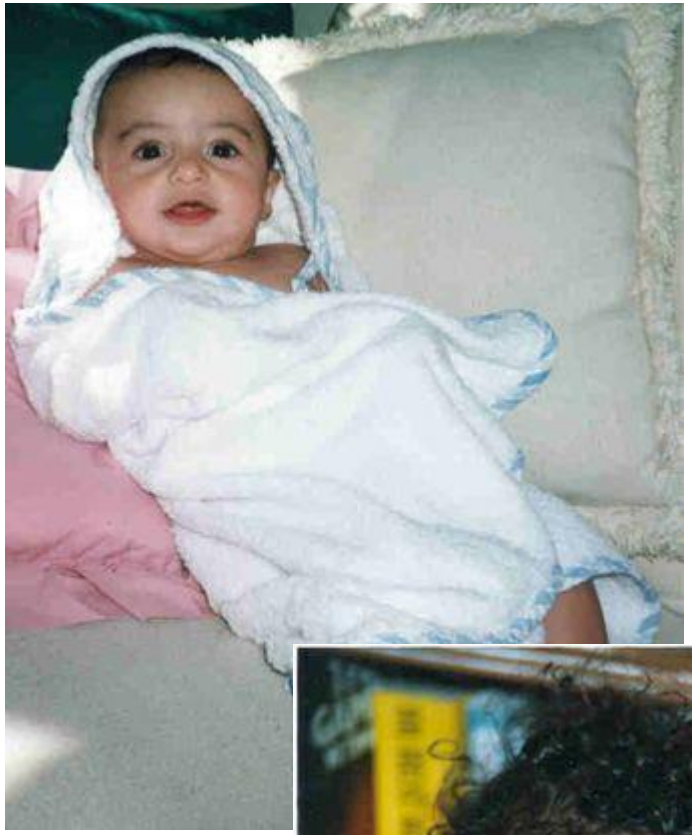
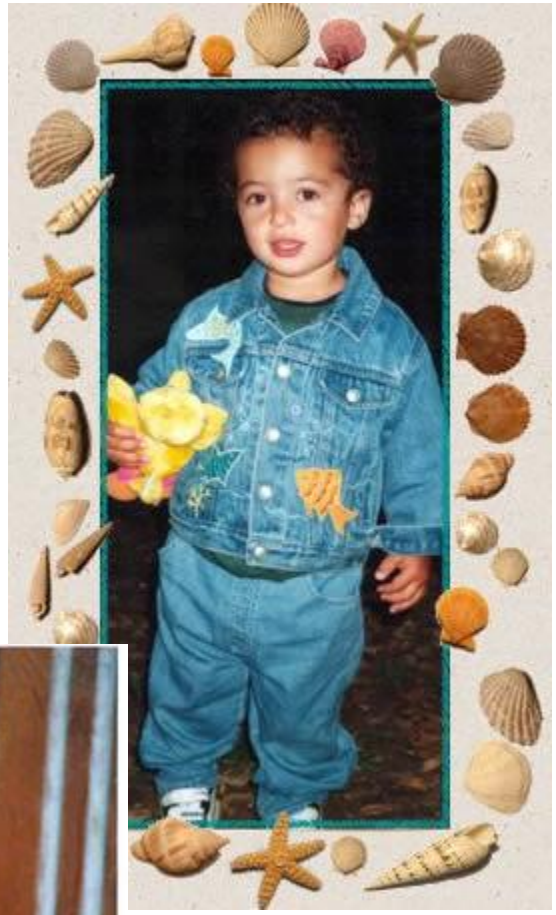
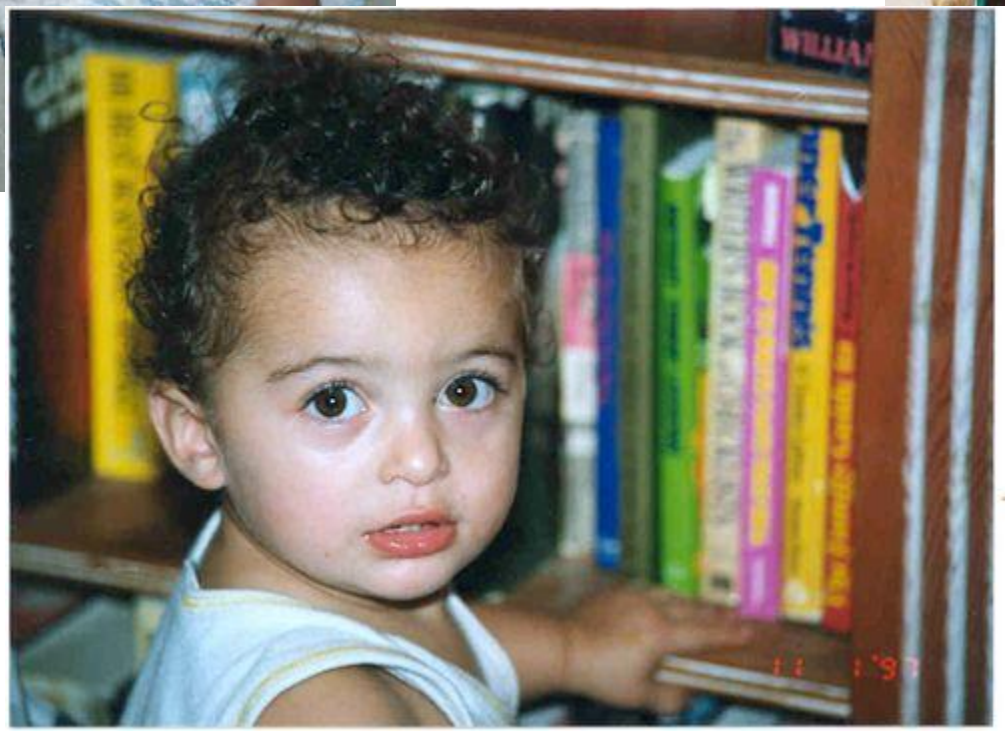
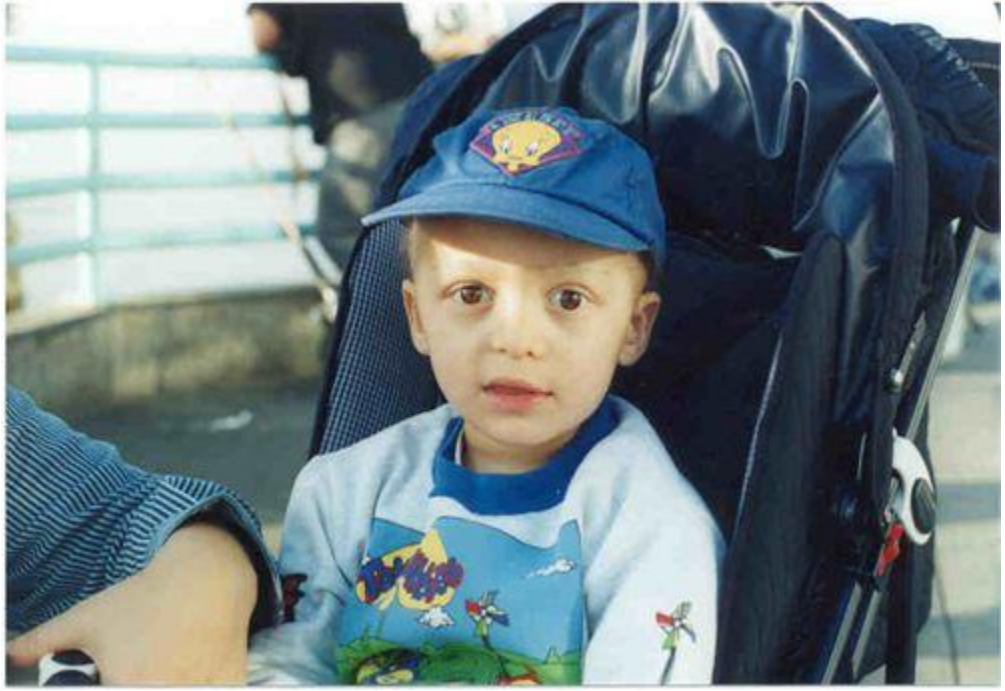
Vaccine Safety and Why Medical Freedom is Necessary

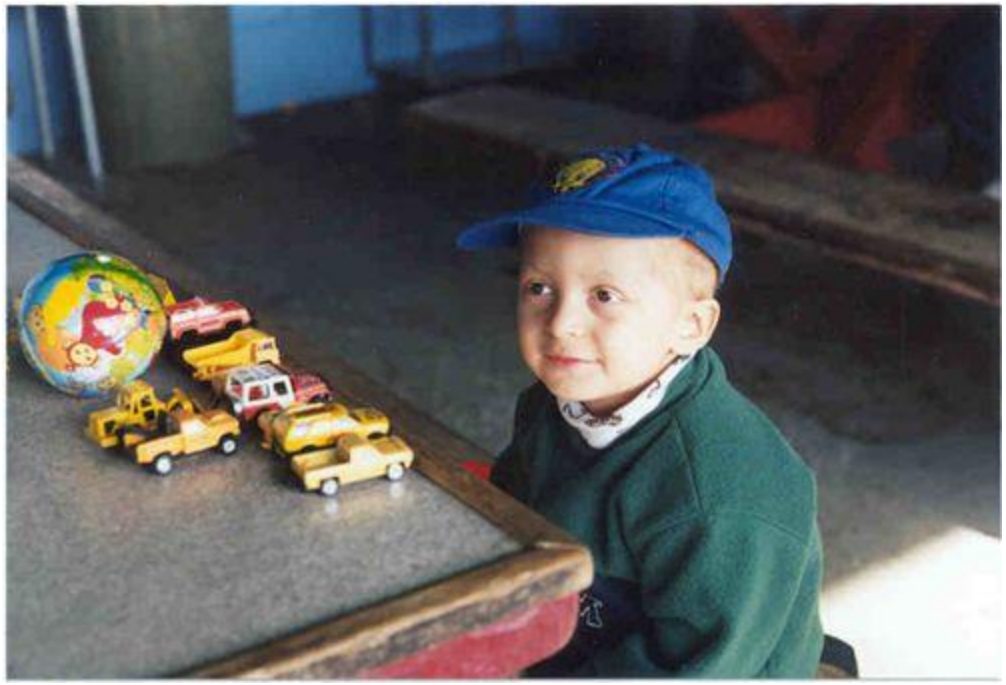
Michael Horwin, MA, JD

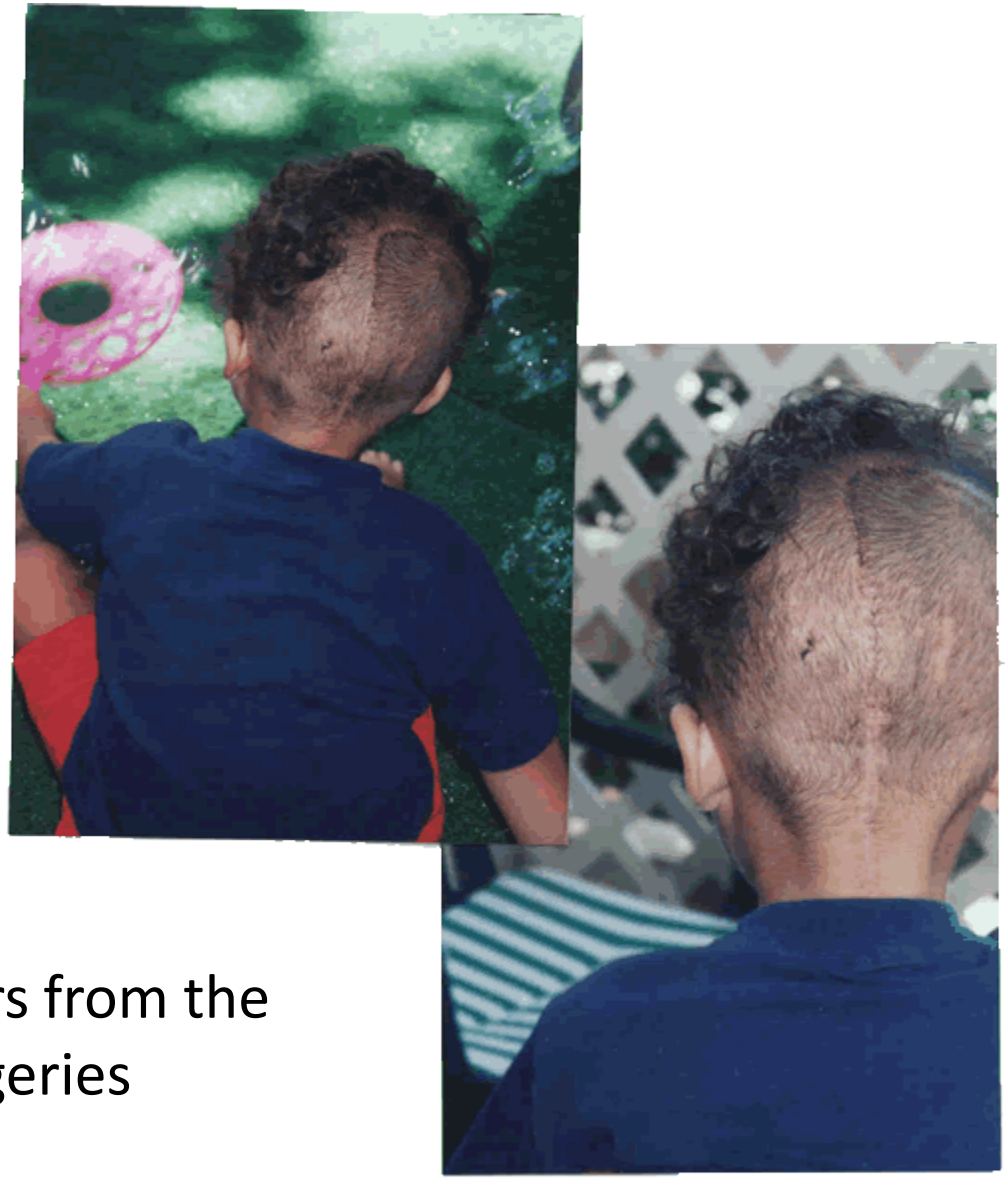






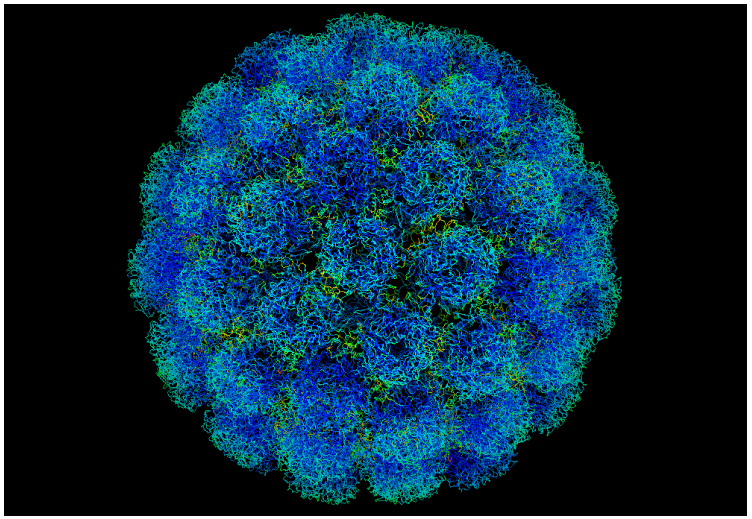






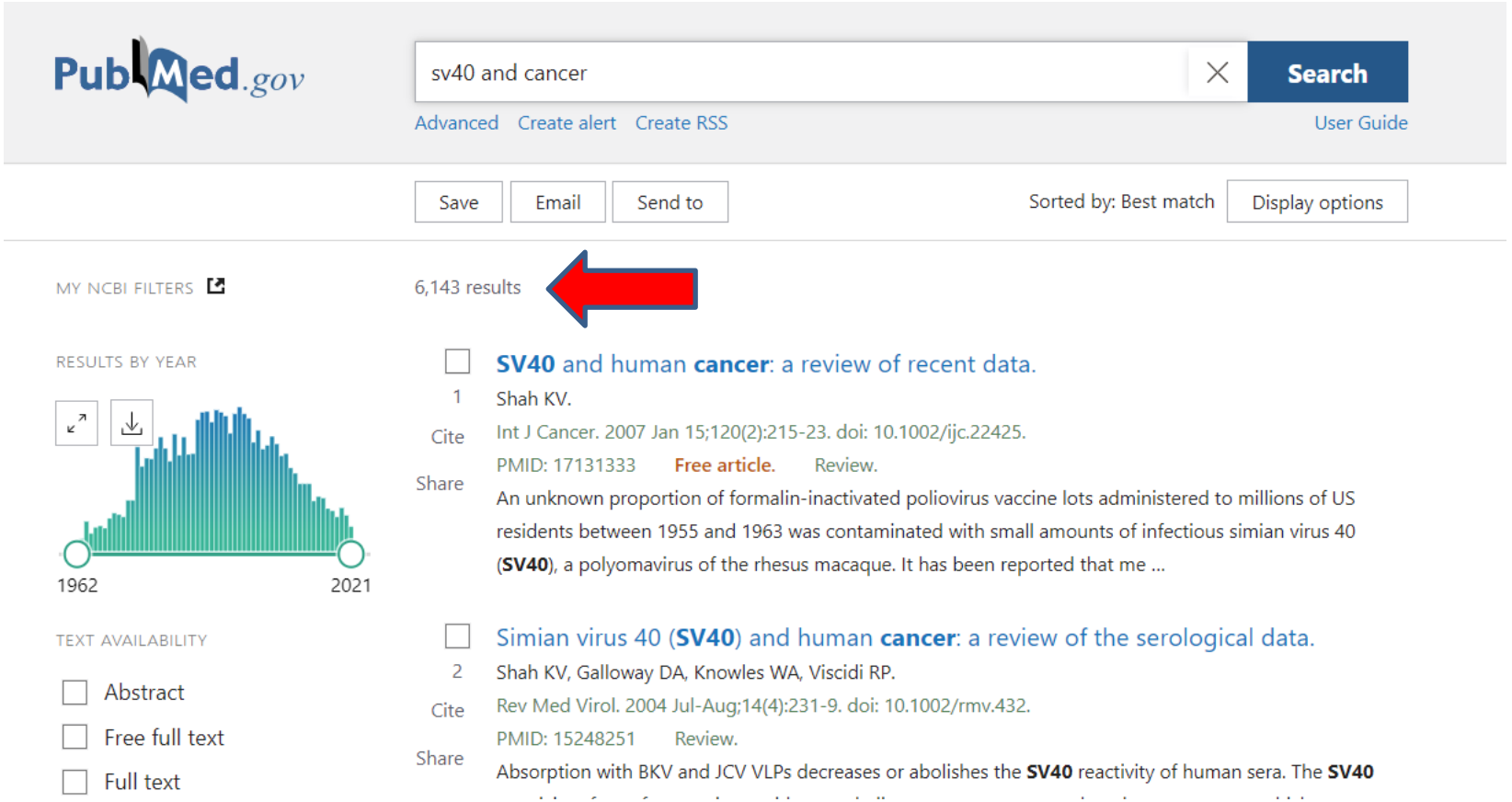
Scars from the surgeries

What is SV40?



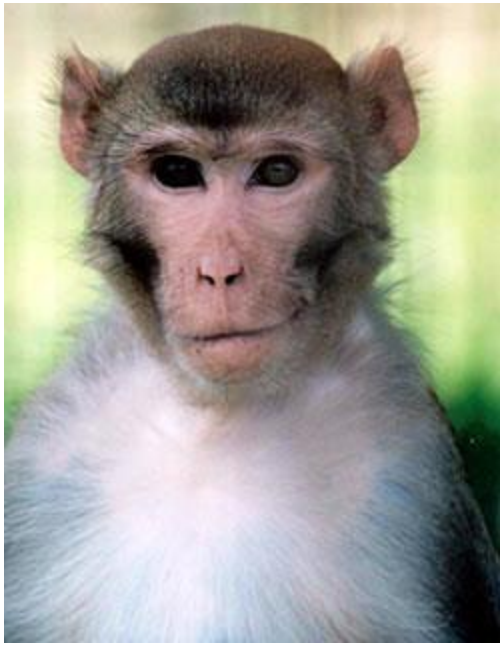
SV40 was the 40th monkey virus found to contaminate polio vaccines. There were hundreds of different viral contaminants in this vaccine. What was special about SV40 is that it was a virus that was highly carcinogenic. It caused various cancers.

What is SV40?



If you do a Pubmed search for “SV40 and cancer” you get over 6,000 medical and scientific articles.

How Did SV40 Get in the Vaccine?



The polio vaccines were made by growing them on monkey kidney cells. The problem is that when you grow a virus for a vaccine on animal cells you end up getting animal viruses mixed into your final product. It this case it was dangerous monkey viruses.

This picture was taken one month before Alexander's death.

Alexander was born on June 7, 1996. Completely healthy.

Oral polio vaccine in November 1997.

August 10, 1998, Alexander was diagnosed with medulloblastoma, a malignant (cancerous) pediatric brain tumor.

He died on January 31, 1999.

The tumor was full of SV40. SV40 is known to cause medulloblastoma.

His cord blood was banked when he was born and was SV40 free which shows that he got the monkey virus after he was born.

His parents were tested and did not carry SV40. We believe it came from his vaccine.

How Did the Health Authorities Deal with SV40 in the Polio Vaccines?

Back in the 1960's the health services knew that SV40 had contaminated the polio vaccines.

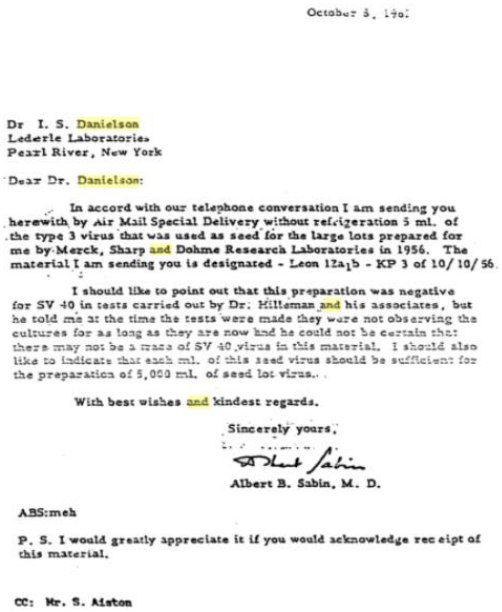
They put in new regulations, but they never required that the contaminated seed stocks be thrown away. The seed stocks were used to make all subsequent oral polio vaccines for decades.

They never required that the contaminated vaccines be thrown away.

They never put in state-of-the-art tests to find SV40 despite the fact that scientists were asking for them.

They did NOT act independently or responsibly in protecting the public. They seemed to be more interested in protecting the pharmaceutical industry.

Historical Document: 1961 Letter from Albert Sabin to Lederle, the company that made the OPV



October 3, 1961

Dr I. S. Danielson
Lederle Laboratories
Pearl River, New York

Dear Dr. Danielson:

In accord with our telephone conversation I am sending you herewith by Air Mail Special Delivery without refrigeration 5 ml. of the type 3 virus that was used as seed for the large lots prepared for me by Merck, Sharp and Dohme Research Laboratories in 1956. The material I am sending you is designated - Leon 12a$_1$b - KP 3 of 10/10/56.

I should like to point out that this preparation was negative for SV 40 in tests carried out by Dr. Hilleman and his associates, but he told me at the time the tests were made they were not observing the cultures for as long as they are now and he could not be certain that there may not be a trace of SV 40 virus in this material. I should also like to indicate that each ml. of this seed virus should be sufficient for the preparation of 5,000 ml. of seed lot virus..

With best wishes and kindest regards.

Sincerely yours,

Albert B. Sabin, M. D.

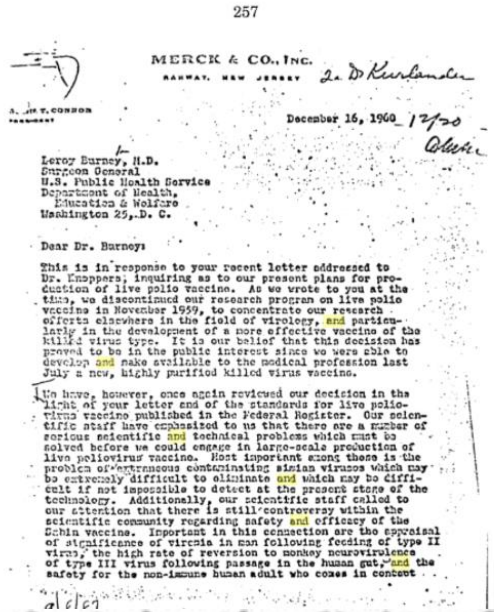
ABS:meh

P. S. I would greatly appreciate it if you would acknowledge receipt of this material.

CC: Mr. S. Aiston

for SV 40 in tests carried out by Dr. Hilleman and his associates, but he told me at the time the tests were made they were not observing the cultures for as long as they are now and he could not be certain that there may not be a trace of SV 40 virus in this material. I should also like to indicate that each ml. of this seed virus should be sufficient for the preparation of 5,000 ml. of seed lot virus..

Historical Document: 1960 Letter from Merck to the Surgeon General



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MERCK & CO., INC.
RAHWAY, NEW JERSEY

Dr. D. Kurlander

December 16, 1960

Leroy Burney, M.D.
Surgeon General
U.S. Public Health Service
Department of Health,
Education & Welfare
Washington 25, D. C.

Dear Dr. Burney:

This is in response to your recent letter addressed to Dr. Knappen, inquiring as to our present plans for production of live polio vaccine. As we wrote to you at the time, we discontinued our research program on live polio vaccine in November 1959, to concentrate our research efforts elsewhere in the field of virology, and particularly in the development of a more effective vaccine of the killed virus type. It is our belief that this decision has proved to be in the public interest since we were able to develop and make available to the medical profession last July a new, highly purified killed virus vaccine.

We have, however, once again reviewed our decision in the light of your letter and of the standards for live poliovirus vaccine published in the Federal Register. Our scientific staff have emphasized to us that there are a number of serious scientific and technical problems which must be solved before we could engage in large-scale production of live poliovirus vaccine. Most important among these is the problem of extraneous contaminating simian viruses which may be extremely difficult to eliminate and which may be difficult if not impossible to detect at the present stage of the technology. Additionally, our scientific staff called to our attention that there is still controversy within the scientific community regarding safety and efficacy of the Sabin vaccine. Important in this connection are the appraisal of significance of viremia in man following feeding of type II virus, the high rate of reversion to monkey neurovirulence of type III virus following passage in the human gut and the safety for the non-immune human adult who comes in contact

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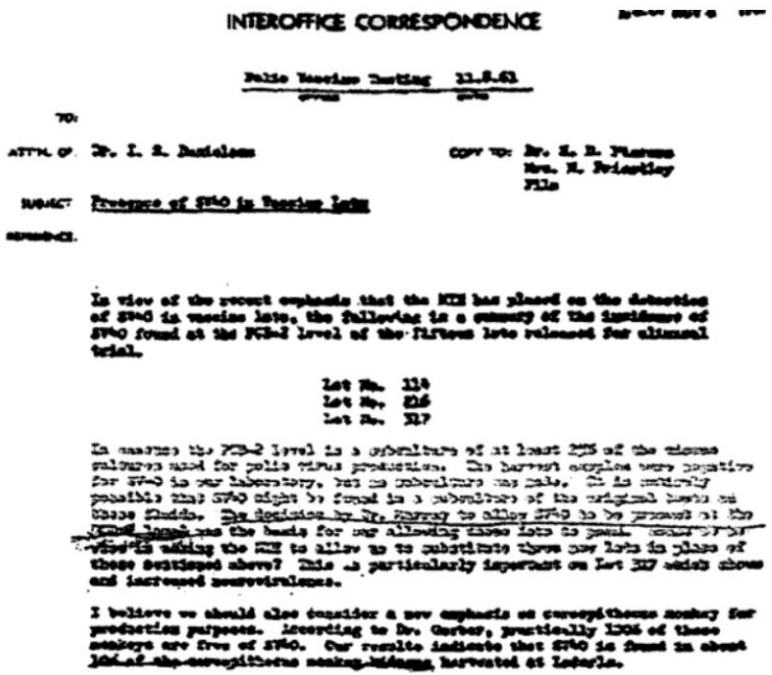
Historical Document: 1961 interoffice memo from Lederle



INTEROFFICE CORRESPONDENCE

Polio Vaccine Testing 11.8.61

TO:

ATTN. OF Dr. I. S. Danielson

COPY TO: Dr. L. D. Plotnow
Mrs. M. Priestley
File

SUBJECT Presence of SV40 in Vaccine Lots

REFERENCE

In view of the recent emphasis that the NIH has placed on the detection of SV40 in vaccine lots, the following is a summary of the incidence of SV40 found at the PCK-2 level of the fifteen lots released for clinical trial.

Lot No. 114
Lot No. 116
Lot No. 117

In testing the PCK-2 level in a subculture of at least 225 of the rhesus kidneys used for polio virus production. The kidneys were negative for SV40 in our laboratory, but no subculture was made. It is entirely possible that SV40 might be found in a subculture of the original lots as these fluids. The decision by Dr. Murray to allow SV40 to be present at the [illegible] level was the basis for our allowing these lots to pass. [illegible] who is asking the NIH to allow us to substitute three new lots in place of those designated above? This is particularly important on Lot 117 which shows and increased neurovirulence.

I believe we should also consider a new emphasis on cercopithecus monkey for production purposes. According to Dr. Gerber, practically 100% of these monkeys are free of SV40. Our results indicate that SV40 is found in about 10% of the cercopithecus monkey kidneys harvested at Lederle.

The decision by Dr. Murray to allow SV40 to be present at the () level was the basis for our allowing these lots to pass…

How Did the Health Authorities Deal with SV40 in the Polio Vaccines?

In Summary:

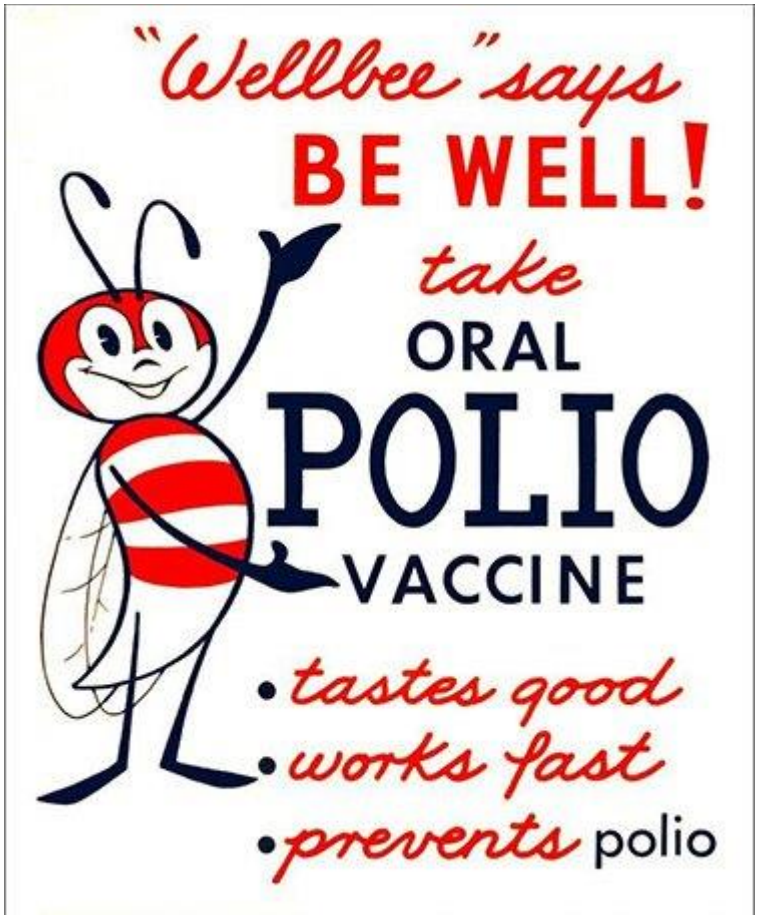
The oral polio vaccine was manufactured by Lederle laboratories.

The vaccine was contaminated with a cancer causing monkey virus called SV40.

In 1994 the company was sold and renamed Wyeth.

Wyeth was bought by Pfizer in 2009.

So let's look at Pfizer now.



How Did the Health Authorities Deal with Covid and Children? Any parallels to SV40?

On Tuesday of this week the FDA's Vaccines and Related Biological Products Advisory Committee voted to approve the use of the Pfizer Covid vaccine in 5 to 11 year old children for Emergency Use. They want to put this experimental injection into our kids.

How Did FDA Committee Do This?

The clinical trials were not double blind. Pfizer didn't follow the gold standard. (But the committee didn't care.)

The clinical trial started in June of this year. Only four months from start to end. Most vaccines are tested for many years. (But the committee didn't care.)

There were no long-term safety studies. Pfizer provided safety data on two groups of children ages 5 to 11. The first group was followed for about two months, the second for only two-and-a-half weeks. (But they didn't care.)

How Did FDA Committee Do This?

For children 5 to 12 years of age, there were 143 Covid-related deaths in the U.S. through Oct. 14, 2021. There are approximately 25 million kids that age in the U.S. Covid deaths is equivalent to .0000057 or .00057% (But that didn't seem to matter.)

Covid morbidity and mortality statistics are based on unreliable and unlicensed PCR tests. In addition, Covid deaths have a comorbidity rate of around 95% which means that for otherwise healthy children, the threat of Covid is almost nil. (But that didn't seem to matter all that much.)

How Did FDA Committee Do This?

But even using their inflated Covid death numbers, about 10 times as many children actually die of cancer. The Covid vaccines contain never before seen chemicals and nanoparticles and were NOT even tested to see if they can cause cancer. (But the committee didn't care.)

The two-month clinical study included 2,268 children ages 5 to 11. This equals .009% of that population in the U.S. Can a study detect all side effects when only one out of every 10,000 children is being represented? No.
(But they didn't care.)

How Did FDA Committee Do This?

According to actual research, children are at a very low risk of infecting other children and adults with Covid. And the idea of injecting healthy kids with experimental products to theoretically protect adults is morally reprehensible.
(But they didn't care.)

How Did FDA Committee Do This?

The Pfizer Covid vaccine caused myocarditis in other age groups. Myocarditis is an inflammation of the heart muscle. Clots can form in your heart, leading to a stroke or heart attack. According to Pfizer, the number of children in the study were "too small to detect any potential risks of myocarditis associated with vaccination." In other words, they don't know if the vaccine will cause myocarditis in children because the study wasn't big enough to tell. And it will NOT be studied until after the vaccine is injected into millions of children. Our kids are the guinea pigs. (But they didn't care about that.)

How Did FDA Committee Do This?

In the clinical trial, all serious adverse events were considered unrelated to the vaccine. (But they didn't care.)

Dr. William Gruber of Pfizer said they did NOT assess whether the vaccine prevents Covid-19 transmission in children. Isn't that important? (But the committee didn't care.)

VAERS is the Vaccine Adverse Events Reporting System. Recent VAERS data released by the CDC included a total of 798,636 reports of side effects (i.e. "adverse events") from all age groups following Covid vaccines, including 16,766 deaths and 117,399 serious injuries between Dec. 14, 2020 and Oct. 8, 2021. (But the committee didn't care.)

How Did FDA Committee Do This?

From the 10/8/2021 release of VAERS data:

Found 798,636 cases where Vaccine is COVID19

Table

Event Outcome	Count	Percent
Death	16,766	2.1%
Permanent Disability	24,805	3.11%
Office Visit	124,397	15.58%
Emergency Room	57	0.01%
Emergency Doctor/Room	89,867	11.25%
Hospitalized	79,446	9.95%
Hospitalized, Prolonged	223	0.03%
Recovered	247,355	30.97%
Birth Defect	555	0.07%
Life Threatening	18,238	2.28%
Not Serious	344,805	43.17%
TOTAL	† 946,514	† 118.52%

† Because some cases have multiple vaccinations and symptoms, a single case can account for multiple entries in this table. This is the reason why the Total Count is greater than 798636 (the number of cases found), and the Total Percentage is greater than 100.

How Did FDA Committee Do This?

VAERS Data revealed:

57 reports of anaphylaxis among 12- to 17-year-olds where the reaction was life-threatening, required treatment or resulted in death — with 96% of cases attributed to Pfizer's Covid vaccine.

529 reports of myocarditis and pericarditis (heart inflammation) with 508 cases attributed to Pfizer's Covid vaccine.

117 reports of blood clotting disorders, with all cases attributed to Pfizer's Covid vaccine.
(But the committee didn't care all that much.)

How Did FDA Committee Do This?

According to Jessica Rose, Ph.D., a VAERS analyst and computational biologist, VAERS does NOT accurately track ALL adverse reactions due to several flaws, including a "huge" backlog of data waiting to be entered into the system — she put the number at "hundreds of thousands."
(But they didn't care.)

A Harvard Pilgrim study showed that the number of VAERS reports is under-reported by a factor of 100. Only about 1% of reports of deaths and injuries from vaccines end up in the database. (But the committee didn't bother to discuss.)

How Did FDA Committee Do This?

Maddie De Garay is a 12-year old girl who volunteered to participate in the Pfizer Covid clinical trial. She was a healthy, straight-A student from Ohio who enjoyed life before receiving her second dose of the Pfizer Covid-19 vaccine. She suffered severe side effects and is now confined to a wheelchair.
(But the committee didn't care to discuss it.)

How Did FDA Committee Do This?

The FDA panel admitted that they have no idea of what side effects may exist from the vaccine if a child is already exposed to Covid. (But they didn't care.)

The FDA doesn't know if the benefits of the vaccine outweigh the risks in children because they don't know either the risks or the benefits. (But they didn't care.)

In 2009, the Justice Department Announced the Largest Health Care Fraud Settlement in Its History.
Pfizer had to Pay $2.3 Billion for Fraudulent Marketing
(But the committee didn't care.)

How Did FDA Committee Do This?

There are safe and effective treatments for Covid such as Ivermectin and Hydroxychloroquine. Since they exist an emergency use vaccine full of experimental technologies and untested ingredients is not necessary.
(But the committee didn't care.)

Some ingredients of the Pfizer vaccine:
((4-hydroxybutyl)azanediyl)bis(hexane-6,1-diyl)bis (2-hexyldecanoate), 2 [(polyethylene glycol)-2000]-N,N-ditetradecylacetamide
1,2-Distearoyl-snglycero-3- phosphocholine

How Did FDA Committee Do This?

As in the case of SV40, our health authorities did NOT act independently or responsibly in protecting the public or our children. They just didn't care! But they did seem to care about Pfizer.

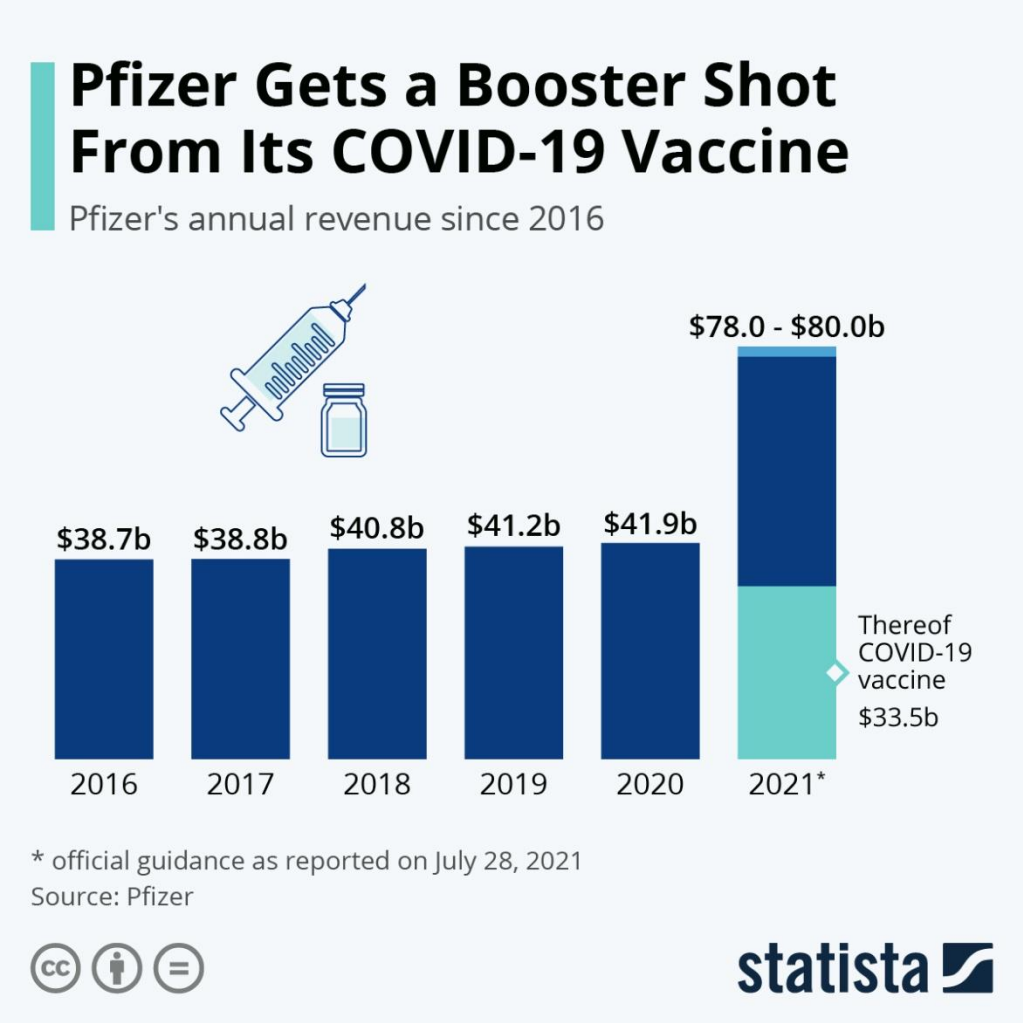
Pfizer's Health Seems Quite Robust

According to Forbes:

"Pfizer expects to generate $33.5 billion in Covid-19 vaccine sales in 2021, up from previous estimates of $26 billion, according to its second quarter earnings reports."

$33,500,000,000.00

Pfizer's Health Seems Quite Robust



Pfizer's Health Seems Quite Robust

According to NBC News:
The Biden administration has purchased 65 million pediatric doses of the Pfizer/BioNTech vaccine — enough to vaccinate an estimated 28 million children.

Why Don’t They Care?

Money

The FDA gets paid by vaccine makers. The FDA actually receives and depends on money from companies like Pfizer in order to function. In 1992, the drug industry started paying the salaries of drug reviewers.

The CDC is in the vaccine business because of its financial licensing interests in various vaccine products and its vaccine patents.

Individuals on the FDA and CDC committees get paid from vaccines. For example, Paul Offit was on the FDA committee. He makes millions from vaccines.

Why Don't They Care?

Group Think

"Groupthink is a phenomenon that occurs when a group of individuals reaches a consensus without critical reasoning or evaluation of the consequences or alternatives. Groupthink is based on a common desire not to upset the balance of a group of people."

No Alternative Viewpoints Allowed

There are no truly independent scientists allowed on these committees.

Cooked Industry Data?

The company with the most to gain, such as Pfizer can, supply their own data and they do.

What's the Antidote?

What's the antidote to a system where medical decisions are being made by people who have their own motives and agenda?

There's only one: Medical Freedom

As long as we can make our own health decisions for ourselves and our children then all the nonsense perpetrated by these corrupt agencies cannot hurt us.

Medical Freedom protects us from the corruption. But without medical freedom we will all be victims of this corruption.

Some Principles

Choice in medical care is a human right that must be protected.

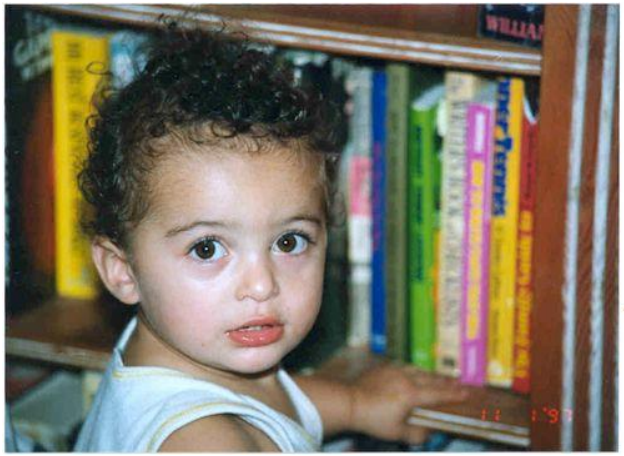
Medical mandates such as vaccine passports or whatever they want to call them should be eliminated because they do not allow for informed consent.

Medical decisions are personal decisions that should be made by the individual with the help of their professional medical provider, not a bureaucrat.

Some Principles

People should always have the right to refuse a medical procedure.

Health decisions for children should be the responsibility of their parents and their doctors, not a bureaucracy. We do NOT co-parent with the government.

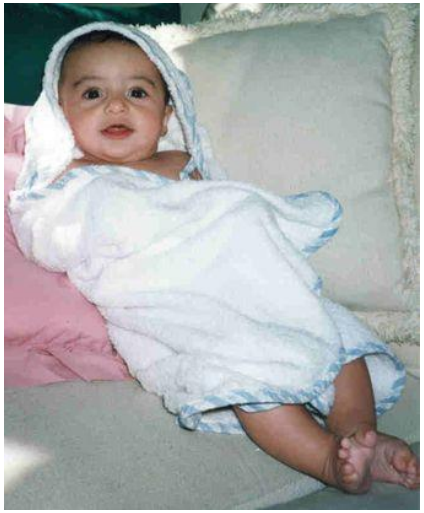


Alexander before the vaccine

Some Principles

“Those who won our independence... valued liberty as an end and as a means. They believed liberty to be the secret of happiness and courage to be the secret of liberty.” – Supreme Court Justice, Louis D. Brandeis

Let us all be courageous in protecting our freedom and our health for ourselves and our children.



Alexander before the vaccine