Judy A. Mikovits, PhD
April 27, 2019

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PLAGUE OF CORRUPTION:

Restoring Faith in the Promise of Science

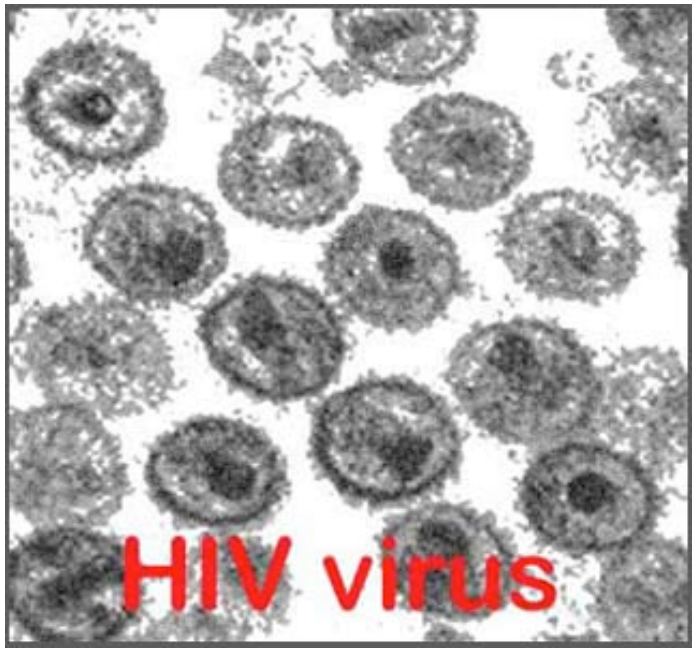
By Kent Heckenlively, J.D.
& Judy Mikovits, Ph.D.

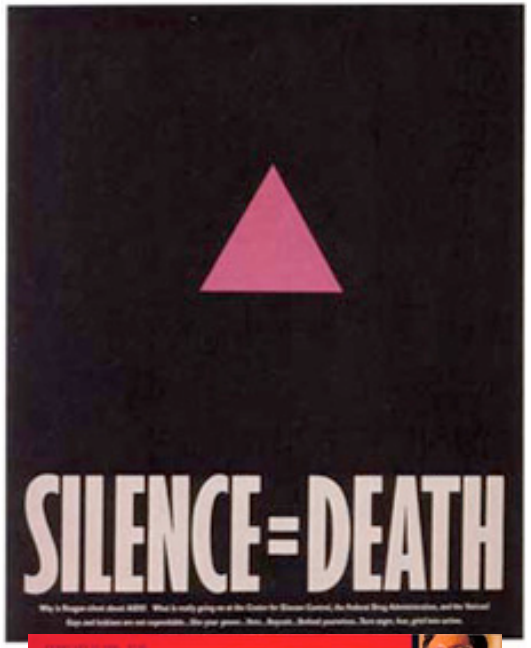
To go against conscience is neither right nor safe.
Therefore, I cannot and will not recant. Here I stand.
I can do no other. God help me, amen.

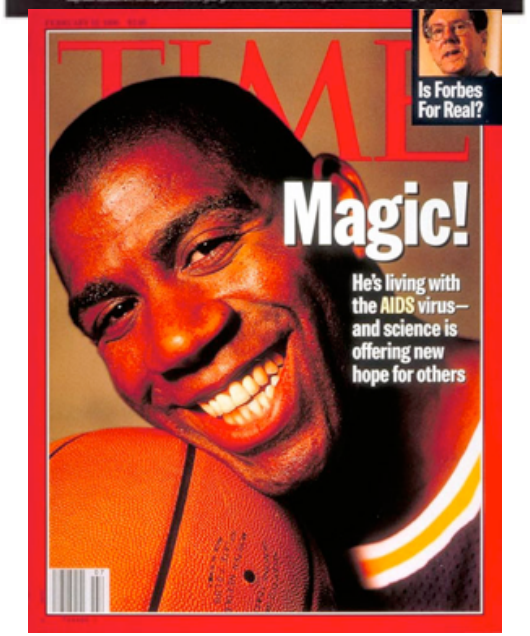
From the movie, *Martin Luther* (1953)

Political Influence on Scientific Research and the Impact it has on us ALL

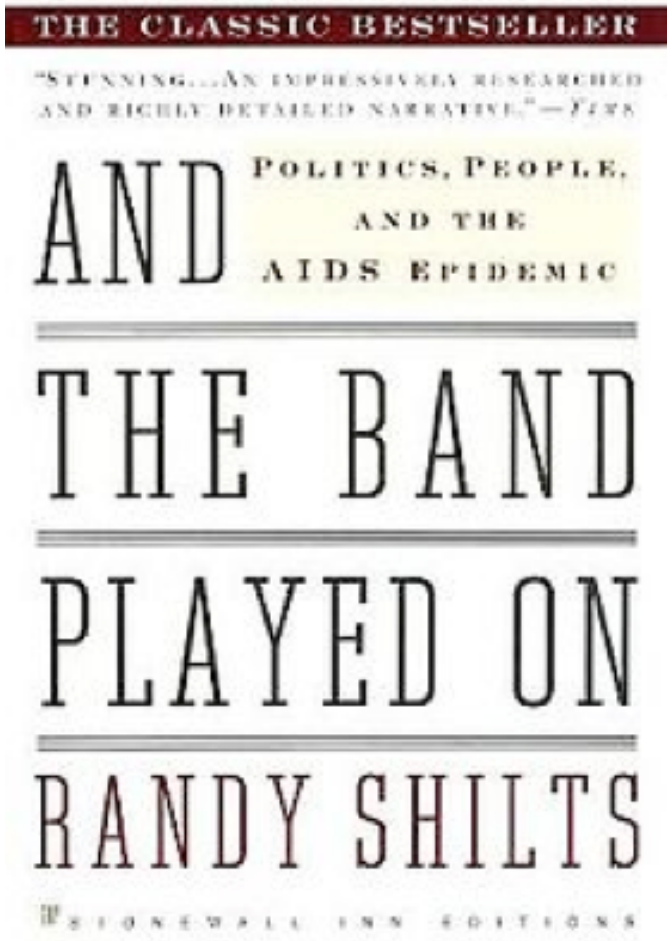
HIV -1 Isolation- 1982



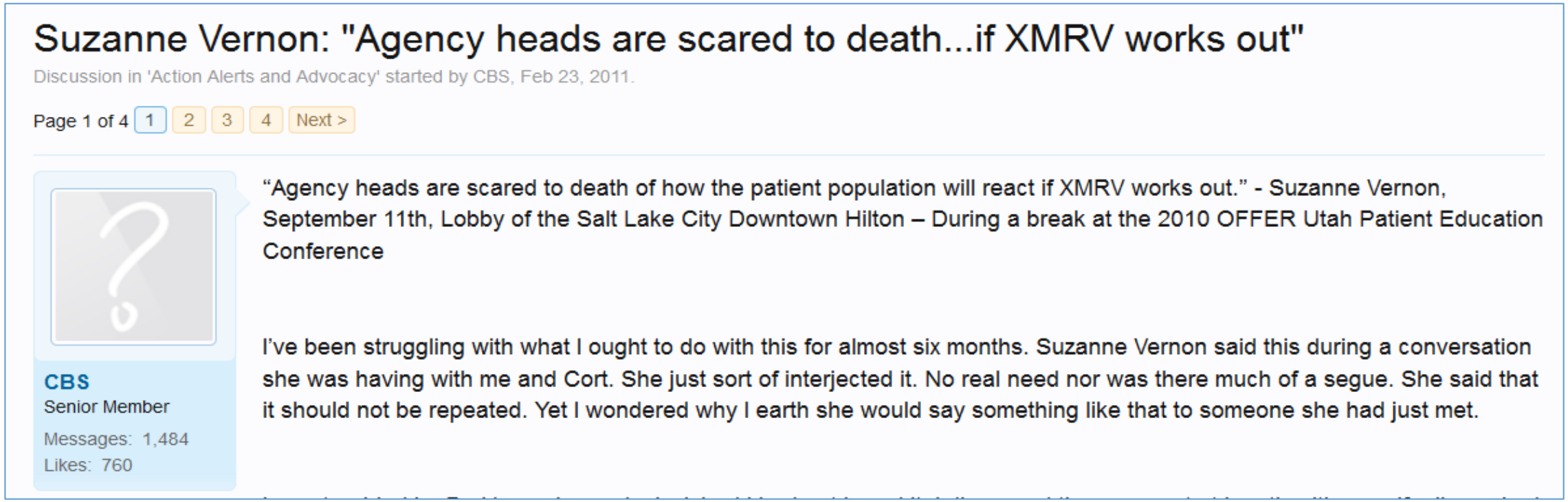




November 7,1991



MANY DEATHS BEFORE ESTABLISHMENT BELIEVED IN RETROVIRAL CAUSE



Suzanne Vernon: "Agency heads are scared to death...if XMRV works out"

Discussion in 'Action Alerts and Advocacy' started by CBS, Feb 23, 2011.

Page 1 of 4 1 2 3 4 Next >

CBS
Senior Member
Messages: 1,484
Likes: 760

"Agency heads are scared to death of how the patient population will react if XMRV works out." - Suzanne Vernon, September 11th, Lobby of the Salt Lake City Downtown Hilton – During a break at the 2010 OFFER Utah Patient Education Conference

I've been struggling with what I ought to do with this for almost six months. Suzanne Vernon said this during a conversation she was having with me and Cort. She just sort of interjected it. No real need nor was there much of a segue. She said that it should not be repeated. Yet I wondered why I earth she would say something like that to someone she had just met.

September 11. 2010

Why? Because in 1991 ONE million Americans were Infected with HIV
in 2010 when studies showed between 10-25 Million Americans were infected with XMRVs

THE BLOOD Supply IS CONTAMINATED with MLV-related viruses!

NYAS Mikovits March29,2011

Summary/Conclusions

- Data suggest there are different strains of Gamma Retroviruses that can infect humans
- Assays that capture the variation of these viruses in the blood supply are the best i.e. Serology and transmission
- Cerus Technologies can inactivate infectious strains of XMRV/HGRVs in Blood Components
- New Disease associations include leukemia, lymphoma and the platelet/megakaryocyte disorder, ITP
- Need more full length sequencing!!!

SCIENTIFIC AMERICAN™

FDA Approval December 1 2014 of Intercept Blood System

Permanent Address: http://www.scientificamerican.com/article/the-intercept-blood-system-rids-blood-donations-of-all-pathogens/

Health » Scientific American Volume 313, Issue 1 » Advances

The INTERCEPT Blood System Rids Blood Donations of All Pathogens

Blood banks begin using the method in donations this summer as the northward spread of chikungunya continues

By Tara Haelle | Jun 16, 2015 |

"*Science* started this and *Science* is going to End This"

John Coffin to Frank Ruscetti November 2010

Failure to Confirm XMRV/MLVs in the Blood of Patients with Chronic Fatigue Syndrome: A Multi-Laboratory Study

Graham Simmons,[1] Simone A. Glynn,[2] Anthony L. Komaroff,[3] Judy A. Mikovits,[4] Leslie H. Tobler,[1] John Hackett Jr.,[5] Ning Tang,[5] William M. Switzer,[6] Walid Heneine,[6] Indira K. Hewlett,[7] Jiangqin Zhao,[7] Shyh-Ching Lo,[8] Harvey J. Alter,[9] Jeffrey M. Linnen,[10] Kui Gao,[10] John M. Coffin,[11] Mary F. Kearney,[12] Francis W. Ruscetti,[12] Max A. Pfost,[4] James Bethel,[13] Steven Kleinman,[14] Jerry A. Holmberg,[15] Michael P. Busch,[1]* for the Blood XMRV Scientific Research Working Group (SRWG)†

12 September 2011; accepted 20 September 2011
Published online 22 September 2011;

Mikovits said she hopes to have full sequences of her new viruses "in a couple of weeks." **–JON COHEN**

VIROLOGY

The Waning Conflict Over XMRV And Chronic Fatigue Syndrome

OTTAWA, CANADA—Less than a day after a new study dealt what many consider a lethal blow to the controversial theory that a newly detected virus, XMRV, is linked to chronic fatigue syndrome (CFS), proponents and skeptics of the theory squared off in a meeting here.

In one corner was Judy Mikovits, research director at the Whittemore Peterson Institute for Neuro-Immune Disease (WPI) in Reno, Nevada, and the main champion of the idea that XMRV and its relatives play a role in CFS. Her opponent, an erstwhile supporter, was heavyweight retrovirologist John Coffin of the Tufts University Sackler School of Graduate Biomedical Sciences in Boston. When Mikovits and Coffin took the stage at the meeting, which was organized by IACFS/ME (an international association devoted to the disease) and attracted 460 researchers and patients, they sat on opposite sides of the lectern. During their introductions, Coffin clasped his hands in front of his mouth, looking like a man in

had asserted—explained the XMRV DNA it found in some patient samples.

In Ottawa, Mikovits came out swinging. But she didn't make the case for XMRV, which stands for xenotropic murine leukemia virus–related virus. Instead, she offered new evidence that people with CFS (known as myalgic encephalomyelitis in some countries) had a virus "highly related" to XMRV.

Unlike the original study that appeared in *Science* that showed entire sequences of XMRV and infection of fresh cells, Mikovits revealed only partial viral sequences that she

Pro and con. Judy Mikovits (*left*) argued for the link between human gammaretro-

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Solution for Agency Heads for 2009 and 2010 XMRV Publications in Elite Journals? ?
Plague of Corruption

Lasker Award Winner , Harvey Alter , Confirms findings

So it's really probably a better term is murine leukemia virus-related viruses which encompasses XMRV so we found this in a very high percentage of the chronic fatigue patients that Dr. Komaroff had sent to us—about 86 percent—and simultaneously found that in about 6.6 percent of our healthy blood donors.

So there was a dramatic association with chronic fatigue syndrome, with the syndrome of chronic fatigue but that's all it is . . . we think basically it confirms the findings of the Whittemore Peterson group.[20]

Detection of an Infectious Retrovirus, XMRV, in Blood Cells of Patients with Chronic Fatigue Syndrome

Vincent C. Lombardi,[1]* Francis W. Ruscetti,[2]* Jaydip Das Gupta,[3] Max A. Pfost,[1] Kathryn S. Hagen,[1] Daniel L. Peterson,[3] Sandra K. Ruscetti,[4] Rachel K. Bagni,[5] Carl Petrow-Sadowski,[6] Bert Gold,[2] Michael Dean,[2] Robert H. Silverman,[3] Judy A. Mikovits[1]

www.sciencemag.org SCIENCE VOL 326 23 OCTOBER 2009

Taken together, these data demonstrate the first direct isolation of infectious XMRV from humans and implicate a role for XMRV infection in the pathogenesis of CFS."

The original abstract of the *Science* article which was published on October, 8, 2009

Detection of MLV-related virus gene sequences in blood of patients with chronic fatigue syndrome and healthy blood donors

Shyh-Ching Lo[a,1], Natalia Pripuzova[a], Binjie Li[a], Anthony L. Komaroff[b], Guo-Chiuan Hung[a], Richard Wang[a], and Harvey J. Alter[a,1]

[a]Tissue Microbiology Laboratory, Division of Cellular and Gene Therapies and Division of Human Tissues, Office of Cellular, Tissue and Gene Therapy, Center for Biologics Evaluation and Research, Food and Drug Administration, Bethesda, MD 20892; [b]Department of Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, MA 02115; and [c]Department of Transfusion Medicine, The Warren Grant Magnuson Clinical Center, National Institutes of Health, Bethesda, MD 20892

Using Lombardi et al. nested PCR methods, *gag* sequences more closely related to polytropic MLV than to XMRV were detected

gag sequences were found in 86.5% of CFS patients' samples drawn in 1991-4 and in 6.8% of control samples

8/9 CFS patients exhibited the same *gag* sequences in blood freshly drawn 15 years later

No mouse mitochondrial DNA could be detected in the samples

Lo et al. presented no evidence of infectious virus

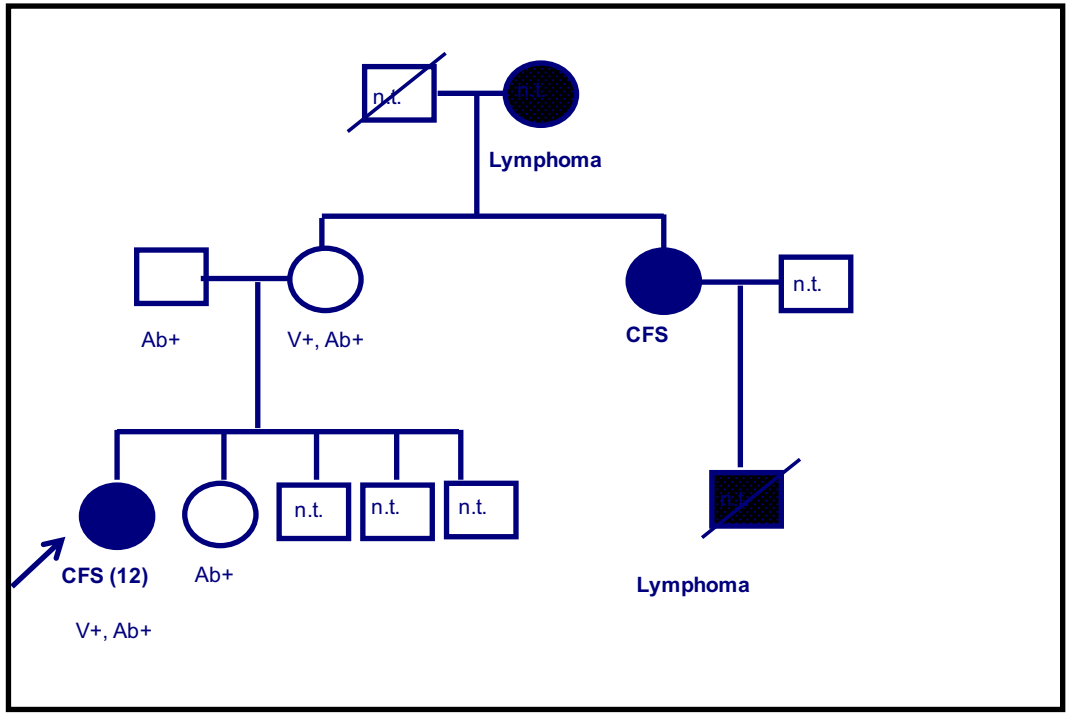
In 2011 Harvey Alter picked the poison to have this 2010 confirmatory study withdrawn

Contamination of the Blood Supply, Additional Strains, Disease Associations Infection in the Worldwide

XMRV/MRVs Detection in Cancer & Blood Diseases in addition to Prostate Cancer and CFS

ID#	XMRV status	Cancer/blood disease
1103	positive	MCL
1109	positive	Thymoma
1118	positive	myelodysplasia
1125	positive	MCL
1186	positive	Lymphoma
1199	positive	Lymphoma
1150	positive	Lymphoma
3818	positive	MCL
1174	positive	Thymoma
1205	positive	lymphoma
1172	positive	MCL
3848	positive	ITP
3827	positive	ITP
1113	positive	CLL
1322	Not tested	MCL
1181	positive	CLL
1188	positive	CLL
1189	positive	MCL
3814	positive	ITP

XMRV in Families of CFS Patients



21st Century Acquired Endocannabinoid Immune Dysfunction: *Unintended?* Consequences of Unsafe Vaccinations & CDC Schedule

Prostate*	Lupus	ME/CFS*
Breast*	Crohn's*	Gulf War Syndrome*
Multiple Myeloma*	Hashimoto's Thyroiditis*	Autism/ASD*
Non Hodgkin's Lymphoma*	Polymyositis	MS*
Chronic Lymphocytic Leukemia*	Sjogren's Syndrome	Parkinson's*
Mantle Cell Lymphoma*	Bechet's Disease*	ALS*
Hairy Cell Leukemia	Primary Billary Cirrhosis*	Fibromyalgia
Bladder*	IBD*	Chronic Lyme Disease*
Coloretal	Psoriasis, Dermatitis	OCD
Kidney*	Diabetes	ADHD
Ovarian*	Cardiovascular Disease*	PTSD
	Neuroendocrine tumors*	Psychosis

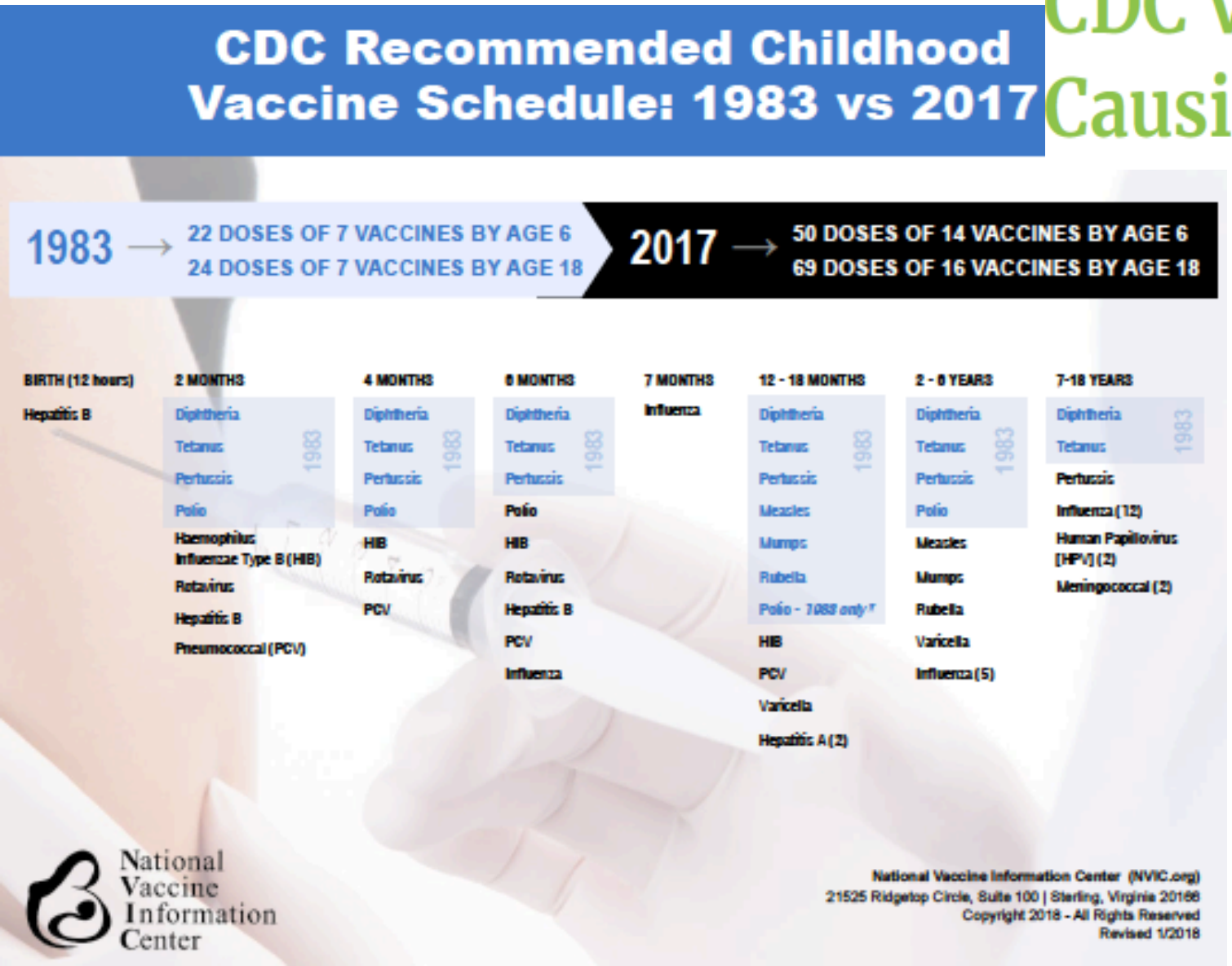
"One of the most widely distributed biological products that frequently involved mice or mouse tissue, at least up to recent years, are vaccines, especially vaccines against viruses . . .

It is possible that XMRV particles were present in virus stocks cultured in mice or mouse cells for vaccine production, and that the virus was transferred to the human population by vaccination

Frontiers in Microbiology published January 2011

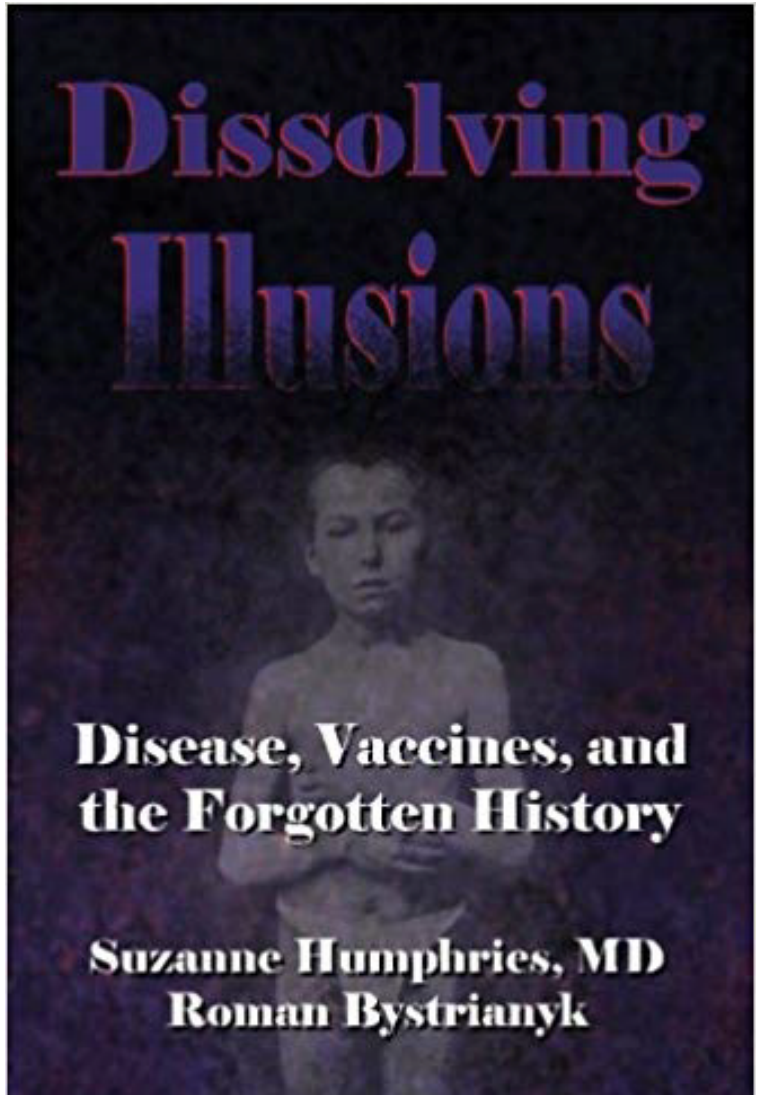
MARCH 19, 2019

Real-Life Data Show that the CDC Vaccine Schedule is Causing Harm



“Before Litigation There was Truth”
Mike Hugo

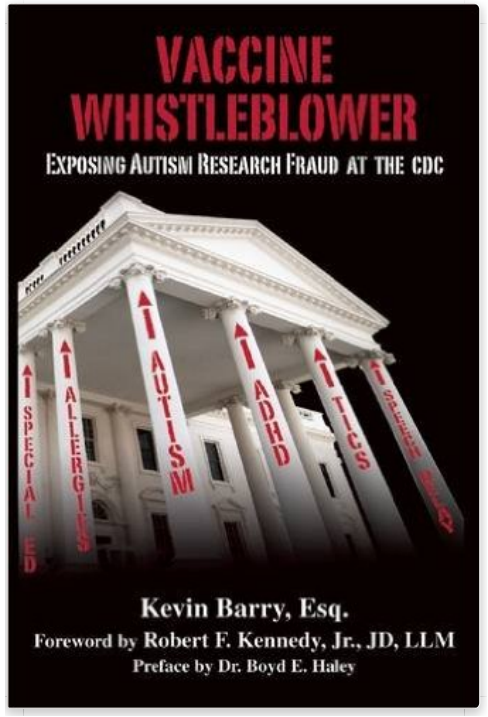
Vaccines didn’t save humanity. Their impact was somewhere between 1-3.5% of the total decline in mortality rates. Improvement in sanitation and standards of living really did (nutrition, living conditions, etc.). Did vaccines contribute to a small decrease of certain acute illnesses? Yes, but their relative benefit is often exaggerated to an extreme, and then used to browbeat, guilt, and scare parents.



Given Protection from Liability by 1986 National Childhood Vaccine Injury Act: Federal Agencies, Pharmaceutical Companies, Mainstream media and Scientific Journals Collude to Censor Truth

Three Main Areas:

1. Vaccine Safety Efficacy
2. Scientific Discovery
3. Alternative Therapies



Vaccine Excipient & Media Summary

Excipients Included in U.S. Vaccines, by Vaccine

Vaccine	Contains	Source: Manufacturer's P.I. Dated
DTaP (Daptacel)	Stainer-Scholte medium, modified Mueller's growth medium, modified Mueller-Miller casamino acid medium (without beef heart infusion), dimethyl 1-beta-cyclodextrin, ammonium sulfate	July, 2012
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	July, 2012
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	July, 2012
DTaP-HepB-IPV (Pediarix)	formaldehyde, gluteraldehyde, 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	August, 2012

"It's troubling to me that in a recent Senate hearing on childhood vaccinations, it was never mentioned that our government has paid out over three billion dollars through a Vaccine Injury Compensation Program for children who have been injured by vaccinations."

-Congressman William Posey on the floor of the US House of Representatives, July 29, 2015.

EBioMedicine 17 (2017) 192–198

Contents lists available at ScienceDirect

EBioMedicine

journal homepage: www.ebiomedicine.com

Research Paper

The Introduction of Diphtheria-Tetanus-Pertussis and Oral Polio Vaccine Among Young Infants in an Urban African Community: A Natural Experiment

Søren Wengel Mogensen [a,1], Andreas Andersen [b,1], Amabelia Rodrigues [a], Christine S Benn [b,c], Peter Aaby [a,b,*]

[a] Bandim Health Project, Indepth Network, Apartado 861, Bissau, Guinea-Bissau
[b] Research Centre for Vitamins and Vaccines (CVIVA), Bandim Health Project, Statens Serum Institute, Artillerivej 5, 2300 Copenhagen S, Denmark
[c] OPEN, Institute of Clinical Research, University of Southern Denmark/Odense University Hospital, 5000 Odense C, Denmark

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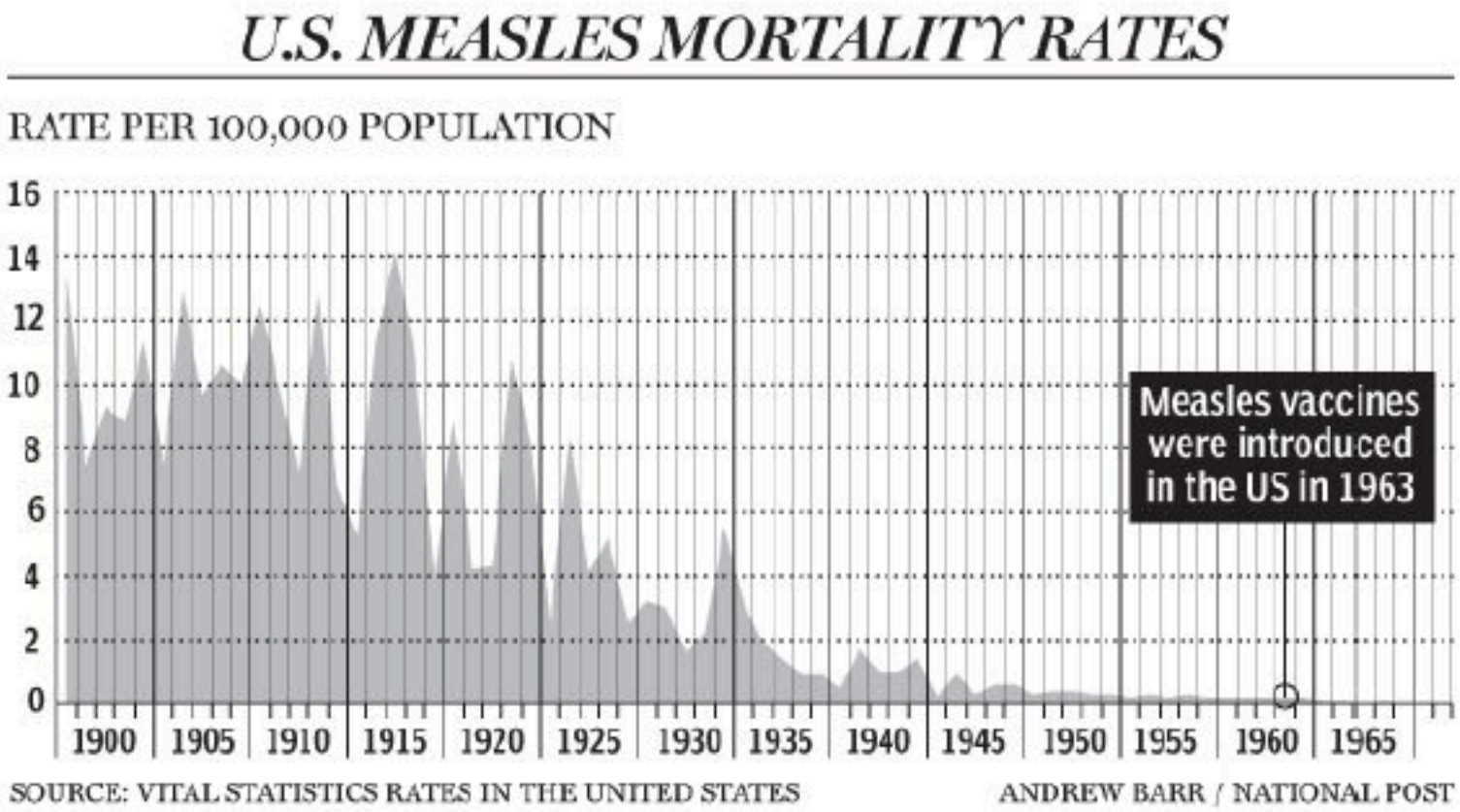
Keywords:
Diphtheria-tetanus-pertussis vaccine
DTP
Measles vaccine
Non-specific effects of vaccines
Oral polio vaccine

ABSTRACT

Background: We examined the introduction of diphtheria-tetanus-pertussis (DTP) and oral polio vaccine (OPV) in an urban community in Guinea-Bissau in the early 1980s.
Methods: The child population had been followed with 3-monthly nutritional weighing sessions since 1978. From June 1981 DTP and OPV were offered from 3 months of age at these sessions. Due to the 3-monthly intervals between sessions, the children were allocated by birthday in a 'natural experiment' to receive vaccinations early or late between 3 and 5 months of age. We included children who were < 6 months of age when vaccinations started and children born until the end of December 1983. We compared mortality between 3 and 5 months of age of DTP-vaccinated and not-yet-DTP-vaccinated children in Cox proportional hazard models.
Results: Among 3–5-month-old children, having received DTP (± OPV) was associated with a mortality hazard ratio (HR) of 5.00 (95% CI 1.53–16.3) compared with not-yet-DTP-vaccinated children. Differences in background factors did not explain the effect. The negative effect was particularly strong for children who had received DTP-only and no OPV (HR = 10.0 (2.61–38.6)). All-cause infant mortality after 3 months of age increased after the introduction of these vaccines (HR = 2.12 (1.07–4.19)).
Conclusion: DTP was associated with increased mortality; OPV may modify the effect of DTP.

Lawrence Solomon: The untold story of measles

Several decades following the vaccine's introduction, the measles death rate rose, largely because the vaccine made adults, expectant mothers and infants more vulnerable



Review

CellPress

Effects of environmental change on zoonotic disease risk: an ecological primer

Agustín Estrada-Peña[1], Richard S. Ostfeld[2], A. Townsend Peterson[3], Robert Poulin[4], and Jo

[1] Department of Animal Pathology, Faculty of Veterinary Medicine, Miguel Servet, 177, 50013-Zaragoza, Spain
[2] Cary Institute of Ecosystem Studies, Millbrook, NY 12545-0129, USA
[3] The University of Kansas Biodiversity Institute, Lawrence, KS 66045-7593, USA
[4] Department of Zoology, University of Otago, Dunedin 9016, New Zealand
[5] SaBio, IREC, Ronda de Toledo s/n, 13071 Ciudad Real, Spain
[6] Center for Veterinary Health Sciences, Oklahoma State University, Stillwater, OK 74078, USA

Trends in Parasitology, April 2014, Vol. 30, No. 4 205

Anatomy of a science study censorship

March 20, 2019 (http://www.ghostshipmedia.com/2019/03/20/how-elsevier-censored-a-sheep-study-that-could-sting-mercks-profits/) Ghost Ship Media (http://www.ghostshipmedia.com/author/gh923ssh1px/)

"*Dear Dr. Luján,*

"I wanted to step in here to say that your manuscript is not being retracted – which implies wrongdoing and could damage your professional reputation," Anne-Marie Pordon, publisher of Pharmacology and Pharmaceutical Sciences titles for Elsevier journals interjected in a heated e-mail exchange between the lead researcher and various editors. *"We are withdrawing the paper, which does not imply misconduct in any way. There will be simply a statement that says "This paper has been withdrawn at the request of the _____" (Authors or Editors in the blank.)"* Pick your poison. You remove it, or we remove it.

"That doesn't mean there isn't another gammaretrovirus to be found. I think enough evidence has been presented that maybe another infectious retrovirus is there. These studies will continue to go on, looking for MLV-related viruses."

John Coffin NIH State of the Knowledge Workshop on ME/CFS from April, 2011

Taken together these data suggest there are
additional human gamma retroviruses which may be involved in the Pathogenesis of neuroimmune disease and cancer!

"The question, which urgently needs to be answered is whether the plague feared by Coffin and Stoye has already arrived, but we do not Recognize it…
THEY SEE WHAT THEY WANT TO SEE AND THAT'S THE ***REAL PLAGUE"***

Plague Chapter 21 p382

What did we find in the MMRV (Priorix Tetra) vaccine?

VACCINEGATE - EN | STAFF CORVELVA | 21 APRILE 2019 | VISITE: 37399

Residual DNA/RNA deriving from cultured cells - Total amount of DNA: 1.7-3.7 µg/dose,
the 80% of which was human (Human fetal DNA / RNA from the MRC-5 cell line). Other amount of DNA: chicken

Adventitious viruses - Human endogenous retrovirus K, Equine infectious anemia virus, Avian leukosis virus, HERV-H/env62
Other microbial contaminants - Proteobacteria, nematode-helminth

Human endogenous retrovirus K - 32 sequences
Equine infectious anemia virus - 2 sequences
Avian leukosis virus - 2 sequences
HERV-H/env62 - 4 sequences
These viruses are known to be adventitious vaccine contaminants and are known to be potentially dangerous, which is why manufacturers are required to verify that they are completely absent from the vaccine.
It follows that this in-depth analysis in this vaccine confirms two nonconformities on efficacy and safety:
The presence of rubella in a very low number of copies (subthreshold)
The presence of potentially dangerous adventitious viruses which certifies that there is no adequate control on vaccines because if there were, these elements would have been detected.

The Name Game and the Immaculate Recombination

How many have we created, John? How many retroviruses are out there Judy Mikovits asking a question to Dr. John Coffin
at the Ottawa IACFS ME/CFS meeting 23 September 2011

Plague Chap 17 p 284

RESEARCH PAPER

Cancer Biology & Therapy 12:7, 617-628; October 1, 2011; © 2011 Landes Bioscience

Frequent detection of infectious xenotropic murine leukemia virus (XMLV) in human cultures established from mouse xenografts

Yu-An Zhang,[1] Anirban Maitra,[2] Jer-Tsong Hsieh,[3] Charles M. Rudin,[4] Craig D. Peacock,[4] Collins Karikari,[2] Rolf A. Brekken,[1] Victor Stastny,[1] Boning Gao,[1] Luc Girard,[1] Ignacio Wistuba,[5] Eugene Frenkel,[6] John D. Minna[1] and Adi F. Gazdar[1,*]

[1]Hamon Center for Therapeutic Oncology Research; [3]Department of Urology; [6]Division of Hematology-Oncology; The University of Texas Southwestern Medical Center at Dallas; Dallas, TX USA; [2]Departments of Pathology and Oncology; The Sol Goldman Pancreatic Cancer Research Center; [4]Department of Oncology; Johns Hopkins University School of Medicine; Baltimore, MD USA; [5]Department of Pathology; MD Anderson Cancer Center; Houston, TX USA

Table 3. Frequent detection of murine leukemia virus (MLV) contamination of non-xenograft human cultures

Characterization of murine leukemia viruses (MLV) detected in human non-xenograft cultures in xenograft culture laboratories

Table 1. Identification of xenotropic murine leukemia viruses (XMLV) and MLV-related viruses in xenograft cell lines

"However, the group also recommended that further studies be undertaken urgently and internationally to put into perspective the very low levels of RT activity found in the vaccines."

4.1. Initial findings

The discovery in 1995 of reverse transcriptase (RT) activity in marketed measles, mumps and rubella (MMR) vaccine raised concerns that the vaccine was contaminated by an unrecognized avian retrovirus with unknown safety implications.

4.2. Background

The usual flow of genetic information is from DNA to RNA. However, the reverse of that process was discovered to be mediated by an RNA-dependent DNA polymerase (reverse transcriptase) that some RNA viruses, such as retroviruses, use to reverse-transcribe their RNA genomes into DNA. That viral DNA can then be integrated into the host genome and replicated, resulting in the production of more RNA virus. RT activity has therefore been used as a biochemical marker for the presence of retroviruses. However, the genes that encode RT are widely distributed in eukaryotic organisms and all reverse transcriptases are evolutionarily related. In addition, cellular DNA-directed DNA polymerases can exhibit some ability to use RNA as a template and reverse-transcribe as well.

Biologicals 42 (2014) 223–236

Contents lists available at ScienceDirect

Biologicals

journal homepage: www.elsevier.com/locate/biologicals

Review

Adventitious agents in viral vaccines: Lessons learned from 4 case studies

John Petricciani [a,*], Rebecca Sheets [b], Elwyn Griffiths [c], Ivana Knezevic [d]

[a] IABS, POB 1925, Palm Springs, CA 92263, USA
[b] Grimalkin Partners, 13401 Norden Drive, Silver Spring, MD 20906, USA
[c] 3 The Farthings, Kingston Upon Thames, Surrey KT2 7PT, UK
[d] Group Lead, Norms and Standards for Biologicals, Department of Essential Medicines and Health Products (EMP) Health Systems and Innovation (HIS) Cluster, WHO L276, Avenue Appia 20, 1211 Geneva 27, Switzerland

ARTICLE INFO

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ABSTRACT

Since the earliest days of biological product manufacture, there have been a number of instances where laboratory studies provided evidence for the presence of adventitious agents in a marketed product. Lessons learned from such events can be used to strengthen regulatory preparedness for the future. We

AND Continues today: creating HOW Many New Diseases?

View full text

Expert Rev Vaccines. 2011 Mar;10(3):355-64. doi: 10.1586/erv.11.7.

New Japanese encephalitis vaccines: alternatives to production in mouse brain.

Halstead SB[1], Thomas SJ.

Author information

Abstract

Japanese encephalitis virus (JEV), a flavivirus maintained in a zoonotic cycle and transmitted by the mosquito Culex tritaeniorhynchus, causes epidemics of encephalitis throughout much of Asia.

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Health

Health

UPDATED: June 13, 2011 NO. 24 JUNE 16, 2011

A Mysterious Disease

People claim to suffer from an infectious condition dubbed HIV-negative AIDS

The Dangers of Xenotransplantation. Nature Medicine 1995

JP Stoye and JM Coffin

- Such viruses are widely distributed in mammalian species including pigs and baboons, potential

  donors for these procedures.
- Since they are inherited in the germ line in the form of proviral DNA,
- They are impossible to remove using the usual methods for deriving pathogen-free animals.

- Implanting an organ carrying a dormant endogenous retrovirus into a patient is equivalent to injecting the patient with live virus.

Xenotransplantation and Primates - Threats Masquerading as Cures.

September 1, 1996

- **Dr. John Coffin*, a leading expert on recombination in viruses, concluded "the infection is a virtually inevitable consequence" of xenotransplantation and "This is a very serious worry because the animals that have been chosen for doing this -- the baboon and the pig -- are both known to carry endogenous viruses, replication competent, but very poorly studied, that are capable of infecting human cells." He further suggested baboon bone marrow experiments could make the HIV-AIDS infection "worse by spreading the host range."**

- Despite scientific skepticism, the FDA supported the clinical experiment to transplant baboon cells into AIDS patient Jeff Getty. Prior to this decision, the FDA convened lengthy hearings of the National Academy of Sciences' Institute of Medicine and its own Biological Response Modifiers Advisory Committee. Dr. Marion Michaels*, from the University of Pittsburgh, told the committee that despite rigorous screening, "the donor organ, the tissue or the accompanying hematopoietic cells can also be the source of infection. Most often these infections are latent organisms and are often clinically silent in the donor."

Isn't Injecting babies and children with mouse viruses capable of infecting human cells the same thing??

U.S. Department of Health and Human Services
Food and Drug Administration
Center for Biologics Evaluation and Research
February 2002

They withdrew in on: Withdrawn May 2015 (Coffin was part of the meetings where they said partners of xeno were not at risk when all previous research said they were. They didn't want it to show that close contact relatives could catch something from a xeno recipient)

Withdrawn - Draft Guidance for Industry: Precautionary Measures to Reduce the Possible Risk of Transmission of Zoonoses by Blood and Blood Products from Xenotransplantation Product Recipients and Their Intimate Contacts

The guidance document entitled **"Draft Guidance for Industry: Precautionary Measures to Reduce the Possible Risk of Transmission of Zoonoses by Blood and Blood Products from Xenotransplantation Product Recipients and Their Intimate Contacts"** was withdrawn on May 8, 2015. Please visit: http://www.gpo.gov/fdsys/pkg/FR-2015-05-06/html/2015-10477.htm for additional information. that link goes to this:

[Federal Register Volume 80, Number 87 (Wednesday, May 6, 2015)]
[Notices]
[Pages 26059-26061]

Current Vaccine schedules compound damage in vulnerable populations with Chronic Disease and Cancer (20-30 Million Americans)

> "Activation of the cellular immune system is important in the pathogenesis of HIV disease, and that fact has given rise to concerns that the activation of the immune system through vaccinations might accelerate the progression of HIV disease . . . If feasible, it is preferable to have patients on antiretroviral therapy (ART) prior to receipt of vaccination." – Accessed May 3, 2013.
>
> UCSF Pediatric Pediatric AIDS Website on HIV and Immunization

- Sterile environments result lack of educated immune systems
- Vaccination schedules result in anergic immune systems that is the inability to mount an immune response to the antigen
- Toxic components exacerbate immune dysfunction resulting in aberrant expression of host endogenous RVs
- Reappearance of disease is BECAUSE of inappropriate vaccinations and the toxic components contained in them

Conclusions

Aberrant evolution of the human genome by:

- Replication competent retroviruses generated laboratories in current vaccines and cell cultures
- Increased zoonosis of novel retroviruses in human population from animal populations.
- That means GMOs and toxins in animals result in compromised immune systems and the expression of endogenous viruses ..eg Bovine leukemia virus
- These retroviruses CAN and have been shown to infect human cells and like HTLV, HIV are passed in milk and other fluids
- The blood supply is contaminated and the vaccines are contaminated as is food supply (milk)

Phytocannabinoids can Enhance Vaccine Efficacy and Reduce Neuroinflammation

www.nature.com/scientificreports

SCIENTIFIC REPORTS

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Transient Cannabinoid Receptor 2 Blockade during Immunization Heightens Intensity and Breadth of Antigen-specific Antibody Responses in Young and Aged mice

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