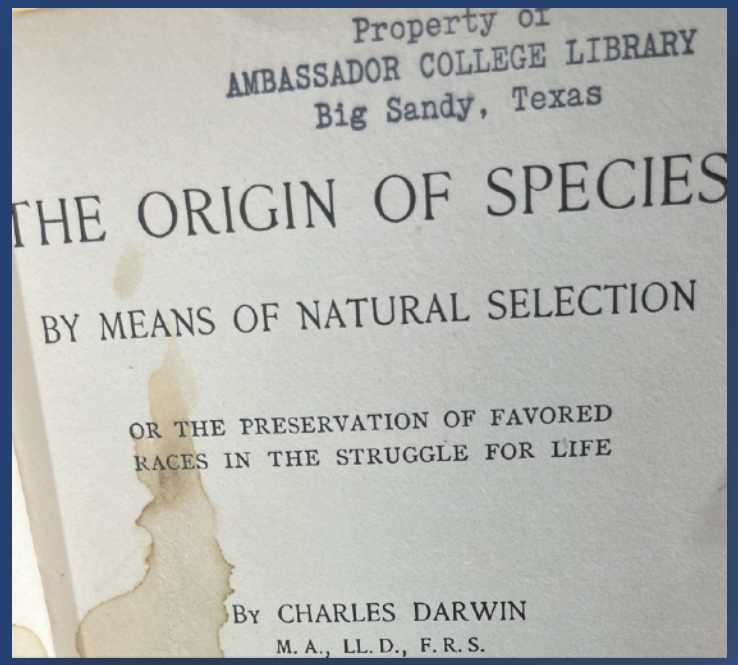
Vaccination is not Immunization It's Extermination

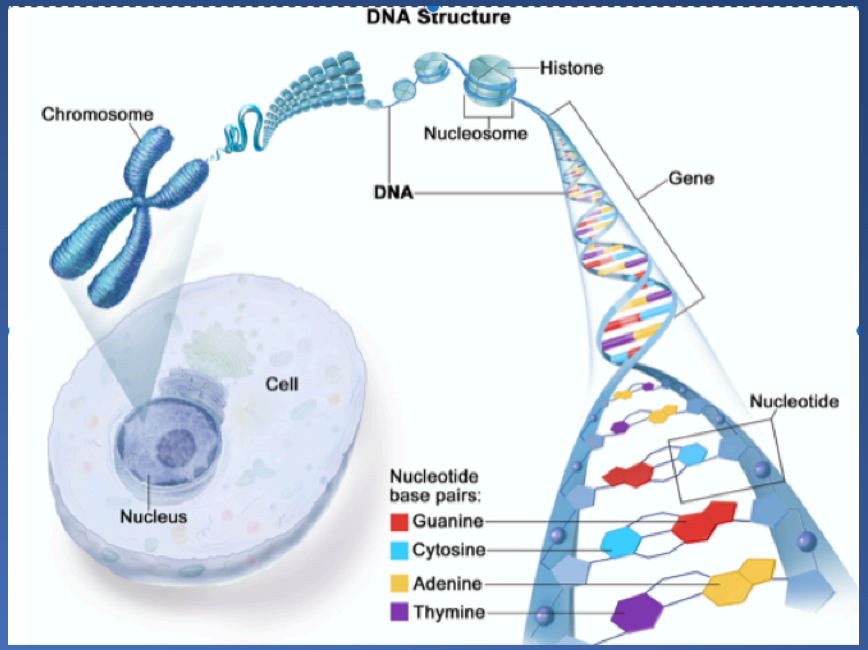
1828 Webster Definition Vaccination: EXTERMINATION of an UNWANTED VARMINT

What are "THEY" Afraid of? COVID VACCINE will crumble the confidence in ALL Vaccines?
We the People will REPENT and turn Back to GOD GIVEN NATURAL IMMUNIYTY



1859

1999



2015

2023 GOD WINS: THE FEAR OF THE LORD is the Beginning of Knowledge but Fools Despise Wisdom & Instruction (PROVERBS 1:7)

If my people , who are called by my name, will humble themselves,
pray & seek my face & turn from their wicked ways, then I hear from heaven, & I will forgive their Sin and will heal their land (2 Chronicles 7:14)

DISCLAIMER: I DON'T WORSHIP SCIENTISM

AIDS RESEARCH AND HUMAN RETROVIRUSES
Volume 36, Number 7, 2020
Mary Ann Liebert, Inc.
DOI: 10.1089/aid.2020.0095

COMMENTARY

Fake Science: XMRV, COVID-19, and the Toxic Legacy of Dr. Judy Mikovits

Stuart J.D. Neil[1] and Edward M. Campbell[2]

Abstract

One cannot spend >5 min on social media at the moment without finding a link to some conspiracy theory or other regarding the origin of SARS-CoV2, the coronavirus responsible for the COVID-19 pandemic. From the virus being deliberately released as a bioweapon to pharmaceutical companies blocking the trials of natural remedies to boost their dangerous drugs and vaccines, the Internet is rife with far-fetched rumors. And predictably, now that the first immunization trials have started, the antivaccine lobby has latched on to most of them. In the last week, the trailer for a new "bombshell documentary" *Plandemic* has been doing the rounds, gaining notoriety for being repeatedly removed from YouTube and Facebook. We usually would not pay much heed to such things, but for retrovirologists like us, the name associated with these claims is unfortunately too familiar: Dr. Judy Mikovits.

SCIENCEINSIDER HEALTH

Fact-checking Judy Mikovits, the controversial virologist attacking Anthony Fauci in a viral conspiracy video

In Plandemic, the former chronic fatigue syndrome researcher makes countless unsubstantiated claims and accusations

8 MAY 2020 • BY MARTIN ENSERINK, JON COHEN

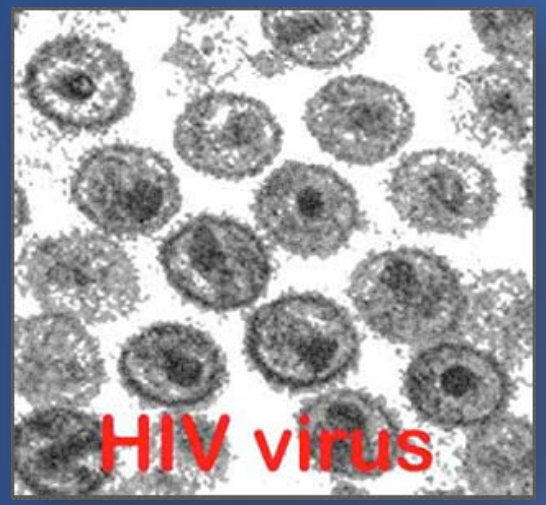
FAITH REQUIRES TRUST
PLANDEMICSERIES.COM

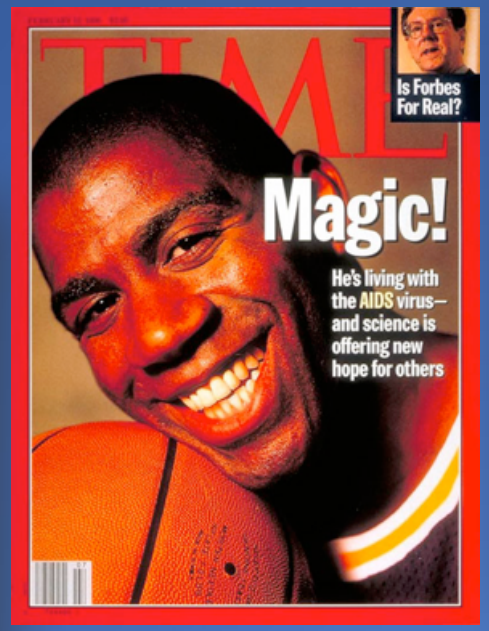
TRUST IN THE LORD WITH ALL YOUR HEART & LEAN NOT ON YOUR ON UNDERSTANDING (PROVERBS 3:5)

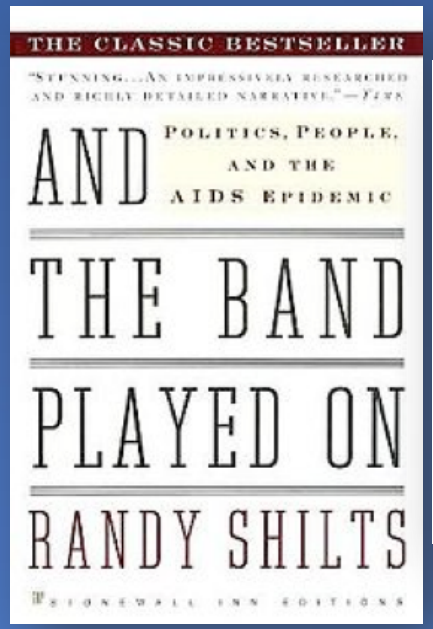
SINCE FAUCI AKA “SCIENCE” IS RESIGNING: A WALK DOWN CRIMES AGAINST HUMANITY LANE

AIDS Plandemic Driven By HBV Vaccine in HIV antibody Positive

LAV Isolation in GRID-1982







AIDS. 2016 Sep 24;30(15):2289-98. doi: 10.1097/QAD.0000000000001202.

Standard vaccines increase HIV-1 transcription during antiretroviral therapy.

Yek C[1], Gianella S, Plana M, Castro P, Scheffler K, García F, Massanella M, Smith DM.

Author information

Abstract

OBJECTIVES: Curative strategies using agents to perturb the HIV reservoir have demonstrated only modest activity, whereas increases in viremia after standard vaccination have been described. We investigated whether vaccination against non-HIV pathogens can induce HIV transcription and thereby play a role in future eradication strategies.

HIV First Fauci GOF 1984
HIV/LAV EM Antibodies DIFFERENT
HIV DID NOT EXIST IN NATURE

November 7,1991 MAGIC TESTS POSITIVE FOR HIV ANTIBODIES
November 14, 1991 MIKOVITS THESIS:

HIV COULD NOT CAUSE AIDS IF INNATE IMMUNITY Healthy

DANGERS OF USE Of ANIMAL RNA,DNA PROTEIN
All Vaccines are GMO Synthetic viruses

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,[1] Sandra K. Ruscetti,[4] Rachel K. Bagni,[5] Cari 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

- XMRV RNA/DNA in 67% of CFS patients tested
- XMRV protein detected in >85% stimulated/dividing T and B cells
- Antibody to XMRV Env detected in >50% CFS patient plasma
- Infectious virus transmitted from >90% CFS patient plasma
- XMRV is a Blood Borne, Infectious Human Retrovirus

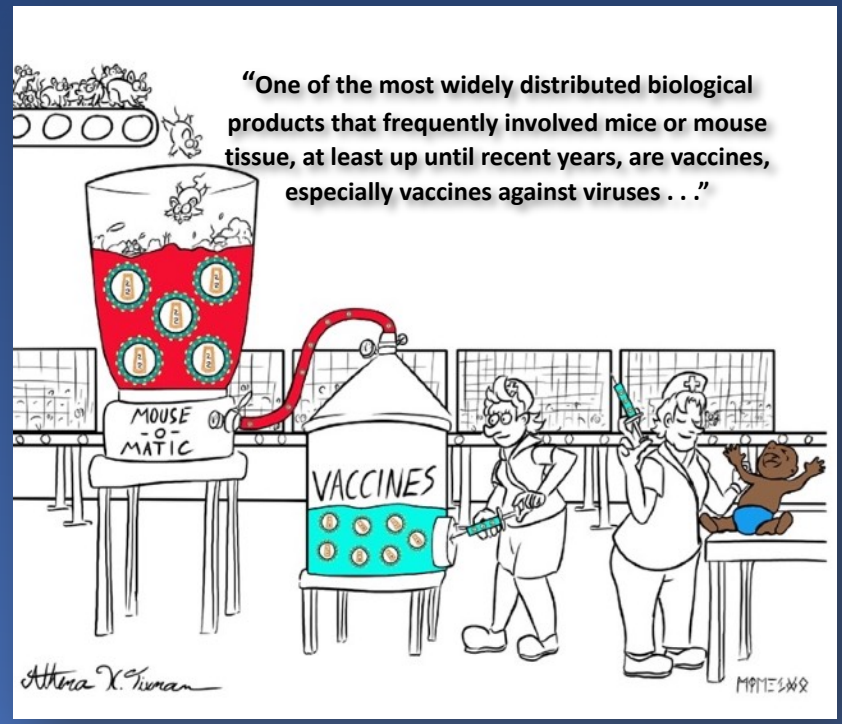
Evidence of XMRV infection in >98% of this cohort
(Mikovits et al Virulence 1:5 1-5 October 2010)

Animal

- Bovine serum (several forms)
- Avian serum - chicken
- Egg protein – ovalbumin
- VERO cell Line – monkey
- Dog kidney cell Line (MDCK)
- Insect cell line

Human cell Lines

- WI-38
- MRC-5
- PER.C6



HOW MANY NEW VIRUSES HAVE WE CREATED John Coffin ??

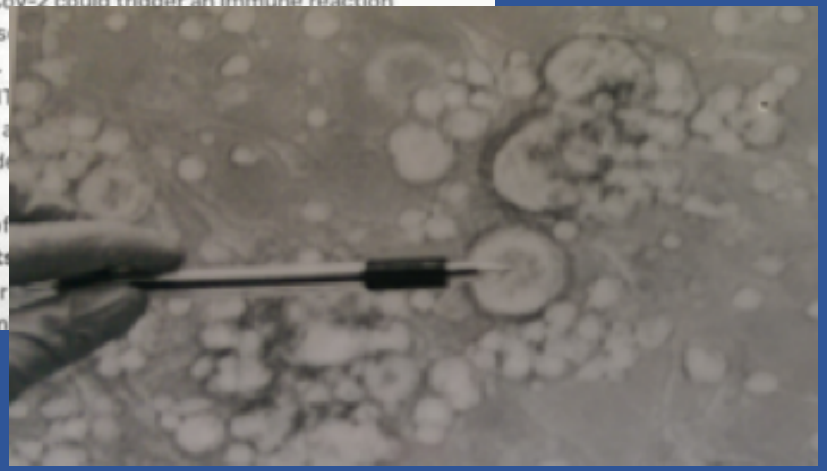
EXPRESSION OF SYNCYTIN IN COVID VACCINE KNOWN to result in female infertility & the Development of AIDS/CANCER

Head of Pfizer Research: Covid Vaccine is Female Sterilization

Health & Money News / December 2, 2020 / News

The vaccine contains a **spike protein (see image) called syncytin-1**, vital for the formation of human placenta in women. If the vaccine works so that we form an immune response AGAINST the spike protein, we are also **training the female body to attack syncytin-1**, which could lead to infertility in women of an unspecified duration.

- The vaccinations are expected to produce antibodies against spike proteins of SARS-CoV-2. However, spike proteins also contain **syncytin-homologous proteins**, which are essential for the formation of the placenta in mammals such as humans. It must be absolutely ruled out that a vaccine against SARS-CoV-2 could trigger an immune reaction against syncytin-1, as otherwis **result in vaccinated women**.
- The mRNA vaccines from BioN **(PEG)**. 70% of people develop means that many people can d to the vaccination.
- The much too short duration o **estimation of the late effect** Nevertheless, BioNTech/Pfizer emergency approval on Decem



Keywords fusion; interferon; SARS-CoV-2; syncytia
Subject Category Immunology
DOI 10.15252/embj.2020106267 | Received 17 July 2020 | Revised 6 October 2020 | Accepted 8 October 2020 | Published online 4 November 2020
The EMBO Journal (2020) 39: e106267

THE EMBO JOURNAL

Syncytia formation by SARS-CoV-2-infected cells

Expression of S without any other viral proteins triggers syncytia formation. Interferon-induced transmembrane proteins (IFITMs), a family of restriction factors that block the entry of many viruses, inhibit S-mediated fusion, with IFITM1 being more active than IFITM2 and IFITM3. On the contrary, the TMPRSS2 serine protease, which is known to enhance infectivity of cell-free virions, processes both S and ACE2 and increases syncytia formation by accelerating the fusion process. TMPRSS2 thwarts the antiviral effect of IFITMs. Our results show that SARS-CoV-2 pathological effects are modulated by cellular proteins that either inhibit or facilitate syncytia formation.

2006: Syncytia in XMRV Infected ME/CFS patient blood

Recombinant origin of SARS-CoV2 in 2004: It's a Retrovirus NOT Coronavirus

Each spike protein snaps together with two others, forming a structure that has a tulip-like shape. A long stem anchors the proteins to the virus, and their top looks like a three-part flower. NYTimes

JOURNAL OF VIROLOGY, Oct. 2004, p. 10628–10635 Vol. 78, No.
0022-538X/04/$08.00+0 DOI: 10.1128/JVI.78.19.10628–10635.2004

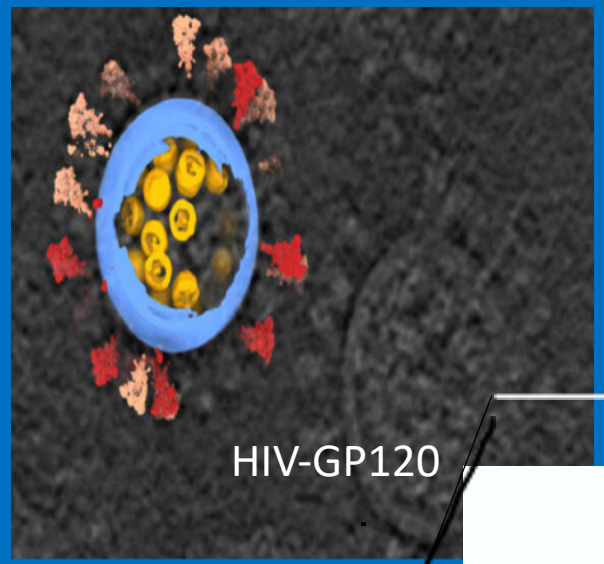
Retroviruses Pseudotyped with the Severe Acute Respiratory Syndrome Coronavirus Spike Protein Efficiently Infect Cells Expressing Angiotensin-Converting Enzyme 2

Michael J. Moore,[1] Tatyana Dorfman,[1] Wenhui Li,[1] Swee Kee Wong,[1] Yanhan Li,[2] Jens H. Kuhn,[1,3] James Coderre,[4] Natalya Vasilieva,[5] Zhongchao Han,[2] Thomas C. Greenough,[4] Michael Farzan,[1]* and Hyeryun Choe[5]*

Partners AIDS Research Center, Brigham and Women's Hospital, and Department of Medicine (Microbiology and Molecular Genetics),[1] and Perlmutter Laboratory, Children's Hospital, and Department of Pediatrics,[5] Harvard Medical School, Boston, and Program in Molecular Medicine, University of Massachusetts Medical School, Worcester,[4] Massachusetts; State Key Laboratory of Experimental Hematology, Institute of Hematology and Hospital of Blood Diseases, Chinese Academy of Medical Sciences and Peking Union Medical College, Tianjin, China[2]; and Department of Biology, Chemistry, Pharmacy, Freie Universität Berlin, Berlin, Germany[3]

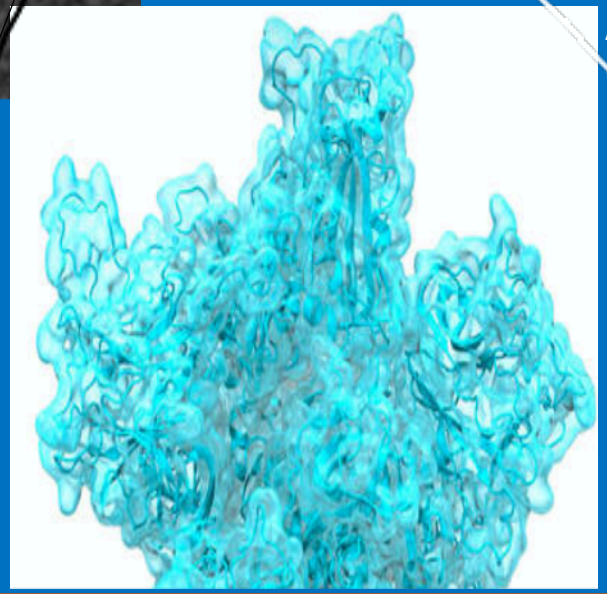
Received 3 February 2004/Accepted 28 May 2004

responses to potential vaccines. Here we show that simian immunodeficiency virus (SIV) pseudotyped with several codon-optimized S-protein variants could efficiently infect Vero E6 cells and HEK293T cells transiently or stably expressing ACE2. One such variant, truncated at its cytoplasmic tail and bearing instead a region of the tail of the human immunodeficiency virus type 1 (HIV-1) envelope glycoprotein (17), was especially efficient at mediating infection. Murine leukemia virus (MLV) pseudotyped with this S-protein variant also infected ACE2-expressing cells more efficiently than MLV pseudotyped with other S-protein variants. We used this sys-



XMRV-Syncytin

SARS-ACE2RBD



COVID19 MRNA vaccines Synthetic Retroviruses: HIV/XMRV/SARS

JVI Journals.ASM.org

4630 jvi.asm.org
Journal of Virology

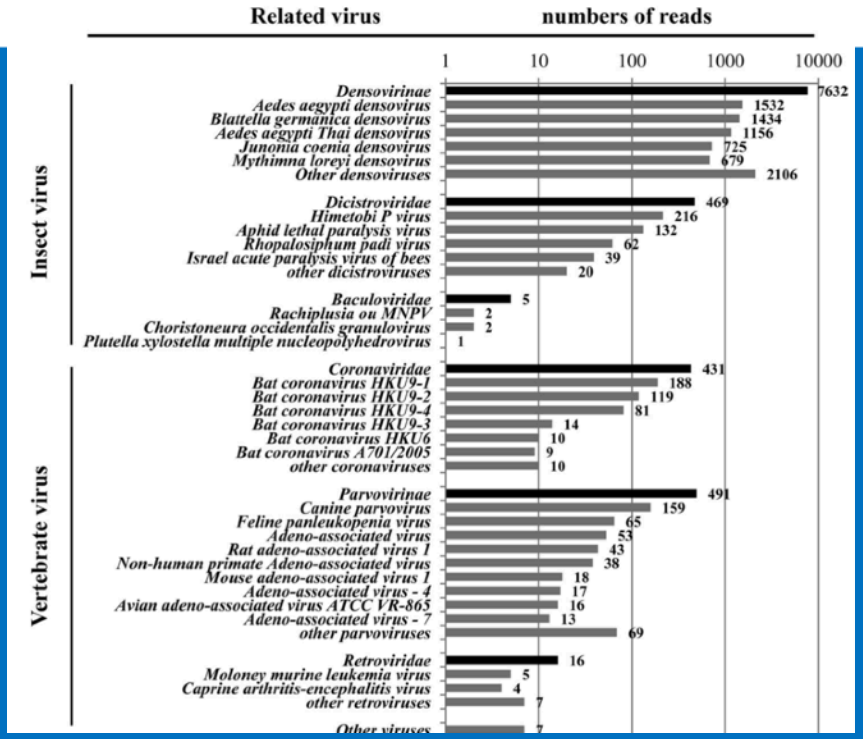
Received 31 October 2011 Accepted 31 January 2012
Published ahead of print 15 February 2012
Address correspondence to Zhengli Shi, zlshi@wh.iov.cn.
Supplemental material for this article may be found at http://jvi.asm.org/.

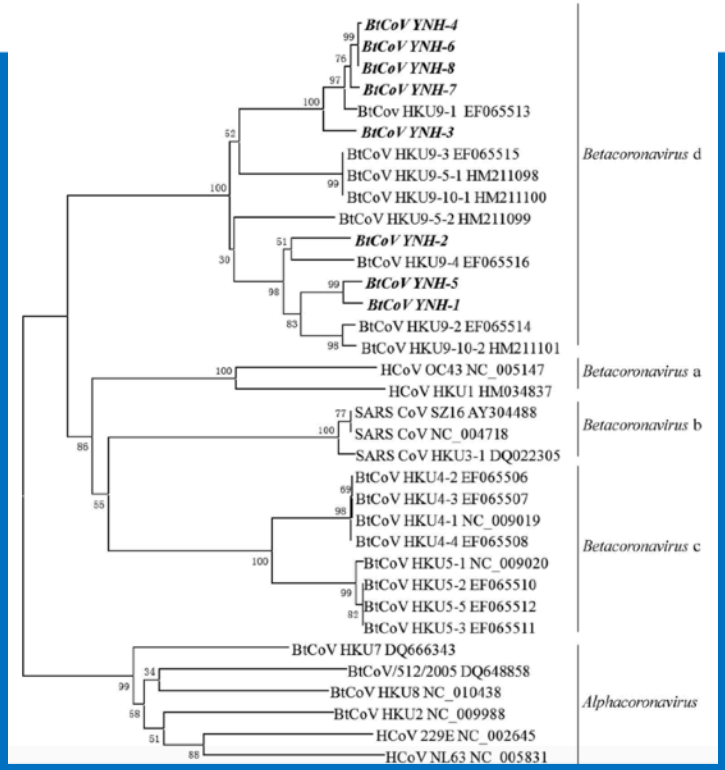
doi:10.1128/JVI.06671-11

Metagenomic Analysis of Viruses from Bat Fecal Samples Reveals Many Novel Viruses in Insectivorous Bats in China

Xingyi Ge,[a] Yan Li,[a] Xinglou Yang,[a] Huajun Zhang,[a] Peng Zhou,[a] Yunzhi Zhang,[b] and Zhengli Shi[a]

State Key Laboratory of Virology, Wuhan Institute of Virology, Chinese Academy of Sciences, Wuhan, China,[a] and Yunnan Institute of Endemic Diseases Control and Prevention, Dali, China[b]





SUPPLEMENTAL.PDF

Metagenomic Analysis of Viruses from the Bat Fecal Samples Reveals Many Novel Viruses in Insectivorous Bats in China

Xingyi Ge[1], Yan Li[1], Xinglou Yang[1], Huajun Zhang[1], Peng Zhou[1], Yunzhi Zhang[2], Zhengli Shi[1*]

Retro-transcribing viruses		
HERV-H/env60	1	*Retroviridae;* unclassified Retroviridae; *Human endogenous retrovirus*
Amphotropic murine leukemia virus	1	
Moloney murine sarcoma virus	1	*Retroviridae; Orthoretrovirinae; Gammaretrovirus*
Xenotropic MuLV-related virus VP62	1	
Moloney murine leukemia virus	5	
Friend murine leukemia virus	1	

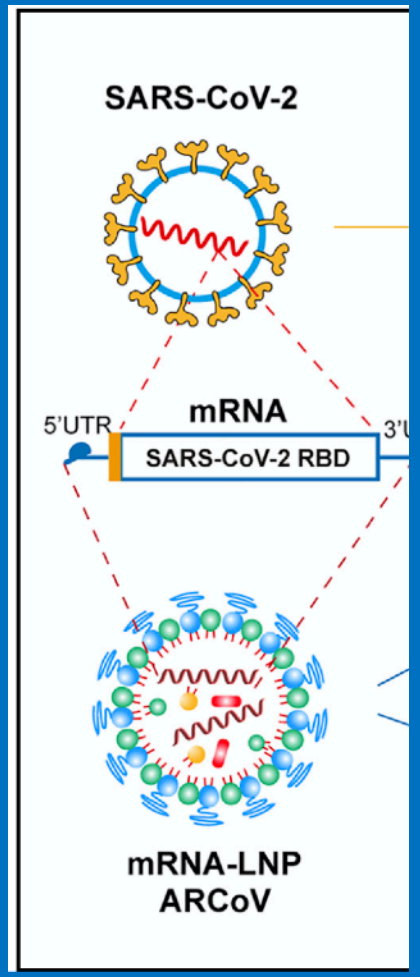
SARS-CoV2 designed to kill the 6% of the victims of 4 Decades of Medical Racism

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

An mRNA Vaccine against SARS-CoV-2 — Preliminary Report

L.A. Jackson, E.J. Anderson, N.G. Rouphael, P.C. Roberts, M. Makhene, R.N. Coler, M.P. McCullough, J.D. Chappell, M.R. Denison, L.J. Stevens, A.J. Pruijssers, A. McDermott, B. Flach, N.A. Doria-Rose, K.S. Corbett, K.M. Morabito, S. O'Dell, S.D. Schmidt, P.A. Swanson II, M. Padilla, J.R. Mascola, K.M. Neuzil, H. Bennett, W. Sun, E. Peters, M. Makowski, J. Albert, K. Cross, W. Buchanan, R. Pikaart-Tautges, J.E. Ledgerwood, B.S. Graham, and J.H. Beigel, for the mRNA-1273 Study Group*



NOTHING in CDC Schedule is a "VACCINE" ALL ARE

1)MANUFACTURED Viruses

- **Bioweapons that activate your own cells to become pathogen manufacturing devices**

2) Synthetic: MRNA viruses (poisons)

- **Snake Venom**

SNAKE NEEDS FANGS

Participants were not screened for SARS- CoV-2 infection by serology/ Antibody or polymerase chain reaction before enrollment.

Antibodies to XMRVs ENV/Spike Reproducibly Detected in 4-6% Population In every single study

Table 1. All XMRV/P-MLV assay results from all laboratories. Abbott-M, Abbott Molecular; Abbott-D, Abbott Diagnostics; WB, whole blood; N/A, not applicable. Boldface entries indicate positive results.

Culture	FDA/Hewlett	0/15	0/10	0/5	**5/5**
	NCI/Ruscetti	**6/15**	**3/10†**	0/5	**5/5**
Serology	Abbott-D	0/15	0/10	0/5	N/A
	CDC	0/15	0/10	0/5	N/A
	NCI/Ruscetti	**8/15**	**3/10**	**2/5†**	N/A
	WPI	**6/15**	**5/10**	**5/5†**	N/A

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

- ADE-Antibody Dependent Enhancement (B-Cell) Adaptive
- Pathogenic Priming (Macrophage) Innate
- ADCC-Antibody Dependent Cellular Cytotoxicity (NK Cells) Innate

TABLE 3 Equivalent levels of XMRV sequences and anti-XMRV antibodies in CFS (chronic fatigue syndrome) patients and matched controls

			CFS/ME cases (n = 147)		Controls (n = 146)	
Lab site	Analysis	Sample	Total studied	No. positive (%)	Total studied	No. positive (%)
CDC	RT-PCR	Plasma	147	0 (0.0)	146	0 (0.0)
FDA	RT-PCR	Plasma	121[a]	0 (0.0)	110[a]	0 (0.0)
	PCR	PBMC	121[a]	0 (0.0)	111[a]	0 (0.0)
Mikovits, Ruscetti, and Hanson	PCR of cultured PBMC	PBMC	117[b]	0 (0.0)	126[b]	0 (0.0)
Mikovits and Ruscetti	Serology	Plasma	147	9 (6.1)	146	9 (6.2)

[a] Numbers represent all samples available for analysis at that site.
[b] Fifty samples (30 cases; 20 controls) were unable to be assayed because at least one of two aliquots from each set of subject PBMC did not grow in tissue culture.

COVID19 is the COVER-UP of Crimes Against Humanity

2017 Flunami: 2004 cloning of 1918 pandemic Influenza A

JOURNAL OF VIROLOGY, Nov. 2004, p. 12462–12470 Vol. 78, No. 22
0022-538X/04/$08.00+0 DOI: 10.1128/JVI.78.22.12462–12470.2004

Novel Origin of the 1918 Pandemic Influenza Virus Nucleoprotein Gene

Ann H. Reid, Thomas G. Fanning, Thomas A. Janczewski, Raina M. Lourens, and Jeffery K. Taubenberger*

Division of Molecular Pathology, Department of Cellular Pathology and Genetics, Armed Forces Institute of Pathology, Rockville, Maryland

Received 30 March 2004/Accepted 7 July 2004

The nucleoprotein (NP) gene of the 1918 pandemic influenza A virus has been amplified and sequenced from archival material. The NP gene is known to be involved in many aspects of viral function and to interact with host proteins, thereby playing a role in host specificity. The 1918 NP amino acid sequence differs at only six amino acids from avian consensus sequences, consistent with reassortment from an avian source shortly before 1918. However, the nucleotide sequence of the 1918 NP gene has more than 170 differences from avian strain consensus sequences, suggesting substantial evolutionary distance from known avian strain sequences. Both the gene and protein sequences of the 1918 NP fall within the mammalian clade upon phylogenetic analysis. The evolutionary distance of the 1918 NP sequences from avian and mammalian strain sequences is examined, using several different parameters. The results suggest that the 1918 strain did not retain the previously circulating human NP. Nor is it likely to have obtained its NP by reassortment with an avian strain similar to those now characterized. The results are consistent with the existence of a currently unknown host for influenza, with an NP similar to current avian strain NPs at the amino acid level but with many synonymous nucleotide differences, suggesting evolutionary isolation from the currently characterized avian influenza virus gene pool.

Vaccine 38 (2020) 350–354

Contents lists available at ScienceDirect

Vaccine

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

Influenza vaccination and respiratory virus interference among Department of Defense personnel during the 2017–2018 influenza season

Greg G. Wolff

Armed Forces Health Surveillance Branch Air Force Satellite, 2510 5th Street, Bldg 840, Wright-Patterson AFB, OH 45433, United States

Respiratory Virus Infections Associated With Receipt of Inactivated Influenza Vaccine

RESEARCH ARTICLE | INFLUENZA

Vaccine-Induced Anti-HA2 Antibodies Promote Virus Fusion and Enhance Influenza Virus Respiratory Disease

Surender Khurana[1], Crystal L. Loving[2], Jody Manischewitz[1], Lisa R. King[1], Phillip C. Gauger[3], Jamie ...

+ See all authors and affiliations

Science Translational Medicine 28 Aug 2013:
Vol. 5, Issue 200, pp. 200ra114
DOI: 10.1126/scitranslmed.3006366

4000 FLU DEATHS/week JAN 2020!

VACCINE AIDS = COVID19

Autoimmune, Autoinflammatory Disease & Cancer *Unintended* Consequences of 3 DECADES LIABILITY FREE VACCINES

Prostate Cancer*	Crohn's Disease*	Gulf War Syndrome*
Breast Cancer *	Hashimoto's Thyroiditis*	Autism / ASD*
Multiple Myeloma*	Polymyositis*	Multiple Sclerosis*
Non-Hodgkins Lymphoma*	Sjogren's Syndrome *	Parkinson's*
Chronic Lymphocytic Leukemia*	Bechet's Disease*	ALS*
Mantle Cell Lymphoma*	Primary Biliary Cirrhosis*	Fibromyalgia*
Hairy Cell Leukemia*	Inflammatory Bowel Disease*	Chronic Lyme Disease*
Bladder Cancer *	Psoriasis, Dermatitis	OCD*
Colorectal Cancer*	Diabetes*	ADHD*
Kidney Cancer *	Cardiovascular Disease*	PTSD*
Ovarian Cancer*	ME / CFS*	Psychosis*
**Neuroendocrine Tumors*	Lupus/SLE*	Rheumatoid Arthritis*

KEY to IMMUNITY is do not defile the TEMPLE of GOD

NEVER GET ANOTHER VACCINE

Call To Action: #stoptheshots

- MORATORIUM on ALL VACCINES on CDC SCHEDULE
- Repeal 1986 National Vaccine Injury Compensation Act
- End all Mandates and Restore Liability to all
- Convict criminals at CDC, FDA, NIH for crimes against humanity
- Eliminate Advisory Committee on Immunization Practices (ACIP)
- Use NIH and CDC & FDA Patent Royalties to Compensate all Victims of this 35 Year Plague Of Corruption

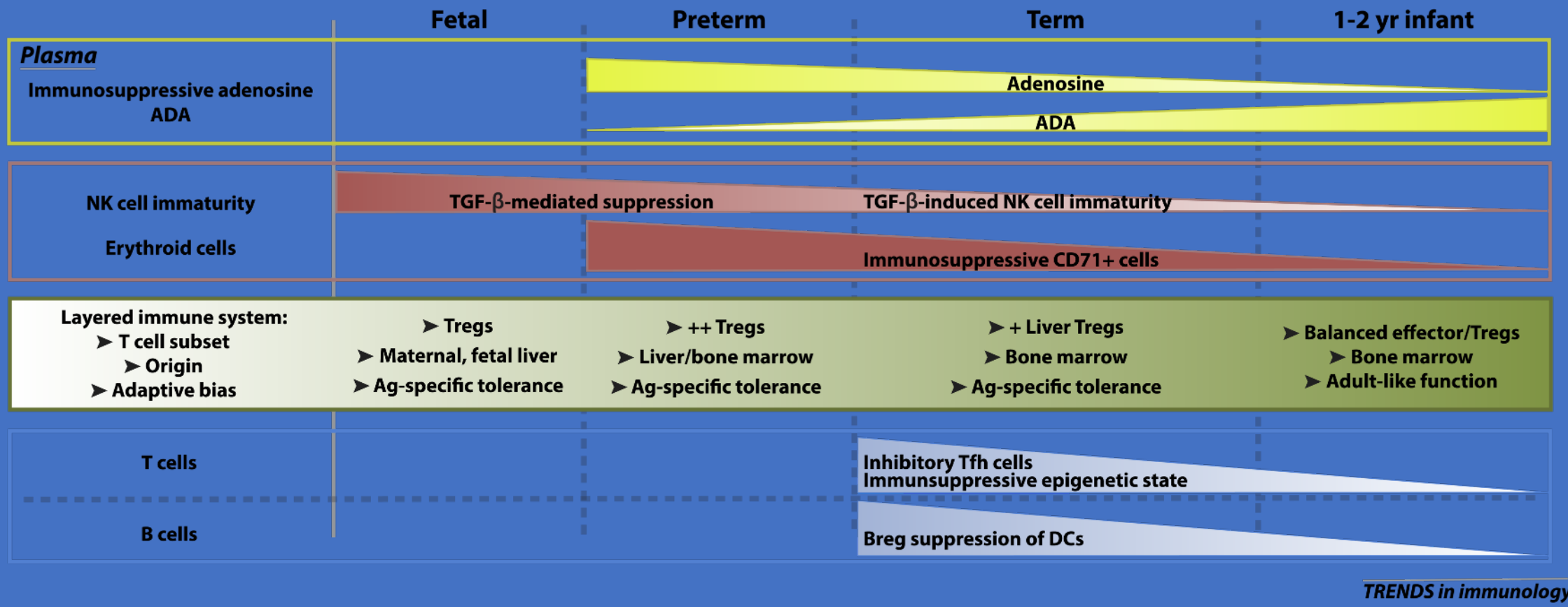
Greg@WriteYourLaws.com

https://writeyourlaws.com/2020/08/the-fda-emergency-use-authorization-emergency-overhaul-act-of-2020/

https://writeyourlaws.com/amendments-to-section-230-eliminating-censorship-from-big-tech-version-2

https://writeyourlaws.com/the-vaccine-manufacturer-full-product-liability-restoration-act-of-2021

Immunity is not static: it changes with age; many unique features in early life



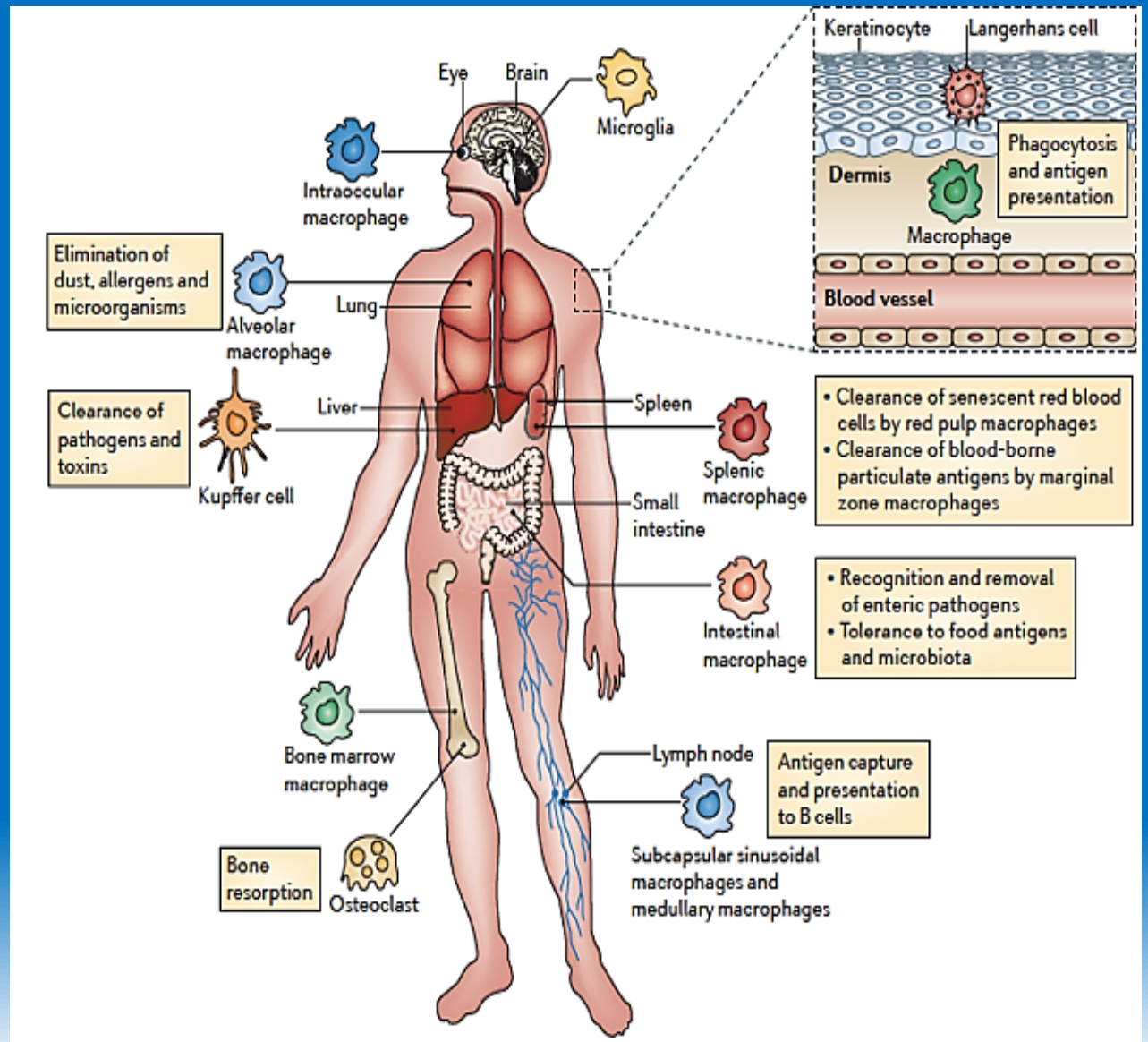
The Brain and The Immune System are inextricably linked from Conception

Monocyte/Macrophage Dysfunction as a Driver of Vaccine AIDS/CANCER

- **Express Purinergic Receptors**:
- **P2XR and P2YR**.
- Express Cannabinoid Receptors
- CB1 & CB2

Tissue Macrophages perform Key Homeostatic Functions Modulated by

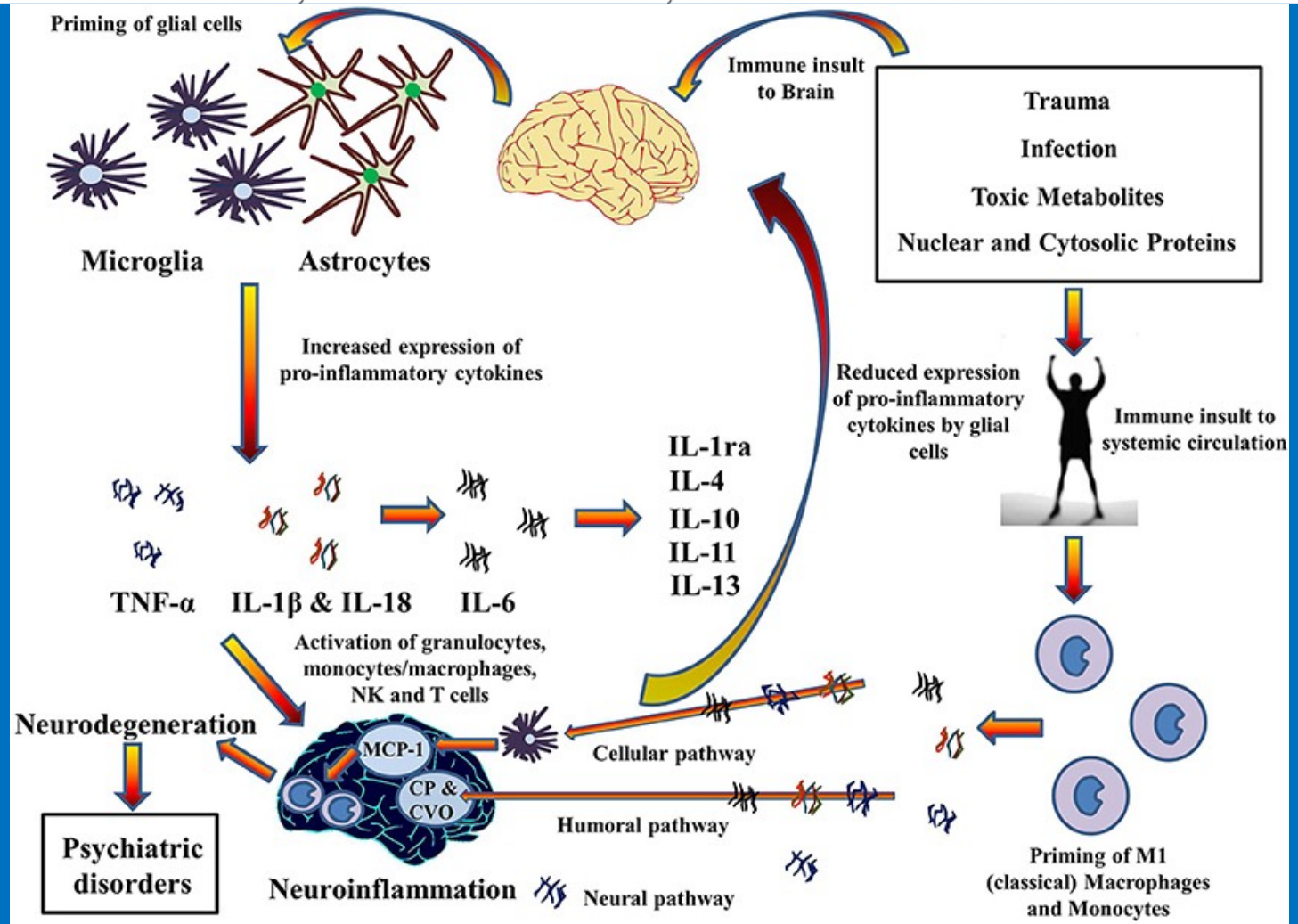
- **Cannabinoids**
- **GcMAF**
- **Suramin**
- **Ivermectin**
- **Vitamin C**
- **DMG**
- **Decitibine (Vidaza)**
- **Peptide T**



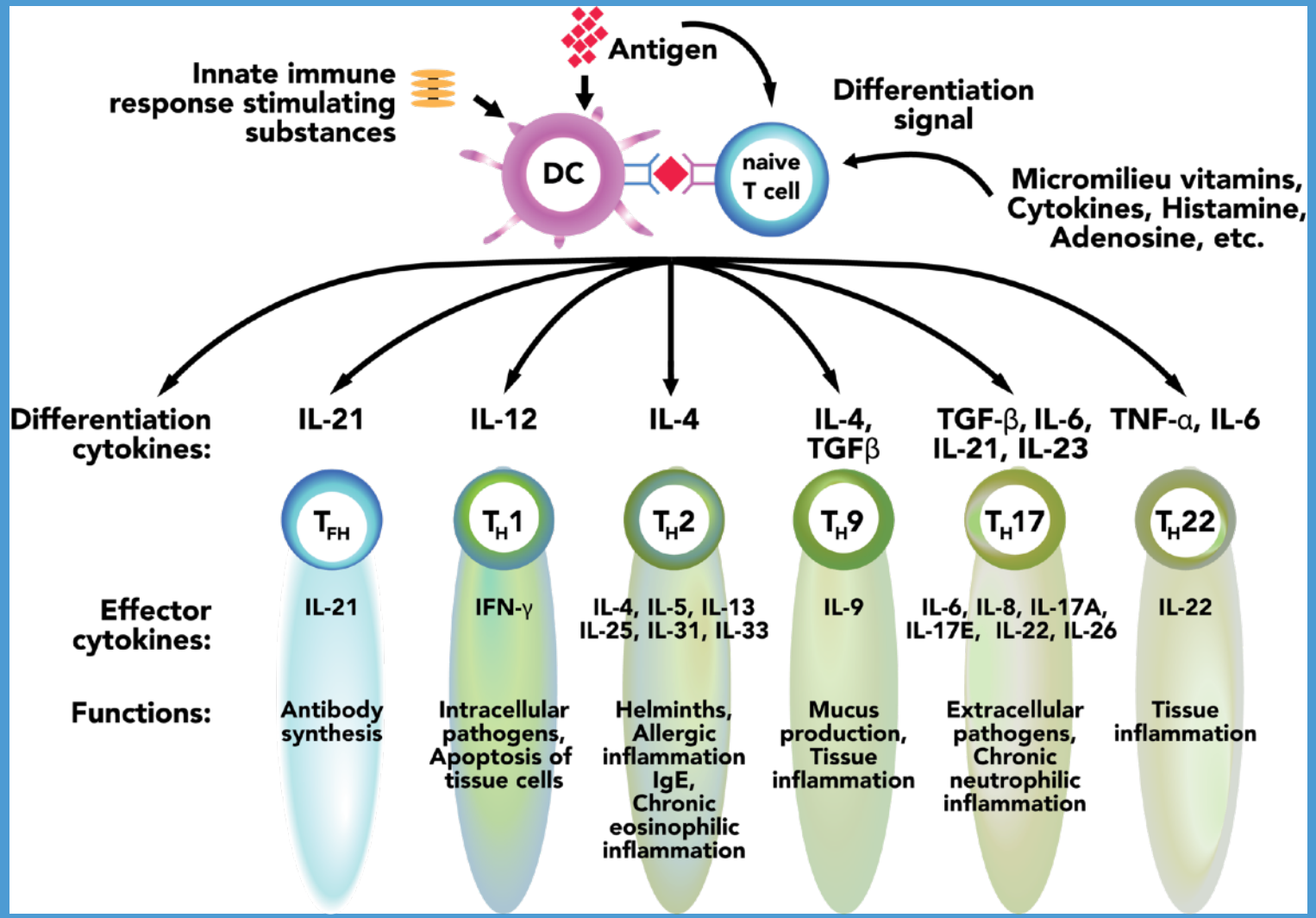
Xenotropic Murine Leukemia Virus-related Virus-associated Chronic Fatigue Syndrome Reveals a Distinct Inflammatory Signature

in vivo *25*: 307-314 (2011)

VINCENT C. LOMBARDI[1], KATHRYN S. HAGEN[1], KENNETH W. HUNTER[4], JOHN W. DIAMOND[2†], JULIE SMITH-GAGEN[3], WEI YANG[3] and JUDY A. MIKOVITS[1]

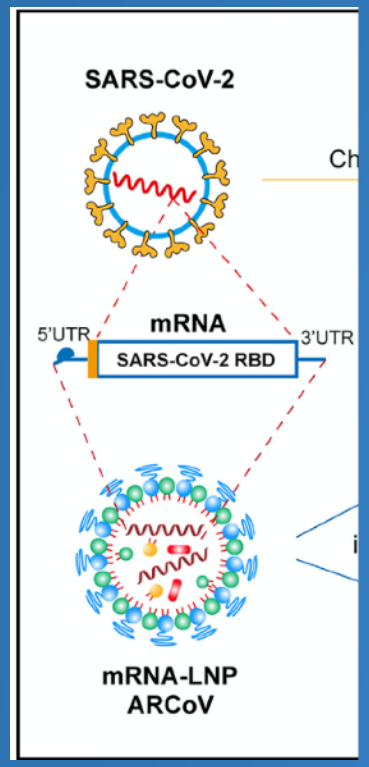


Inappropriate Activation of the cellular Immune system is important in the pathogenesis of human Retrovirus Associated Disease



Every Experimental injection Bypasses The Innate Immune System

Stop Shedding /Transmission: XMRVs (all prior Vaccines)& Synthetic Lipid Nano Particles (COVID 19 shot)

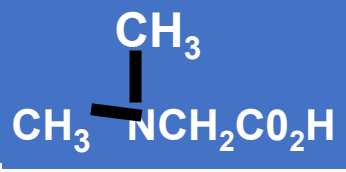


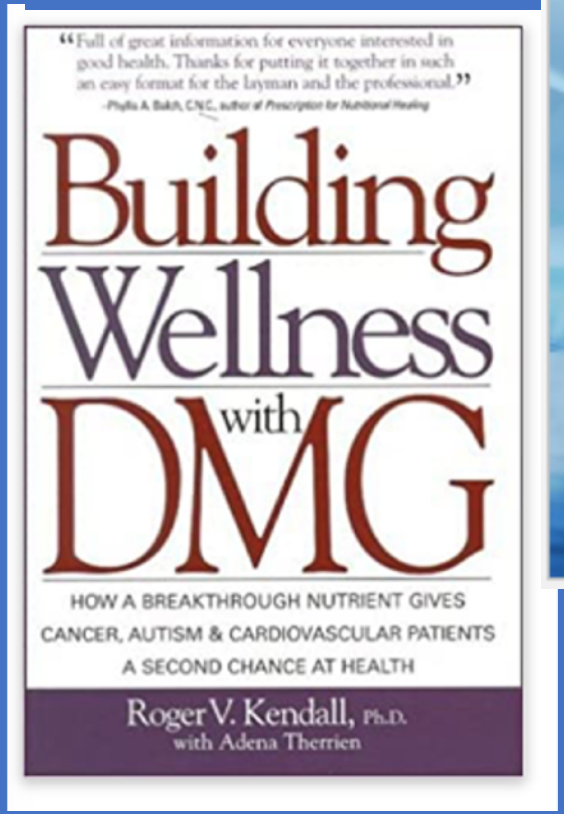
Detoxing that synthetic Lipid Nano Particle
(SARS-CoV2 virus & COVID Vaccine)

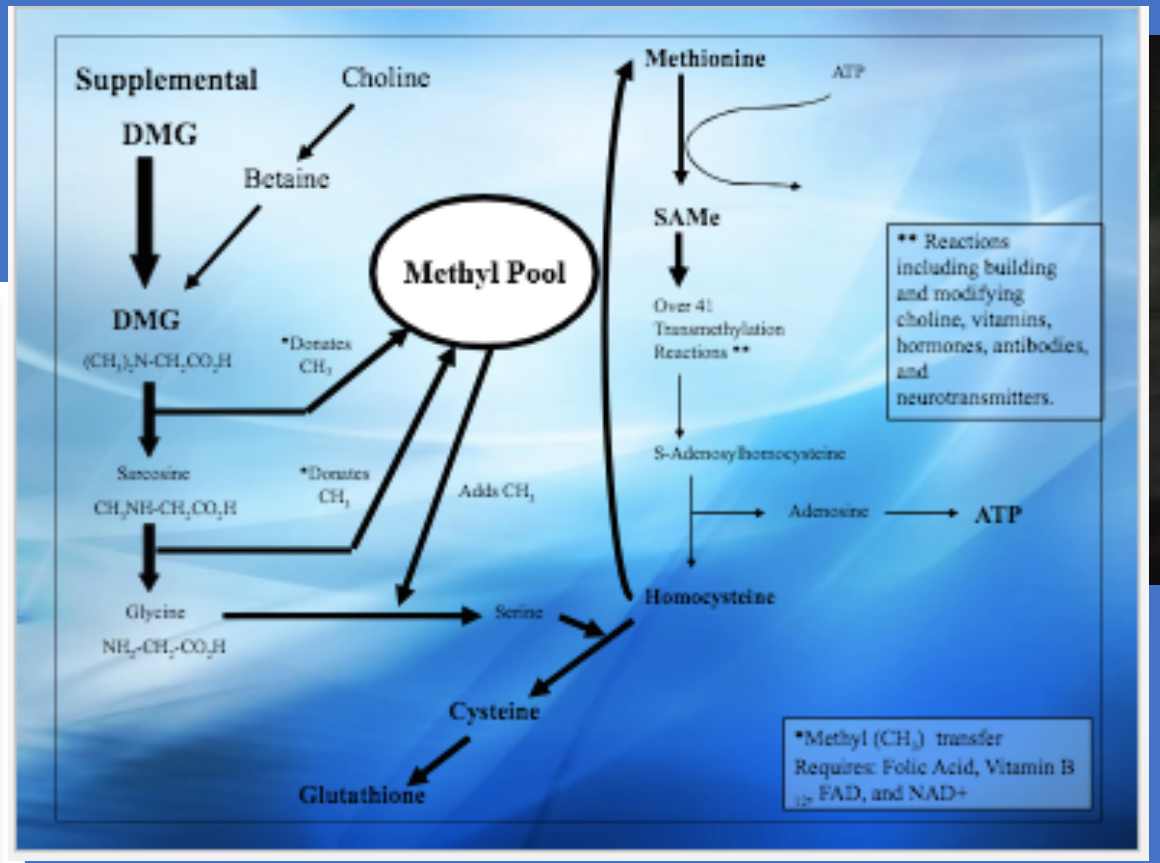
- Ozone therapies
- Specialized Pro resolving mediators
- Chlorine Dioxide, MMS, CDS

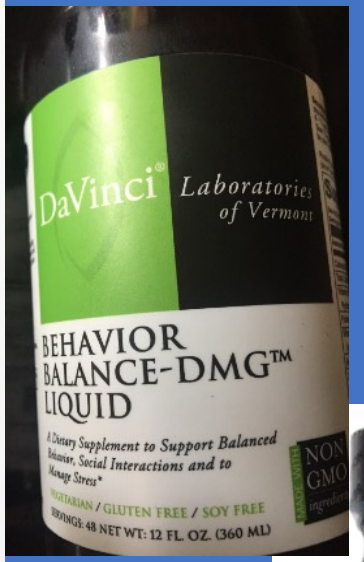
DiMethylGlycine: Silence Reactivated viruses

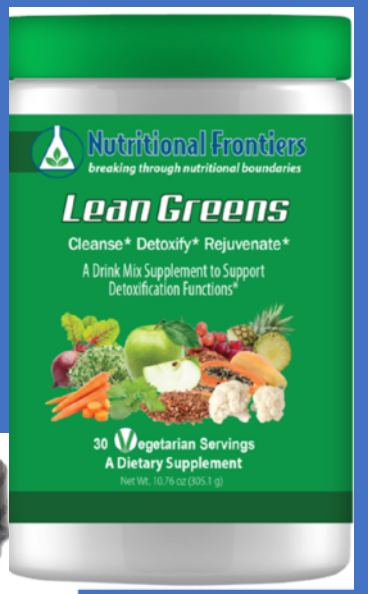
Nutrition's Best Kept secret for strengthening Genomic Pathways and Preventing Disease







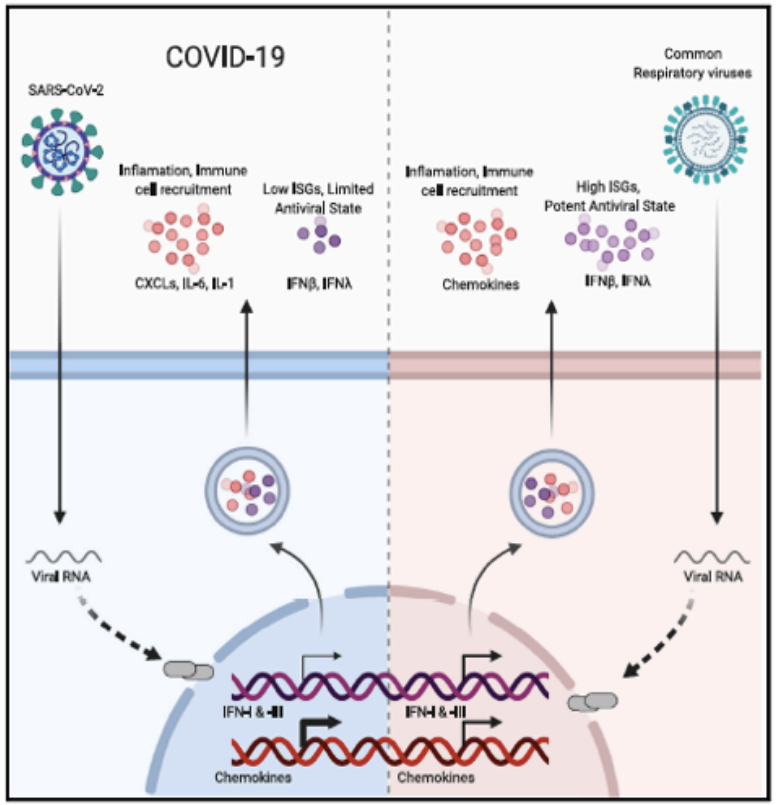




- Important nutrient found in low levels in our food
- As a Key Nutrient DMG PROTECTS OUR GENES

Imbalanced IFN Response to RNA Viruses Drives Development of Autoimmune, Autoinflammatory Disease & Cancer

Graphical Abstract



Authors

Daniel Blanco-Melo, Benjamin E. Nilsson-Payant, Wen-Chun Liu, ..., Jean K. Lim, Randy A. Albrecht, Benjamin R. tenOever

Correspondence

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In Brief

In comparison to other respiratory viruses, SARS-CoV-2 infection drives a lower antiviral transcriptional response that is marked by low IFN-I and IFN-III levels and elevated chemokine expression, which could explain the pro-inflammatory disease state associated with COVID-19.

Poisons (ADJUVANTS): Aluminum, LPS (ENDOTOXIN), Xenoestrogens, Arsenic in Vaccines food & water
Cripple Innate Immune responses= VACCINE AIDS

Lupus is an autoimmune inflammatory disease in which the body produces auto antibodies antibodies causing the immune system to affect the skin, joints, blood and kidneys.

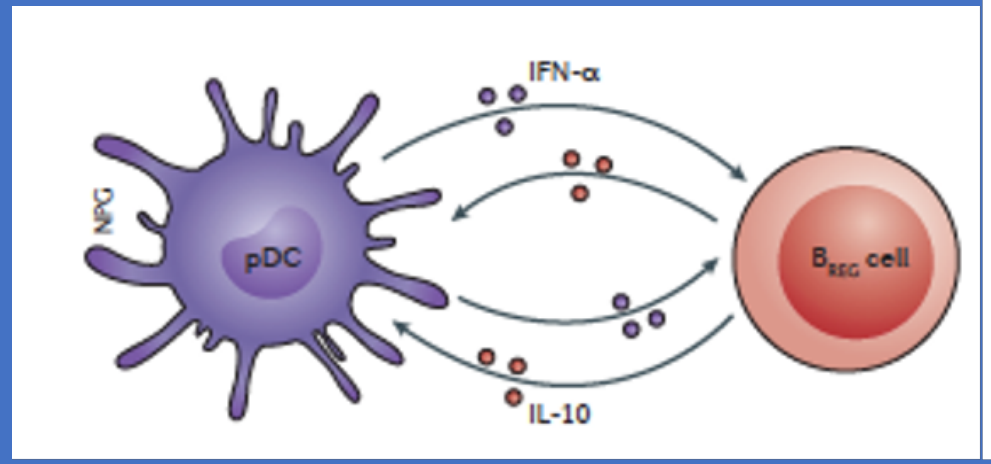
Symptoms include:

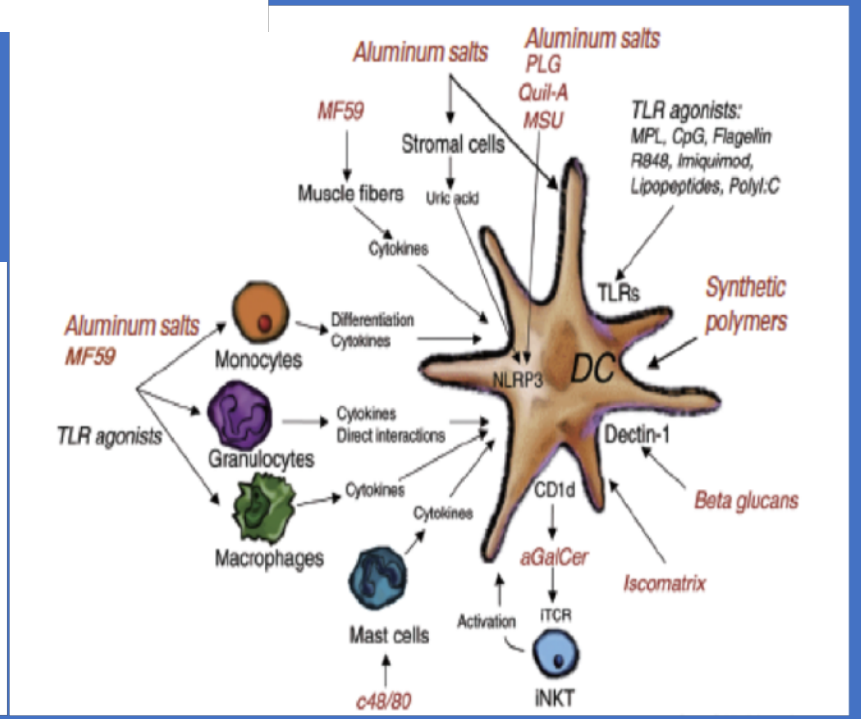
- Skin rashes/ Inflammation
- Arthritis/ Joint Pain
- Extreme Fatigue
- Anemia/ Blood Disorders
- Kidney Damage
- Immune Disorder
- Antinuclear Antibodies
- CVID: Antibody Deficiency

Nature Reviews Rheumatology | Published online 24 Mar 2016; doi:10.1038/nrrheum.2016.43

SYSTEMIC LUPUS ERYTHEMATOSUS

Compromised pDC–B_{REG} cell crosstalk

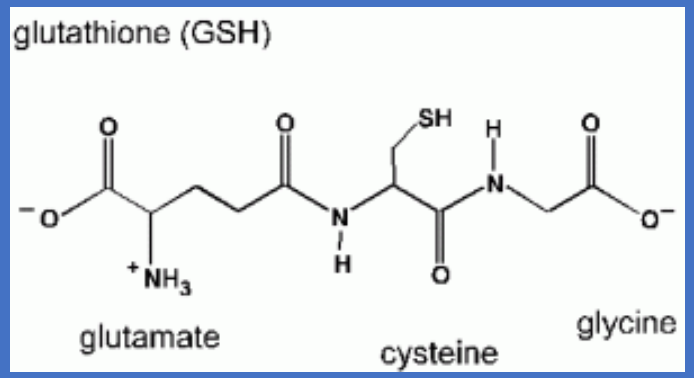


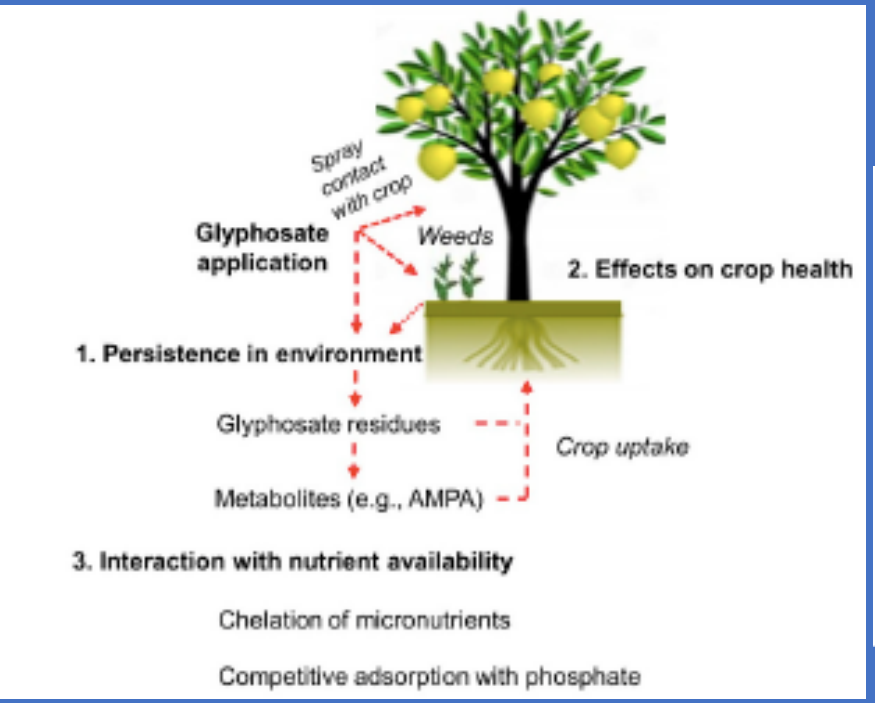


Unintended Consequences of Inappropriate Innate Immune Activation?

Glyphosate: Damages Key GOD GIVEN antioxidant Glutathione

Produced by the liver, glutathione is made up of three amino acids: Lcysteine, glycine, and L-glutamate





American Chemical Society
Public Health Emergency Collection
Public Health Emergency COVID-19 Initiative

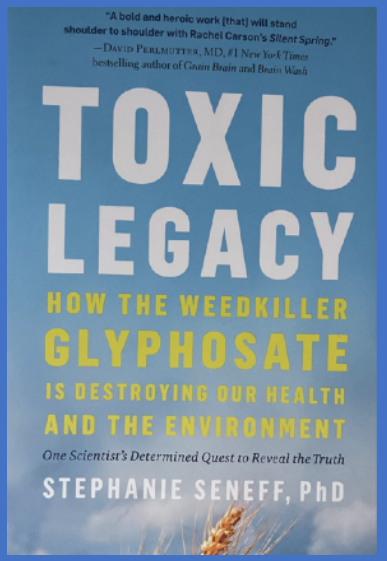
ACS Infect Dis. 2020 May 28 : acsinfecdis.0c00288.
Published online 2020 May 28. doi: 10.1021/acsinfecdis.0c00288
PMCID: PMC7263077
PMID: 32463221

Endogenous Deficiency of Glutathione as the Most Likely Cause of Serious Manifestations and Death in COVID-19 Patients

Alexey Polonikov*

Author information · Article notes · Copyright and License information Disclaimer

Endogenous glutathione deficiency appears to be a crucial factor enhancing SARS-CoV-2-induced oxidative damage of the lung and, as a result, leads to serious manifestations, such as acute respiratory distress syndrome, multiorgan failure, and death in COVID-19 patients. When the antiviral activity of GSH is taken into account, individuals with glutathione deficiency seem to have a higher susceptibility for uncontrolled replication of SARS-CoV-2 virus and thereby suffer from an increasing viral load. The severity of clinical manifestations in COVID-19 patients is apparently determined by the degree of impaired redox homeostasis attributable to the deficiency of reduced glutathione and increased ROS production. This assumption can be supported by our findings. In particular, COVID-19 patients with moderate and severe illness had lower levels of glutathione, higher ROS levels, and greater redox status (ROS/GSH ratio) than COVID-19 patients with a mild illness. Long-term and severe manifestations of COVID-19 infection in one of our patients with marked glutathione deficiency suggest that the degree of glutathione decrease correlates negatively with viral replication rate and that an increasing viral load exacerbates oxidative damage of the lung. This finding suggests that the virus cannot actively replicate at higher levels of cellular glutathione, and therefore, milder clinical symptoms are observed with lower viral loads.

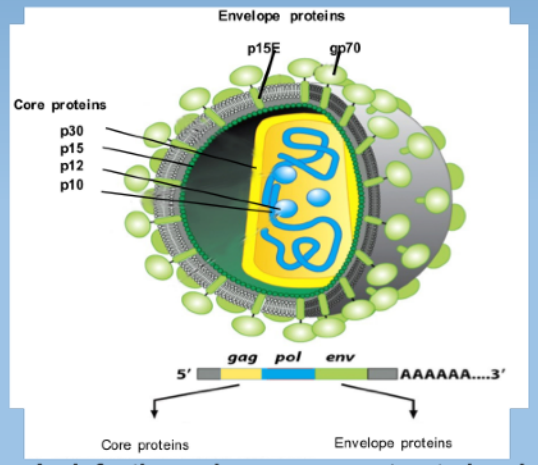


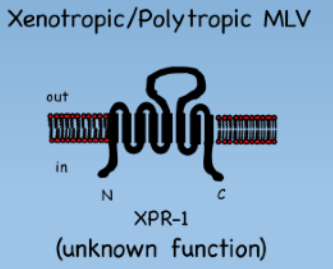
Glyphosate in our soil -> our Food is Contaminated !!

XMRV is pivotal because WE DETECTED VIRAL PROTEINS & ANTIBODY

- Evidence of infection in families with diagnoses: ASD, CFS, Chronic Lyme disease, prostate cancer and EVERY study found antibodies 4-6% in US “healthy controls”.. that is 20 million Americans at risk of Developing Vaccine AIDS = LONG HAUL COVID!!
  - ANTIBODY Test identified XMRV ENV/Spike Syncytin pathology
    - Including infection of brain microglia
    - Infection and dysregulation of gut tight junctions
    - Vasculitis
    - Inflammatory dysfunction: cytokine/chemokine
    - autoimmunity

Xenotropic Murine Leukemia Virus-Related Virus (XMRV)





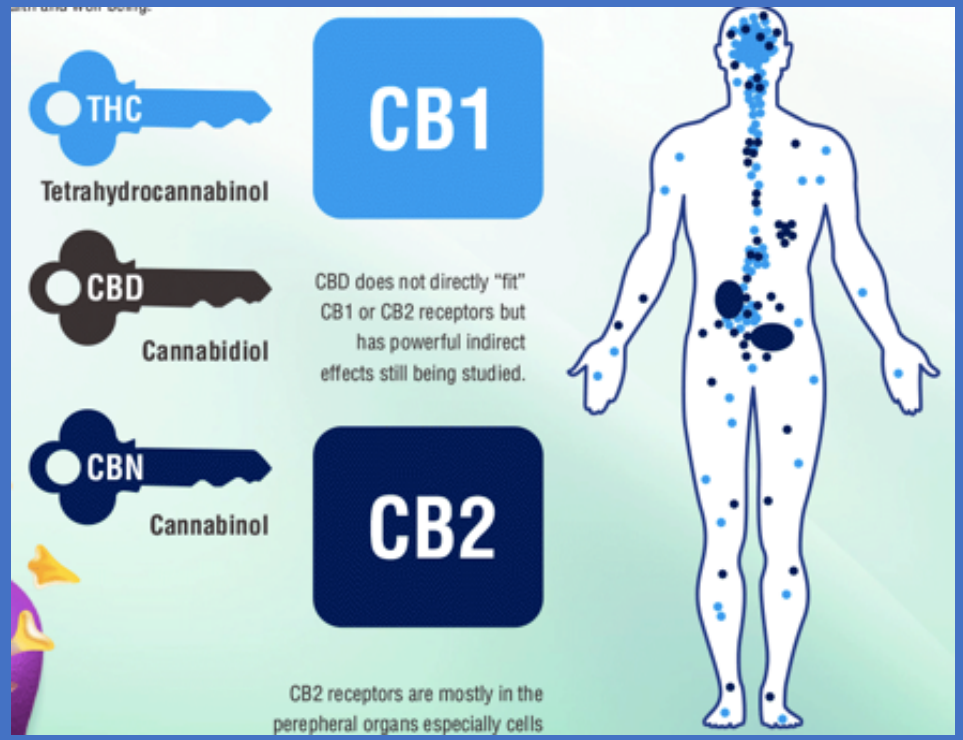
Like mouse xenotropic MuLV, XMRV uses the Xpr-1 receptor to enter cells (Dong et al., PNAS, 2007)

An infectious clone was constructed and sequenced and found to be a novel gammaretrovirus (Dong et al., PNAS, 2007)

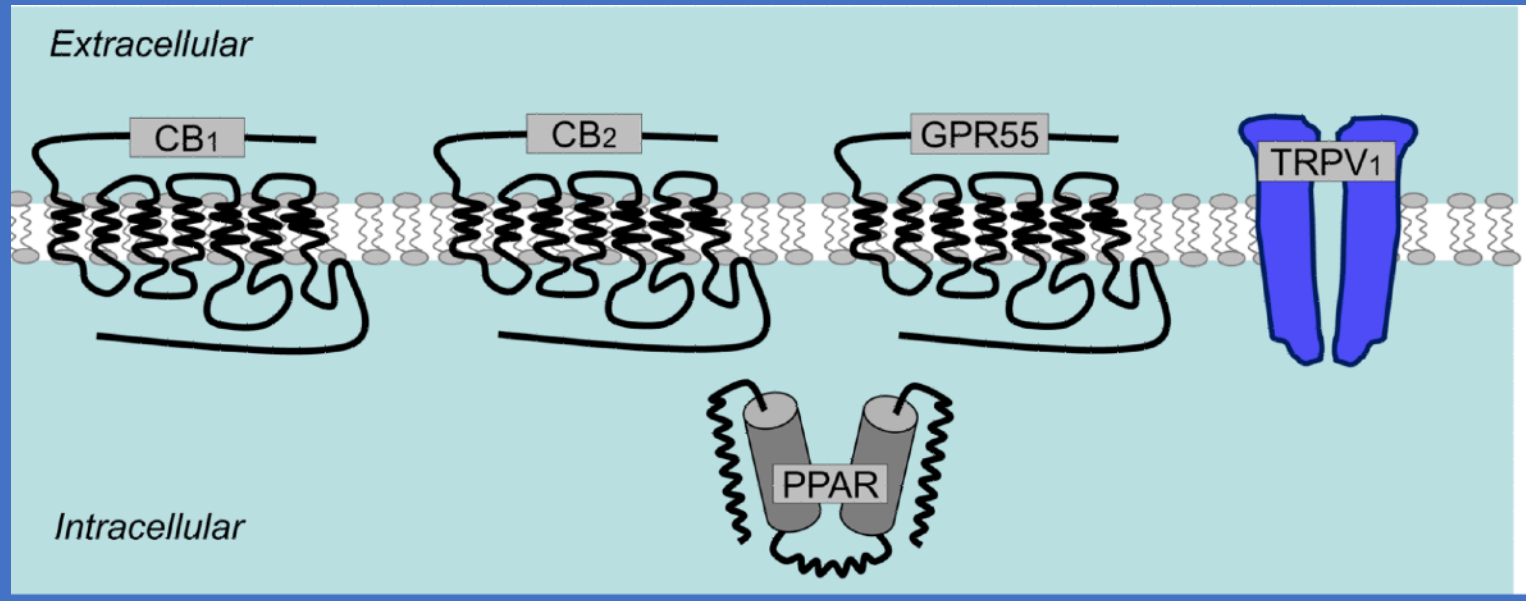
XMRV proviral integration occurs preferentially in CpG islands: gene promoters (Kim et al., JVirol, 2008)

The Human Endocannabinoid System (eCS)
GOD GIVEN DIMMER SWITCH ON INFLAMMATION

A signaling system that helps to modulate all other physiological, behavioral, and energetic processes in the body.



Glia. 2010 July ; 58(9): 1017–1030

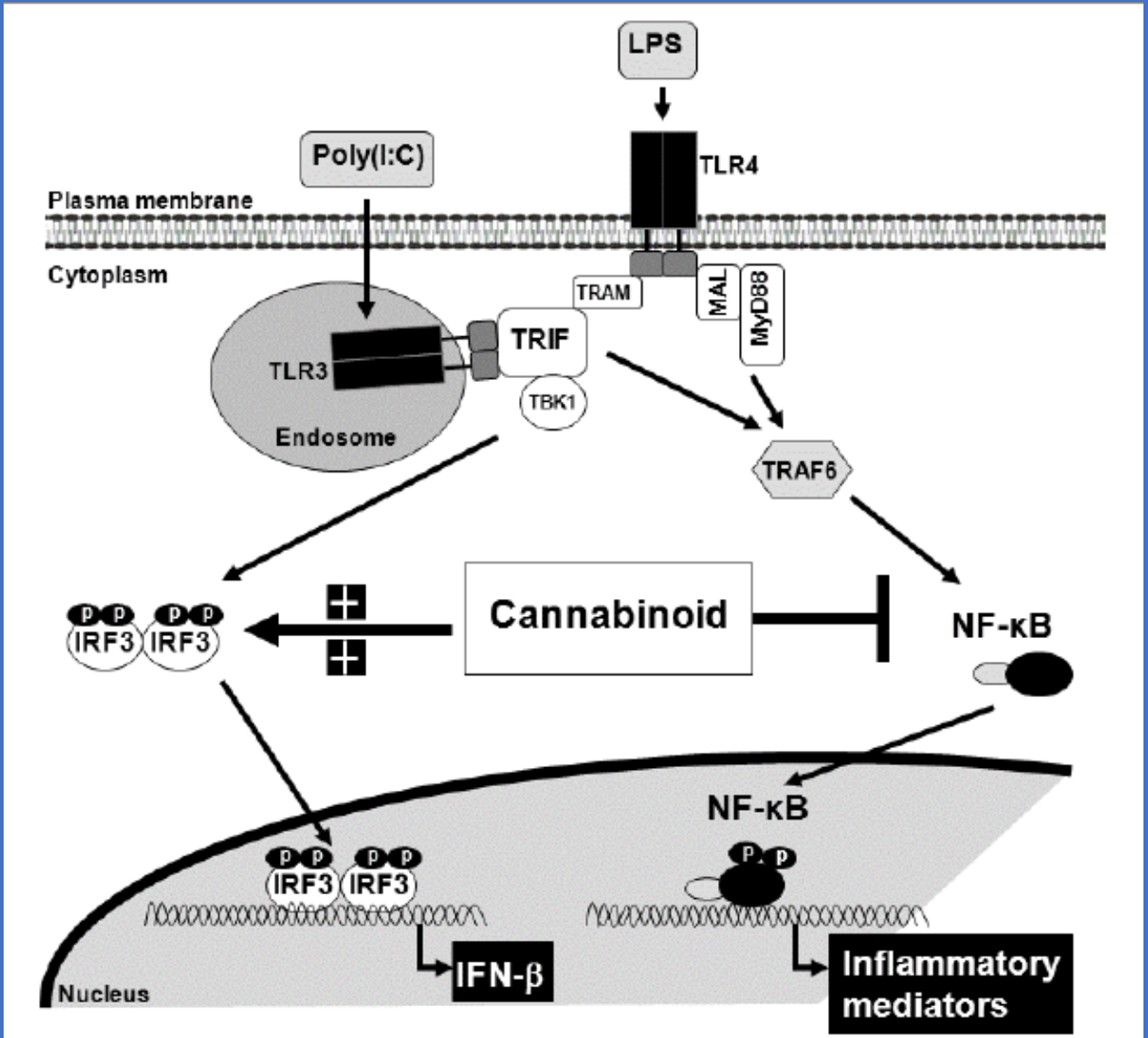


Anxiety
Depression
Sleep Disorders
Pain
Itch
Wound healing

- ***neuroprotection & plasticity***
- ***immunity & inflammation***
- ***apoptosis &carcinogenesis***
- ***pain and emotional memory***
- ***Supports detoxification:***
  - ***repairs Fibrosis***
  - ***fatty Liver disease***

Cannabinoids are Anti-Viral and Reduce inflammation

THE DIMMER SWITCH ON THE FLAME



CANNABIS is NOT a DRUG! IT'S Food!! Nourish CELLS
ALL Plants (HEMP & CANNABIS) Removed from US 1938!

Drug Metabolism Reviews
http://informahealthcare.com/dmr
ISSN: 0360-2532 (print), 1097-9883 (electronic)
Drug Metab Rev, 2014; 46(1): 86–95
© 2014 Informa Healthcare USA, Inc. DOI: 10.3109/03602532.2013.849268
informa healthcare

REVIEW ARTICLE

Exogenous cannabinoids as substrates, inhibitors, and inducers of human drug metabolizing enzymes: a systematic review

GOD GIVEN LIPID/FAT SIGNALING SYSTEM in EVERY Cell MEMBRANE

Viruses cause Disease by Dysregulating Key Immune molecules modulated by the eCS
The Dimmer switch of Inflammation: Putting out the Fire

Comparison of cellular gene expression in Ebola-Zaire and Ebola-Reston virus-infected primary human monocytes

C. Xiang[1], H. Young[2], H. Alterson[1], D. Reynolds[2], M. Bittner[2], Y. Chen[2], G. Gooden[1], Y. Jiang[1], P. Meltzer[2], J. Trent[1], J. Mikovits[2] & K. Anderson[1]

[1]Virology Division, US Army Medical Research Institute of Infectious Diseases, [2]Division of Basic Sciences, NCI-FCRDC, Frederick, Maryland 21702, USA [3]Laboratory of Cancer Genetics, NHGRI-NIH, Bethesda, Maryland 20892, USA

Ebola viruses are filamentous, enveloped, nonsegmented RNA viruses. Although most Ebola viruses, notably Ebola-Zaire virus, are highly infectious for primates and can cause severe haemorrhagic diseases, Ebola-Reston virus does not cause serious disease in humans. Microarray technology was employed to compare cellular gene responses to Ebola-Zaire and Ebola-Reston virus infection of primary human monocytes, the early targets of Ebola-Zaire virus infection. We found that approximately 200 of 1,400 human genes on the array exhibited changes in expression in response to Ebola-Zaire virus infection after 24 hours. Most affected genes were upregulated in their level of expression, including cytokine and chemokine genes (IL-1, IL-1, IL-6, IL-8, IL-15, MIP-1, MIP-1 and TNF), genes involved in regulation of cell cycle or apoptosis and other genes involved in signal transduction. The gene expression profile from Ebola-Reston–infected monocytes was totally different from that observed with Ebola-Zaire virus. The results from northern-blot or ribonuclease protection assays confirmed the array data. The possible influence of differences in cellular gene expression observed between Ebola-Zaire and Ebola-Reston viruses on the ability of these viruses to cause diseases will be discussed.

Research (/tags/106) In-Press Preview (/tags/113) Cell biology (/tags/16) Immunology (/tags/25) Free access | 10.1172/JCI122462 (https://doi.org/10.1172/JCI122462)

TGFβ-induced epigenetic deregulation of SOCS3 facilitates STAT3-signaling to promote fibrosis

Clara Dees, Sebastian Pötter, Yun Zhang, Christina Bergmann, Xiang Zhou, Markus Luber, Thomas Wohlfahrt, Emmanuel Karouzakis, Andreas Ramming, Kolja Gelse, Akihiko Yoshimura, Rudolf Jaenisch, Oliver Distler, Georg Schett, and Jörg H.W. Distler

First published January 28, 2020 - More info

Abstract

Fibroblasts are key-effector cells in tissue remodeling. They remain persistently activated in fibrotic diseases, resulting in progressive deposition of extracellular matrix. Although fibroblast activation maybe initiated by external factors, prolonged activation can induce an “autonomous”, self-maintaining pro-fibrotic phenotype in fibroblasts. Accumulating evidence suggests that epigenetic alterations play a central role to establish this persistently activated pathologic phenotype of fibroblasts. We demonstrated that in fibrotic skin of patients with systemic sclerosis (SSc), a prototypical idiopathic fibrotic disease, transforming growth factor-β (TGFβ) induced the expression of DNA-

Phytocannabinoids Oral Immunization adjuvants

www.nature.com/scientificreports

SCIENTIFIC REPORTS

OPEN

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

Received: 31 August 2016
Accepted: 12 January 2017
Published: 17 February 2017

Emmanuel Dotsey[1], Irina Ushach[1], Egest Pone[2], Rie Nakajima[1], Algis Jasinskas[1], Donovan A. Argueta[3], Andrea Dillon[3], Nicholas DiPatrizio[3], Huw Davies[1], Albert Zlotnik[4], Peter D. Crompton[5] & Philip L. Felgner[1]

BJP British Journal of Pharmacology

DOI:10.1111/j.1476-5381.2011.01238.x
www.brjpharmacol.org

Themed Issue: Cannabinoids in Biology and Medicine, Part I

REVIEW

Taming THC: potential cannabis synergy and phytocannabinoid-terpenoid entourage effects

Ethan B Russo

GW Pharmaceuticals, Salisbury, Wiltshire, UK

Correspondence
Ethan Russo, MD, 20402 81st Avenue SW, Vashon, WA 98070, USA. E-mail: ethanrusso@comcast.net

Keywords
cannabinoids; terpenoids; essential oils; THC; CBD; limonene; pinene; linalool; caryophyllene; phytotherapy

Received
19 November 2010
Revised
29 December 2010
Accepted
12 January 2011

Oral and Topical Immunization: Z-Stack Plus

Beta-caryophyllene is a dietary cannabinoid

Jürg Gertsch*†, Marco Leonti‡§, Stefan Raduner*§, Ildiko Racz¶, Jian-Zhong Chen‖, Xiang-Qun Xie‖, Karl-Hein
Meliha Karsak¶, and Andreas Zimmer¶

*Institute of Pharmaceutical Sciences, Department of Chemistry and Applied Biosciences, Eidgenössische Technische Hochschule (ETH) Zurich
Switzerland; ‡Dipartimento Farmaco Chimico Tecnologico, University of Cagliari, 01924 Cagliari, Italy; ¶Department of Molecular Psychiatry,
Bonn, 53115 Bonn Germany; and ‖Department of Pharmaceutical Sciences, University of Pittsburgh, Pittsburgh, PA 15260

β-Caryophyllene, A Natural Dietary CB2 Receptor Selective Cannabinoid can be a Candidate to Target the Trinity of Infection, Immunity, and Inflammation in COVID-19

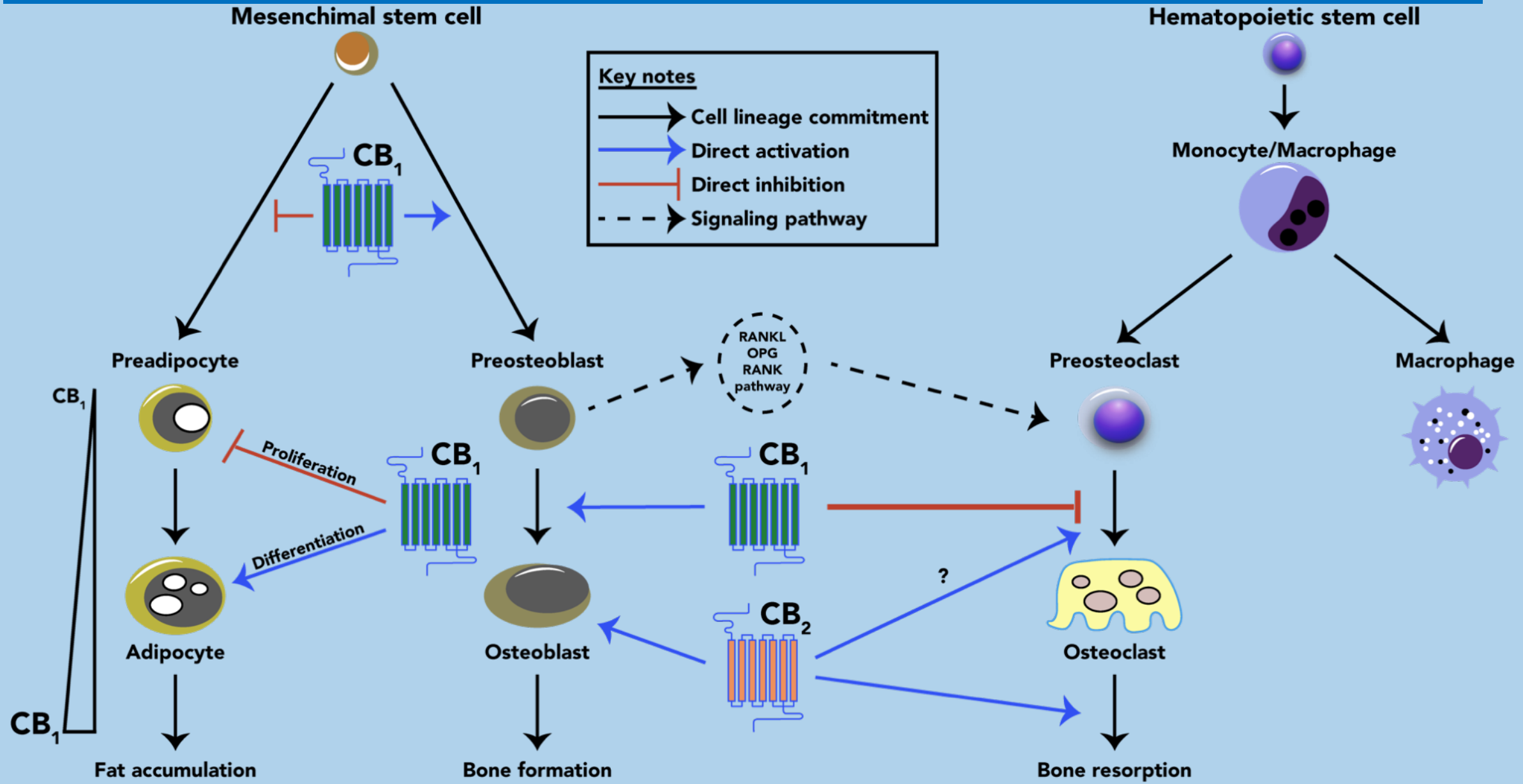
Niraj Kumar Jha[1†], Charu Sharma[2†], Hebaallah Mamdouh Hashiesh[3], Seenipandi Arunachalam[3], MF Nagoor Meeran[3], Hayate Javed[4], Chandragouda R. Patil[5], Sameer N. Goyal[6] and Shreesh Ojha[3*]

Beta-caryophyllene enhances wound healing through multiple routes

Sachiko Koyama, Anna Purk, Manpreet Kaur, Helena A. Soini, Milos V. Novotny, Keith Davis, C. Cheng Kao, Hiroaki Matsunami, Anthony Mescher

Published: December 16, 2019 • https://doi.org/10.1371/journal.pone.0216104

CB2 Dysregulation is associated with Chronic Inflammation Driving cardiovascular disease, Obesity & Cancer



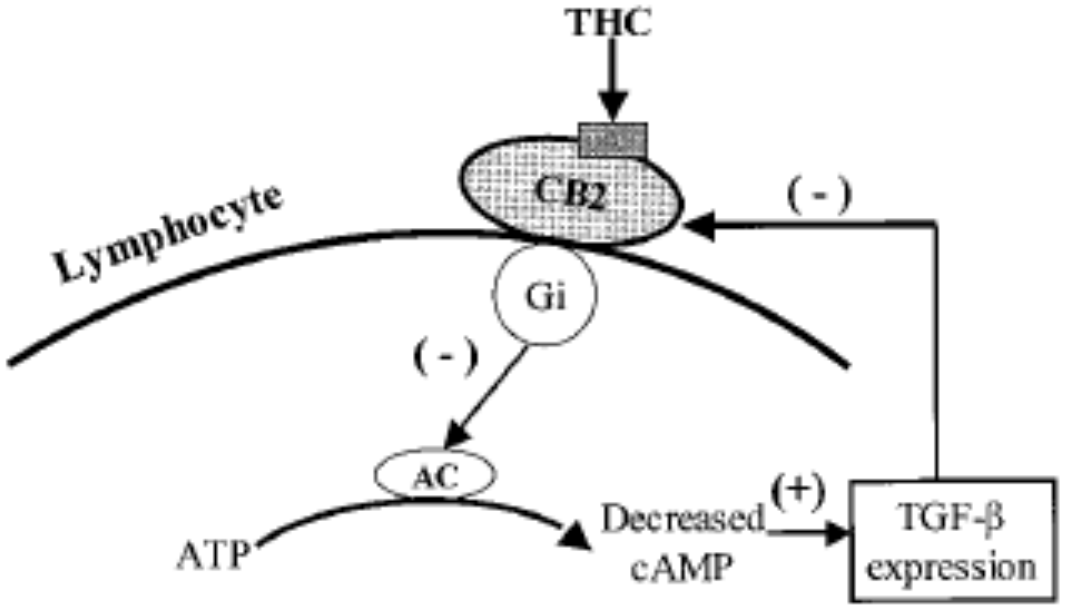
Trends in Neurosciences

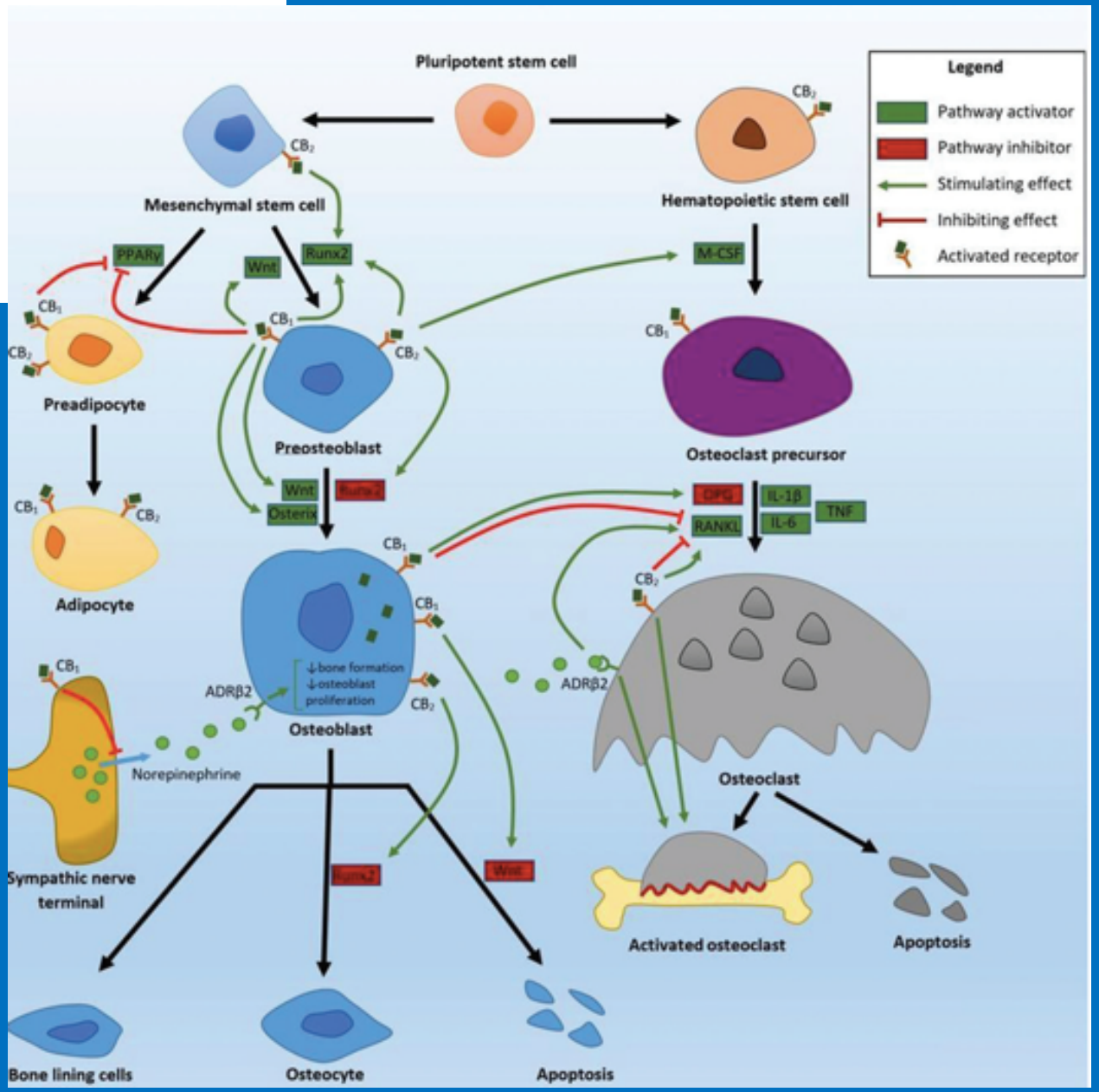
CellPress
REVIEWS

Review

Ion Channel Functions in Early Brain Development

Richard S. Smith[1,*] and Christopher A. Walsh[1,*]





Cannabinoids regulate MINERALS in Immune Cells via endocannabinoid System Receptors

- A downside of activation of MINERALS is the dysregulation endogenous microbes
- OUR SOILS ARE DEPLETED OF MINERALS

TRPV1/2	Ca^{2+}/Na^{+}	PM	Heat (fever?), low pH, mechanical stress		Mono, macro	Degranulation, phagocytosis, cytokine production
TRPC3/6	Ca^{2+}/Na^{+}	PM	PLC activation (DAG), PIP_2		T, B, NK cells, neutro	Chemotaxis, degranulation
TRPM2	Ca^{2+}/Na^{+}	PM, lys	H_2O_2, NAADP, cADPR		T, B, neutro, mast cells, DC	Cytokine production, degranulation
Magnesium						
TRPM6	$Mg^{2+}>Ca^{2+}$	PM		Inhibited by $[Mg^{2+}]_i$	Gut, kidney, hematopoietic (not T cells)	Unknown in immune cells
TRPM7	$Mg^{2+}>Ca^{2+}$	PM	Unknown (BCR, TCR?) PIP_2 (?)	Inhibited by $[Mg^{2+}]_i$	Ubiquitous	T cell development, T and B cell proliferation, cytokine production

Review

CellPress

Divalent cation signaling in immune cells

Benjamin Chaigne-Delalande and Michael J. Lenardo

Molecular Development of the Immune System Section, Lymphocyte Molecular Genetics Unit, Laboratory of Immunology, National Institute of Allergy and Infectious Diseases, National Institutes of Health, Bethesda, MD 20892, USA

Divalent cations of two alkaline earth metals Ca^{2+} and Mg^{2+} and the transition metal Zn^{2+} play vital roles in the immune system, and several immune disorders are associated with disturbances of their function. Until re-

Trends in Immunology July 2014, Vol. 35, No. 7

VIRUSES/POSIONS: Loss of TASTE AND SMELL
Lack of Minerals, Essential Amino acids, Phytocannabinoids

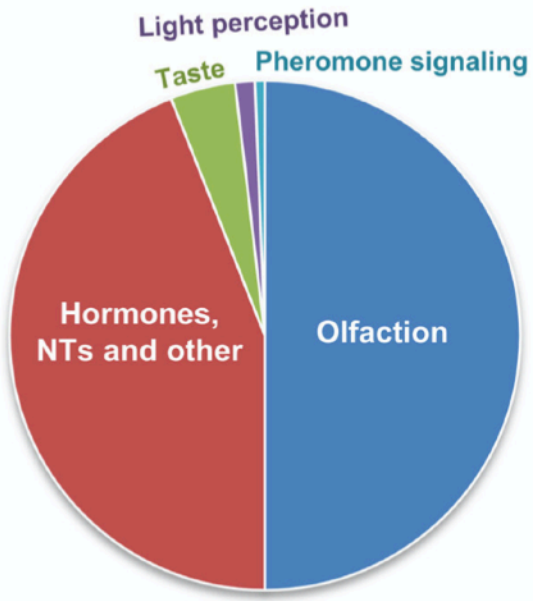
frontiers in Aging Neuroscience

REVIEW
published: 24 April 2019
doi: 10.3389/fnagi.2019.00089

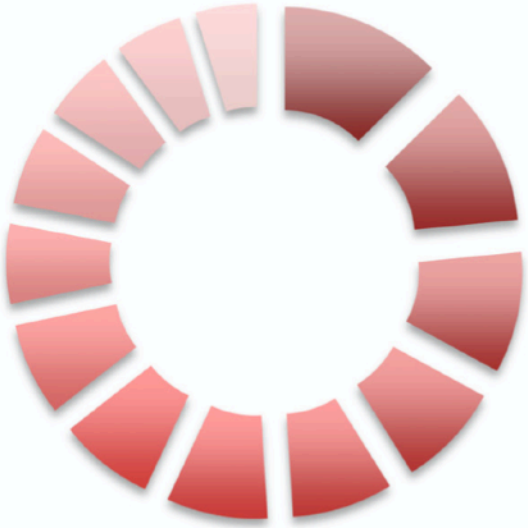
$G_{i/o}$-Protein Coupled Receptors in the Aging Brain

Patrícia G. de Oliveira[†], Marta L. S. Ramos[†], António J. Amaro[2], Roberto A. Dias[†‡] and Sandra I. Vieira[†]*

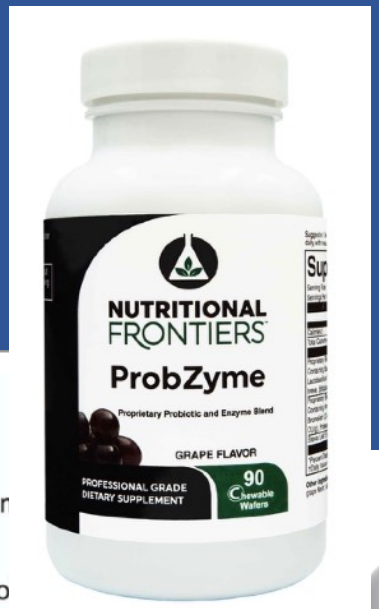
A GPCRs main functions



B Tissue distribution of $G_{i/o}$-coupled GPCRs



- Bone r
- Brain
- Gastro
- Lung
- Male tissues
- Female tissues
- Endocrine tissues
- Kidney & Urinary bladder
- Skin
- Liver and gallbladder
- Muscle tissues
- Pancreas
- Adipose and soft tissues



Nitric oxide is a soluble gas that is continually being made from arginine in endothelial cells. Endothelial cells comprise a layer of cells inside the lining of our blood vessels.

Pharmacogn. Res.
ORIGINAL ARTICLE

Vitamin D_3, L-Arginine, L-Citrulline, and Antioxidant Supplementation Enhances Nitric Oxide Bioavailability and Reduces Oxidative Stress in the Vascular Endothelium – Clinical Implications for Cardiovascular System

Hazem Dawoud, Tadeusz Malinski

Department of Chemistry and Biochemistry, Nanomedical Research Laboratories, Ohio University, Athens, Ohio, USA

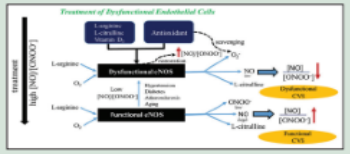
ABSTRACT

Background: Nitric oxide (NO) is a crucial signaling molecule which regulates the blood flow and prevents the adhesion of blood components to the vascular wall. A deficiency in bioavailable NO concentration is associated with the dysfunction of endothelial NO synthase (eNOS) and/or an increase in oxidative stress. The deficiency of bioavailable NO is a common denominator of several cardiovascular diseases, including diabetes, atherosclerosis, and hypertension. **Materials and Methods:** We used a nanomedical technology to elucidate the balance between bioavailable NO and oxidative stress (peroxynitrite $ONOO^-$) in human umbilical vein endothelial cells (HUVECs) treated with a supplement containing L-arginine, L-citrulline, Vitamin D_3, and antioxidants. Nanosensors, with a diameter of 200–300 nm, are capable of measuring *in situ* NO and peroxynitrite ($ONOO^-$) concentrations produced by single endothelial cells. **Results:** The ratio of the concentration of cytoprotective NO [NO] to the concentration of cytotoxic peroxynitrite [$ONOO^-$] was used to estimate the efficiency of eNOS. HUVECs incubated with L-citrulline, L-arginine, and Vitamin D_3 increased the [NO]/[$ONOO^-$] ratio by 25%, while in the presence of antioxidants, the increase was 15%. The synergistic effect between the mix of L-arginine, L-citrulline, Vitamin D_3, and antioxidants was a favorable increase of the overall [NO]/[$ONOO^-$] ratio by 50%. **Conclusion:** The findings of the study presented here clearly indicate that L-arginine, L-citrulline, and Vitamin D_3 can significantly alter the function of the endothelium and NO production, in a favorable manner, while pointedly reducing $ONOO^-$ – the main component of oxidative stress. This effect can be significantly potentiated in the presence of antioxidants.

Key words: Antioxidant, endothelium, L-arginine, L-citrulline, nitric oxide, peroxynitrite, Vitamin D_3

SUMMARY

- Nanomedical studies were used to elucidate the role of a mixture of Vitamin D_3, L-arginine, L-citrulline, and several antioxidants in the improvement of nitric oxide production and the reduction of oxidative stress in human endothelial cells. It appears that the combination of natural products can effectively improve endothelial function by about 50% and has shown that, on cellular models, it could potentially be used to improve the endothelial function in cardiovascular diseases.



Abbreviations Used: HUVECs: Human umbilical vein endothelial cells; $O_2^{\cdot-}$: Superoxide; HBSS: Hank's balanced salt solution; EC: Endothelial cell; Cal: Calcium ionophore; CVD: Cardiovascular disease; eNOS: Endothelial nitric oxide synthase.

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Nanomedical Research Laboratories, Ohio University, 350 West State Street, Athens, Ohio, USA.
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Website: www.phcogres.com
Quick Response Code:

INTRODUCTION

Nitric oxide (NO) is a gaseous molecule that is generated by the NO synthase (NOS) enzyme. NO is synthesized from two substrates: L-arginine (non-essential amino acid) and oxygen.[1,2] This synthesis occurs through NOS in a five-electron transfer oxidation of L-arginine to L-citrulline. NOS is located in the membrane of endothelial cells, and its synthesis is stimulated by calcium flux.[3,4] In the cardiovasculature, the calcium flux is triggered by a mechanical process (shear stress)[5] and chemical stimuli such as acetylcholine, norepinephrine, angiotensin II, and many others.[6,7]

NO can react rapidly with many biological components, including superoxide ($O_2^{\cdot-}$), Fe (III) of hemoglobin, guanylate cyclase, and many others.[8–10] Therefore, the measurement of reactive "free" NO is a challenging problem. In our laboratories, we are able to perform measurements of bioavailable NO produced by a single endothelial cell in different segments of the cardiovascular system, such as capillary vessels, aorta, and heart. Maximal NO concentrations vary significantly, depending on the location of the endothelial cells – with the lowest concentrations in the small capillary (about 80 nM) and the highest in the endocardium of the heart (about 2.0 μM).[11] The level of NO concentration depends largely on the velocity and type of blood flow (laminar vs. turbulent).[12–14]

Cite this article as: Dawoud H, Malinski T. Vitamin D_3, L-Arginine, L-Citrulline, and antioxidant supplementation enhances nitric oxide bioavailability and reduces oxidative stress in the vascular endothelium – Clinical implications for cardiovascular system. Phcog Res 2020;12:17-23.

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Functional characterization of nutraceuticals using spectral clustering: Centrality of caveolae-mediated endocytosis for management of nitric oxide and vitamin D deficiencies and atherosclerosis

OPEN ACCESS

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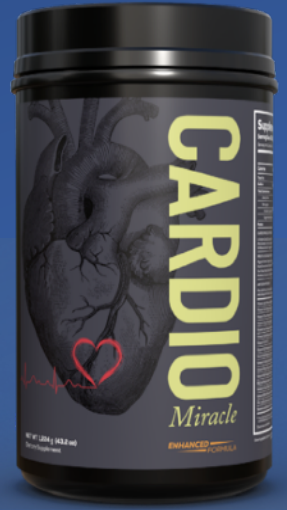
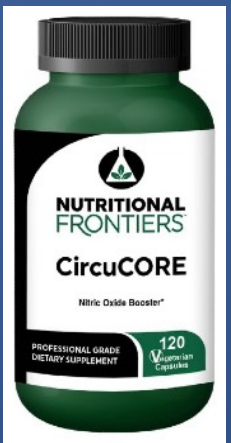
SCAN ME





Provide the building blocks to support nitric oxide formation enhance overall circulation, including heart health and erectile dysfunction.

AMERICAN SOCIETY FOR MICROBIOLOGY | Antimicrobial Agents and Chemotherapy®

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100 Years of Suramin

Natalie Wiedemar,[a,b] Dennis A. Hauser,[a,b] Pascal Mäser[a,b]

SURAMIN, THE FRUIT OF EARLY MEDICINAL CHEMISTRY

SURAMIN AS AN ANTIPARASITIC DRUG

SURAMIN AS AN ANTIVIRAL AGENT

SURAMIN AGAINST CANCER

SURAMIN AS AN ANTIDOTE

Three of the many biological activities of suramin support its potential use as a protective agent: the inhibition of thrombin, the inhibition of phospholipase A2, and the inhibition of purinergic signaling.

FURTHER POTENTIAL USES OF SURAMIN

TABLE 1 Diseases and pathogens susceptible to suramin

Disease and/or pathogen	Activity in[a]: Cell culture	Animal model	Patient
Parasitic infections			
T. b. rhodesiense HAT	X	X	X
T. brucei gambiense HAT	X	X	X
Surra, *T. evansi*	X	X	NA
River blindness, *O. volvulus*	X	X	X
T. cruzi	X		
Leishmania spp.	X		
P. falciparum	X		
Viral infections			
Hepatitis virus	X	X	X
AIDS, HIV	X		X
Herpes simplex virus	X	X	
Chikungunya virus	X	X	
Enterovirus 71	X	X	
Dengue virus	X		
Zika virus	X		
Ebola virus	X		
Neoplastic diseases			
Non-small cell lung cancer	X	X	
Breast cancer	X	X	
Bladder cancer	X	X	
Brain tumors	X	X	
Prostate cancer	X	X	X
Other			
Snakebite	X	X	
Arthritis	X	X	
Autism	NA	X	X

OVERVIEW

Potential therapeutic targets in the rapidly expanding field of purinergic signalling

Clin Med JRCPL2002;2:45–53

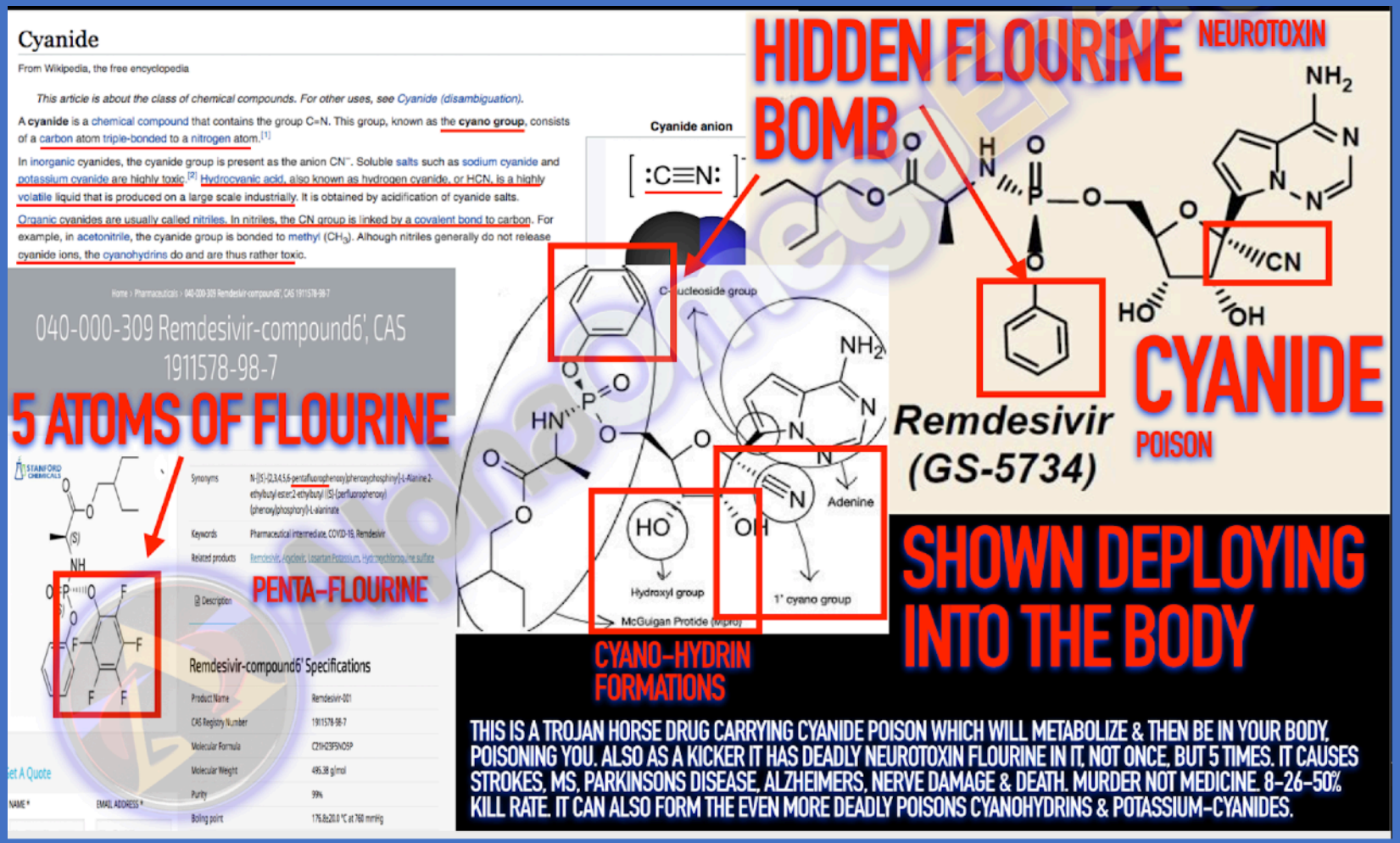
G Burnstock

The role of P2X3 receptors in nociception and a new hypothesis concerning purinergic mechanosensory transduction in visceral pain will be considered, as will the therapeutic potential of purinergic agonists or antagonists for the treatment of supraventricular tachy- cardia, cancer, dry eye, bladder hyperactivity, erectile dysfunction, osteoporosis, diabetes, gut motility and vascular disorders.

Key Points

- Purinergic signalling is widespread and acts through three families of receptors: P1 receptors for adenosine; P2X ligand-gated ion-channel receptors and P2Y G-protein-coupled receptors for ATP, ADP and UTP
- $P2Y_{12}$ receptors on platelets mediate ADP-induced aggregation; the $P2Y_{12}$ receptor antagonist clopidogrel (Plavix) is a very promising antithrombotic agent, especially when combined with aspirin
- $P2X_3$ receptors are located in nociceptive sensory nerves, and selective antagonists are being developed against the initiation of visceral pain
- $P2Y_2$ agonists and antagonists, which modulate mucus secretion, are being developed to treat cystic fibrosis, chronic obstructive pulmonary disease and 'dry eye' conditions
- Purinergic agonists and antagonists are being developed to modulate cell proliferation, migration, differentiation and death for treating tumours, osteoporosis and vascular conditions such as hypertension, atherosclerosis and restenosis, and to regulate angiogenesis
- Adenosine is being used successfully as a diagnostic tool and for reversing supraventricular tachycardia. The antiasthmatic drug theophylline acts through antagonism of the adenosine receptor, and adenosine-receptor antagonists are being explored for the treatment of Parkinson's disease
- The therapeutic potentials of purinergic compounds in kidney failure, glaucoma, osteoporosis, wound healing, detrusor instability, erectile function and arthritis are also being considered

Fauci's Trojan horse Poisons Remdesivir and Paxlovid "Ivermectin" with Cyanide & Fluorine



SYNCYTIN: ONLY One Component of Snake Venom additional components/toxins in Food, Water "Drugs"

FEBS Letters 436 (1998) 256–258 FEBS 20902

Enhancement and inhibition of snake venom phosphodiesterase activity by lysophospholipids

ScienceDirect

Estuarine, Coastal and Shelf Science
Volume 219, 5 April 2019, Pages 161-168

Microplastic pollution in commercial salt for human consumption: A review

Diogo Peixoto [a], Carlos Pinheiro [a], João Amorim [a], Luís Oliva-Teles [a, b], Lúcia Guilhermino [a, c], Maria Natividade Vieira [a, b]

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Outline | Share | Cite

https://doi.org/10.1016/j.ecss.2019.02.018

Highlights

- Plastics as marine debris are the new addition to the list of global threats.
- Marine pollution will undoubtedly lead to the contamination of sea products.
- Microplastics in salts might pose a threat to human food safety and health.
- Microplastics sorb contaminants and transfer them to salt and other products.

Bayer Request for Additional Information and Attestation
Regarding Religious Exemption / Accommodation Form (Covid-19 Vaccine)

Bayer requires additional information to further consider your request for a religious exemption/accommodation. Please complete this form and attestation and submit it to accommodations_US@bayer.com.

Your request appears to be principally based upon your objection to the use of fetal cell lines in the testing, research, or development of the COVID-19 vaccine and/or your belief concerning the purity of the body. The information reported on this form will serve to validate your understanding of fetal cell use in common medicines and consumer products, and aid in assessing the sincerity of your professed religious belief.

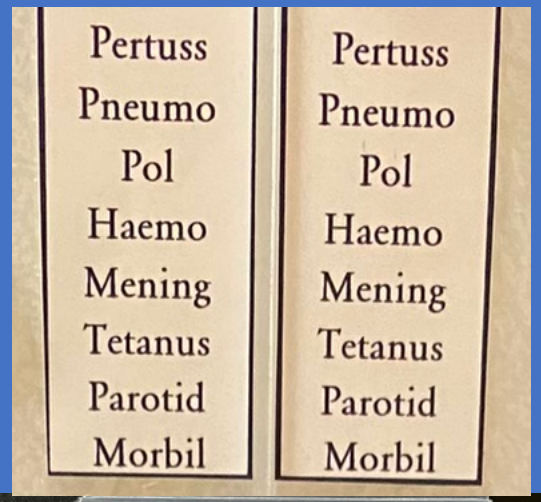
The following is a non-exhaustive list of common medicines and products that have used fetal cells in testing, research, and/or development.[1]

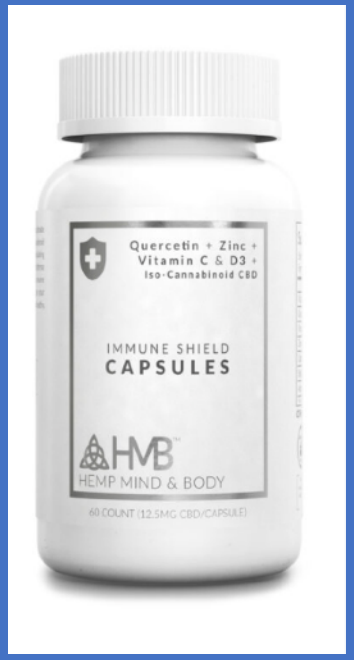
Acetaminophen	Enbrel	Maalox	Sudafed
Acetylsalicylic Acid (ASA)	Ex-Lax, Zocor	Metformin/Glucophage	Suphedrine
Advil	Havrix	Motrin	Toprol
Albuterol	Hydroxychloroquine	Mucinex	Tums
Aleve	Ibuprofen	Pepto Bismol	Tylenol
Amlodipine/Norvasc	Ivermectin	Preparation H	Varilrix
Aspirin	Levothyroxine	Prilosec OTC/Zegrid	Zoloft
Azithromycin	Lidocaine	Robitussin/Delsym	Zostavax
Benadryl	Lipitor	Senokot	
Claritin	Losartan/Cozaar	Simvastatin	

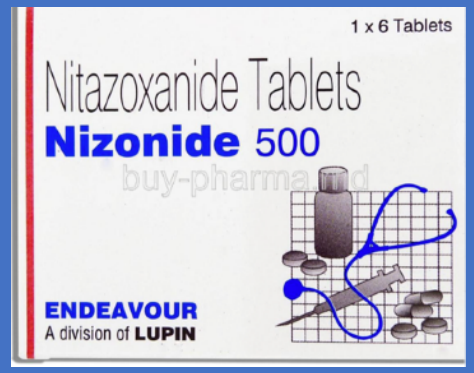
To be Completed by Individual Requesting the Accommodation	
Full Name:	Click here to enter name.
Employee or Contractor ID #	
Email:	Click here to enter email.

Please state whether your religious objection to the COVID-19 vaccine is equally applicable to the above medicines and other products that used fetal cells in testing, research, and/or development. If not, please explain why.	Click here to enter text.
If your religious objection to the COVID-19 vaccine is equally applicable to medicines and products that used fetal cells in testing, research, and/or development, please state whether you abstain from using all such medicines and products. If not, please	Click here to enter text.

Proven Effective Oral Immunization strategies for XMRVs Including SARSCoV2







Dear Dr. Judy, I am excited to report that thanks to my jabbed neighbors and friends I have now developed sufficient immunity and ample antibodies. Look at my value: 417! Thanks to your protocol I am not afraid of the vaccinated. I take my Immune Formulation 200, Cardio Miracle, Prolean Greens DMG & Paximune regularly. Natural immunity for the win! It truly works!

AFFIDAVIT OF Proof of IMMUNITY
EXEMPTION ON RELIGIOUS GROUNDS FROM Vaccination/Inoculations

Date: ____/____/20____

Governing Authority Name (business issuing mandate): _____
Address: _____

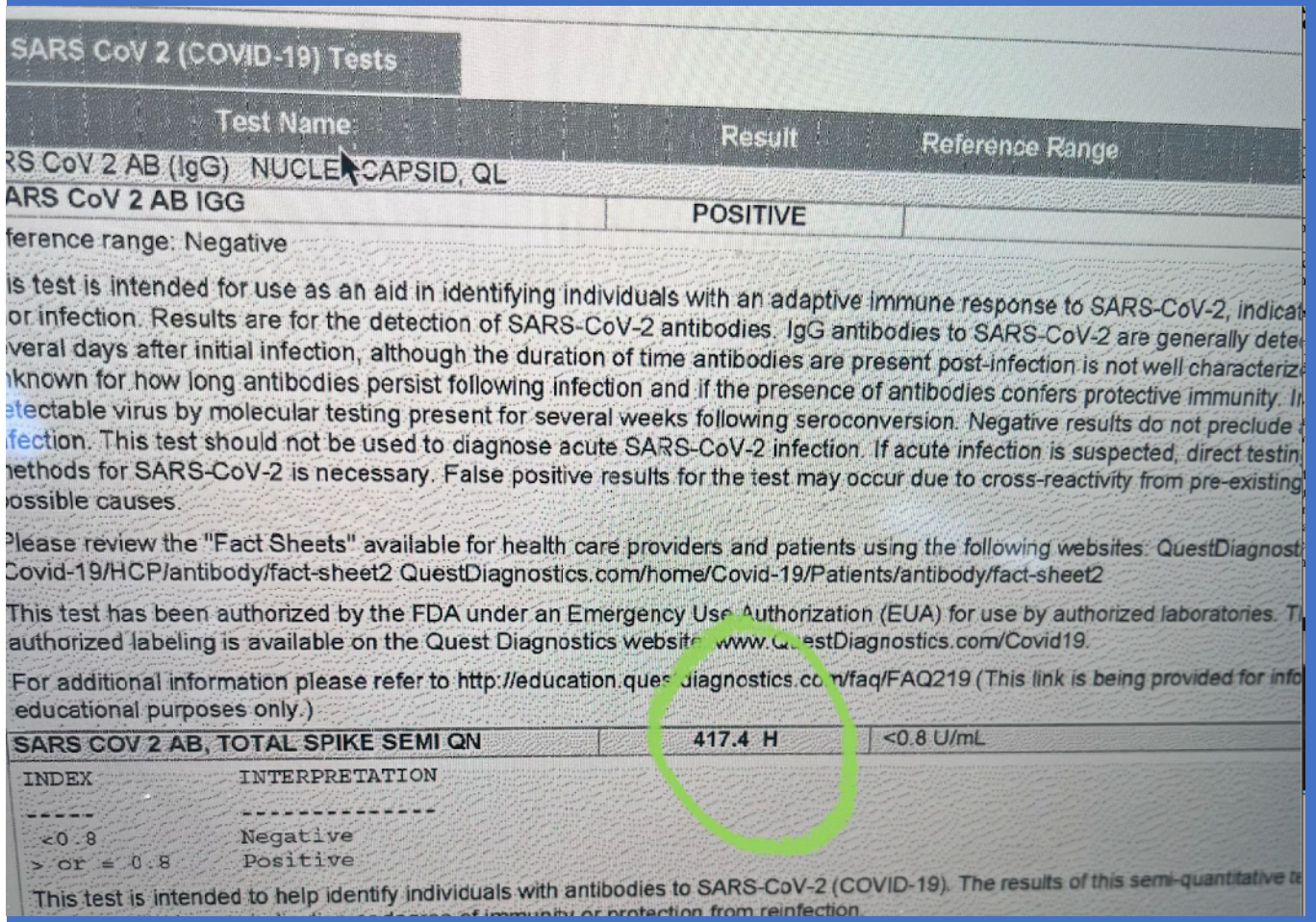
RE: Religious Exemption from Vaccination Requirements. Declaration of God Given IMMUNITY

I, (Name) _____, the undersigned do hereby swear and affirm that I am a member of a recognized religious organization, and that the immunizations required by (Governing Authority Name) _____, are contrary to my religious tenets and practices. On this basis, as no vaccination on the CDC schedule has been safety tested as acknowledged by the CDC in 2019 and EUA COVID19 inoculations/immunization requirements violate my right to freely exercise my religion as guaranteed by the First Amendment of the Constitution of The United States of America, I am asserting my rights to an exemption by (Governing Authority Name) _____ from EUA Inoculation and nasal Swab PCR test requirements as I am immunized/vaccinated by virtue of the robust natural immunity I acquired when I recovered from COVID19.

I also am immunized according to my religious beliefs as I regularly take oral booster. My immunization strategy **exceeds** FDA and CDC mandated standards, which were recently changed such that the vaccination need not provide immunity.

I qualify for this exemption based on the First Amendment of the United States Constitution and *42 U.S. Code § 2000a - Prohibition against discrimination or segregation in places of public accommodation*, which states "All persons shall be entitled to the full and equal enjoyment of the goods, services, facilities, privileges, advantages, and accommodations of any place of public accommodation, as defined in this section, without discrimination or segregation on the ground of race, color, religion, or national origin." Failure to uphold 42 U.S. Code § 2000a may be met with legal action.

Thank you in advance


SARS CoV 2 (COVID-19) Tests

Test Name	Result	Reference Range
RS CoV 2 AB (IgG) NUCLEOCAPSID, QL		
ARS CoV 2 AB IGG	POSITIVE	

ference range: Negative

is test is intended for use as an aid in identifying individuals with an adaptive immune response to SARS-CoV-2, indicat or infection. Results are for the detection of SARS-CoV-2 antibodies. IgG antibodies to SARS-CoV-2 are generally dete veral days after initial infection, although the duration of time antibodies are present post-infection is not well characteriz known for how long antibodies persist following infection and if the presence of antibodies confers protective immunity. I tectable virus by molecular testing present for several weeks following seroconversion. Negative results do not preclude fection. This test should not be used to diagnose acute SARS-CoV-2 infection. If acute infection is suspected, direct testin ethods for SARS-CoV-2 is necessary. False positive results for the test may occur due to cross-reactivity from pre-existing ossible causes.

Please review the "Fact Sheets" available for health care providers and patients using the following websites: QuestDiagnost Covid-19/HCP/antibody/fact-sheet2 QuestDiagnostics.com/home/Covid-19/Patients/antibody/fact-sheet2

This test has been authorized by the FDA under an Emergency Use Authorization (EUA) for use by authorized laboratories. T authorized labeling is available on the Quest Diagnostics websit www.QuestDiagnostics.com/Covid19.

For additional information please refer to http://education.ques diagnostics.com/faq/FAQ219 (This link is being provided for info educational purposes only.)

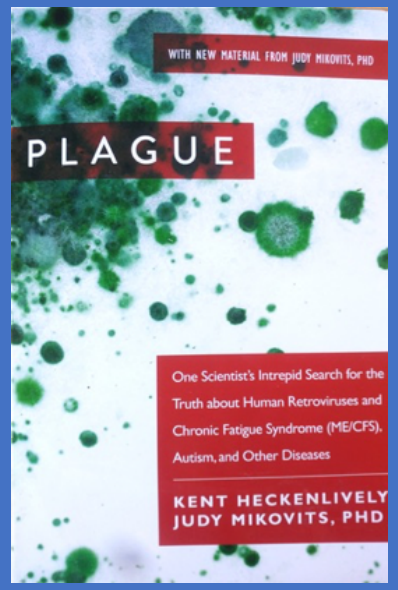
SARS COV 2 AB, TOTAL SPIKE SEMI QN	417.4 H	<0.8 U/mL

INDEX	INTERPRETATION
<0.8	Negative
> or = 0.8	Positive

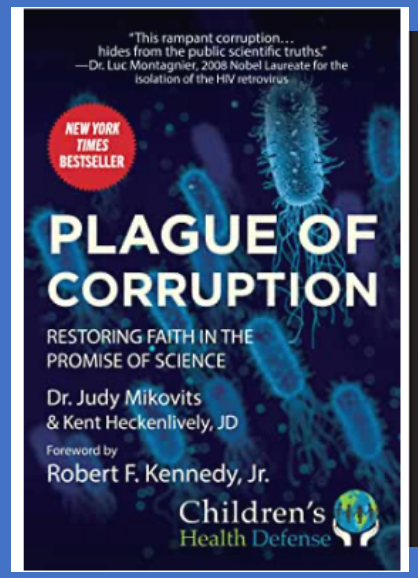
This test is intended to help identify individuals with antibodies to SARS-CoV-2 (COVID-19). The results of this semi-quantitative t

GOD's People are destroyed from lack of Knowledge (Hosea 4:6)

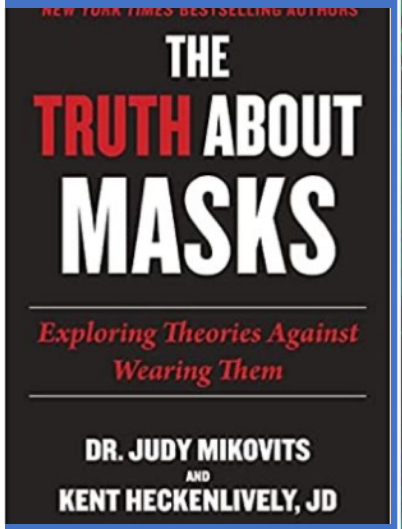
2014



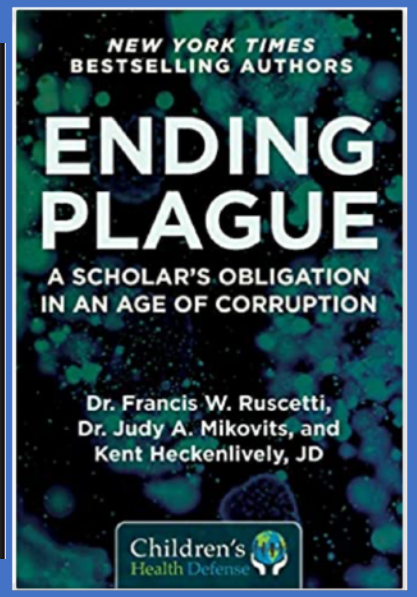
(James 1:19-22) 2017



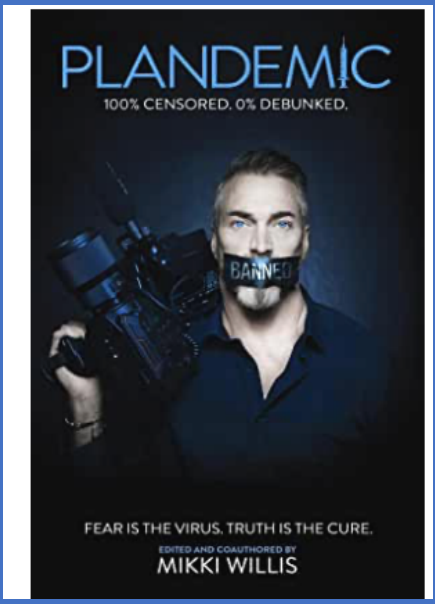
2020 (Psalm 91)



2020 1(Cor 3:18)



2021(Ephesians 5:11)



2021(2 Chronicles 7:14)

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