.com/mthfr-mutation/

http://www.vacfacts.info/mthfr-genetics-autism-and-disease.html

**Association between MTHFR gene polymorphisms and the risk of autism spectrum disorders: a meta-analysis.**

https://www.ncbi.nlm.nih.gov/m/pubmed/23653228/

MTHFR:

http://www.thefamilythathealstogether.com/vaccine-contraindications-six-people-not-vaccinated/

Many with autoimmune diseases often have the MTHFR mutation and other genetic mutations that make it so that people have difficulty detoxing these toxins out of the body. These genetic mutations increase the risk of developing some kind of autoimmune disorder from them or a long list of other injuries. GSTM1, HLA, and CBS are others.

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2843136/

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2722448/

http://jid.oxfordjournals.org/content/198/1/16.full

http://www.march-against-monsanto.com/the-truth-about-vaccines-facts-to-know-before-innoculating-your-child/

Autism and MTHFR

http://www.jpands.org/vol9no4/boris.pdf

http://www.vacfacts.info/mthfr-genetics-autism-and-disease.html

MTHFR and medications

https://myjourneywithmthfr.wordpress.com/2013/04/28/medications-that-could-be-dangerous-to-someone-with-mthfr/

CBS, HLA, Gardasil

http://sanevax.org/four-year-analysis-adverse-reactions-gardasil-hpv-vaccine/

GSTM1
Livewello

https://livewello.com/library/glutathione-s-transferases-gstm1-gstt1-gstp1-genes?id=6737166583988224

"These are the slides from a presentation that discusses concerns with MTHFR and Mercury vaccines like DPT and DTaP causing autism... it's pretty in-depth."

http://www.nationalacademies.org/hmd/~/media/4B8DAC4AD18F432283E67D91DB81F49B.ashx

How to read the book "Vaccines and Autoimmunity" & more information:

https://720d2911-a-62cb3a1a-s-sites.googlegroups.com/site/fdfhdf53/edc42/Vaccines-and-Autoimmunity.pdf?attachauth=ANoY7cqyvHUP7ezYAzy3VF9N3YXMsFnwh9YX8q_MuQCKskEcrO-tNWShSqNG7nKazg5rMT7XZoAWs5mfXmuBfJuf8UL6eEdX_1PYeBxY4a81pvWwWNpGQKjomdqIa8BKuW0oLHevv0XxEB7XuUSW-mlp-_wsZB1YzC3857zbiv_G3WIpkzSO_qCK4b0Ooy2gOdFY5DYgNLMzrLRR0FRcunlbHxL-bt8sPt-WEa_kTeNWOwYxP3akNRw%3D&attredirects=0

Online library
http://onlinelibrary.wiley.com/book/10.1002/9781118663721

Synopsis
http://cdn2.hubspot.net/hubfs/519118/Vaccines_Autoimmunity_draft3.pdf?t=1451507339294

Interview with Shoenfeld

http://bmcmedicine.biomedcentral.com/articles/10.1186/1741-7015-11-118

Article by Shoenfeld

http://maurizioproietti.eu/wp-content/uploads/2015/08/Vaccination-and-Autoimmunity.pdf

To read:

https://pdfs.semanticscholar.org/a69b/f88358f2fc057b1597cf8e5a868ed38c4e47.pdf

http://smjournals.com/vaccines/download.php?file=fulltext/smvvj-v1-1011.pdf

http://vaccinesafetycouncilminnesota.org/wp-content/uploads/2012/01/Mechanisms-of-aluminum-adjuvant-toxicity-and-autoimmunity-in-pediatric-populations.pdf

http://www.cmsri.org/wp-content/uploads/2015/08/1810-Soriano-Who-might-be-at-risk.-Pharmacol-Res-2014.pdf

https://www.ima.org.il/FilesUpload/IMAJ/0/95/47610.pdf

Studies to read

https://www.ncbi.nlm.nih.gov/m/pubmed/20193633/

Interview with Shoenfeld

http://bmcmedicine.biomedcentral.com/articles/10.1186/1741-7015-11-118

Article by Shoenfeld

http://maurizioproietti.eu/wp-content/uploads/2015/08/Vaccination-and-Autoimmunity.pdf

MTHFR

MTHFR - you will NOT know if your child has this and badly reacts to a vaccine until it's too late!

What is the 1# Risk Factor in Vaccine Injuries that the CDC will never tell you?

The public has not been informed about treatable genetic risk factors and life saving treatment options now available to everyone.

30% to 50% of the US population has a genetic polymorphism called MTHFR. This is a genetic defect that limits the bodies ability to converte folate into the active form utilize by the body. Many metabolic reactions are dependent on this methylated forms of folate.

Most people know folic acid is used in fordification, added to many processed foods. But, the best source of "folate" is found in green organic vegetables. If you have the MTHFR genetic mutation in your family, it's very important to identfy this genetic weekness early in life.

Note: The synthetic form of Folic acid added to most prenatal vitamins and may be a risk factor in pregnant women who have the MTHFR mutations.

Did you know Methylation is the holy grail to the human Metabolism, needed for detoxification of heavy metals, including toxic adjuvants now being added to most vaccine.

Individuals with MTHFR mutations have Methylation insufficiencies and impaired detoxification metabolism. The Methylation pathway is crucial for the production of Glutathione -- The Body's Master Anti-Oxidant. Individuals with the MTHFR defects are more susceptible to toxins as they have limited amounts Glutathione.

MTHFR MUTATION - IS A TREATABLE CONDITION

The first step in treating MTHFR mutations is to test for the MTHFR mutations. Genetic testing is the only reliable method for identifying this genetic risk factor.

"The treatment of MTHFR mutations is often a two-pronged approach. First, supplemental methylfolate and methylcobalamin directly address dysfunction in methylation pathways. Second, it is important to adopt appropriate lifestyle habits to down-regulate epigenetic expression of MTHFR mutations."

Note: Dosing requirements for methylfolate vary from person to person.

http://www.mthfrtreatment.com/index.html#.V_x0kVMVDVI

Another way to identify if you have MTHFR genetic risk factors is to ask your Doctor to test homocysteine as homocysteine levels will be elevated in people who have the MTHFR gene mutations.

Homocysteine is produced when the amino acid Methionine is broken down in the body. Elevated homocysteine levels increased the risk for hardening of the arteries, heart attack and stroke.

Why is MTHFR the 1# Risk Factor in Vaccine Injuries?

Many lifes have been destroy by toxic adjuvants being added to vaccines. We must understand vaccines are toxic by design, that's how they work

Vaccine adjuvants is a substance added to a vaccine to increase the body's immune response to the vaccine.

Many of these substance are neurotoxins and are destructive to nerve tissue. Common additives in vaccines are Aluminum phosphate, Thimerosal, Phenoxyethanol, Formaldehyde and Polysorbate 80 to name a few.

Vaccine manufactures are not required to use the same high standard testing as other pharmaceutical drugs. Vaccine are not double blind, placebo controlled tested.

If you think vaccines are safe I hope you enjoy being lied to, because vaccine adjuvants have never been proven safe.

Individuals who have the MTHFR mutations has limited detoxification abilties. The MTHFR genetic mutation may be the most significant risk factor in vaccine Injuries.

- Pediatric Doctors need to start testing for treatable genetic risk factors before any vaccine protocols are introduce into susceptible individuals.

Pregnant mothers also need to be made aware of genetic risk factors associated with Methylation insufficiencies and vaccine injury.

Note: The Glyphosate herbicide used in GMO Foods also depletes many essential co-factors needed for Methylation and may contribute to Methylation insufficiencies.

Please share this life saving information and check out mthfr.net for more details about MTHFR

Methylation Made Easy Part 1 of 4 - Overview Here:

https://www.youtube.com/watch?v=o4uqEDK6BvM

Glutathione. The Body's Master Anti-Oxidant
http://www.mcvitamins.com/glutathione.htm

MTHFR Mutations Explained In Plain English
https://www.youtube.com/watch?v=K1AlLRjcUlI

Food allergies & vaccines:

source author unknown

I hear alot about the science of vaccines. How can you study and declare vaccines safe when you don't even know all the ingredients?

It has been known for over 100 years that if you inject food into an animal, that you produce a food allergy. Food allergies were unknown until vaccines. I traced the history of food allergies and they follow the history of injections exactly. First allergy milk. First injection had cow serum in it. Peanut oil was added to injections in 1919 and the first nut allergy was discovered in 1920.

Wild animals don't have food allergies unless vaccinated. People in countries without modern medicine available to them do not have food allergies. (That is used as the reasoning behind the hygiene theory of food allergies. See http://www.medscape.com/viewarticle/842500)

And food allergies are epidemic now. And they are serious ones. People die from them. A friend's daughter, if she smells fish cooking, has to immediately go to the hospital oshe will die.

It's been kept a secret where these allergies have come from. But it is from injections which include vaccines.

The pharmaceutical companies are allowed to "self-affirm generally recognized as safe" ingredients. What this means is - they decide they want to use GMO soy oil as a pharmaceutical ingredient. They pay for a study using the skin prick test. They pay for some experts to review the study. Then they are free to use GMO soy as a pharmaceutical ingredient. NOTHING appears on the package insert. NOTHING is submitted to the FDA. GMO soy becomes a trade secret, protected by international trade law.

Now there is a big difference between eating GMO soy, doing a skin prick test with GMO soy oil, and injecting GMO soy oil along with a vaccine adjuvant which increases the body's immune response to anything injected along with it. You can search patents on line for vaccine adjuvants and you will find every food oil known to man listed as a possible ingredient.

Not only are people seriously allergic to peanuts, but you can find people who are allergic to every single ingredient found in vaccines.

http://barbfeick.com/vaccinations

Then you can look at bovine serum. The cause of Mad Cow disease cannot be removed from the serum. They monitor the source and hopefully choose healthy cattle.

OK... but the "prions" supposedly cause Mad Cow disease. We don't know for sure. We also don't know if there are other prions in the serum that could be causing other diseases. There could be something like that - that causes diabetes.

http://www.nvic.org/vaccines-and-diseases/Diabetes/juvenilediabetes.aspx

http://barbshealthblog.blogspot.com/2016/10/vaccines-are-direct-cause-of-diabetes.html

Do we know for sure? No. But diabetes is epidemic. So we should just take our chances?

Is that what we do with our children? Experiment on them?

Is it really OK for pharmaceutical companies to "self affirm GRAS" ingredients and not submit anything to the government? Is it OK for pharmaceutical companies to have secret ingredients in vaccines? Is it OK for your doctor and pharmacist to not know if peanut oil is an ingredient in a product that is going to be used on a child with peanut allergy?

Our children are sicker than ever. The medical community says "cause unknown". Are we really that stupid?

"Children and Youth With Disabilities. In 2013–14, the number of children and youth ages 3–21 receiving special education services was 6.5 million, or about 13 percent of all public school students. Among students receiving special education services, 35 percent had specific learning disabilities."

http://nces.ed.gov/programs/coe/indicator_cgg.asp

"May 3, 2011 - The number of people with asthma continues to grow. One in 12 people (about 25 million, or 8% of the population) had asthma in 2009, compared with 1 in 14 (about 20 million, or 7%) in 2001. More than half (53%) of people with asthma had an asthma attack in 2008. More children (57%) than adults (51%) had an attack."

http://www.cdc.gov/vitalsigns/asthma/

"Fifteen years ago, type 2 diabetes in children was almost unheard of. In June, researchers from the SEARCH for Diabetes in Youth study released data showing that type 2 diabetes in 10- to 19-year-olds had increased 21 percent between 2001 and 2009. "

http://www.diabetesforecast.org/2012/nov/more-kids-than-ever-have-type-2-diabetes.html?

"Food allergies:
"This potentially deadly disease affects 1 in every 13 children (under 18 years of age) in the U.S. That's roughly two in every classroom.
"According to a study released in 2013 by the Centers for Disease Control and Prevention, food allergies among children increased approximately 50% between 1997 and 2011."

https://www.foodallergy.org/facts-and-stats

"Even with the decrease, however, about 1,500 infants died of SIDS in 2014, making it the leading cause of death in infants 1 to 12 months old in the United States. (1)

"The cause of SIDS remains a true mystery. Current research is focused on trying to hone in on genetic or physiological causes. However, even though we do not know the cause of SIDS, there are well established methods that can be taken to minimize the risk."

http://acsh.org/news/2016/08/18/in-the-middle-of-the-night-sids-recommendations-go-out-the-window

"The fact that the peak age for SIDS is 2–4 months, which coincides with the introduction of 11 shots containing 16 vaccines (within the US immunization schedule), is so obvious a cause for concern,"

http://kellybroganmd.com/driving-epidemic-sudden-infant-death-sids/

"The kind the best prevalence tracking system for severe autism – the California Department of Developmental Services – says has increased 21% among children born in just the 5 years from 2002 to 2006.

"I'm talking about children who, for example, will never be able to take care of themselves, who need one-on-one classroom aides, who cannot speak for themselves, who are 12 years old and still wear diapers. I'm talking about the child with autism in so much pain that he beats himself bloody, whose parents had to institutionalize him.

"I'm talking about the children and adults with autism who also have co-occurring conditions: gastro-intestinal problems, sleep issues, seizure disorders, metabolic disorders, motor problems.

"Study or no study, you can't help but see it with your own eyes.
"Long waiting lists for outpatient behavioral services.
"Long waiting lists to see a good developmental pediatrician.
"Long waiting lists for respite services.
"Long waiting lists for adult day programs
"Skyrocketing special education costs.
"They can tell us there's no autism epidemic but we all live it. Every day."

http://www.safeminds.org/blog/2015/08/20/face-it-there-is-an-autism-epidemic/

"The National Institutes of Health (NIH estimates up to 23.5* million Americans suffer from autoimmune disease and that the prevalence is rising.

"Autoimmune disease is one of the top 10 leading causes of death in female children and women in all age groups up to 64 years of age."

https://www.aarda.org/autoimmune-information/autoimmune-statistics/

Dangers of Tylenol and vaccines:

Tylenol (Acetaminophen) depletes glutathione levels in the body, which are essential for detoxification. Vaccines have alum adjuvants and other ingredients (See attached link to see specific vaccine ingredients and adverse reactions associated). If you give your child tylenol before or after vaccines, they can't process these toxins and they become even more susceptible to autism and other vaccine injury, as these peer reviewed scientific studies affirm.

Acetaminophen (paracetamol) use, measles-mumps-rubella vaccination, and autistic disorder: the results of a parent survey

"This preliminary study found that acetaminophen use after measles-mumps-rubella vaccination was associated with autistic disorder."

http://www.ncbi.nlm.nih.gov/pubmed/18445737

The acetaminophen metabolite N-acetyl-p-benzoquinone imine (NAPQI) inhibits glutathione synthetase in vitro; a clue to the mechanism of 5-oxoprolinuric acidosis?
http://www.ncbi.nlm.nih.gov/pubmed/27086508

Acetaminophen decreases intracellular glutathione levels and modulates cytokine production in human alveolar macrophages and type II pneumocytes in vitro.

http://www.ncbi.nlm.nih.gov/pubmed/15878691

https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1

Dtap/Tdap/aP background:

**Dtap/TDap/aP is not a live vaccine, but has been shown by the FDA to cause the vaccinated to become asymptomatic carriers whenever exposed to the bacteria for both pertussis and diphtheria, thus spreading the illness without knowing. The Dtap/TDap vaccine is associated with the most adverse reactions recorded.**

**Pertussis carrier:**
**http://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm376937.htm**

**Diphtheria carrier: http://www.cdc.gov/diphtheria/clinicians.html**

**Pertussis epidemic despite high levels of vaccination coverage with acellular pertussis vaccine.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/24216286/**

**"Further, we show that aP vaccination impedes host immunity against B. parapertussis-measured as reduced lung inflammatory and neutrophil responses. Thus, we conclude that aP vaccination interferes with the optimal clearance of B. parapertussis and enhances the performance of this pathogen. Our data raise the possibility that widespread aP vaccination can create hosts more susceptible to B. parapertussis infection."**

**http://www.ncbi.nlm.nih.gov/m/pubmed/20200027/**
**Acellular pertussis vaccines protect against disease but fail to prevent infection and transmission in a nonhuman primate model**

"The observation that aP, which induces an immune response mismatched to that induced by natural infection, fails to prevent colonization or transmission provides a plausible explanation for the resurgence of pertussis and suggests that optimal control of pertussis will require the development of improved vaccines."

**http://m.pnas.org/content/111/2/787.abstract**

**"Our unvaccinated and under-vaccinated population did not appear to contribute significantly to the increased rate of clinical pertussis. Surprisingly, the highest incidence of disease was among previously vaccinated children in the eight to twelve year age group."**

**http://www.ncbi.nlm.nih.gov/pubmed/22423127**

**Vaccinations create more powerful and virulent strains of bacteria and viruses. The reason for the current whooping cough outbreak. Read more here from the CDC"Vaccination against 2 avian viruses, the Marek disease virus, and the infectious bursal disease virus, were associated with the emergence of more virulent strains (33). An important role of host immunity in selecting for virulence is also suggested by the co-evolution of the myxomatosis virus and rabbits (34). Furthermore, immune pressure was shown to select for more virulent Plasmodium chabaudi parasites in mice (35). Based on mathematical modeling, vaccines designed to reduce pathogen growth rate and/or toxicity may result in the evolution of pathogens with higher levels of virulence.**

**(36)."http://wwwnc.cdc.gov/eid/article/15/8/08-1511_article.htm**

**"After the fifth dose of DTaP, the odds of acquiring pertussis increased by an average of 42% per year."**

**http://www.nejm.org/doi/full/10.1056/NEJMoa1200850#discussion**

**Whooping Cough Study May Offer Clue on Surge**

http://www.nytimes.com/2013/11/26/health/study-finds-vaccinated-baboons-can-still-carry-whooping-cough.html

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**Acellular pertussis vaccines protect against disease but fail to prevent infection and transmission in a nonhuman primate model**

http://www.pnas.org/content/111/2/787.abstract

**98% Vaccinated Involved in Whooping Cough Outbreak**
http://www.activistpost.com/2015/02/98-vaccinated-involved-in-whooping.html

**This study shows that efficacy of the DTaP falls rapidly. At 2 years post-vaccination, it's just 75%. By 5 years, it's down to 11.9%.**

**http://www.ncbi.nlm.nih.gov/pubmed/24903664**

**Pertussis 53 - 64% effective in adolescents and adults:**

**http://www.ncbi.nlm.nih.gov/pubmed/23873919**
**-the pertussis vaccine doesn't work, and any immunity from it wanes**

**Importantly, we demonstrate that acellular vaccine antigen-encoding genes are evolving at higher rates than other surface protein-encoding genes. This was true even prior to the introduction of pertussis vaccines but has become more pronounced since the introduction of the current acellular vaccines. The fast evolution of vaccine antigen-encoding genes has serious consequences for the ability of current vaccines to continue to control pertussis.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/25489002/**

**Tetanus is anaerobic, so it dies when it hits oxygen. Bleeding from an injury is a good thing. Tetanus is more common where farm animal feces is present. If tetanus is suspected, an immune globulin shot can be administered. Getting a tetanus shot at the time of injury is a moot point. It would take several weeks to create antibodies, and if tetanus were present, death could occur within a week. Tetanus is extremely rare.**

**Severe tetanus in immunized patients with high anti-tetanus titers**
**https://www.ncbi.nlm.nih.gov/m/pubmed/1565228/**

**Adverse event reports after tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccines in pregnant women.**

**https://www.ncbi.nlm.nih.gov/m/pubmed/22727350/**

**https://www.sciencedaily.com/releases/2015/06/150624071018.htm**

**Is pertussis actually reemerging? Insights from an individual-based model**

**http://www.scielo.br/scielo.php?script=sci_arttext&pid=S0102-311X2001000300005&lng=en&nrm=iso&tlng=en**

https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1

**Articles:**

**http://vaccinechoicecanada.com/in-the-news/a-not-so-perfect-vaccine/**

**Whooping cough on the rise in San Diego, even for the vaccinated – KUSI.com – KUSI News – San Diego CA – News, Weather, PPR**
http://www.kusi.com/story/25336329/whooping-cough-on-the-rise-in-san-diego-even-for-the-vaccinated

.

**Tetanus Vaccine Causes a New disease known as antiphospholipid Syndrome**

http://healthimpactnews.com/2013/tetanus-vaccine-causes-a-new-disease-known-as-antiphospholipid-syndrome/

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**"Mass sterilization": Kenyan Doctors Find Anti-fertility Agent in UN Tetanus Vaccine**

http://healthimpactnews.com/2014/mass-sterilization-kenyan-doctors-find-anti-fertility-agent-in-un-tetanus-vaccine/

DTap/Tdap Vaccine Insert Information:

Diphtheria, Pertussis, Tetanus (DtaP)- Given at 2 months, 4 months, 6 months, 15-18 months, 4-6 years

Diphtheria, Pertussis, Tetanus (Tdap or Td) -Given at 11-12 years, at 19-65+ Substitute 1-time dose of Tdap for Td booster; then boost with Td every 10 yrs, also given to pregnant women

**Vaccine Ingredients:**

Dtap (Infanrix)- formaldehyde, glutaraldehyde, aluminum hydroxide, polysorbate 80, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium

Dtap (Daptacel)- aluminum phosphate, formaldehyde, glutaraldehyde, 2-phenoxyethanol, Stainer-Scholte medium, modified Mueller's growth medium, modified Mueller-Miller casamino medium (without beef heart infusion), dimethyl-1-beta-cyclodextrin, ammonium sulfate

Dtap+IPV (Kinrix)- formaldehyde, glutaraldehyde, aluminum hydroxide, vero (monkey kidney) cells, calf serum, lactalbumin hydrolysate, polysorbate 80, neomycin sulfate, polymyxin B, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium

Dtap+Hep B+IPV (Pediarix)- formaldehyde, glutaraldehyde, aluminum hydroxide, aluminum phosphate, lactalbumin hydrolysate, polysorbate 80, neomycin sulfate, polymyxin B, yeast protein, calf serum, Fenton medium (containing bovine extract), modified Latham medium (derived from bovine casein), modified Stainer-Scholte liquid medium, Vero (monkey kidney) cells

Dtap+IPV+Hib (Pentacel)- aluminum phosphate, polysorbate 80, formaldehyde, glutaraldehyde, bovine serum albumin, 2-phenoxyethanol, neomycin, polymyxin B sulfate, Mueller's growth medium, Mueller-Miller casamino acid medium (without beef heart infusion), Stainer-Scholte medium (modified by the addition of casamino

acids and dimethyl-beta-cyclodextrin), MRC-5 (human diploid) cells, CMRL 1969 medium (supplemented with calf serum), ammonium sulfate, medium 199

Tdap (Boostrix)- formaldehyde, glutaraldehyde, aluminum hydroxide, polysorbate 80 (tween 80), Latham medium derived from bovine casein, Fenton medium containing a bovine extract, Stainer-Scholte liquid medium

Tdap (Adacel)- aluminum phosphate, formaldehyde, glutaraldehyde, 2-phenoxyethanol, ammonium sulfate, Stainer-Scholte medium, dimethyl-beta cyclodextrin, modified Mueller's growth medium, Mueller-Miller casamino acid medium (without beef heart infusion)

**Adverse Reactions from the package inserts:**

Dtap (Infanrix)- Bronchitis, cellulitis, respiratory tract infection, Lymphadenopathy, thrombocytopenia, anaphylactic reaction, hypersensitivity, encephalopathy, headache, hypotonia, syncope, ear pain, cyanosis, apnea, cough, angioedema, erythema, pruritus, rash, urticaria, fatigue, injection site induration, injection site reaction, Sudden infant death syndrome (SIDS)

Dtap (Daptacel)- temp above 105 degrees within 48 hours, collapse or shock-like state, persistent/ inconsolable crying lasting more than 3 hours, seizures with or without fever, Guillain- Barre syndrome, anorexia, drowsiness, vomiting, lymphadenopathy, cyanosis, nausea, diarrhea, injection site pain/ swelling/ module/ mass/ cellulitis/ abscess, hypersensitivity, allergic reaction, anaphylactic reaction, convulsions, febrile convulsions, partial seizures, syncope, screaming

Dtap+IPV (Kinrix)- temp above 105 degrees, collapse or shock-like state, persistent/ inconsolable crying lasting more than 3 hours, seizures with or without fever, cellulitis, constipation, foreign body trauma, injection site vesicles, syncope, anaphylactoid reactions, anaphylaxis, angioedema, urticaria, apnea, lymphadenopathy, thrombocytopenia

Dtap+Hep B+IPV (Pediarix)- pyrexia, gastroenteritis, bronchiolitis, Sudden infant death syndrome (SIDS), convulsive disorder, congenital immunodeficiency with sepsis, neuroblastoma, By chance alone some cases of SIDS can be expected to follow receipt of pertussis- containing vaccines. Chronic illness, asthma, febrile and afebrile seizures, infantile spasms, jaundice, meningitis, lichen planus

Dtap+IPV+Hib (Pentacel)- dehydration, gastroenteritis, asthma, pneumonia, death due to suffocation, head trauma, sudden infant death syndrome (SIDS), neuroblastoma, cyanosis, vomiting, nausea, meningitis, rhinitis, viral infection, decreased appetite, depressed level of consciousness, screaming, apnea, cough, erythema, skin discoloration, pallor

Tdap (Boostrix)- hypersensitivity, anaphylaxis, coma, decreased level of consciousness, prolonged seizures, Guillain- Barre syndrome, brachial neuritis, syncope, progressive or unstable neurologic disorders, Arthus- type hypersensitivity, headache, fatigue, gastrointestinal symptoms, fever, lymphadenitis, allergic reaction such as anaphylactic and anaphylactoid reactions, myocarditis, injection site mass, back pain, convulsions (with and without fever), encephalitis, facial palsy, loss of consciousness

Tdap (Adacel)- headache, body ache, tiredness, chills, sore and swollen joints, nausea, lymph node swelling, diarrhea, vomiting, rash, anaphylactic reaction, hypersensitivity reaction, Guillain- Barre syndrome, brachial neuritis, facial palsy, convolutions, syncope, myositis, myocarditis, pruritus, extensive limb swelling

Vitamin k shot:
Via Todd Wilke

Do you know why vitamin K is pushed on parents and their children?

Because pharmaceutical companies don't like to lose money, doctors don't like to be questioned, the American Academy of Pediatrics dare not change its recommendations. It's just a whole lot easier to give every child vitamin K to prevent the bleeding that could occur from the harm caused by our birth procedures, vaccinations, medications, and unnecessary medical interventions. It just makes complete sense to further assault a newborn's body with an insane level of synthetic vitamin, some antifreeze, and a substance derived from coal-tar that bears a threat of death that warrants a black-box warning.

STUDY shows CHILDHOOD LEUKEMIA RISK from SYNTHETIC Vit K VACCINES given at birth..
( BUT NATURAL VIT K ---orally ok. )
There was a significant association (p = 0.002) --with intramuscular vitamin K --- (odds ratio 1.97, 95% confidence interval 1.3 to 3.0) when compared with oral vitamin K or no vitamin K.

http://www.bmj.com/content/305/6849/341.abstract?ijkey=69c0133cec7e49e8537b087ca33aa20713b41c9d&keytype2=tf_ipsecsha

Here are other reasons why Vit K shots are not safe. But ORAL Vit K at birth is safe and good to ingest:)

http://healthimpactnews.com/2014/the-high-risks-of-vitamin-k-shot-for-your-newborn-baby/

Between four and twelve weeks, we give babies twelve vaccines including a second dose of Hep B, and two doses of DtaP, IPV, Hib, and PCV, all of which can cause vasculitis and brain encephalitis that can induce a hemorrhage. Our children also get

two doses of a live rotavirus vaccine that can shed, infect others, and cause hemorrhagic enteritis and thrombocytopenic purpura (a bleeding disorder). Have you read your child's vaccine inserts?

http://www.immunize.org/packageinserts/

BOX WARNING!!!!

https://dailymed.nlm.nih.gov/dailymed/drugInfo.cfm?setid=e8808230-2c44-44c6-8cab-8f29b6b34051

They are trying to prevent bleeding
BUT THEY Are NOT worried about all the vaccines that they want to give That can cause bleeding/hemorrhage!!! REALLY
SMH

More good info
http://www.livingwhole.org/synthetic-vitamin-k-shot/

You choose ! Your choice!

http://www.bmj.com/content/305/6849/341.abstract?ijkey=69c0133cec7e49e8537b087ca33aa20713b41c9d&keytype2=tf_ipsecsha

Vit k - doctor info
Author of collected info unknown

Vit K... Interesting points made about Vitamin K at birth by a brilliant pediatrician who remains anonymous:

"You know what "synthetic vitamin K" enthusiasts don't understand? The thought that babies (and all animals for that matter) have lower levels of vitamin K at birth for a beneficial, protective, reason. I'm just going to throw these "common sense-based" thoughts out there but let's consider them:

and BTW this goes for the Drops as well as the Shot

First, in order to absorb vitamin K we have to have a functioning biliary and pancreas system. Your infant's digestive system isn't fully developed at birth which is why we give babies breast milk (and delay solids) until they are at least 6-months-old, and why breast milk only contains a small amount of highly absorbable vitamin K. Too much vitamin K could tax the liver and cause brain damage (among other things). As baby ages and the digestive tract, mucosal lining, gut flora, and enzyme functions develop, baby can process more vitamin K. Low levels of vitamin K at birth just…makes…sense. ???

Secondly, cord blood contains stem cells, which protect a baby against bleeding and perform all sorts of needed repairs inside an infant's body. Here's the kicker, in order for a baby to get this protective boost of stem cells, cord-cutting needs to be delayed and the blood needs to remain thin so stem cells can easily travel and perform their functions. Imagine that, baby has his/her own protective mechanism to prevent bleeding and repair organs…that wasn't discovered until after we started routinely giving infants vitamin K injections.

Third, a newborn might have low levels of vitamin K because it's intestines are not yet colonized with bacteria needed to synthesize it and the "vitamin K cycle" isn't fully functional in newborns. It makes sense then to bypass the gut and inject vitamin K right into the muscle right? Except baby's kidneys aren't fully functional either.

Fourth, babies are born with low levels of vitamin K compared to adults, but this level is still sufficient to prevent problems; vitamin K prophylaxis isn't necessarily needed.

Finally, several clinical observations support the hypothesis that children have natural protective mechanisms that justify their low vitamin K levels at birth . I don't know about you, but we should probably figure out why that is before we "inject now and worry about it later."

Do you know why vitamin K is pushed on parents and their children? Because pharmaceutical companies don't like to lose money, doctors don't like to be questioned, the American Academy of Pediatrics dare not change its recommendations."

"Since 1985, the medical profession has known that oral vitamin K raises blood levels 300 - 9,000 times higher. The injectable vitamin K, results in vitamin K levels 9,000 times thicker than adults blood.

Baby's blood thickened with vitamin K, causes a situation where stem cells have to move through sludge, not nicely greased blood vessels full of blood which can allow stem cells easy acess to anywhere. Maybe one day it will dawn on the medical profession that not only are cord blood stem cells important and useful to the newborn baby, but that stem cells need to thin blood for a reason."

"Any fetus which gets being wrung out like a wet towel while travelling down a narrow drain pipe, can incur damage in any part of the body, including in the brain, and needs an in-built fix-it. And stem cells cross the brain blood barrier. In fact, stem cells can go ... anywhere!!! Amazing don't you think. God's design has solutions for situational problems. Three solutions, actually. The second is the fact that naturally, in the first few days, a baby's blood clotting factors are lower than normal.

But ... pediatricians consider this a ... "defect" ... so want to give vitamin K which results in blood nearly 100 times thicker than an adult's. This vitamin K injection, so they say ... (like they say immediate cord clamping is safe, and normal, and delayed cord clamping is an unproven intervention) ... is because the baby wasn't designed right, and if you don't give a vitamin K injection, the baby "could bleed to death".

It's not for nothing that the vitamin K syringe, sits right alongside that cord clamp and the scissors!

But there is an unanswered question:

"Why are blood clotting factors in babies low in the first few days after birth? Why has a baby got much thinner blood as a result?"

Might a logical hypothesis be, that thinner blood allows freer and quicker access of cord blood stem cells to any part of the body damaged during birth? After all, why should stem cells have to fight through a baby's blood which is now 100 times thicker than any adult's, courtesy of another needle?"

MMR/MMRV Vaccine Insert Information:

Measles, Mumps, Rubella (MMR)- Given at 12-15 months, 4-6 years, 1 or 2 doses at 19-50

Vaccine ingredients:

MMR (MMR-II)- recombinant human albumin, neomycin, sorbitol, hydrolyzed gelatin, chick embryo cell culture, WI-38 human diploid lung fibroblasts, Medium 199, minimum essential medium, phosphate
Live virus vaccine that has to potential to shed.

MMRV (ProQuad)- sucrose, hydrolyzed gelatin, sorbitol, monosodium L-glutamate, sodium phosphate dibasic, human albumin, sodium bicarbonate, potassium phosphate monobasic, potassium chloride, potassium phosphate dibasic, neomycin, bovine calf serum, chick embryo cell culture, WI-38 human diploid lung fibroblasts, MRC-5 cells
Live virus vaccine that has the potential to shed.

**Adverse Reactions from the package inserts**

MMR (MMRII)- Panniculitis, atypical measles, fever, syncope, headache, dizziness, malaise, irritability, vasculitis, pancreatitis, diarrhea, vomiting, parotitis, nausea, diabetes belitus, thrombocytopenic, purpura, regional lymphadenopathy, leukocytosis, anaphylaxis, anaphylactoid, pneumonia such as angioneurotic edema, bronchospasm, arthritis, arthralgia, myalgia, encephalitis, encephalopathy, measles inclusion body encephalitis (MIBE), subacute sclerosing panencephalitis (SSPE),

Guillain- Barre syndrome, febrile seizures, febrile seizures, ataxia, polyneuritis, polyneuropathy, ocular palais, paresthesia, ear nerve deafness, death

MMRV (Proquad)- fever, irritability, measles-like rash, varicella-like rash, rash, upper respiratory infection, viral exanthema, diarrhea, rubella-like rases, rhinorrhea, atypical measles, candidiasis, cellulitis, herpes zoster, infection, influenza, measles, respiratory infection, varicella (vaccine strain), anaphylaxis, pneumonia, afebrile seizures, aseptic meningitis, Bell's Palsy, dizziness, ear pain, nerve deafness, bronchial spasms, eczema, death

Studies and articles:

**Malignant Mumps In MMR Vaccinated Children**

http://www.greenmedinfo.com/blog/vaccines-dont-work-malignant-mumps-mmr-vaccinated-children-12

**The Vaccinated Spreading Measles: WHO, Merck, CDC Documents Confirm**

http://vaccineimpact.com/2015/the-truth-about-measles-the-mainstream-media-is-suppressing/

**California Measles: 85% of Those Contracting It Are Fully Vaccinated**

**http://www.activistpost.com/2014/06/california-measles-85-of-those.html**

**A Population-Based Study of Measles, Mumps, and Rubella Vaccination and Autism — NEJM**

http://www.nejm.org/doi/full/10.1056/NEJMoa021134

MMR

https://books.google.com/books?id=CRuY3HmoG4wC&pg=PT136&lpg=PT136&dq=An+Excerpt+from+the+Original+M.M.R.+Drug+Insert&source=bl&ots=g6h4pYFY_O&sig=In7Alt-w-PI6UjGGGwcTiAgLYzU&hl=en&sa=X&ved=0ahUKEwiLo9SUrcPQAhVJ3WMKHXBqB1gQ6AEIXjAJ#v=onepage&q=An%20Excerpt%20from%20the%20Original%20M.M.R.%20Drug%20Insert&f=false

http://currenthealthscenario.blogspot.com/2015/09/original-mmr-package-insert-has-autism.html?m=1

Many children with autism, who have been vaccinated with MMR, have abnormally elevated levels of Measles antibodies and 90% of those also create

AUTOANTIBODIES, suggesting a strong association between MMR and central nervous system autoimmunity in autism.

"Because many autistic children harbor elevated levels of measles antibodies, we conducted a serological study of measles-mumps-rubella (MMR) and MBP autoantibodies. Using serum samples of 125 autistic children and 92 control children, antibodies were assayed by ELISA or immunoblotting methods. ELISA analysis showed a significant increase in the level of MMR antibodies in autistic children. Immunoblotting analysis revealed the presence of an unusual MMR antibody in 75 of 125 (60%) autistic sera but not in control sera. This antibody specifically detected a protein of 73-75 kD of MMR. This protein band, as analyzed with monoclonal antibodies, was immunopositive for measles hemagglutinin (HA) protein but not for measles nucleoprotein and rubella or mumps viral proteins. Thus the MMR antibody in autistic sera detected measles HA protein, which is unique to the measles subunit of the vaccine. Furthermore, over 90% of MMR antibody-positive autistic sera were also positive for MBP autoantibodies, suggesting a strong association between MMR and CNS autoimmunity in autism. Stemming from this evidence, we suggest that an inappropriate antibody response to MMR, specifically the measles component thereof, might be related to pathogenesis of autism."

https://www.ncbi.nlm.nih.gov/m/pubmed/12145534/

http://www.cbsnews.com/news/doctor-explains-why-he-lets-kids-avoid-the-measles-vaccine/

Resource:

https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1

Glyphosate in vaccines and glyphosate poisoning:

Dr. Stephanie Seneff is a senior research scientist at MIT.

"There is a clear explanation for why MMR would cause autism. In fact, there have been a series of papers published by Prof. Singh in Utah showing that autistic kids tend to have very high levels of antibodies to measles hemagglutinin protein (i.e., the vaccine "took" VERY WELL). However, those who have those very high antibody levels also usually have antibodies to myelin basic protein in the brain as well. This protein is in the myelin sheath of the nerve fibers, so the child basically has acquired an autoimmune reaction to its own brain as a consequence of an overly strong reaction to hemagglutinin. I believe the reason it is happening more these days than it used to (autism rates are sharply on the rise) is because of synergistic toxicity with glyphosate, the active ingredient in the pervasive herbicide, Roundup. Roundup is all over the food supply due to pervasive Roundup Ready crops as well

as crops like wheat and sugar cane being sprayed with Roundup right before the harvest. Chronic glyphosate exposure leads to leaky gut, leaky brain barrier, and weakened general immunity. This makes it necessary to over-react to the measles protein (high antibody levels) and then the measles virus gets into the brain and the brain develops autoantibodies through molecular mimicry (the hemagglutinin protein resembles the myelin basic protein). Worse than this, there might actually BE glyphosate in the MMR vaccine, because the live virus is grown on gelatin derived from the ligaments of pigs fed a heavy dose of glyphosate in their feed. My recent research together with Anthony Samsel proposes that glyphosate can get into proteins by mistake in place of glycine. If glyphosate got into the measles hemagglutinin, it would make it much more allergenic and also difficult to break down (get rid of)."

**Glyphosate poisoning.**

http://www.ncbi.nlm.nih.gov/m/pubmed/15862083/

**Concerns over use of glyphosate-based herbicides and risks associated with exposures: a consensus statement.**

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4756530/

**Concerns over use of glyphosate-based herbicides and risks associated with exposures: a consensus statement.**

http://www.ncbi.nlm.nih.gov/m/pubmed/26883814/?i=1&from=glyphosate%20toxicity%20cancer

**Glyphosate: environmental contamination, toxicity and potential risks to human health via food contamination.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27541149/?i=2&from=glyphosate%20toxicity

**[Glyphosate and its formulations--toxicity, occupational and environmental exposure].**

http://www.ncbi.nlm.nih.gov/m/pubmed/24502134/?i=9&from=glyphosate%20toxicity%20cancer

**Potential toxic effects of glyphosate and its commercial formulations below regulatory limits.**

http://www.ncbi.nlm.nih.gov/m/pubmed/26282372/?i=6&from=/23756170/related

5 vaccines that contain carcinogenic glyphosate

https://truthkings.com/5-vaccines-contain-carcinogenic-glyphosate/

**Glysphosate in Childhood Vaccines.**

http://www.momsacrossamerica.com/glyphosate_in_childhood_vaccines

http://www.thehealthtrainers.net/what-s-the-deal-on-vaccinations--.html

The gut and vaccines - Studies

"In healthy subjects, only 0.3% of orally administered aluminum is absorbed via the GI tract, and the kidneys effectively eliminate aluminum from the human body. Only when the GI barrier is bypassed, such as by intravenous infusion or in the presence of advanced renal dysfunction, does aluminum have the potential to accumulate. As an example, with intravenously infused aluminum, 40% is retained in adults and up to 75% is retained in neonates."
"If a significant aluminum load exceeds the body's excretory capacity, the excess is deposited in various tissues, including bone, brain, liver, heart, spleen, and muscle. This accumulation causes morbidity and mortality through various mechanisms."

https://www.ncbi.nlm.nih.gov/pubmed/26948677

*Doctors who explain clearly why vaccines aren't safe or effective. Version 2.0.

1. Dr. Nancy Banks - http://bit.ly/1Ip0aIm
2. Dr. Russell Blaylock - http://bit.ly/1BXxQZL
3. Dr. Shiv Chopra - http://bit.ly/1gdgh1s
4. Dr. Sherri Tenpenny - http://bit.ly/1MPVbjx
5. Dr. Suzanne Humphries - http://bit.ly/17sKDbf
6. Dr. Larry Palevsky - http://bit.ly/1LLEjf6
7. Dr. Toni Bark - http://bit.ly/1CYM9RB
8. Dr. Andrew Wakefield - http://bit.ly/1MuyNzo
9. Dr. Meryl Nass - http://bit.ly/1DGzJsc
10. Dr. Raymond Obomsawin - http://bit.ly/1G9ZXYl
11. Dr. Ghislaine Lanctot - http://bit.ly/1MrVeUL
12. Dr. Robert Rowen - http://bit.ly/1SIELeF
13. Dr. David Ayoub - http://bit.ly/1SIELve
14. Dr. Boyd Haley PhD - http://bit.ly/1KsdVby
15. Dr. Rashid Buttar - http://bit.ly/1gWOkL6
16. Dr. Roby Mitchell - http://bit.ly/1gdgEZU
17. Dr. Ken Stoller - http://bit.ly/1MPVqLI
18. Dr. Mayer Eisenstein - http://bit.ly/1LLEqHH

19. Dr. Frank Engley, PhD - http://bit.ly/1OHbLDI
20. Dr. David Davis - http://bit.ly/1gdgJwo
21. Dr Tetyana Obukhanych - http://bit.ly/16Z7k6J
22. Dr. Harold E Buttram - http://bit.ly/1Kru6Df
23. Dr. Kelly Brogan - http://bit.ly/1D31pfQ
24. Dr. RC Tent - http://bit.ly/1MPVwmu
25. Dr. Rebecca Carley - http://bit.ly/K49F4d
26. Dr. Andrew Moulden - http://bit.ly/1fwzKJu
27. Dr. Jack Wolfson - http://bit.ly/1wtPHRA
28. Dr. Michael Elice - http://bit.ly/1KsdpKA
29. Dr. Terry Wahls - http://bit.ly/1gWOBhd
30. Dr. Stephanie Seneff - http://bit.ly/1OtWxAY
31. Dr. Paul Thomas - http://bit.ly/1DpeXPf
32. Many doctors talking at once - http://bit.ly/1MPVHOv
33. Dr. Richard Moskowitz - http://bit.ly/1OtWG7D
34. Dr. Jane Orient - http://bit.ly/1MXX7pb
35. Dr. Richard Deth - http://bit.ly/1GQDL10
36. Dr. Lucija Tomljenovic - http://bit.ly/1eqiPr5
37. Dr Chris Shaw - http://bit.ly/1IlGiBp
38. Dr. Susan McCreadie - http://bit.ly/1CqqN83
39. Dr. Mary Ann Block - http://bit.ly/1OHcyUX
40. Dr. David Brownstein - http://bit.ly/1EaHl9A
41. Dr. Jayne Donegan - http://bit.ly/1wOk4Zz
42. Dr. Troy Ross - http://bit.ly/1IlGlNH
43. Dr. Philip Incao - http://bit.ly/1ghE7sS
44. Dr. Joseph Mercola - http://bit.ly/18dE38I
45. Dr. Jeff Bradstreet - http://bit.ly/1MaX0cC
46. Dr. Robert Mendelson - http://bit.ly/1JpAEQr
47. Dr. Garth Nicolson - http://bit.ly/1OQVJsF
48. Dr. Marc Girard - http://bit.ly/1iw0smT
49. Dr. Charles Richet - http://bit.ly/1G5GG7j
50. Dr. Zac Bush - http://bit.ly/1LS19OZ
51. Dr. Judy Mikovits - http://bit.ly/1IseF05

Many more doctors testifying that vaccines aren't safe or effective, in these documentaries....
1. Vaccination - The Silent Epidemic - http://bit.ly/1vvQJ2W
2. The Greater Good - http://bit.ly/1icxh8j
3. Shots In The Dark - http://bit.ly/1ObtC8h
4. Vaccination The Hidden Truth - http://bit.ly/KEYDUh
5. Vaccine Nation - http://bit.ly/1iKNvpU
6. Vaccination - The Truth About Vaccines - http://bit.ly/1vlpwvU
7. Lethal Injection - http://bit.ly/1URN7BJ
8. Bought - http://bit.ly/1M7YSlr
9. Deadly Immunity - http://bit.ly/1KUg64Z
10. Autism - Made in the USA - http://bit.ly/1J8WQN5

11. Beyond Treason - http://bit.ly/1B7kmvt
12. Trace Amounts - http://bit.ly/1vAH3Hv
13. Why We Don't Vaccinate - http://bit.ly/1KbXhuf
14. Autism Yesterday - http://bit.ly/1URU2A7

*Vaccine Related Videos, Documentaries, & Audio Clips

(https://www.scribd.com/doc/258899259/Vaccine-Related-Videos-Documentaries-Audio-Clips)

*Why Do Parents Refuse to Vaccinate Their Children?
Dr. Sam Eggertsen, MD

http://bit.ly/1O13BUe

*Why Do Doctors Push Vaccines?
Dr. Janet Levatin, MD

http://tenpennyimc.com/2011/12/24/why-do-doctors-push-vaccines/

*One Nurse's Story - I have seen the cover up

https://www.facebook.com/OccupyProhibition/photos/a.751344764969797.1073742541.317596638344614/751345011636439/?type=3&theater

*Former Sergeant of Police Chris Savage explains how police and doctors who are brainwashed to blame parents for vaccine injury and death

https://www.youtube.com/watch?v=GnVkJmoXDqc

*Scientists Against Vaccines - Hear From Those Who Have Done The Research

http://www.organiclifestylemagazine.com/scientists-against-vaccines-hear-from-those-who-have-done-the-research

*Money Talks - Conflicting Interests Between Healthcare Establishment, Pharmaceutical Corporations, & the Push to Vaccinate

https://www.facebook.com/OccupyProhibition/photos/pb.317596638344614.-2207520000.1448286801./752259758211631/?type=3&theater

*Links to Resources - Vaccine & Contemporary Eugenics Related FAQ-Memes

https://www.facebook.com/notes/occupy-prohibition/vaccine-faq-memes-contemporary-eugenics/701201553317452

Peer reviewed 1000 studies:

http://vaccine-injury.info/pdf/vaccinepeerreview.pdf

http://www.scribd.com/mobile/doc/83004210/CDC-MMR-MMR-V-MMRV-Seizure-Rates

http://www.scribd.com/mobile/doc/33874512/Merck-Vitamin-K-Package-Insert-Aquamephyton-PI

http://www.dailymail.co.uk/health/article-376203/Former-science-chief-MMR-fears-coming-true.html

Autism Spectrum Disorders and Aluminum Vaccine Adjuvants
Lucija Tomljenovic, Russell L. Blaylock, Christopher A. Shaw

Abstract
Impaired brain function, excessive inflammation, and autoimmune manifestations are common in autism. Aluminum (Al), the most commonly used vaccine adjuvant, is a demonstrated neurotoxin and a strong immune stimulator. Hence, adjuvant Al has the necessary properties to induce neuroimmune disorders. Because peripheral immune stimuli in the postnatal period can compromise brain development and cause permanent neurological impairments, the possibility that such outcomes could also occur with administration of Al vaccine adjuvants needs to be considered. In regard to the risk of adjuvant toxicity in children, the following should be noted: (i) children should not be viewed as "small adults" as their unique physiology makes them more vulnerable to toxic insults; (ii) in adult humans Al adjuvants can cause a variety of serious autoimmune and inflammatory conditions including those affecting the brain, yet children are routinely exposed to much higher amounts of Al from vaccines than adults; (iii) compelling evidence has underscored the tight connection between the development of the immune system and that of the brain. Thus, it appears plausible that disruptions of critical events in immune development may also play a role in the establishment of neurobehavioral disorders; (iv) the same immune system components that play key roles in brain development appear to be targeted for impairment by Al adjuvants. In summary, research data suggests that vaccines containing Al may be a contributing etiological factor in the increasing incidence of autism.

http://link.springer.com/referenceworkentry/10.1007%2F978-1-4614-4788-7_89

And what do multiple vaccines combined with aluminum adjuvants do? They over-activate the brains microglia cells and with resulting long term levels of brain inflammation. The studies and the data are all there on the page that I gave you

Journal of American Nutraceutical Association 6: 21-35, 2003.

Interaction of Cytokines, Excitotoxins, and Reactive Nitrogen and Oxygen Species in Autism Spectrum Disorders, Russell Blaylock, MD* Medical Director, Advanced Nutritional Concepts Ridgeland, Mississippi

ABSTRACT

There is growing and compelling evidence that excessive peripheral as well as central immune activation of brain microglia can result in alterations in brain growth and connectivity during rapid brain growth, the so-called "brain growth spurt." A considerable amount of evidence, presented in this paper, demonstrates the deleterious effects of immune factors, such as cytokines, chemokines, and excitotoxins, when present in excess. The interaction between excitotoxicity, ROS and RNS injury and immune dysfunction is discussed. It is concluded that excessive activation of the brain's immune system during critical growth periods can occur when vaccines are given as combination vaccines, using schedules that are too close together or by the use of certain live viruses in the vaccines.

http://www.nutrimedical.com/news.jhtml?method=view&news.id=1084

J Toxicol. 2014; 2014: 491316.
Published online 2014 Oct 2. doi: 10.1155/2014/491316

Aluminum-Induced Entropy in Biological Systems: Implications for Neurological Disease
Christopher A. Shaw, 1 , 2 , 3 ,* Stephanie Seneff, 4 Stephen D. Kette, 5 Lucija Tomljenovic, 1 John W. Oller, Jr., 6 and Robert M. Davidson 7

Abstract
Over the last 200 years, mining, smelting, and refining of aluminum (Al) in various forms have increasingly exposed living species to this naturally abundant metal. Because of its prevalence in the earth's crust, prior to its recent uses it was regarded as inert and therefore harmless. However, Al is invariably toxic to living systems and has no known beneficial role in any biological systems. Humans are increasingly exposed to Al from food, water, medicinals, vaccines, and cosmetics, as well as from industrial occupational exposure. Al disrupts biological self-ordering, energy transduction, and signaling systems, thus increasing biosemiotic entropy. Beginning with the biophysics of water, disruption progresses through the macromolecules that are crucial to living processes (DNAs, RNAs, proteoglycans, and proteins). It injures cells, circuits, and subsystems and can cause catastrophic failures ending in death. Al forms toxic complexes with other elements, such as fluorine, and interacts negatively with mercury, lead, and glyphosate. Al negatively impacts the central nervous system in all species that have been studied, including humans. Because of the global impacts of Al on water dynamics and biosemiotic systems, CNS disorders in humans are sensitive indicators of the Al toxicants to which we are being exposed.

1.2. The Toxic Effects of Aluminum as a Vaccine Adjuvant

Read more:
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4202242/

These are research papers from scientists, doctors, pathologists, and immunologists, but these are very interesting research documents. Some actually from the CDC, which is ironic. The first one actually. Vaccine safety division of the CDC. If autism (brain damage) doesn't happen, then why are there some many research papers like this?

http://www.scribd.com/mobile/doc/220807175/124-Research-Papers-Supporting-the-Vaccine-Autism-Link

Great Books:

Dr. Judy Mikovits

http://www.organiclifestylemagazine.com/vaccines-retroviruses-dna-and-the-discovery-that-destroyed-judy-mikovits-career

http://www.plaguethebook.com

Dr. Mary's Monkeys

http://www.myneworleans.com/New-Orleans-Magazine/July-2007/Dr-Marys-monkey/

http://doctormarysmonkey.com

Dissolving Illusions by Dr. Suzanne Humphries MD
http://www.dissolvingillusions.com

Hep B Vaccine:

**Acute Disseminated Encephalomyelitis following Vaccination against Hepatitis B in a Child: A Case Report and Literature Review**

**http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4960329/**

**Hepatitis B vaccination of male neonates and autism diagnosis, NHIS 1997-2002**

**http://www.ncbi.nlm.nih.gov/pubmed/21058170**

**Acute Disseminated Encephalomyelitis following Vaccination against Hepatitis B in a Child: A Case Report and Literature Review**

**http://www.ncbi.nlm.nih.gov/m/pubmed/27478662/**
**I**
**Acute Disseminated Encephalomyelitis following Vaccination against Hepatitis B in a Child: A Case Report and Literature Review**

**http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4960329/**

**Failure of neonatal hepatitis B vaccination: the role of HBV-DNA levels in hepatitis B carrier mothers and HLA antigens in neonates**

**http://www.sciencedirect.com/science/article/pii/S0168827805804945**

**SEVEN-YEAR STUDY OF HEPATITIS B VACCINE EFFICACY IN INFANTS FROM AN ENDEMIC AREA (SENEGAL)**

**http://www.thelancet.com/journals/lancet/article/PIIS0140-6736(86)90543-X/abstract**

**Hepatitis B vaccine induces apoptotic death in Hepa1–6 cells**

**http://link.springer.com/article/10.1007/s10495-011-0690-1#/page-1**

**The persistence of anti-HBs antibody and anamnestic response 20 years after primary vaccination with recombinant hepatitis B vaccine at infancy.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/25483689/**

**Follow-up of six blood donors highlights the complementary role and limitations of hepatitis C virus antibody and nucleic acid amplification tests.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/12823724/**

**Vaccine induced immunologic memory for hepatitis B surface antigen: implications for policy on booster vaccination**

**http://www.sciencedirect.com/science/article/pii/0264410X9600062X**

**Ten-year neonatal hepatitis B vaccination program, the Netherlands, 1982–1992: protective efficacy and long-term immunogenicity**

**http://www.sciencedirect.com/science/article/pii/S0264410X97000807**

**SEVEN-YEAR STUDY OF HEPATITIS B VACCINE EFFICACY IN INFANTS FROM AN ENDEMIC AREA (SENEGAL)**

**http://www.sciencedirect.com/science/article/pii/S014067368690543X**

**Hepatitis B vaccine and liver problems in U.S. children less than 6 years old, 1993 and 1994**

**http://www.ncbi.nlm.nih.gov/m/pubmed/10230847/**

**Adverse events associated with hepatitis B vaccine in U.S. children less than six years of age, 1993 and 1994.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/11164115/**

**A one year followup of chronic arthritis following rubella and hepatitis B vaccination based upon analysis of the Vaccine Adverse Events Reporting System (VAERS) database.**

**http://www.ncbi.nlm.nih.gov/m/pubmed/12508767/**

**"After reviewing the literature, we observed that complications seen after Hepatitis B vaccination are sudden infant death syndrome, multiple sclerosis, chronic fatigue syndrome, idiopathic thrombocytopenic purpura, vasculitis optic neuritis, anaphylaxis, systemic lupus erytymatosus, lichen planus and neuro-muscular disorder. "**
**Hepatitis B vaccination and associated oral manifestations: a non-systematic review of literature and case reports**

**http://www.ncbi.nlm.nih.gov/m/pubmed/25506472/**

"In it, the Court ruled that the victim, an adult female, had contracted a form of demyelinating disease and MS, and eventually died, after receiving the Hepatitis B vaccine series. It was just the most recent case in a rash of rulings in the omnibus proceeding dealing with hepatitis B vaccine and "demyelinating diseases such as transverse myelitis (TM), Guillain-Barré syndrome (GBS), chronic inflammatory demyelinating disease (CIDP), and multiple sclerosis (MS)," according to court papers."

http://maxresistance.com/vaccines-causing-multiple-sclerosis-nurse-sues-over-h1n1-shot/

http://www.ageofautism.com/2009/02/vaccine-court-hepatitis-b-shot-causes-ms.html

Hepatitis B Vaccine Insert Information:

Hepatitis B- Given at Birth, 2 months, 6-18 months, 3 doses at 19-65+ if high risk, given to pregnant women at high risk

**Vaccine Ingredients:**

Hib+Hep B (Comvax)- yeast (vaccine contains no detectable yeast DNA), nicotinamide adenine dinucleotide, hemin chloride, soy peptone, dextrose, mineral salts, amino acids, formaldehyde, potassium aluminum sulfate, amorphous aluminum hydroxyphosphate sulfate, sodium borate, phenol, ethanol, enzymes, detergent

Hep A+Hep B (Twinrix)- formalin, yeast protein, aluminum phosphate, aluminum hydroxide, amino acids, phosphate buffer, polysorbate 20, neomycin sulfate, MRC-5 human diploid cells

Hep B (Engerix-B)- aluminum hydroxide, yeast protein, phosphate buffers

Hep B (Recombivax)- yeast protein, soy peptone, dextrose, amino acids, mineral salts, potassium aluminum sulfate, amorphous aluminum hydroxyphosphate sulfate, formaldehyde

**Adverse Reactions from the package inserts:**

Hib+Hep B (Comvax)- irritability, somnolence, crying (unusual, high pitched, prolonged for more than 4 hours), anorexia, vomiting, fever over 103 degrees, diarrhea, upper respiratory infection, rash, rhinorrhea, respiratory congestion, cough, anaphylaxis, angioedema, urticaria, seizures, pruritus, edema, syncope, arthritis

Hep A+Hep B (Twinrix)- headache, fatigue, diarrhea, nausea, fever, vomiting, upper respiratory tract infection, anorexia, agitation, insomnia, dizziness, migraine, paresthesia, somnolence, syncope, vertigo, erythema, rash, sweating, back pain, myalgia, lymphadenopathy, dysgeusia, photophobia, hypotension, herpes zoster, meningitis, Bell's palsy, convulsions, Guillain- Barre syndrome, multiple sclerosis, neuritis, earaches, tinnitus, palpitations, hepatitis, jaundice, eczema, lichen planus, arthritis, chills

Hep B (Engerix-B)- herpes zoster, meningitis, thrombocytopenia, anaphylaxis, hypersensitivity syndrome, encephalitis, encephalopathy, migraine, multiple sclerosis, neuritis, neuropathy, Guillain- Barre syndrome, Bell's palsy, optic neuritis, seizures, syncope, earaches, tinnitus, vertigo, palpitations, asthma, eczema, alopecia, lichen planus, purpura, arthritis, muscular weakness, abnormal liver function tests, anorexia, insomnia

Hep B (Recombivax)- fatigue, weakness, high fever, nausea, diarrhea, pharyngitis, upper respiratory infection, sweating, chills, diminished appetite, rhinitis, influenza,

cough, vertigo/ dizziness, paresthesia, pruritus, rash, angioedema, myalgia, back pain, lymphadenopathy, insomnia, earaches, dysuria, hypotension, elevation of liver enzymes, constipation, Guillain- Barre syndrome, multiple sclerosis, seizures, febrile seizures, herpes zoster, encephalitis, eczema, arthritis, extreme pain, lupus-like syndrome

**Reference:**

**https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1**

The Flu:
NATURAL VS. SYNTHETIC - YOU DECIDE!

Here are ways to avoid the flu. You don't need a shot with potential adverse reactions and dangerous elements/ingredients in
It to do so. Prevention is always best. These studies can give insight into natural prevention vs. synthetic.

NATURAL PREVENTION:

Elderberry Syrup:

**Randomized study of the efficacy and safety of oral elderberry extract in the treatment of influenza A and B virus infections.**

https://www.ncbi.nlm.nih.gov/m/pubmed/15080016/?i=2&from=elderberry%20syrup

**The effect of Sambucol, a black elderberry-based, natural product, on the production of human cytokines: I. Inflammatory cytokines.**

https://www.ncbi.nlm.nih.gov/m/pubmed/11399518/?i=2&from=elderberry%20syrup%20flu

General vitamins:

**Effects of a nutritional supplement on the immune response and cytokine production in free-living Chilean elderly.**

https://www.ncbi.nlm.nih.gov/m/pubmed/15449576/?i=4&from=b12%20flu

Vitamin c and Zinc studies:

**Red ginseng and vitamin C increase immune cell activity and decrease lung inflammation induced by influenza A virus/H1N1 infection.**

https://www.ncbi.nlm.nih.gov/m/pubmed/26898166/?i=3&from=vitamin%20c%20flu

**A new mechanism of vitamin C effects on A/FM/1/47(H1N1) virus-induced pneumonia in restraint-stressed mice.**

https://www.ncbi.nlm.nih.gov/m/pubmed/25710018/?i=8&from=vitamin%20c%20flu

**In Silico Analysis to Compare the Effectiveness of Assorted Drugs Prescribed for Swine flu in Diverse Medicine Systems.**

https://www.ncbi.nlm.nih.gov/m/pubmed/24799734/?i=10&from=vitamin%20c%20flu

**Effect of oral gavage treatment with ZnAL42 and other metallo-ion formulations on influenza A H5N1 and H1N1 virus infections in mice.**

https://www.ncbi.nlm.nih.gov/m/pubmed/17626596/?i=3&from=zinc%20effective%20against%20flu

**Low pH gel intranasal sprays inactivate influenza viruses in vitro and protect ferrets against influenza infection.**

https://www.ncbi.nlm.nih.gov/m/pubmed/17509128/?i=4&from=zinc%20effective%20against%20flu

Sulfate (MSM) Study:

**Glycans on influenza hemagglutinin affect receptor binding and immune response.**

https://www.ncbi.nlm.nih.gov/m/pubmed/19822741/?i=2&from=sulfate%20effective%20against%20flu

The common cold can be helped by the above suggested vitamins/minerals/plants. Colloidal silver nasal spray can also be of help.

**Efficacy of a new medical device based on colloidal silver and carbossimetyl beta glucan in treatment of upper airways disease in children.**

https://www.ncbi.nlm.nih.gov/m/pubmed/21946445/

SYNTHETIC PREVENTION:

Flu shot information:

"There is no safe level of mercury and no one has actually shown there is a safe level and I would say mercury is a very toxic substance."--Dr Friberg MD Ph.D. former head of toxicology WHO.

Here are some things in the insert not on the store's or doctor's information sheet:
1. The single dose vial contains mercury (thimerisol) at ≤1mcg (This is called a "trace amount" by the industry.) (The multi-vial contains 25 mcg per shot)
2. People with egg allergies are contraindicated.
3. "Safety and effectiveness have not been established in pregnant women, nursing mothers and children under four. There are no adequate and well-controlled studies in pregnant women. This vaccine should be used during pregnancy only if clearly needed. It is not known whether fluvarin is excreted in human milk."
4. "Fluvarin has not been evaluated for carcinogenic or mutagenic potential, or for impairment of fertility."
5. "Antibody response is low in the geriatric population."
6. "Serious reactions, including anaphylactic shock, have been observed."
7. "There are no data to assess the concomitant administration of flu vaccine with other vaccines."
8. "The vaccine has been associated with an increased frequency of Guillain-Barre syndrome."
9. "In some studies, fluvarin protected up to 50% of subjects."

Flu Vaccine Insert Information

Vaccine Ingredients:

Influenza (yearly)- Starting at 6 months and up, also given to pregnant women

Influenza (Fluzone: standard, high dose)- formaldehyde, octylphenol ethoxylate (Triton X-100), gelatin (standard trivalent formulation only), thimerosal (multi-dose vial only), egg protein, phosphate buffers, sucrose

Influenza (Fluvirin)- nonylphenol ethoxylate, thimerosal (multi-dose only in prefilled syringe), polymyxin, neomycin, beta-propiolactone, egg proteins

Influenza (Flulaval)- thimerosal, a-tocopheryl hydrogen succinate, polysorbate 80, formaldehyde, sodium deoxycholate, ovalbumin

Influenza (Flumist)- ethylenediaminetetraacetic acid (EDTA), monosodium glutamate, hydrolyzed porcine gelatin, arginine, sucrose, dibasic potassium phosphate, monosodium potassium phosphate, gentamicin sulfate, egg protein

*Live virus vaccine that has the potential to shed. - Taken off US Market

Adverse Reactions from the package inserts:

Influenza (Fluzone: High dose)- myalgia, malaise, headache, fever, thrombocytopenia, lymphadenopathy, anaphylaxis, ocular hyperemia, Guillain-Barre syndrome, convulsions, Bell's palsy, brachial neuritis, syncope (shortly after vaccination), dizziness, rhinitis, cough, wheezing, chest pain, vomiting, nausea, diarrhea, chills

Influenza (Fluvirin)- pain, headache, fatigue, fever, sweating, arthralgia, myalgia, malaise, sore throat, chills, nausea, cough, wheezing, stroke, hypersensitivity reactions, vasculitis, loss of appetite, confusion, febrile seizures, Guillain- Barre syndrome, urticaria, rash, cellulitis, encephalopathy

Influenza (Flulaval)- Guillain- Barre syndrome, syncope (fainting), headache, fatigue, malaise, sore throat, cough, chills, facial swelling, diarrhea, lymphadenopathy, eye pain, photophobia, dysphagia, chest pain, allergies, anaphylaxis, angioedema, rhinitis, laryngitis, cellulitis, arthritis, dizziness, tremors, seizures, limb paralysis, insomnia, bronchospasm, throat tightness

Influenza (FluMist)- wheezing, runny nose, decreased appetite, lethargy, chills, high fever, pericarditis, genetic disorders, nausea, vomiting, diarrhea, hypersensitivity, Guillain- Barre syndrome, Bell's Palsy, meningitis, vaccine- associated encephalitis, rash

Reference:

https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1

Peer reviewed scientific studies:

"Influenza vaccines have a modest effect in reducing influenza symptoms and working days lost. There is no evidence that they affect complications, such as pneumonia, or transmission. WARNING: This review includes 15 out of 36 trials funded by industry (four had no funding declaration). An earlier systematic review of 274 influenza vaccine studies published up to 2007 found industry funded studies were published in more prestigious journals and cited more than other studies independently from methodological quality and size. Studies funded from public sources were significantly less likely to report conclusions favorable to the vaccines. The review showed that reliable evidence on influenza vaccines is thin but there is evidence of widespread manipulation of conclusions and spurious notoriety of the studies. The content and conclusions of this review should be interpreted in light of this finding."

http://www.ncbi.nlm.nih.gov/m/pubmed/20614424/

"Increased risk of noninfluenza respiratory virus infections associated with receipt of inactivated influenza vaccine."

http://www.ncbi.nlm.nih.gov/m/pubmed/22423139/

"Increased Risk of Noninfluenza Respiratory Virus Infections Associated With Receipt of Inactivated Influenza Vaccine"

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3404712/

**Association between the 2008–09 Seasonal Influenza Vaccine and Pandemic H1N1 Illness during Spring–Summer 2009: Four Observational Studies from Canada**

http://journals.plos.org/plosmedicine/article?id=10.1371/journal.pmed.1000258

http://www.nvic.org/NVIC-Vaccine-News/April-2016/cdc-admits-flu-shots-fail-half-the-time.aspx

https://liveto110.com/flu-vaccines-are-toxic/

The flumist live virus vaccine was causing a mutation that was causing severe flu in mice, that's why the CDC doesnt recommend it any longer, not to mention it was shedding the flu virus to others.." we found that the FluMist vaccine backbone could regain virulence to cause severe disease in mice. The revertant also regained virulence and caused significant disease in mice, with severity comparable to that caused by a wild type 2009 H1N1 pandemic virus."

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**Insight: Evidence grows for narcolepsy link to GSK swine flu shot**

http://mobile.reuters.com/article/idUSBRE90L07H20130122

**AS03 Adjuvanted AH1N1 Vaccine Associated with an Abrupt Increase in the Incidence of Childhood Narcolepsy in Finland**

http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0033536#close

**Increased Incidence and Clinical Picture of Childhood Narcolepsy following the 2009 H1N1 Pandemic Vaccination Campaign in Finland**

http://journals.plos.org/plosone/article?id=10.1371/journal.pone.0033723

Thimerisol/Mercury brain damage:

http://personalhealthdiary.co/dementia-now-striking-people-in-their-40s-as-mercury-from-vaccines-causes-slow-degenerative-brain-damage/

http://www.royalrife.com/flu_shots.html

Pregnancy:

Flu Vaccine and miscarriage increase
http://Flu Vaccine and miscarriage increase

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3888271/

Maternal Transfer of Mercury on Developing Fetus

http://www.tandfonline.com/doi/full/10.1080/02772248.2012.724574#.VHEJAIvF_pc

2 vaccines cause increase in fetal deaths in 2009/2010

http://www.ncbi.nlm.nih.gov/pubmed/23023030

http://prn.fm/flu-vaccine-and-pregnant-women/

http://community.babycenter.com/post/a38354068/flu_shot_4250_increase_in_fetal_death_reports

Our bodies are amazing. Even getting an illness is part of our bodies journey to transformation and total health. Dr. Stephanie Seneff (Senior Research Scientist MIT) made this fairly new discovery.

She says we live in a symbiotic relationship with all the other species, even the pathogens. The Flu virus goes into the muscles cells and reprograms them to hand over their sulfate. The flu virus delivers the sulfate to your blood. The sulfate cleans your blood by killing off the weak cells allowing growth for new cells. In effect, the flu virus is rescuing your blood from a future meltdown (more serious illnesses, perhaps cancer).

The potential meltdown was there before getting the flu. It's much like an overcrowded Forrest that catches fire to thin and clear out the debris and weak trees. It's a natural process. Depending on what we eat, if we exercise, and if we take supplements to fortify our forest (blood) keeping only the strongest trees (cells) to begin with. Getting sick with the flu isn't bad, it's actually good for your body. You are cleaning house. So this flu season (Fall), think about this. Pretty cool!

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http://prn.fm/flu-vaccine-and-pregnant-women/

http://community.babycenter.com/post/a38354068/flu_shot_4250_increase_in_fetal_death_reports

- Hib (PedVaxHib brand only) – 225 micrograms per shot.
- Hepatitis B – 250 micrograms.
- DTaP – depending on the manufacturer, ranges from 170 to 625 micrograms.
- Pneumococcus – 125 micrograms.
- Hepatitis A – 250 micrograms.
- HPV – 225 micrograms.
- Pentacel (DTaP, HIB and Polio combo vaccine) – 330 micrograms.
- Pediarix (DTaP, Hep B and Polio combo vaccine) – 850 micrograms.
-

http://www.askdrsears.com/topics/health-concerns/vaccines/vaccine-faqs

http://www.scribd.com/mobile/doc/83004210/CDC-MMR-MMR-V-MMRV-Seizure-Rates

http://www.scribd.com/mobile/doc/33874512/Merck-Vitamin-K-Package-Insert-Aquamephyton-PI

http://www.dailymail.co.uk/health/article-376203/Former-science-chief-MMR-fears-coming-true.html

Autism Spectrum Disorders and Aluminum Vaccine Adjuvants
Lucija Tomljenovic, Russell L. Blaylock, Christopher A. Shaw

Abstract
Impaired brain function, excessive inflammation, and autoimmune manifestations are common in autism. Aluminum (Al), the most commonly used vaccine adjuvant, is a demonstrated neurotoxin and a strong immune stimulator. Hence, adjuvant Al has the necessary properties to induce neuroimmune disorders. Because peripheral immune stimuli in the postnatal period can compromise brain development and cause permanent neurological impairments, the possibility that such outcomes could also occur with administration of Al vaccine adjuvants needs to be considered. In regard to the risk of adjuvant toxicity in children, the following should be noted: (i) children should not be viewed as "small adults" as their unique physiology makes them more vulnerable to toxic insults; (ii) in adult humans Al adjuvants can cause a variety of serious autoimmune and inflammatory conditions including those affecting the brain, yet children are routinely exposed to much higher amounts of Al from vaccines than adults; (iii) compelling evidence has underscored the tight connection between the development of the immune system and that of the brain. Thus, it appears plausible that disruptions of critical events in immune development may also play a role in the establishment of neurobehavioral disorders; (iv) the same immune system components that play key roles in brain development appear to be targeted for impairment by Al adjuvants. In summary, research data suggests that vaccines containing Al may be a contributing etiological factor in the increasing incidence of autism.

http://link.springer.com/referenceworkentry/10.1007%2F978-1-4614-4788-7_89

And what do multiple vaccines combined with aluminum adjuvants do? They over-activate the brains microglia cells and with resulting long term levels of brain inflammation. The studies and the data are all there on the page that I gave you

Journal of American Nutraceutical Association 6: 21-35, 2003.
Interaction of Cytokines, Excitotoxins, and Reactive Nitrogen and Oxygen Species in Autism Spectrum Disorders, Russell Blaylock, MD* Medical Director, Advanced Nutritional Concepts Ridgeland, Mississippi

ABSTRACT

There is growing and compelling evidence that excessive peripheral as well as central immune activation of brain microglia can result in alterations in brain growth and connectivity during rapid brain growth, the so-called "brain growth spurt." A considerable amount of evidence, presented in this paper, demonstrates the deleterious effects of immune factors, such as cytokines, chemokines, and excitotoxins, when present in excess. The interaction between excitotoxicity, ROS and RNS injury and immune dysfunction is discussed. It is concluded that excessive activation of the brain's immune system during critical growth periods can occur when vaccines are given as combination vaccines, using schedules that are too close together or by the use of certain live viruses in the vaccines.

http://www.nutrimedical.com/news.jhtml?method=view&news.id=1084

J Toxicol. 2014; 2014: 491316.
Published online 2014 Oct 2. doi: 10.1155/2014/491316

Aluminum-Induced Entropy in Biological Systems: Implications for Neurological Disease
Christopher A. Shaw, 1 , 2 , 3 ,* Stephanie Seneff, 4 Stephen D. Kette, 5 Lucija Tomljenovic, 1 John W. Oller, Jr., 6 and Robert M. Davidson 7

Abstract
Over the last 200 years, mining, smelting, and refining of aluminum (Al) in various forms have increasingly exposed living species to this naturally abundant metal. Because of its prevalence in the earth's crust, prior to its recent uses it was regarded as inert and therefore harmless. However, Al is invariably toxic to living systems and has no known beneficial role in any biological systems. Humans are increasingly exposed to Al from food, water, medicinals, vaccines, and cosmetics, as well as from industrial occupational exposure. Al disrupts biological self-ordering, energy transduction, and signaling systems, thus increasing biosemiotic entropy. Beginning with the biophysics of water, disruption progresses through the macromolecules that are crucial to living processes (DNAs, RNAs, proteoglycans, and proteins). It injures cells, circuits, and subsystems and can cause catastrophic failures ending in death. Al forms toxic complexes with other elements, such as fluorine, and interacts

negatively with mercury, lead, and glyphosate. Al negatively impacts the central nervous system in all species that have been studied, including humans. Because of the global impacts of Al on water dynamics and biosemiotic systems, CNS disorders in humans are sensitive indicators of the Al toxicants to which we are being exposed.

1.2. The Toxic Effects of Aluminum as a Vaccine Adjuvant

Read more:
http://www.ncbi.nlm.nih.gov/pmc/articles/PMC4202242/

These are research papers from scientists, doctors, pathologists, and immunologists, but these are very interesting research documents. Some actually from the CDC, which is ironic. The first one actually. Vaccine safety division of the CDC. If autism (brain damage) doesn't happen, then why are there some many research papers like this?

http://www.scribd.com/mobile/doc/220807175/124-Research-Papers-Supporting-the-Vaccine-Autism-Link

HPV Vaccine information:

There has never been a study that demonstrated the HPV vaccines reduced incidence of cancer. The vaccine was fast tracked simply on the basis that it created antibodies to the HPV strain. There are studies that show the vaccine actually increases your chance of many other cancer causing viruses.

On the relationship between human papilloma virus vaccine and autoimmune diseases.

https://www.ncbi.nlm.nih.gov/m/pubmed/24468416/

Premature ovarian failure 3 years after menarche in 16 year old girl following HPV vaccination.

http://casereports.bmj.com/.../2012/bcr-2012-006879.abstract

"We retrospectively described a case series including 18 girls (aged 12–24 years) referred to our “Second Opinion Medical Network” for the evaluation of “neuropathy with autonomic dysfunction” after HPV vaccination. All girls complained of long-lasting and invalidating somatoform symptoms (including asthenia, headache, cognitive dysfunctions, myalgia, sinus tachycardia and skin rashes) that have developed 1–5 days (n = 11), 5–15 days (n = 5) and 15–20 days (n = 2) after the vaccination. These cases can be included in the recently described immune dysfunction named autoimmune/inflammatory syndrome induced by adjuvants (ASIA)."

https://link.springer.com/article/10.1007/s12026-016-8820-z

"958 hospitalizations
+ 19,351 ED (emergency room visits)
= 20,309 serious adverse events"

So 20,000 of 195,000 girls who got HPV vaccines
---- OR 10% ----
had a SERIOUS enough reaction that it warranted a trip to the ER within 42 days of vaccination.

"Rates of AEFI [adverse reactions] after HPV immunization in Alberta are low and consistent with types of events seen elsewhere."

http://www.sciencedirect.com/science/article/pii/S0264410X16002036

Efficacy of Quadrivalent HPV Vaccine against HPV Infection and Disease in Males.

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3495065/

**Human papilloma virus vaccine and primary ovarian failure: another facet of the autoimmune/inflammatory syndrome induced by adjuvants.**

http://www.ncbi.nlm.nih.gov/m/pubmed/23902317/

**Adverse events following HPV vaccination, Alberta 2006-2014.**

http://www.ncbi.nlm.nih.gov/m/pubmed/26921782/

**New Concerns about the Human Papillomavirus Vaccine**
*American College of Pediatricians – January 2016*

http://www.acpeds.org/the-college-speaks/position-statements/health-issues/new-concerns-about-the-human-papillomavirus-vaccine

**Pancreatitis after human papillomavirus vaccination: a matter of molecular mimicry.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27421720/?i=6&from=Gardasil

**Human papillomavirus vaccine and systemic lupus erythematosus.**

http://www.ncbi.nlm.nih.gov/m/pubmed/23624585/

**Orthostatic intolerance and postural tachycardia syndrome as suspected adverse effects of vaccination against human papilloma virus.**

http://www.ncbi.nlm.nih.gov/m/pubmed/25882168/

**Vaccinations and secondary immune thrombocytopenia with antiphospholipid antibodies by human papillomavirus vaccine.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27312165/

**Is Chronic Fatigue Syndrome/Myalgic Encephalomyelitis a Relevant Diagnosis in Patients with Suspected Side Effects to Human Papilloma Virus Vaccine?**

http://medcraveonline.com/IJVV/IJVV-01-00003.pdf

**Orthostatic intolerance and postural tachycardia syndrome as suspected adverse effects of vaccination against human papilloma virus**

http://www.sciencedirect.com/science/article/pii/S0264410X15004375

**Behavioral abnormalities in female mice following administration of aluminum adjuvants and the human papillomavirus (HPV) vaccine Gardasil.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27421722/

**Severe somatoform and dysautonomic syndromes after HPV vaccination: case series and review of literature.**

http://www.ncbi.nlm.nih.gov/m/pubmed/27503625/

**A link between human papilloma virus vaccination and primary ovarian insufficiency: current analysis.**

http://www.ncbi.nlm.nih.gov/m/pubmed/26125978/?i=5&from=%2F23902317%2Frelated

**On the relationship between human papilloma virus vaccine and autoimmune diseases.**

https://www.ncbi.nlm.nih.gov/m/pubmed/24468416/

**Postural tachycardia syndrome following human papillomavirus vaccination**

http://onlinelibrary.wiley.com/doi/10.1111/ene.12272/abstract;jsessionid=7704D99DE252DB80E59332AA817C6A15.f02t02

21 page package insert from the manufacturer:

https://www.merck.com/product/usa/pi_circulars/g/gardasil_9/gardasil_9_pi.pdf

**Peripheral Sympathetic Nerve Dysfunction**

http://www.ncbi.nlm.nih.gov/m/pubmed/?term=Peripheral+sympathetic+nerve+dysfunction+in+adolescent+Japanese+girls+following

Mainstream news sources:

http://www.cbsnews.com/news/gardasil-researcher-speaks-out/

http://www.usatoday.com/story/news/nation/2013/01/13/hpv-testing-false-negatives/1830897/

https://www.acpeds.org/the-college-speaks/position-statements/health-issues/new-concerns-about-the-human-papillomavirus-vaccine

HPV Vaccine Insert Information:

Human papillomavirus (HPV)- Given at 11-26 years (3 doses)

HPV Vaccine Ingredients:

HPV (Cervarix)- vitamins, amino acids, lipids, mineral salts, aluminum hydroxide, sodium dihydrogen phosphate dihydrate, insect cell and viral protein, 3-0-desacyl-4' Monophosphoryl lipid

HPV (Gardasil)- yeast protein, vitamins, amino acids, mineral salts, carbohydrates, amorphous aluminum hydroxyphosphate sulfate, L-histidine, polysorbate 80, sodium borate

HPV (Cervarix) - extreme pain, headache, nasopharyngitis, influenza, dizziness, upper respiratory infection, dysmenorrhea, chlamydia infection, pharyngitis, injection site bruising, vaginal infection, back pain, UTI, arthritis, celiac disease, dermatomyositis, diabetes, hyperthyroidism, hypothyroidism, inflammatory bowel disease, psoriasis, death, allergies, vitiligo, seizures

HPV (Gardasil)- headache, fever, nausea, arthritis, death, autoimmune thyroiditis, celiac disease, inflammatory bowel disease, multiple sclerosis, nephritis,

pigmentation disorder, diabetes, chills, fatigue, malaise, autoimmune disease, seizures, cellulitis, deep venous thrombosis

https://docs.google.com/document/d/1BQOqgpgA_UlkYNmyVeQcgs79RO-m5-85h2TNMcnhqHo/mobilebasic?pli=1

Are Vaccines safe during pregnancy? Here are few articles, scientific research studies, and videos about this subject.

Blaylock -
https://www.facebook.com/669084493234593/videos/701407633335612/

https://vactruth.com/2012/11/23/flu-shot-spikes-fetal-death/

http://journeyboost.com/2014/11/13/flu-shots-in-pregnancy-2/

http://www.vaccinationinformationnetwork.com/vaccinations-during-pregnancy-resource-page-2/

http://www.modernalternativepregnancy.com/2015/11/16/flu-vaccines-unsafe-and-ineffective-for-pregnant-women/

Dr. Paul video warning for pregnant moms
https://m.youtube.com/watch?v=VoY6vXEMsU8

**Comparison of VAERS fetal-loss reports during three consecutive influenza seasons: was there a synergistic fetal toxicity associated with the two-vaccine 2009/2010 season?**

http://www.ncbi.nlm.nih.gov/m/pubmed/23023030/

**Adverse event reports after tetanus toxoid, reduced diphtheria toxoid, and acellular pertussis vaccines in pregnant women.**

http://www.ncbi.nlm.nih.gov/m/pubmed/22727350/

**Comparison of VAERS fetal-loss reports during three consecutive influenza seasons - Was there a synergistic fetal toxicity associated with the two-vaccine 2009/2010 season?**

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3888271/

**Administration of aluminium to neonatal mice in vaccine-relevant amounts is associated with adverse long term neurological outcomes.**

http://www.ncbi.nlm.nih.gov/pubmed/23932735

**Prenatal environmental exposures, epigenetics, and disease**

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3171169/

**Maternal transfer of mercury to the developing embryo/fetus: is there a safe level?**

http://www.tandfonline.com/doi/abs/10.1080/02772248.2012.724574?journalCode=gtec20#/doi/abs/10.1080/02772248.2012.724574?journalCode=gtec20

**Tdap**

http://www.thehealthyhomeeconomist.com/tdap-vaccine-pushed-on-pregnant-women-despite-fetal-risks/

https://m.youtube.com/watch?v=c3EEuMGhCwA

https://cogforlife.org/2009/12/20/pregnant-women-and-the-h1n1-vaccine/

http://articles.mercola.com/sites/articles/archive/2013/11/10/vaccination-during-pregnancy.aspx

http://paulthomasmd.com/2015/11/09/should-i-get-the-tdap-while-pregnant-does-my-newborn-need-the-hepatitis-b-vaccine/

VAERS has about 18 reports of fetal deaths following TDaP, and a handful of pre-term labor reports shortly following TDaP. You can do a search on CDC VAERS wonder site for the full list of reports.

**Repeat Tdap Vaccination and Adverse Birth Outcomes**

http://jama.jamanetwork.com/mobile/article.aspx?articleid=2504806

Natural healing vitamins/minerals/plants:

"All that man needs for health and healing has been provided by God in nature, the challenge of science is to find it."
- Philippus Theophrastrus Bombast that of Aureolus ~ Paracelsus (1493-1541)

Immune System:

Vitamin C - Immune boosting, detox.
Vitamin B12 - Immune System building, Cognition, Brain health.
Vitamin B Complex - Immune System building, Cognition, Brain health.

Full Spectrum Minerals - Immune System Health
Elderberry Syrup - Immune System Booster
Echinacea - Immune System booster.
Goldenseal - Immune System booster.
Garlic - Immune System booster.
Ginger - Immune System booster, digestive aide.
Beta Glucan - Immune System booster.
Oil of Oregano - Immune System booster.
Ashwaganda, immune tonic safe for everyday use. Good for stress relief.
Chinese Goldenthread - Purify blood, detox, & immune system repair

Anti-Inflammatory:

Turmeric - Anti-inflammatory, digestive.
Molecularly distilled fish oil/Krill oil/Cod Liver Oil - Arthritis, Fibromyalgia, brain, mood/depression, and heart health.
Wobenzym - Anti-inflammatory, joint health, environmental toxins, blood clots, and heart health.
Zinc - Anti-inflammatory, immune system booster, free radical fighter, hormonal imbalance, and cancer help.
CBD Oil - Anti-inflammatory, cell regeneration, cancer help,

Hair, Skin, & Nails:

Silica - Hair, skin, nails.
Biotin - Hair, skin, nails.
Magnesium Oil - Skin.
Emu Oil - Skin, sunburn.
Lavender Oil - Scars, stress, skin irritation.
Coconut Oil - Skin, hair, digestion, and immunity.
Tea Tree Oil - Skin blemishes
MSM - Skin

Joints/Bones/Body:

Glucosamine Chondroitin - Joints health.
MSM - Anti-inflammatory, connective tissues, scar tissue, Skin, muscles, and newly discovered help for getting over the flu faster.
L-Glutamine - Amino Acid that improves protein metabolization.
Calcium - Bone strength and development.
L-Theanine - Relaxation, and healthy vascular function.
Boron - Bone density.

Muscles:

Chamomile - Muscle pain and soreness.

Tart Cherries - muscle relaxer, anti-inflammatory, & anti-oxidant.
Peppermint - Muscles relaxer, backaches, leg pain, & tension headaches.
Cayenne Pepper - Reduce muscle pain, stiffness, & inflammation.
Epsom Salt - Relaxes the nervous system, removes toxins, & helps with pain and inflammation.
Valerian - Muscle spasms
Arnica - Anti-inflammatory & improves blood circulation.
Lavender - Reduces pain, swelling & inflammation.
Passion Flower - Muscle spasms & joint soreness.
Raspberry Leaf Tea - Muscle pain & cramps
Magnesium oil - Topical for muscle pain & cramps.

Organ/Body Function:

Milk Thistle - Liver function in humans and dogs (great for hangovers from what I've been told)
Chromium Picolinate - Insulin, uptake of glucose into cells.
Dandelion - Kidney Health.
Cayenne - blood pressure, metabolism boost, lowers cholesterol.
Bilberry - Diarrhea, eye problems, varicose veins, poor circulation and even cancer .
Apple Vinegar Cider - Diabetes, cancer, heart health, high cholesterol, and weight loss.

Digestive:

Digestive enzymes - Digestive health and also inflammation.
DGL - Licorice Root Extract that aids digestion and treats stomach complaints, including heartburn and indigestion.
Probiotics - Gut health, Immune system booster.
Marshmallow root tea - Acid reflux and heartburn.

Allergies:

Quercetin - Natural Antihistamine (seasonal allergies an allergic reactions)
Nettle, Stinging Leaf (tea) - Allergies.

Detox/Antioxidate/Radiation:

Vitamin C - Detox, and Immune system booster and repair
DMG (N-Dimethylglycine) - improves oxygen utilization, detoxification, cell protection, immune system modulation, and physical performance.
Glutathione - Super Antioxident, stress, and injuries.
King Chlorella - Cleaning out environmental toxins/heavy metals.
Iodine (liquid kelp) - Protection against radiation.
Beet Root - Body detox.
CoQ10 - Antioxidant, heart health, anti-aging.

Green Tea - Antioxidant.
Activated Charcoal - Detox.
Bentonite Clay - Heavy Metal Detox.
Diatomaceous Earth - Heavy Metal Detox.
Cilantro - Heavy Metal Detox.
Organic Citrus peels - Heavy Metal Detox.
Spirulina - Heavy Metal Detox.
Garlic - Heavy Metal Detox.

**Oil of oregano - Purify blood, detox, & immune system repair.**
**Echinacea - Purify blood, detox, & immune system repair.**
**Goldenseal - Purify blood, detox, & immune system repair.**
**Chinese Goldenthread - Purify blood, detox, & immune system repair.**
**Milk Thistle - Liver Detox**

Depression/Brain/Migraines:

Vitamin D - Metal health, Immune Boosting, scar healing, Bones.
Lithium Orotate - Mood Stabilizer.
Vitamin B-6 - Neurological Health.
Magnesium - Migraines.
B2 - Migraines.

Energy:

Pantothenic Acid - (Vitamin B-5) generation of energy from fat, carbohydrates and proteins.
Eleuthero root - Stimulant.

Sleep:

Melatonin - Sleep aide.
Valerian - Sleep aide.
Chamomile - Relaxation, sleep aide.
Tart Cherry Juice - Sleep, gout, and illness prevention.

Essential Oils:

Tea Tree: Good for fungus, acne blemishes, and skin fungal infections like athlete's foot.

Lavender: Works on bruises, cuts, and skin irritation too. Good stress reliever too (Sleep/Depression).

Calendula: Used to reduce the appearance of acne scars. You can also put a drop in your bath water to soothe psoriasis.

Chamomile: Used as a tea or oil for relaxation.

Peppermint: “Peppermint purifies and stimulates the mind. It also can increase mental alertness,” Also good for indigestion.

Frankincense: Relaxation, heal bug bites, scars, depression, inflammation, immunity, and awareness.

Oregano: This oil has naturally antibacterial qualities, which help to fight colds and other sicknesses.

Lemon: “Lemon oil can be used not only to detox the body but it can also help with acne." This is also good for increasing focus and concentration. As a bonus, it can help keep fleas away when used on your pets.

Grapefruit: It’s a natural antiseptic, good for fatigue, and you can add it to your homemade household cleansers to keep your home safe and clean.

Eucalyptus: It has many antibacterial properties and has been known to stimulate the immune system, helps with colds, allergies, and nasal congestion.

Lemongrass: Used as aromatherapy to relieve muscle pain, externally to kill bacteria, ward off insects, and reduce body aches, and internally to help your digestive system.

Thieves: Supports Healthy Immune System and great as deodorant.

Olive leaf extract: Natural antiviral and immune booster.

Without Prejudice or Disadvantage: Thoughts on history and our human codes of ethics.

"Rightful liberty is unobstructed action according to our will within limits drawn around us by the equal rights of others. I do not add 'within the limits of the law' because law is often but the tyrant's will, and always so when it violates the rights of the individual."

-Thomas Jefferson

The most basic right that anyone should have, is what happens to their own physical body, or their child's in regard to health decisions. I find it ironic that the medical and scientific community can demean the average parents intelligence in regard to medical decisions, yet on the television every day we see pharmaceutical ads aimed at the average person, enticing them to ask their doctor to write a prescription. This

is extremely hypocritical. We aren't allowed to ask questions about the true science behind our medicine, but we can ask for a drug we see on TV without hesitation.

Taking away rights, like going to school because of non-compliance with a medical, bodily mandate, is the start to medical tyranny. Using terms like "The Greater Good" to justify these actions, does not justify anything. If you want to take certain medications, and get certain shots, that is your right. You, and the government do not have any right to tell anyone else what they must do with their own physical bodies, even out of fear for yourself or your children.

Fear is a strong word, and a strong emotion. Many times fear is unfounded as well. The book "Dissolving Illusions" by Suzanne Humphries explains in detail the history of vaccines and what really stopped the spread of these terrible diseases. History is the evidence that so many have forgotten, nor bother to read about. Fear drives good people to do things that can change the world into a place no one wants to live in. It happens through quiet steps and strategy by those who want power over others, and they use fear and propaganda to get what they want. Our country is at an intersection. A wrong turn can be difficult to correct, and can have unintended consequences for us all.

The Universal Declaration on Bioethics and Human Rights was adopted by the United Nations Educational, Scientific, and Cultural Organisation (UNESCO) in 2005, and is an important step in the search for global minimum standards in biomedical research, and clinical practice.

The Declaration has important principles relating to bioethics, such as part of Article 6 - Informed Consent (Article 6.1).

"Any preventive, diagnostic and therapeutic medical intervention is only to be carried out with the prior, free and informed consent of the person concerned, based on adequate information. The consent should, where appropriate, be express and may be withdrawn by the person concerned at any time and for any reason without disadvantage or prejudice."

"Without disadvantage or prejudice" are important words. Telling families that their children must have a certain amount of shots by certain dates, or they cannot attend public school, would be considered both a disadvantage, and prejudice. This would also be true for anyone who is being strong armed for their job, or other.

This is the definition of informed consent: Informed consent means the patient's right of self-decision can be effectively exercised when the patient possesses enough information to make an informed choice, and that the patient should make his or her own determination about treatment.

We have doctors who can't tell patients what the ingredients of a vaccine are, or the adverse reactions that could happen, even though the package inserts clearly state

all of those. They are also unaware they need to warn the recently vaccinated that they can infect others for quite awhile after having been given a shot. That is also listed in the vaccine inserts. Yet these doctors recommend and administer something they know very little about. Many gain financial bonuses by doing so. This is clearly a conflict of interest, and their lack of understanding on the most basic vaccine knowledge makes them incapable of giving adequate information in regard to "Informed Consent".

We fought WWII because the Nazis were trying to erase certain people they felt were unequal in value to others. They felt they owned those people, and could do whatever they wanted with them. They were used in horrible medical experiments without their permission, and then thrown away. They were simply considered collateral damage for the bigger picture. This seems like an extreme in comparison, and that we could never get to that point again, but we can. It starts softly, and once the steps are in motion, and time moves forward, people start to accept the deceit. When there is no liability, no one to put a company, country, or state in check, those in power will take until there is nothing left to take.

Every person has the right to their own physical body. If they pay taxes, they have the right to go to schools that use those tax dollars, or work wherever they wish without prejudice or disadvantage. All should be afforded the equal right to benefits by the government or State if they turn down a medical recommendation. Fear is not a reason to take another human beings rights away. Medicine can be considered an "experiment" simply because we are all unique, and will not all react the same way to just about anything you can think of, so choice should always exist. We are not inert objects. We are living, breathing, intelligent creatures who deserve a voice. So we must all ask ourselves if we are on the side of the true greater good, or if we will allow fear to rule us.

Reference Links to important documents in regard to informed consent and medical freedom:

The Universal Declaration of Bioethics and Human Rights: (Article 6:1 - Consent)

http://www.svri.org/sites/default/files/attachments/2016-01-13/2060-IBC.Henk_.Bioethics_Declaration_(October_2005)_(3).doc

http://www.unesco.org/new/en/social-and-human-sciences/themes/bioethics/bioethics-and-human-rights/

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2598251/

Geneva Conventions:

https://www.law.cornell.edu/wex/geneva_conventions

United States Constitution:

https://www.archives.gov/founding-docs

Nuremberg Code:

https://history.nih.gov/research/downloads/nuremberg.pdf

United States Holocaust Memorial Museum.

https://www.ushmm.org/information/exhibitions/online-features/special-focus/doctors-trial/nuremberg-code

http://portal.unesco.org/en/ev.php-URL_ID=31058&URL_DO=DO_TOPIC&URL_SECTION=201.html

http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2598251/

Doctors: Need help finding one?

http://www.vaclib.org/basic/health/novaxdoctors.htm

http://naturallynicole.com/naturally-nicoles-complete-list-of-vaccine-choice-doctors-by-state/

http://birthofanewearth.blogspot.com/2014/09/doctors-who-are-friendly-to-anti.html?m=1

http://www.newsmax.com/t/newsmax/article/646935

List of resources to locate alternative medical providers

Many have switched to Naturopaths, Homeopaths, Holistic MD's, or Chiropractors
Sometimes Osteopaths are more open; and sometimes Family Practice Doctors

American Association of Naturopathic Physicians
206-298-0126
Referral number: 206-298-0125
www.naturopathic.org

American Holistic Medical Association
703-556-9245
www.holisticmedicine.org

American Naturopathic Medical Association
702-897-7053

http://anma.net

Holistic Pediatric Association
707-237-5312

http://www.hpakids.org

http://www.novaxdoctors.webs.com/

Functional medicine doctors are often less likely to push vaccines.

https://www.functionalmedicine.org/practitioner_search...

Another source is the International Chiropractic Pediatric Association:

http://www.icpa4kids.org/locator/index.php

If you're not sure about chiropractic and how it may help your child,
you can find answers here:

http://www.icpa4kids.org/why.htm

http://groups.yahoo.com/group/AP_Doctor_Referral/

http://www.mothering.com/discussions/

click on finding your tribe

Holistic Moms Network
http://www.holisticmoms.org/
click on "chapters' in the column on the right of the page

AANP - American Association of Naturopathic Physicians:
Natural...
NATUROPATHIC.ORG
Websites with lists of "vaccine friendly"

doctors:

http://naturallynicole.com/naturally-nicoles-complete-list.../

http://www.askdrsears.com/.../find-vaccine-friendly-doctor-ne...

https://vactruth.com/2016/06/21/how-to-find-a-physician/

Med exemption help:

http://avoiceforchoice.org/second-opinion/

Alternate forms:

https://www.facebook.com/permalink.php?story_fbid=238779579893806&id=232581190513645

https://www.facebook.com/notes/rights-resources-for-vaccine-freedom/physician-acknowledgment-and-guarantee/1177938115611955

http://media.wix.com/ugd/c2b39a_31c10e06ed7c6ffd555759157e9e8838.doc?dn=ALTERNATE%2BDR%2BVACCINE%2BREFUSAL%2BFORM.doc

**The contract:**

A Statement by the Physician I, (Physician's name)....., do hereby state I have advised the parent(s) of (child's name) ..... that in my professional opinion the child should be given (vaccine drug or other) ..... (Manufacturer's name) ..... (Serial number).....(Batch number).....
I have this day (Day) ..... (Month) ..... (Year) ..... administered this medication after advising the parents that the child is at little or no risk from the treatment. I hereby do agree that should the child at any time suffer or develop any permanent condition deleterious or injurious to their health as a result of this treatment then I will pay any and all costs relating to the care and treatment of this child for the rest of its natural life.
I further agree that if my earnings are insufficient to meet those costs I will sell my home, my business and all my material possessions to put the proceeds towards meeting those costs.
..... Signature of Physician ..... Occupational Position
..... Signature of Witness: Parent or Other ..... Name position of witness

Religious Exemption information:

Religious exemption letter

We, _____ and _____, as the parents of _____, are exercising our rights under the US constitution to receive Religious Exemption from vaccination due to our genuine and sincere beliefs which are contrary to the practices which are herein required.

The following are ways in which these violations manifest themselves in the vaccinations recommended by the Center for Disease Control.
The use of cells, cellular debris, protein, and DNA from willfully aborted human children found in Adenovirus, Polio, Dtap/Polio/HiB Combo, Hep A, Hep A/Hep B

Combo, MMR, MMRV Pro Quad, Rabies, Varicella, and the Shingles vaccines violate the very basic commands found in Exodus 20:13 and Deuteronomy 5:15 which instructs us to not murder. The following ingredients were derived from no fewer than 107 human souls who were sacrificed for social reasons and then used in past and ongoing vaccine research: PER C6, HEK293, WI-38 (RA 27/3), WI-1, WI-2, WI-3, WI-4, WI-5, WI-6, WI-7, WI-8, WI-9, WI-10, WI-11, WI-12, WI-13, WI-14, WI-15, WI-16, WI-17, WI-18, WI-19, WI-20, WI-21, WI-22, WI-23, WI-24, WI-25, WI-26, WI-27, WI-38, WI-44, and MCR-5 plus many other ingredients obtained from human children not required to be listed by FDA guidelines. Supporting vaccinations and vaccination developments is an endorsement of the sacrifice of those and the continuing sacrifice of other human souls.

Genesis 4:1, 17 and Jeremiah 1:5 demonstrate that the deceased children used in the aforementioned vaccinations were recognized by God as human souls from the point of conception in the same way that we, as parents, recognized our child as a human from the moment we were aware of his/her presence in his/her mother's wombs.

Genesis 1:27 - 28, 4:1, 2 Kings 17:17-18, Psalm 22:10-11, 106: 35, 37-38, 113:7-9, 127:3, 139:13-16, Amos 1:13, Matthew 18:1-4, and Matthew 19:13-15 are just a few verses that illustrate the aforementioned children as blessings from God that are valued and loved by him, their Creator, in whose image they were created and that their killing is condemned and causes God's destructive anger to burn against their murderers and those complicit in those murders.
Exodus 20:13, Leviticus 18:21 & 20:2-5, Deuteronomy 5:13, 12:30-32, 18:10, 2 Kings 16:3, and Psalm 106:38 illustrate that all child sacrifice is condemned with no exception clauses allowing for the greater good or public exception clauses found anywhere in the sacred scriptures.

1 Corinthians 6:19-20 and 10:31 remind us that we are to regard our bodies a temples of God's Holy Spirit and that we are to honor God, our Creator and possessor of our very bodies by not defiling them. Notwithstanding the presence of socially sacrificed human cells and debris in vaccinations, we firmly believe that the presence of neurotoxins, hazardous substances, attenuated viruses, animal cells, foreign DNA, albumin from human blood, carcinogens, and chemical wastes is in strict violation of our imperative to treat our bodies as holy temples of the very Spirit of God.

Genesis 9:4, Leviticus 17:10-11, 17:14, Deuteronomy 12:23, Acts 15:20 and 29 informs us that blood represents the life force of human and animal species and that human blood was to be kept pure under all circumstances and free from contaminants such as animal cells, parts, and blood.
We thank you for respectfully adhering to our first amendment rights guaranteed as citizens of the United States of America by her great Constitution and reinforced on a state level by the fourteenth amendment without prejudice.

The US Supreme Court held in Frazee V. Illinois Dept. of Security, 489 US 829, that a religious belief is subject to protection even though no religious group espouses such beliefs or the fact that the religious group to which the individual professes to belong may not advocate or require such belief. This ruling is also reflected in Title VII of the Civil Rights Act of 1964 as amended Nov. 1, 1980; Part 1605.1- Guidelines on Discrimination Because of Religion.

Sincerely,

_____

More:

Someone posted this a while back. Great info.

The use of cells, cellular debris, protein, and DNA from willfully aborted human children found in Adenovirus, Polio, Dtap/Polio/HiB Combo, Hep A, Hep A/Hep B Combo, MMR, MMRV Pro Quad, Rabies, Varicella, and the Shingles vaccines violate the very basic commands found in Exodus 20:13 and Deuteronomy 5:17 which instructs us to not murder. The following ingredients were derived from no fewer than 107 human souls who were sacrificed for social reasons and then used in past and ongoing vaccine research: PER C6, HEK293, WI-38 (RA 27/3), WI-1, WI-2, WI-3, WI-4, WI-5, WI-6, WI-7, WI-8, WI-9, WI-10, WI-11, WI-12, WI-13, WI-14, WI-15, WI-16, WI-17, WI-18, WI-19, WI-20, WI-21, WI-22, WI-23, WI-24, WI-25, WI-26, WI-27, WI-38, WI-44, and MCR-5 plus manmj MN kkkkkj

Any other ingredients obtained from human children not required to be listed by FDA guidelines. Supporting vaccinations and vaccination developments is an endorsement of the sacrifice of those and the continuing sacrifice of other human souls.

Genesis 4:1 and Jeremiah 1:5 demonstrate that the deceased children used in the aforementioned vaccinations were recognized by God as human souls from the point of conception in the same way that we, as parents, recognized our child as a human from the moment we were aware of his/her presence in his/her mother's wombs.

Genesis 1:27 - 28, 4:1, 2 Kings 17:17-18, Psalm 22:10-11, 106: 37-40, Matthew 18:1-4, and Matthew 19:13-15 are just a few verses that illustrate the aforementioned children as blessings from God that are valued and loved by him, their Creator, in whose image they were created and that their killing is condemned and causes God's destructive anger to burn against their murderers and those complicit in those murders.

Exodus 20:13, Leviticus 18:21 & 20:2-5, Deuteronomy 12:30-32, 18:10, 2 Kings 16:3, and Psalm 106:37-40 illustrate that all child sacrifice is condemned with no exception clauses allowing for the greater good or public exception clauses found anywhere in the scriptures.

1 Corinthians 6:19-20 and 10:31 remind us that we are to regard our bodies a temples of God's Holy Spirit and that we are to honor God, our Creator and possessor of our very bodies by not defiling them. Notwithstanding the presence of socially sacrificed human cells and debris in vaccinations, we firmly believe that the presence of neurotoxins, hazardous substances, attenuated viruses, animal cells, foreign DNA, albumin from human blood, carcinogens, and chemical wastes is in strict violation of our imperative to treat our bodies as holy temples of the very Spirit of God.

Genesis 9:4, Leviticus 17:10-11, 17:14, Acts 15:20 and 29 informs us that blood represents the life force of human and animal species and that human blood was to be kept pure under all circumstances and free from contaminants such as animal cells, parts, and blood.

Conspiracy Theory:

Fascinating History: Thinking there is a plan to deceive us does not make you crazy. In fact, it makes you intelligent. History has shown us that some of the craziest conspiracy theories, were actually true.

Here's a list of conspiracy theories that were proven true (hence - now history). Operation Mockingbird isn't in this list, but totally worth looking up. It's about CIA controlled media output (1950-60's). This hasn't actually stopped, it's just changed ownership.

COINTELPRO , a vast and extreme domestic program of monitoring, infiltration, and targeting (lots of evidence seems to pretty strongly indicate it included killings as well) of activists and political groups in the U.S. during the late 1950s through 1960s/70s.

Operation Northwoods was a dramatic one, involving plans to launch a series of terrorist attacks against U.S. targets and blame it on Cuba, to promote a U.S. invasion of Cuba to overthrow Castro in the early 1960s. Who planned it? The Pentagon, and the Joint Chiefs of Staff even signed off on it. Yikes.

The Watergate scandal was derided as a bunch of made-up political attacks and lies against Nixon and the GOP etc., when it was being investigated and media reports were making claims and so on. So I guess this counts as something once considered a conspiracy theory but eventually proven to be true.

The October Surprise related to operations by supporters of Ronald Reagan to assure his election in 1980, including contacts with Iran to convince them to hold the hostages until after the election (to deny Carter a last-minute political "win" that could swing the election his way, thus "October Surprise") and laid the groundwork for the subsequent Iran-Contra scandal. The evidence proving this is not even a big secret (among other things, it includes Bill Casey's credit card bill during his travel on meetings to arrange the deals), but it's still not well known to most of the public despite the evidence being overwhelming and glaringly obvious.

Iran-Contra is a big one, and the true depths of it are rarely appreciated by most people, actually, in part the origins mentioned above related to the hostages in Iran. And on one more related note...

Rex 84 is one of the least-known but most shocking government conspiracies in U.S. history, and one created in part by Oliver North. Many people don't even realize this is a documented part of history, but it was very real indeed. Basically, it was the plan to suspend the U.S. Constitution and enact martial law, round up people considered "subversive," and defend the government against any mass uprising of the citizens in opposition to something like an unpopular foreign war for example. Rex 84 was part of the broader Operation Garden Plot, which is the U.S. military plan to respond to civil unrest in the U.S.

Most people are shocked by the extent to which such plans involve erasure of civil liberties, rounding up of subversives or even potentially large numbers of certain ethnic groups (some of the plan is based on plans to target black citizens during the 1950s and 1960s, for example), to impose a complete military dictatorship, and the extent to which the plans remain in place today under different names and with updates by the DHS and FEMA (this isn't just paranoid talk, the Senate Appropriations Committee has explicitly mentioned DHS's role in Garden Plot and retention of the model plans in the modern era, openly stating such in public published documents).

Business Plot - an attempt to assassinate FD Roosevelt and install military dictatorship over USA, orchestrated by Wall Street.

Operation Ajax - the reason we have problems with Iran. Removed democratically elected leader for oil-friendly dictator.

P2 - foiled attempt by exiled and irregular Freemasons to take over Italy (current prime minister is former member)

MKULTRA - CIA experiments in mind control, primarily for interrogation.

Tuskegee Experiment - Studies and experiments on impoverished black men with syphilis, by denying them treatment or informing them of the disease.

Operation Snow White- largest infiltration of US in history. Scientologists infiltrated all levels of many federal agencies and erased information about themselves, and pointed away from illicit and criminal activities. Damage still unknown.