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Dear Friends,

Welcome to the Holistic Child Health Newsletter.

The goal of my holistic pediatric practice, and of this newsletter, is to inform and support parents to become more empowered as the primary health care providers for their children.



Extending far beyond Western medicine's conventional treatments, holistic medicine and mindful parenting allow us to boost children's natural immunity, support their optimum health and wellness, safely heal any illnesses, and prevent disease-without dangerous side effects. Holistic medicine provides us with the tools to nurture the physical, emotional, social and spiritual health of your children.

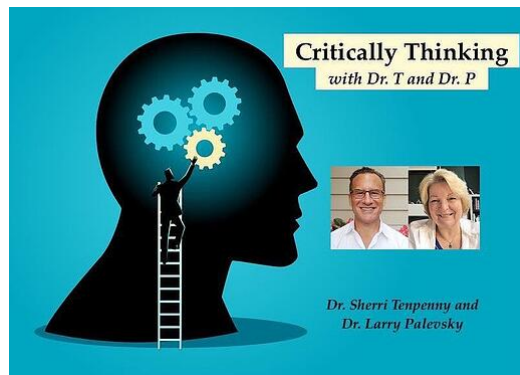
Over the coming months and years, I intend to use this newsletter to address some of your concerns as parents, share information that you might want to add to your knowledge base, and inform you of important issues and current events that are happening in the news, and in your area.

Together we can heal the whole child. Naturally.

Yours in Health,

Lawrence B. Palevsky, MD

CRITICAL THINKING WITH DR T & DR P



Watch & Listen to the weekly podcast
CRITICALLY THINKING with Dr T & Dr P

Open debate and discussion on topics such as vaccines, medicine, science, health, and true wellness have disappeared in today's polarized environment.

In this podcast, "Critically Thinking" Dr. Sherri Tenpenny and Dr. Larry Palevsky challenge the status quo with raw and intelligent discussions designed to restore your curiosity and satisfy your thirst for truth. You will discover answers and possibilities you didn't know existed.

Listen in each week for up-to-date reviews on many different topics that affect your life and the lives of those you love.

To Sign Up for Each Week's Podcast, Click Here:

https://uso2web.zoom.us/webinar/register/WN_HFMeoHUFRjiDwWYIOzdfWg

To Listen To/Watch Each Week's Podcast, Click The Relevant Episode Below:

Week 1

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-1_PH6SFYKmosAIAkg.html

Week 2

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-2_iro7uXMpkZiPg2r.html

Week 3

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-3_N5lMFFGuEEb1kEo.html

Week 4:

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-4_TjkC5BntSkxfv3F.html

Week 5

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-5_JRgdj07FRRncLFs.html

Week 6

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-6_H6sEIpnidHtkwLG.html

Week 7

https://disseminate.tv/watch/critically-thinking-with-dr_EZ8sgssMdlTyW1Z.html

Week 8

<https://disseminate.tv/watch/2d3W2fL1nJCPQ7K>

Week 9

https://disseminate.tv/watch/critically-thinking-with-dr_SxLlenursUmkkqT.html

Week 10

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-10_2lrPwq6zTP7t6t7.html

Week 11

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-11_qafDRZr21xwJraz.html

Week 12

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-12_owiWrwBeHJlbyrU.html

Week 13

<https://disseminate.tv/watch/gFGTtr8qfXtOHGI>

Week 14

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-14_F1UasXuBvTYPesb.html

Week 15

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-15_KFWEEc8nTgXMz2z.html

Week 16

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-16_gN2rLta5zECRive.html

Week 17

https://disseminate.tv/watch/critically-thinking-with-dr-t-and-dr-p-episode-17_EWEKWWU6uczWDma.html

ACTION ALERT



From Autism Action Network

ACTION NEEDED!

Act Now: Tell Your Lawmakers Not to Mandate The Flu Shot For All NJ Students! - Stand for Health Freedom

<https://standforhealthfreedom.com/action/act-now-tell-your-lawmakers-not-to-mandate-the-flu-shot-for-all-nj-students/>

TAKE ACTION



From Autism Action Network**ACTION NEEDED - NEW YORK!****Cuomo Considering Mandatory Flu Shots by Executive Order**

New York State Health Commissioner Howard Zucker said on Wednesday that he is considering following the example of Massachusetts and order all daycare, pre-school, K-12, and college students to get annual flu shots. Connecticut and New Jersey are considering similar orders.

Currently, flu shots are required to attend pre-school and daycare in New York City but nowhere else in the State. Connecticut and New Jersey require flu shots for pre-school and daycare. Massachusetts will become the first state that requires flu shots to attend K-12 school or college. All those states allow religious exemptions.

Zucker is the guy who Gov. Cuomo claims said it was OK to put people with Covid-19 into nursing homes while disregarding all the federal guidelines to prevent the spread of the infection, and now they both refuse to say how many nursing home residents died as the result of their folly. These are the guys who said we had to temporarily shut down to "flatten the curve" so hospitals would not be overwhelmed with Covid-19 cases. That was 180 days ago, hospitals have single digit numbers of Covid-19 cases, and we are still substantially shut down.

The ostensible reason Zucker cited for this new forced vaccination campaign was his concern that unless minors get flu shots now the demand for flu tests later will somehow interfere with Covid-19 testing. They are willing to continue to erode parental rights to make medical decisions our children and compel a medical procedure of questionable safety and efficacy simply to avoid the possibility of busy labs.

Normally adding a new vaccine mandate would require passing laws in the Assembly and Senate followed by approval by the Governor. Just like with any law. But since the legislature gave dictatorial control of the State to Cuomo last March 3, before they crept off to hide, the Governor can order forced flu shots with a decree.

Before you take action on these issues if you value the information and advocacy provided to you by the Autism Action Network please make it possible for us to do so by supporting our one fundraising event of the year we do in conjunction with the Autism Community Walk, here: <https://grouprev.com/autismcommunitywalk2020->

[AutismActionNetwork](#) which is TENTATIVELY scheduled for Sunday, October 18 in Belmont Lake State Park in Babylon, NY. Cuomo willing.

Use this link to send messages to Cuomo, Zucker and our State Senator and Assemblymember letting them know you oppose any type of forced flu shot.

Click Here: <https://cqrcengage.com/achamp/app/write-a-letter?3&engagementId=509256>

Call Cuomo and Zucker and let them know you oppose this scheme:

Gov. Andrew Cuomo, (518) 474-8390

Health Commissioner Howard Zucker, (518) 474-2011

Call your State Senator and Assemblymember. They will want your votes on November 3, so call them and ask them if they will do anything to protect your right to choose whether or your children are forced to get flu shots.

A US Department of Defense Study released earlier this year, the only study done so far on the topic, found that the flu shot INCREASES the risks from coronavirus by 36%. "Receiving influenza vaccination may increase the risk of other respiratory viruses, a phenomenon known as "virus interference... 'vaccine derived' virus interference was significantly associated with coronavirus..." Again, this is not about public health, this is about money and power.

<https://www.sciencedirect.com/science/article/pii/S0264410X19313647?via%3Dihub>

Glaxo, Sanofi, CSL and Astra Zeneca make all the flu shots sold in the US, with approximately 6,000,000 children in New York between 6 months and 18 years, and college students, and the federal Vaccines for Children program selling flu vaccines at around \$13.50 per dose, the New York flu shot market for young people is worth about \$81 million per year, not including the administration fees charge by pediatricians and other flu shot retailers.

According to multi-decade studies of flu shot efficacy by the prestigious Cochrane Collaboration, inactivated flu vaccines (flu shots) in children age 3-16 reduce the risk of influenza in children from 30% to 11% on average. In some years, like 2017, flu shots have no measurable efficacy at all, even in the best-case scenarios devised by the flu shot manufacturers. There is insufficient data according to the Cochrane Collaboration to even say what affect flu shots have on children under 2, yet they would be required to get the shots.

Some flu shots in the US (approximately 20%) are still made using thimerosal, an obsolete preservative based on toxic mercury, that is banned in most of the world, but not the US. In 2008, New York banned the use of thimerosal in shots given to pregnant women and children under three, but the Cuomo administration has illegally suspended the law for past several years misusing language in the bill that allows the state to suspend the law if there is a declared shortage of mercury-free flu shots. There has never been a declared shortage, but Zucker suspends the law anyway. An internal CDC study showed 700% more autism among children who received thimerosal-containing vaccines in infancy. And studies have shown 20% higher autism rates among children whose mothers received pre-natal flu shots in the first trimester.

Please share this message with friends and family, and please share on social networks while we still can.

SPECIAL ALERT



Two Vaccine Mandates Coming Up....

Virginia Health Commissioner Wants to Mandate COVID-19 Vaccine

<https://thevaccinereaction.org/2020/08/virginia-health-commissioner-wants-to-mandate-covid-19-vaccine/>

and

Massachusetts Public Health Officials Require Annual Flu Shots for Child Care and All Grades Through College

<https://thevaccinereaction.org/2020/08/massachusetts-public-health-officials-require-annual-flu-shots-for-child-care-and-all-grades-through-college/>

COVID 19



Dr Andrew Kaufman Destroys the Covid 19 Myth

<https://bit.ly/3cCo83C>

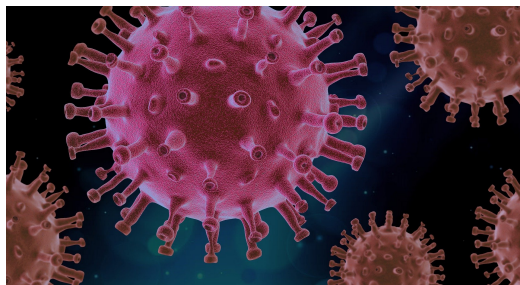
Community and World United, We Say No!

<https://bit.ly/3mW1qv0>

These Are The Good Old Days (NOW Are You Scared?)

https://brandnewtube.com/watch/these-are-the-good-old-days-now-are-you-scared_6TdtD55Qb1yTQBM.html

THE VIRUS



More Experts Point to SARS-CoV-2 Being Created in a Lab

<https://articles.mercola.com/sites/articles/archive/2020/09/04/coronavirus-chimera.aspx>

Chief Science Officer for Pfizer Says "Second Wave" Faked on False-Positive COVID Tests, "Pandemic is Over"

<https://hubpages.com/politics/Pfizer-Chief-Science-Officer-Second-Wave-Based-on-Fake-Data-of-False-Positives-for-New-Cases-Pandemic-is-Over>

Covid Second Wave is Happening All Over Europe, Except in Sweden

<https://www.organiclifestylemagazine.com/covid-second-wave-is-happening-all-over-europe-except-in-sweden>

THE VACCINE



AstraZeneca Made a Bone-Headed Decision to Not Be Transparent With American Public: Chris Meekins

<https://bit.ly/3kKuCU2>

Polls Find Many Americans Do Not Want COVID-19 Vaccine as Soon as It's Licensed

<https://bit.ly/342T38u>

AstraZeneca Halts, Resumes COVID-19 Vaccine Trial After Serious Neurological Event

<https://bit.ly/2HwgNKp>

President Trump Uses 9/11 PREP Act History to Authorize Pharmacists To Administer COVID Vaccines To Kids 3 and Older

<https://bit.ly/2FWSxkx>

Makers of Experimental COVID-19 Vaccines Lobby EU for Liability Protection

<https://thevaccinereaction.org/2020/08/makers-of-experimental-covid-19-vaccines-lobby-eu-for-liability-protection/>

Yale Investigates How To Guilt-Trip People Into Taking Controversial COVID Vaccine

<https://www.lifesitenews.com/news/yale-investigates-how-to-guilt-trip-people-into-taking-controversial-covid-vaccine>

2021: The Year of the Deadliest Vaccine

<https://www.lewrockwell.com/2020/09/gary-d-barnett/2021-the-year-of-the-deadliest-vaccine/>

C-19 TECHNOLOGY & TECHNOCRACY



The Pressing Dangers of Technocracy

<https://bit.ly/2EBxF1g>

A DARPA-Funded Implantable Biochip to Detect COVID-19 Could Hit Markets by 2021

<https://www.mintpressnews.com/darpa-covid-19-vaccine-implant-mrna/271287/>

LOCKDOWN LUNACY



It's Now Looking Like the Lockdowns May Have Been a Huge Mistake

<https://nypost.com/2020/09/06/its-now-looking-like-the-lockdowns-may-have-been-a-huge-mistake/>

The Failed Experiment of Covid Lockdowns

<https://trendmacro.com/system/files/20200901TrendMacroLuskinWSJ.pdf>

VACCINE VIEW



ICAN Files Formal Petition Demanding That The FDA Withdraw It's Licensure of Hepatitis B Vaccines.

ACTION REQUESTED!

ICAN, through its attorneys, has provided the U.S. Department of Health & Human Services (HHS) and the Food and Drug Administration (FDA) numerous opportunities over the past three years to provide proof it licensed the vaccines for Hepatitis B based on clinical trials that reviewed safety for more than five days after injection.

All that HHS and FDA have provided in response are a litany of excuses -- not science. ICAN's attorneys have therefore filed a petition to revoke or suspend the license for these vaccines until a proper clinical trial is conducted.

The Centers for Disease Control and Prevention (CDC) childhood vaccine schedule recommends universal vaccination of all infants with a Hepatitis B vaccine at birth, 1-month of age, and 6-months of age. There are only two Hepatitis B vaccines licensed for administration to newborns: Engerix-B and Recombivax HB.

In 2017, a supporter of ICAN told us that the clinical trials the FDA relied upon to license these vaccines only reviewed safety for a few days after injection. ICAN found this claim incredible. It assumed the claim was likely false.

It seemed improbable because Congress mandated that the FDA only license drugs that have been proven to be "safe and effective" in a clinical trial, and five days of post-injection safety data would be patently insufficient to demonstrate safety. This is why drugs, such as Enbrel, Lipitor, Belviq, and Botox, typically given to adults, had safety review periods of 6.6 years, 4.8 years, 2 years, and 51 weeks respectively, and each was tested against a placebo control group. The FDA even states that the clinical trial relied upon for licensure is typically "1 to 4 years" in duration.

If anything, the safety review period in a clinical trial for a vaccine given to babies and toddlers should be longer since autoimmune, neurological, and developmental disorders will often not be diagnosed until after babies are at least a few years old. Indeed, a 2019 review of 306 pediatric studies, authored by researchers at the FDA and Duke University, explained that, compared to licensing a drug for adults, "data on drug efficacy and safety in children may require an additional 6 years."

Putting this all together, it seemed like science fiction to claim the Hepatitis B vaccines were licensed by the FDA based on only a few days of post-injection safety data. That was, until ICAN reviewed the package inserts for Engerix-B and Recombivax HB issued by their manufacturers and approved by the FDA. To ICAN's

amazement, the package inserts stated that safety in these clinical trials was only reviewed for a few days post-injection into babies.

Therefore, on October 12, 2017, ICAN sent a letter to HHS demanding that it "list and provide the safety data relied upon when recommending babies receive the Hepatitis B vaccine on the first day of life." On January 18, 2018, HHS (in a response reviewed and approved by the FDA) responded by letter in which it failed to provide any clinical trial supporting the safety of either Hepatitis B vaccine that reviewed safety for more than a week post-injection.

After careful review of every single study HHS cited, ICAN responded on December 31, 2018, in a letter which pummeled HHS for failing to provide a single clinical trial reviewing the safety of the Hepatitis B vaccines for more than a week post-injection to babies. ICAN even provided HHS and the FDA another opportunity to provide any such data, including sending a follow-up letter repeating this demand on March 12, 2020. HHS and FDA still have not provided any such data.

ICAN's legal team also formally submitted a Freedom of Information Act request to the FDA demanding a copy of the clinical trials relied upon to license Recombivax HB and received a 1,264 page production of clinical trial reports for this vaccine which confirmed it only reviewed safety for a few days post-injection into babies. ICAN's legal team made a similar request to the FDA for Engerix-B, but this time only requested clinical trials that reviewed safety for more than a week post-injection. The FDA has refused to respond to this request and ICAN's legal team has filed a federal lawsuit against the FDA demanding it respond to this request – more on that lawsuit in a future legal update.

Given the foregoing, it has come time to take formal action. ICAN's legal team, therefore, on September 4, 2020, formally filed a petition to the FDA demanding that the licensure of the Hepatitis B vaccines be revoked or suspended until their safety, as required by law, is determined in a properly designed clinical trial of sufficient duration.

Please submit a comment to the FDA today in support of the petition! Click Here and then on the COMMENTS button: <https://beta.regulations.gov/document/FDA-2020-P-1857-0001>

There may be some uncertainty as to what is required under federal law to determine that a product is "safe" prior to licensure, but what is clear is that five days cannot possibly be sufficient to meet that requirement. Hence, if the FDA

refuses to grant or timely respond to ICAN's petition, we intend to sue the FDA in federal court to demand that the license for these vaccines be revoked or suspended until a proper clinical trial is conducted.



Results of ICAN Lawsuit Against FDA Reveals That Merck's Chicken Pox Vaccine, Varivax, Should Never Have Been Licensed

Merck sells the first and only chicken pox vaccine (VARIVAX) sold in the United States. VARIVAX is produced by growing chicken pox virus on the [cell strain from aborted fetal tissue](#) and was licensed by the FDA in 1995 for people aged 12 months and older.

Prior to FDA licensure of a new experimental vaccines, such as VARIVAX, are expected to undergo long-term placebo-controlled clinical trials with typically tens of thousands of participants to assure their safety.

To evaluate whether Merck met any of these criteria, ICAN, through its attorneys, headed by Aaron Siri, demanded that the FDA produce copies of all the clinical trial reports relied upon to license Merck's chicken pox vaccine. After the FDA stonewalled ICAN for 14 months, failing to produce even a single document, [ICAN sued the FDA in federal court](#) to receive the clinical trial reports for this vaccine.

In this lawsuit, the FDA finally capitulated and began its production of the requested clinical trial reports, the last of which were recently received. A total of 10,796 pages were produced encompassing all the clinical trial reports submitted by Merck to get the VARIVAX license approved. A complete copy of the production is available here. <https://www.dropbox.com/s/pr2afi1mw7uy36m/Productions%201-9%20Combined.pdf?dl=0>

What do these documents show? They show that this product should never have been licensed.

Only one trial relied upon to license this vaccine included a placebo control. Buried deep in the reports, however, it admits that this "placebo" is an injection of

lyophilized stabilizer containing approximately 45 mg of neomycin per milliliter. This is not a placebo! A placebo is an inert substance like saline -- it is not another drug! Worse, this trial only contained 956 children, of which 491 received VARIVAX and 465 received the alleged "placebo." Compounding its fake placebo and miniscule number of participants, these children were only monitored for clinical complaints for 56 days and then for an additional 14 days thereafter for serious adverse reactions.

Nonetheless, even in this shoddy, underpowered study, four serious adverse events (including seizure, ulcer, and viral enteritis) were reported in the group of children receiving VARIVAX and none were reported in the group receiving the alleged placebo. That differential was deemed acceptable by the FDA because Merck's paid investigator did not consider these serious adverse events to be related to VARIVAX.

Nine months after this study began, the children from the "placebo" group were administered VARIVAX. This assured that any differences in the longer-term reaction profile between those that received VARIVAX and those that received the "placebo" in this miniscule trial could never be uncovered.

There was also one study that involved just 61 children in which 32 received VARIVAX and 29 received nothing. At the 8-week mark, the control group (the children who did not receive an injection) were vaccinated with VARIVAX. During that 8-week period, the children receiving VARIVAX had triple the rate of temperatures above 102° F, nearly double the rate of ear infections, 50% increased rate of upper respiratory illness, and 33% increased rate of systemic clinical complaints.

The rest of the trials produced by the FDA were not blinded, not randomized, nor did they include a control group. These trials could therefore not provide scientifically valid data to support the product's safety for licensure.

The FDA's basis for licensing VARIVAX is incredible when considering that: (i) states mandate by law that millions of children receive VARIVAX every year; (ii) Merck cannot be sued for most injuries caused by this product under [federal law](#); and (iii) Merck's sales of this product, alone or in combination with another of its products, was [\\$2.275 billion in 2019](#). But yet, the FDA licensed this product based on one fake "placebo" controlled trial, with less than 1,000 children, that reviewed safety for only around 2 months, and then vaccinated the "placebo" group at nine months!

The FDA should be ashamed to call itself a regulatory agency. ICAN will be taking additional formal legal action regarding the lack of safety relied upon to license this product.

FOLLOW DR PALEVSKY ON FACEBOOK



Join Dr Palevsky on his Facebook page today....browse important articles, scientific papers, and medical information shared for your interest, with added comments from the doctor. Click on the icon above and please feel free to share the link.

ABOUT DR PALEVSKY

Dr. Palevsky is a NYS licensed pediatrician, who utilizes a holistic approach to children's wellness and illness. Dr. Palevsky received his medical degree from the NYU School of Medicine in 1987, completed a three-year pediatric residency at The Mount Sinai Hospital in NYC in 1990, and served as a pediatric fellow in the ambulatory care out-patient department at Bellevue Hospital, NYC, from 1990-1991. Since 1991, his clinical experience includes working in pediatric emergency and intensive care medicine, in-patient and out-patient pediatric medicine, neonatal intensive care medicine, newborn and delivery room medicine, and conventional, holistic and integrative pediatric private practice.



Dr. Palevsky is a diplomate of the American Board of Integrative Holistic Medicine, and Past-President of the American Holistic Medical Association. He received his pediatric board certification in 1990, and passed his pediatric board recertification exams in 1997, 2004, and 2011.

In his current practice in Northport, Long Island and Manhattan, NYC, Dr. Palevsky offers well-child examinations, consultations and educational programs to families and practitioners in the areas of preventive and holistic health; childhood development; lifestyle changes; nutrition for adults, infants and children; safe, alternative treatments for common and difficult to treat acute and chronic pediatric and adult conditions; vaccination controversies; mindful parenting; and rethinking the medical paradigm. Additionally, he teaches holistic integrative pediatric & adolescent medicine to parents, and medical and allied health professionals, both nationally & internationally, and is available for speaking engagements worldwide.

For more information, or to contact Dr. Palevsky go to: www.drpalevsky.com

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If you would like to contact us, please feel free to do so.
We are happy to hear from you.

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