SPIKE DETOX PROTOCOLS INITIAL SITUATION

- 1. What are we facing
- 2. Successful strategies
- 3. Patient results





ONGOING RESEARCH AND DEVELOPMENT FEEDBACKING WITH RESULTS FROM PRACTICE











PRE-CLINICAL AND IN-VITRO LABORATORY



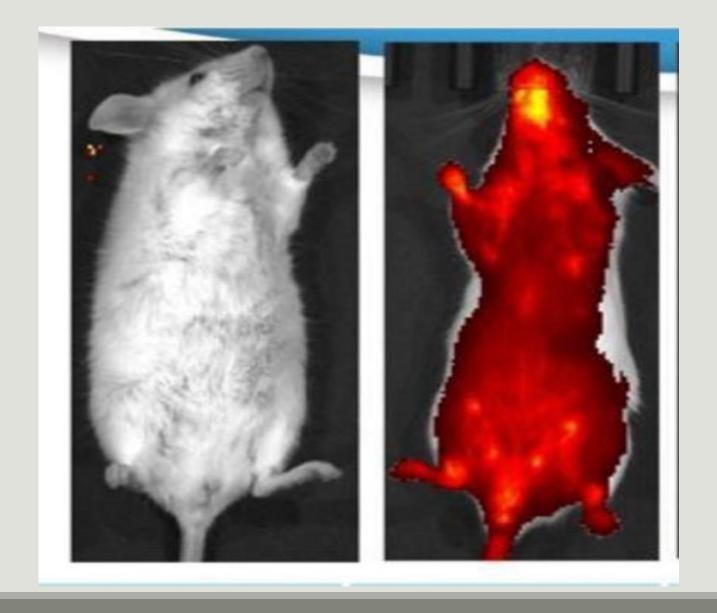
We conducted several hundred in silico, In vitro, pre-clinical and clinical studies on various combinations of molecules to arrive at our Protocol.

A large number of peer reviwed publications came out of this group so far.





ANIMAL 72 HOURS AFTER INJECTION OF SPIKE PROTEIN





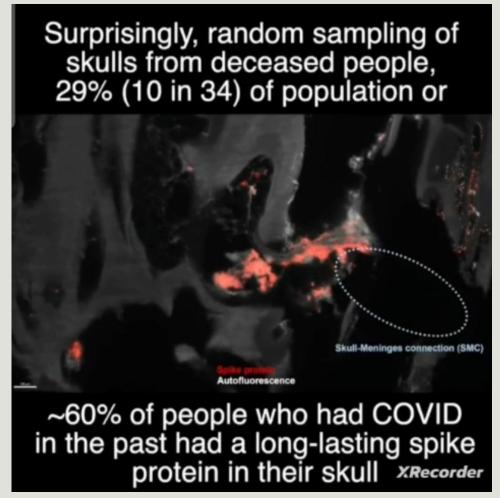


60% of all!!! Sars CoV 2 infected have persistent spike in the bone marrow of the skull and god knows where else. This has to be

removed

SARS-CoV-2 Spike Protein
Accumulation in the Skull-MeningesBrain Axis: Potential Implications for
Long-Term Neurological
Complications in post-COVID-19

Zhouyi Rong, Hongcheng Mai, Saketh Kapoor, Victor G. Puelles, Jan Czogalla, Julia Schädler,







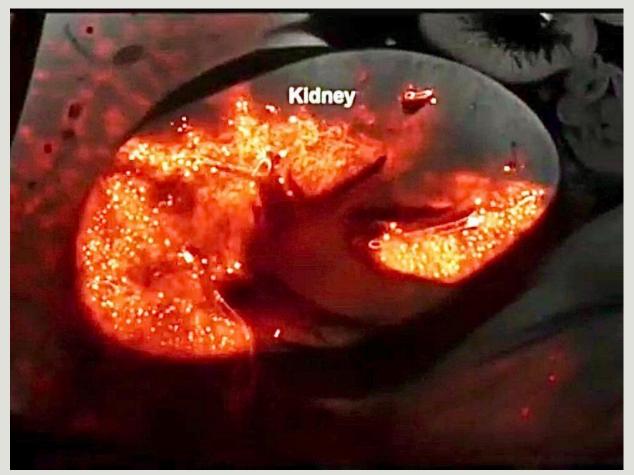








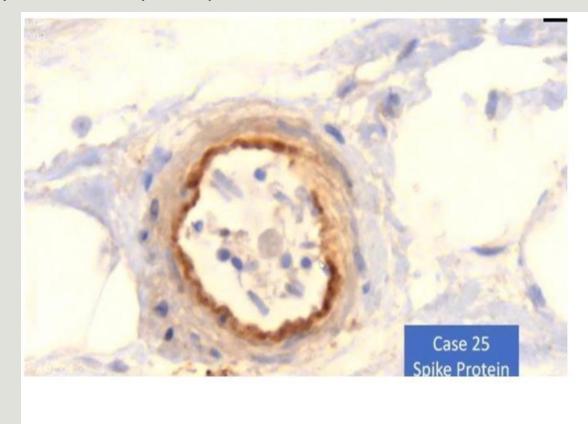








Synthetic Spike protein in arterioles:



Schuss Doutlinger Automaia

Reutlinger Autopsie/Histologie-Studie Impfnebenwirkungen und -Todesfälle

8 Kooperierende Pathologen/ Biologen international 25 Todesfälle / 3 Proben von Lebenden

- 15 Fälle ausgewertet Stufe1: Routine Histologie
- 1 Fall Stufe 2: Spezialmethoden

7 Männer, 8 Frauen; 28 bis 95 Jahre alt Tod 7 Tage bis 6 Monate nach letzter Injektion Impfstoffe:

Comirnaty/Pfizer-BioNTech 8, Moderna 2, Janssen 1, Astra-Zeneca 2, unbekannt 2

VEDICINALS°

Reutlinger Histologie-Studie - Vortrag von Prof. Dr. med. Arne Rurkhardt am 21.01.2022

Spite 11 v

And we find it not only on the capillaries, but also on the small arterioles. Here ["Case 25 Spike Protein"] this clear positive reaction and here an interruption in the inner vessel wall layers. And you see here inside again the sloughed off endothelia and single inflammatory cells.

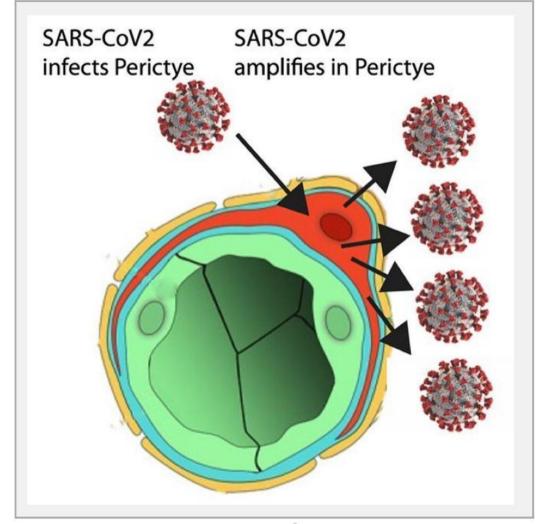
Figure depicts SARS-CoV-2 spreading through blood vessels (green) to infect pericytes (red), which amplify infection and can spread infection

to other cell types in the brain.

3D "assembloid" shows how SARS-CoV-2 infects brain cells

Peer-Reviewed Publication

UNIVERSITY OF CALIFORNIA - SAN DIEGO





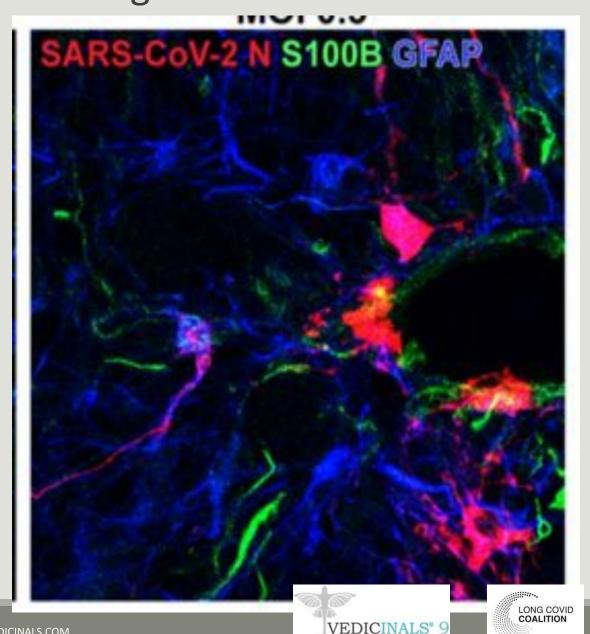


Mature and precursor astroglial cells indicate high infection.

Tropism of SARS-CoV-2 for Developing Human Cortical Astrocytes

Madeline G. Andrews, Tanzila Mukhtar, Ugomma C. Eze, Camille R. Simoneau, Yonatan Perez, Mohammed A. Mostajo-Radji, Shaohui Wang, Dmitry Velmeshev, Jahan Salma, G. Renuka Kumar, Alex A. Pollen, Elizabeth E. Crouch, Melanie Ott, Arnold R. Kriegstein

doi: https://doi.org/10.1101/2021.01.17.427024



Malfunctioning of glymphatic drainage; a network of lymphatic vessels that clear waste from the central nervous system (CNS), mostly during sleep.1 This results in the abnormal accumulation of fluid, waste products, and inflammatory cytokines in the brain parenchyma, leading to nervous system symptoms like fatigue, depression, myalgias, anosmia, and in particular, brain fog.2

MINI REVIEW

Glymphatics and brain fog - the post-COVID-19 phenomenon

Mulazim Hussain Bukhari^{1*}, Shumaila Liaqat², Nadia Naseem³





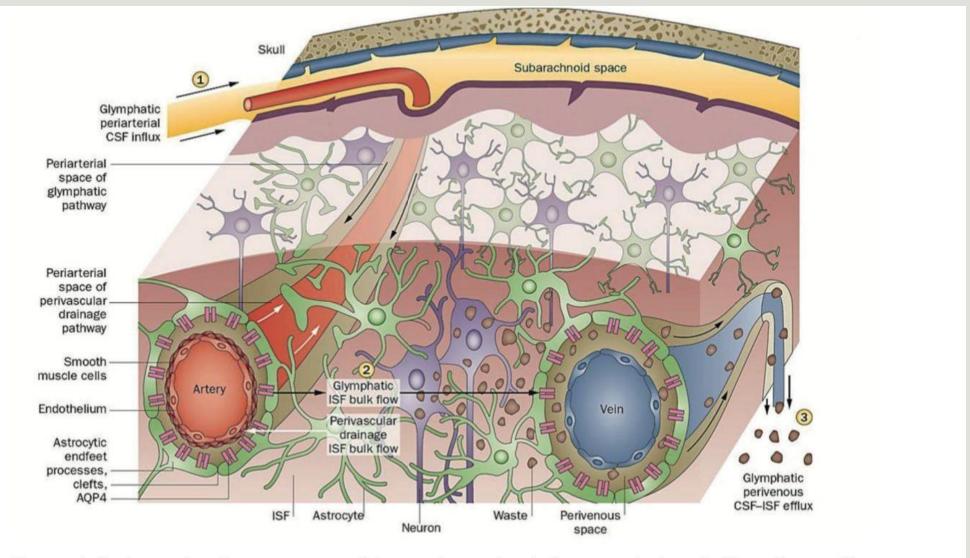


Figure 1. Perivascular clearance comprising perivascular drainage and glymphatic pathways.19





NEURO INFLAMMATION



Research Article

Neuroscience

Recapitulation of pathophysiological features of AD in SARS-CoV-2-infected subjects

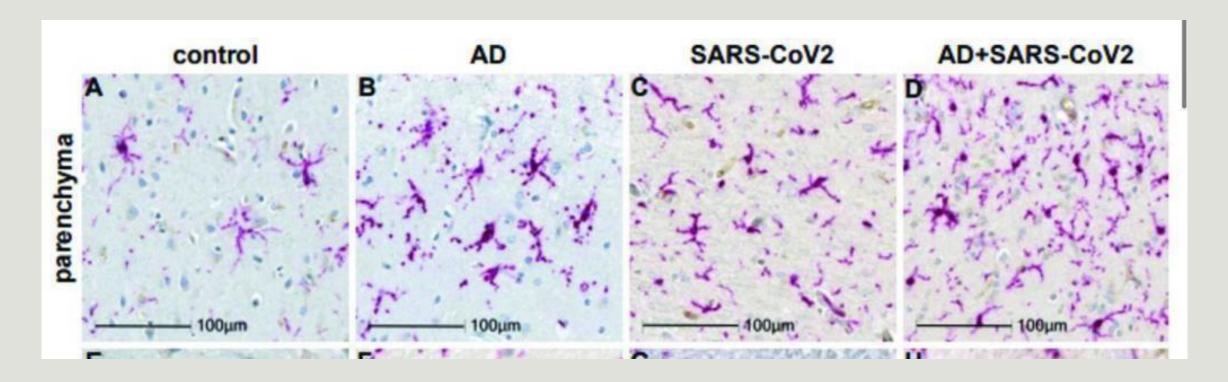
Elizabeth Griggs, Kyle Trageser ... Giulio Maria Pasinetti et al.

Jul 7, 2023 · https://doi.org/10.7554/eLife.86333 👌 💿





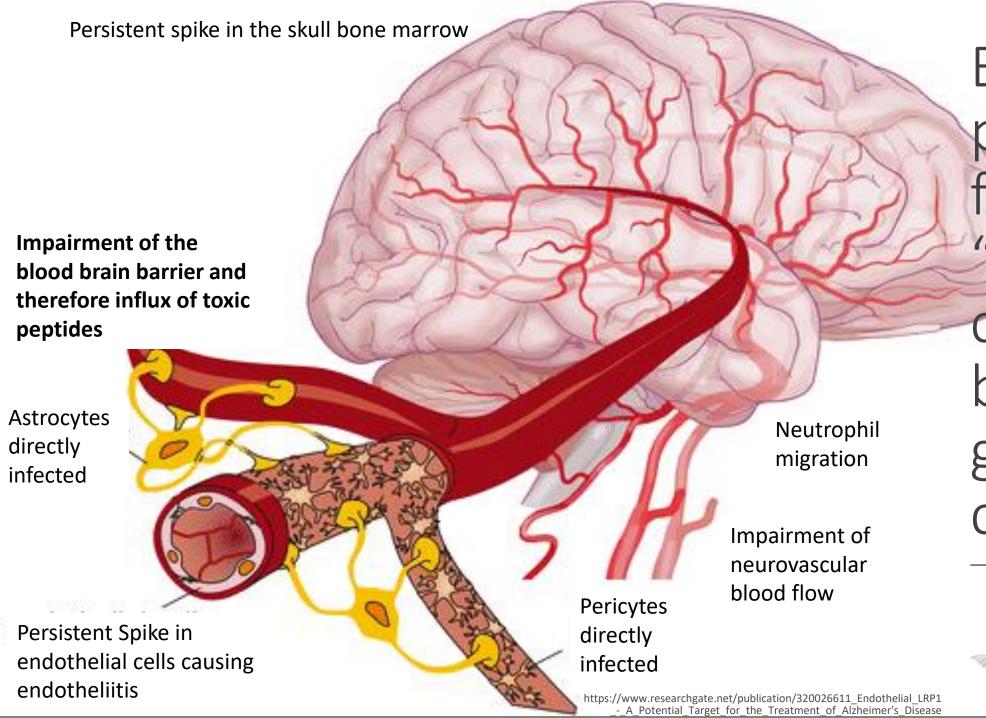
SARS-CoV-2 generates a similar neuroinflammatory environment in neurodegenerative disorders like AD.



Microgliosis and nodular lesions in neurological controls, SARS-CoV-2, Alzheimer's disease (AD), and SARS-CoV-2-infected AD individuals. Level of microglial activation.







Brain periphery forming a "pressure cooker" by blocking glymphatic cleansing





Inside the "pressure cooker"

Prionprotein misfolding

Amyloid-beta

Tau proteins

α-Synuclein

TDP 43

Lewy-Body Formation

> Filopodia formation

Microgliosis

Astrogliosis

Syncytia formation

Senescence

Demyelination

Ferroptosis

Dopaminergic **Neurons**

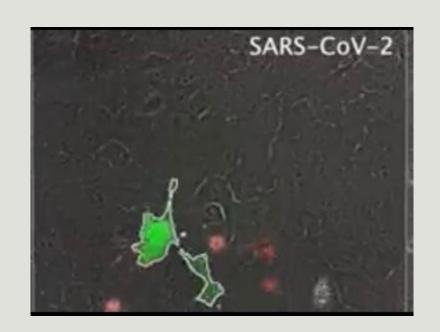
Mitochondrial dysfunction

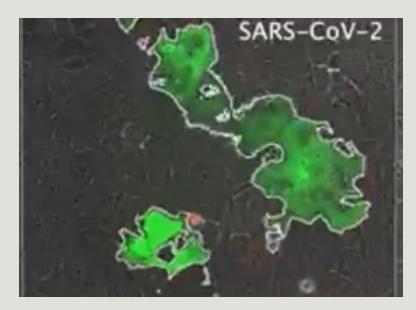


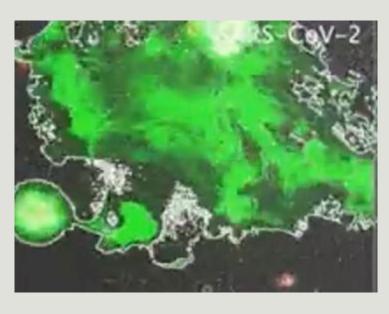


Alzheimer's disease, ALS, Parkinson's disease, Multiple Sklerosis, Prion diseases

Senescent cells / Syncytia are harboring Spike and or persistent viruses. Our protocols prevent the formation of cell fusion by Spike proteins which is important during detox as organ protection!

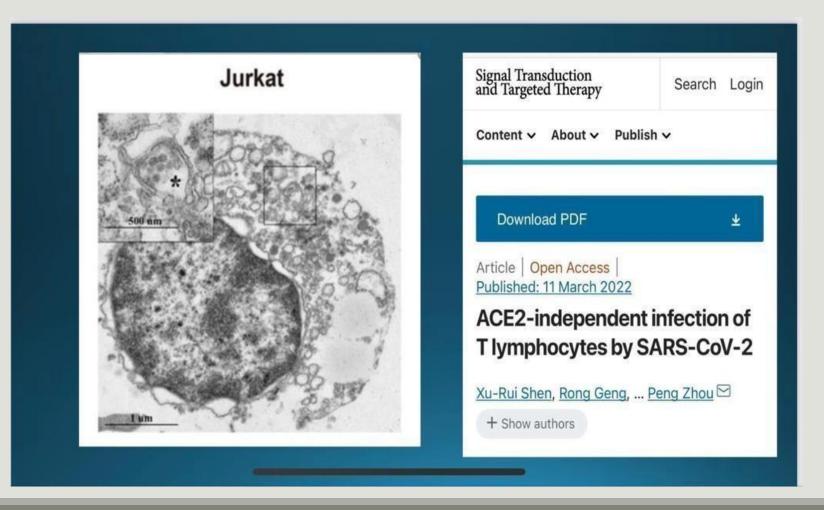










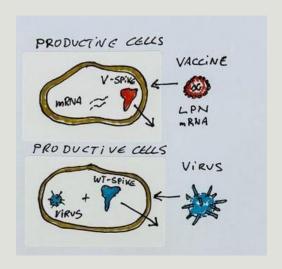


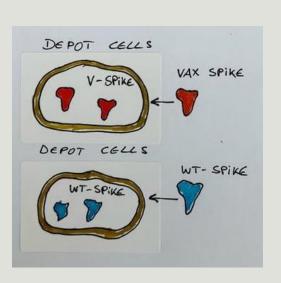
This will result in IMMUNODEFICIENCY!

This will impair the ability to control infections and cancer proliferation









The task:

In all cells the presence of virus and or spike is blocking autophagy and triggers a senescent state.

Phase 1

Override the blockade of the autophagy machanisms.

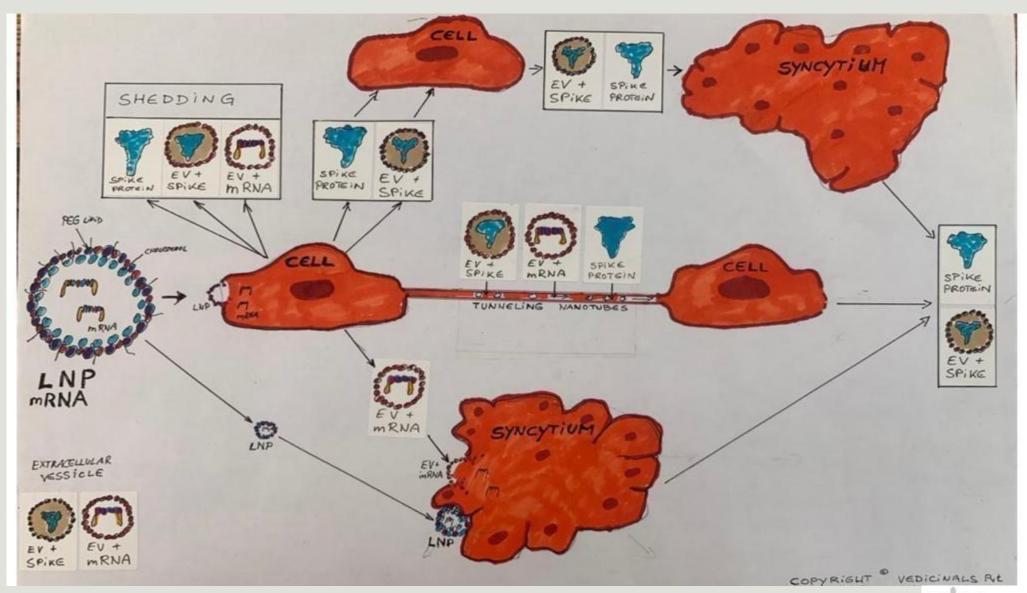
Phase2

Break up senescent cells. Stop viral spreading and new cell entry and reproduction. Stop spike binding to host cell receptors use chelators that can bind to spike.





VACCINE LNP ENTRY MECHANISMS AND SUBSEQUENT PROPAGATION OF SYNTHETIC SPIKE PROTEINS

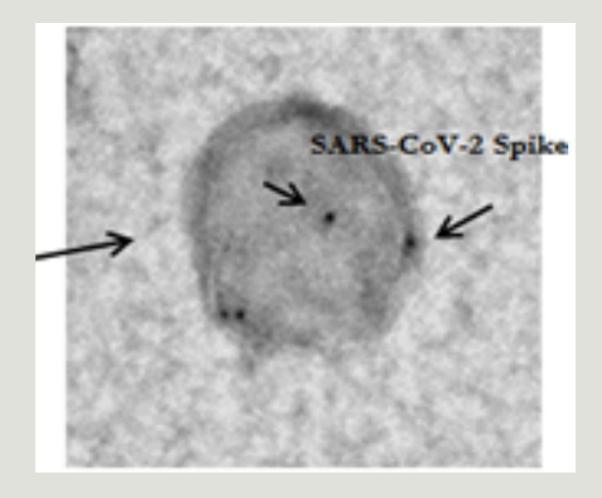






RESEARCH ARTICLE | NOVEMBER 15 2021





Transmission electron microscopy images of SARS-CoV-2 spike Ag on exosomes from control exosomes from control and vaccinated individuals. Arrows indicate SARS-CoV-2 spike-positive exosomes.







Review

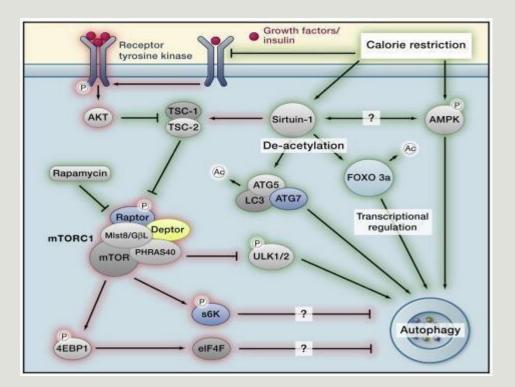
Cell Clearing Systems as Targets of Polyphenols in Viral Infections: Potential Implications for COVID-19 Pathogenesis

Fiona Limanaqi ¹, Carla Letizia Busceti ², Francesca Biagioni ², Gloria Lazzeri ¹, Maurizio Forte ², Sonia Schiavon ³, Sebastiano Sciarretta ^{2,3}, Giacomo Frati ^{2,3} and Francesco Fornai ^{1,2,*}

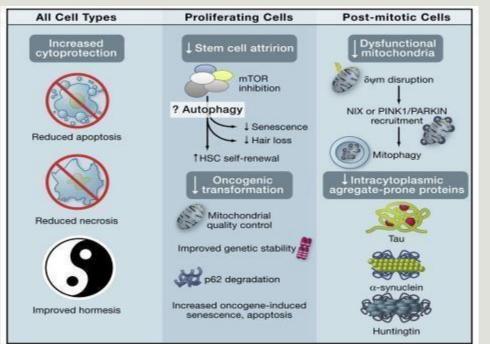
RAGE/TLR4

Figure 5. Summary of the potential mechanisms underlying SARS-CoV-2-induced alterations of Figure 5. Summary of the potential mechanisms underlying SARS-CoV-2-induced alterations of **autophagy** and (immuno-)proteasome.





The Regulation of Autophagy and Life Span





Autophagy may increase organismal fitness by inhibiting cell death, reducing oncogenic transformation, or increasing hormesis, both in quiescent and dividing cells (left). In addition, autophagy may contribute to life span extension through distinct mechanisms in post mitotic (middle) and proliferating cells (right). HCS, hematopoietic stem cell. Vedicinals-9 molecules are known promotors of autophagy which can save the injured cells.





Senolytics - Break up of senescent cells and / or syncytia cell conglomerates

Liberating persistent / intracellular spike and viruses



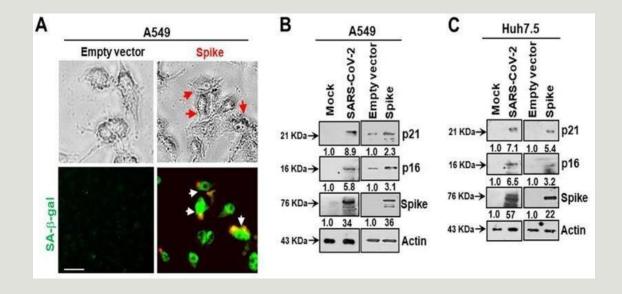




SARS-CoV-2 Spike Protein Induces Paracrine Senescence and Leukocyte Adhesion in Endothelial Cells

Authors: Keith Meyer D, Tapas Patra D,

Virus-infected or spiketransfected human epithelial cells exhibited an increase in senescence, with a release of senescence-associated secretory phenotype (SASP)related inflammatory molecules.



Virus-infected or spike proteinexpressing A549 (B) or Huh7.5 (C) cells also exhibited induction of senescence markers p21 and p16, as well as SARS-CoV-2 spike protein expression.

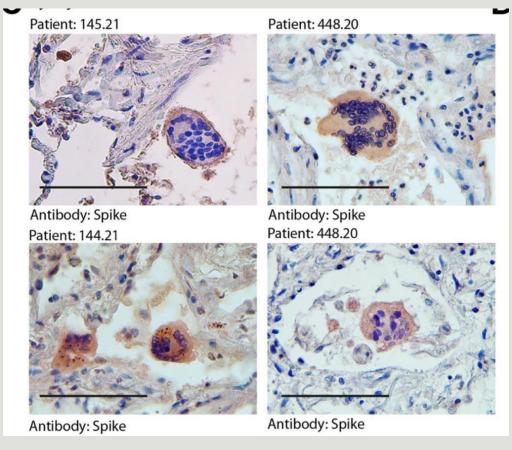




Persistent SARS-CoV-2 infection in patients seemingly recovered from COVID-19

Rossana Bussani, Lorena Zentilin, Ricardo Correa,

Despite apparent virological remission, lung pathology was similar to that observed in acute COVID-19 individuals, including microand macro-vascular thrombosis (67% of cases), vasculitis (24%), squamous metaplasia of the respiratory epithelium (30%), frequent cytological abnormalities and syncytia (67%), and the presence of dysmorphic features in the bronchial cartilage (44%).

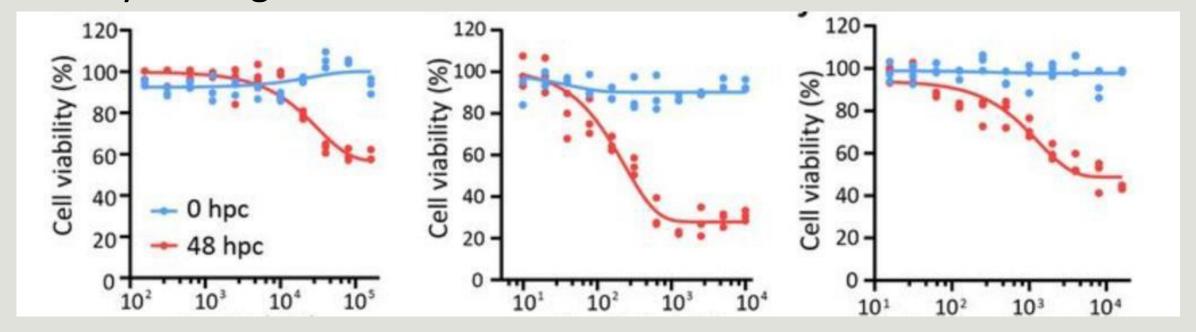


Spike-positive syncytial cells.





Furthermore, SARS-2-S syncytia could be selectively killed by senolytic drugs







Potential of green tea EGCG in neutralizing SARS-CoV-2 Omicron variant with greater tropism toward the upper respiratory tract

Zhichao Zhang a^1 , Meng Hao b^1 , Xiangchun Zhang c^1 , Yufeng He b^1 , Xiongsheng Chen $a^2 \ge a^2$, Ethan Will Taylor $a^1 \ge a^2$, Jinsong Zhang $a^1 \ge a^2$.



(EGCG) and its derivatives including theaflavin-3,3'-di-O-digallate (TFDG) strongly inactivated the conventional SARS-CoV-2 by **binding to the receptor binding domain (RBD)** of the S-protein.









Phenolic compounds disrupt spike-mediated receptor-binding and entry of SARS-CoV-2 pseudo-virions

Anna Goc, Conceptualization, Formal analysis, Investigation, Methodology, Project administration, Supervision, Validation, Visualization, Writing – original draft, Writing – review & editing,* Waldemar Sumera, Formal analysis, Investigation, Methodology, Validation,

Here, we provide experimental evidence that, among 56 tested polyphenols, including plant extracts, brazilin, theaflavin-3,3'-digallate, and curcumin displayed the highest binding with the receptor-binding domain of spike protein







Anti-Entry Activity of Natural Flavonoids against SARS-CoV-2 by Targeting Spike RBD

by

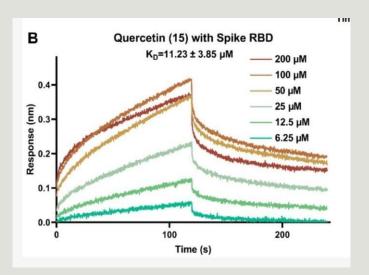
B Jie-Ru Meng

B Jiazheng Liu

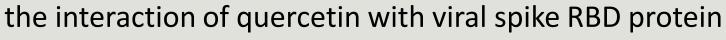
R Lu Fu

R

8 Tong Shu ², 8 Lingzhi Yang ²,







https://www.mdpi.com/1999-4915/15/1/160

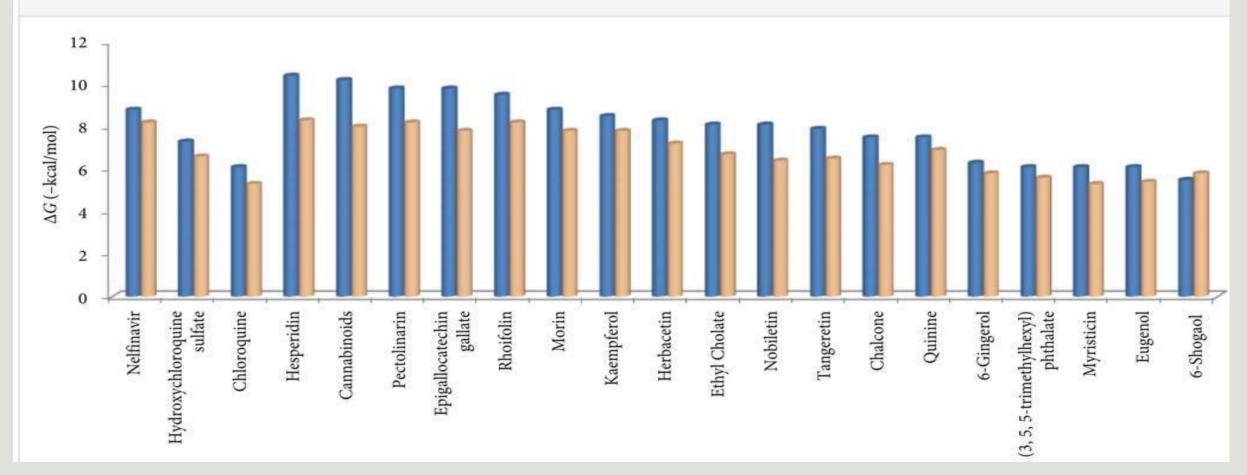






BLUE IS SPIKE BINDING!

Histogram showing the binding energy value ΔG (-kcal/mol) of S protein and M^{pro} with several inhibitor compound candidates.



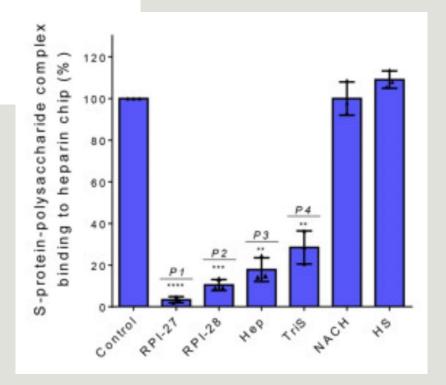




Sulfated polysaccharides effectively inhibit SARS-CoV-2 in vitro

Paul S. Kwon, #1,2 Hanseul Oh, #3
Seok-Joon Kwon, #1 Weihua Jin, 1,4

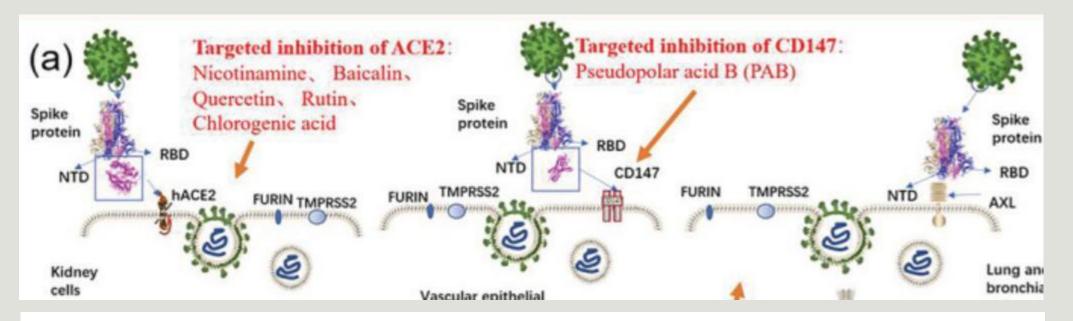
Our results reveal that specific sulfated polysaccharides bind tightly to the S-protein of SARS-CoV-2 in vitro, which suggests that they can act as decoys to interfere with S-protein binding to the heparan sulfate coreceptor in host tissues

















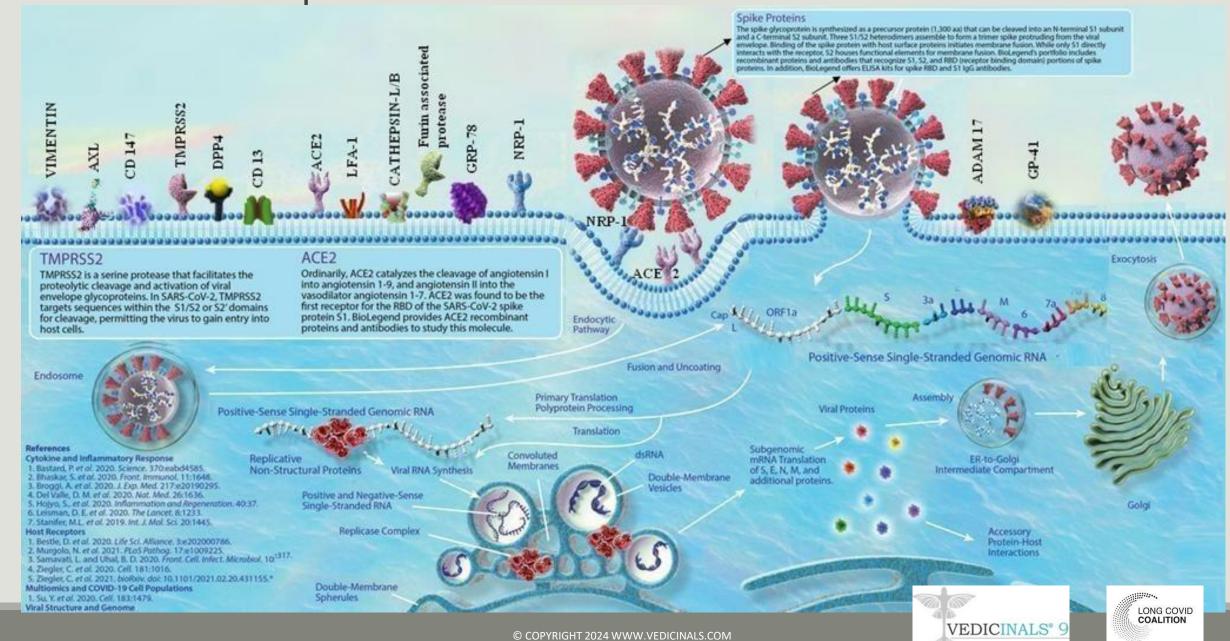
Promising natural products against SARS-CoV-2: Structure, function, and clinical trials

Yan Zhao, ¹ Shanshan Deng, ² Yujiao Bai, ² Jinlin Guo, ³ Guoyin Kai, ^{⊠ 4} Xinhe Huang, ^{⊠ 1} and Xu Jia ^{⊠ 2}



VEDICINALS® 9

Host cell receptors for Sars-CoV-2



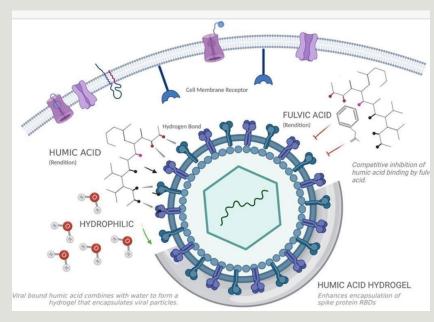
Gut detox from Spike and Viruses

Clinical review of humic acid as an antiviral: Leadup to translational applications in clinical humeomics



David C. Socol^{1,2*}

Humic acid binds to viral spike protein receptor binding domains (RBD) and inhibits viral fusion with target cell membrane receptors. The hydrophilic properties of the humic acid molecule attract water to form a hydrogel which encapsulates spike protein RBD and suspends the viral lifecycle. In the presence of fulvic acid, humic acid's potential to bind spike protein RBDs is impaired, which is the molecule's primary mechanism of action









Advanced Humeomics LLC, Beverly Hills, CA, United States

² SocolMD, Beverly Hills, CA, United States

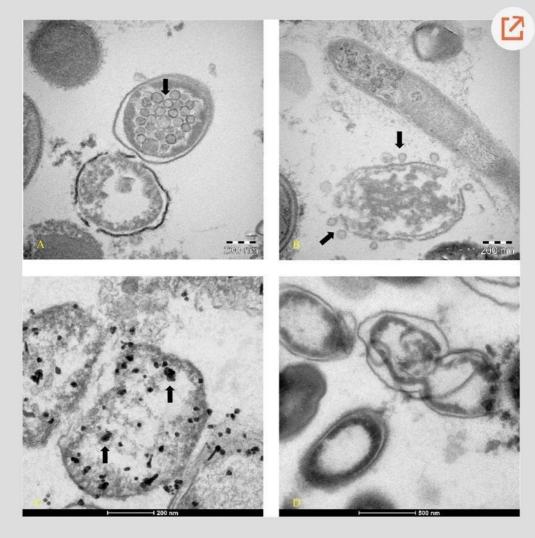


Figure 2. Transmission electron microscopy (postembedded immunogold).

SARS-CoV-2 has a dual mechanism: it infects human cells but first infects bacterial cells in our microbiome!!

Consequence – release of toxins that go into the blood stream, disturbs the CNS and gut-brain axis

The first report on detecting SARS-CoV-2 inside human fecal-oral bacteria: A case series on asymptomatic family members and a child with COVID-19 [version 1; peer review: 1 approved with reservations]

✓ Carlo Brogna (b)¹, Simone Cristoni², [...]
 Marina Prisco⁶, Marina Piscopo⁶ +

https://f1000research.com/articles/11-135

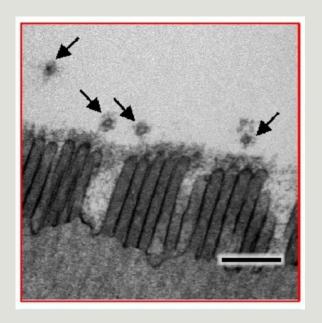


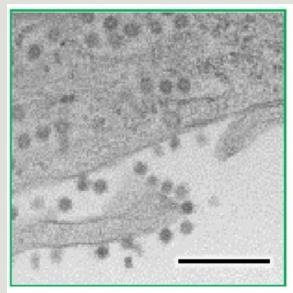


Article Open access | Published: 18 July 2023

Effective SARS-CoV-2 replication of monolayers of intestinal epithelial cells differentiated from human induced pluripotent stem cells

Shohei Minami, Naomi Matsumoto, ... Shintaro Sato ™





The transmission electron microscopic analysis of IEC#17 monolayers infected with SARS-CoV-2. IEC#17 cells were seeded on Transwell membranes





The spike protein of SARS-CoV-2 induces heme oxygenase-1: Pathophysiologic implications

Raman Deep Singh ^a, Michael A. Barry ^b,

Anthony J. Croatt ^a, Allan W. Ackerman ^a,

Joseph P. Grande ^c, Rosa M. Diaz ^d, Richard G. Vile ^d

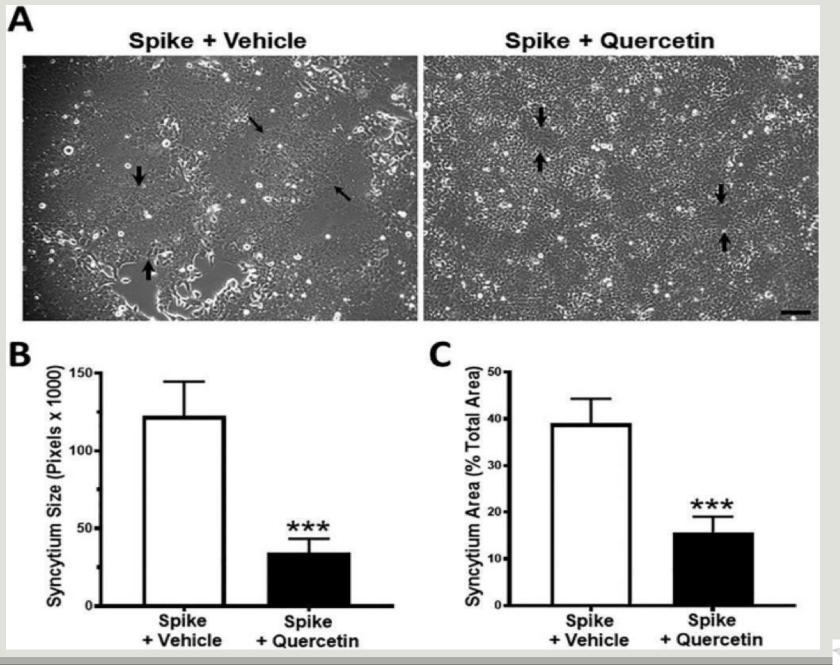
, Anupam Agarwal ^e, Karl A. Nath ^a

SARS-CoV-2 spike protein expression in kidney cells results in syncytia formation with cellular sloughing

 Quercetin reduces syncytia size and spike protein expression





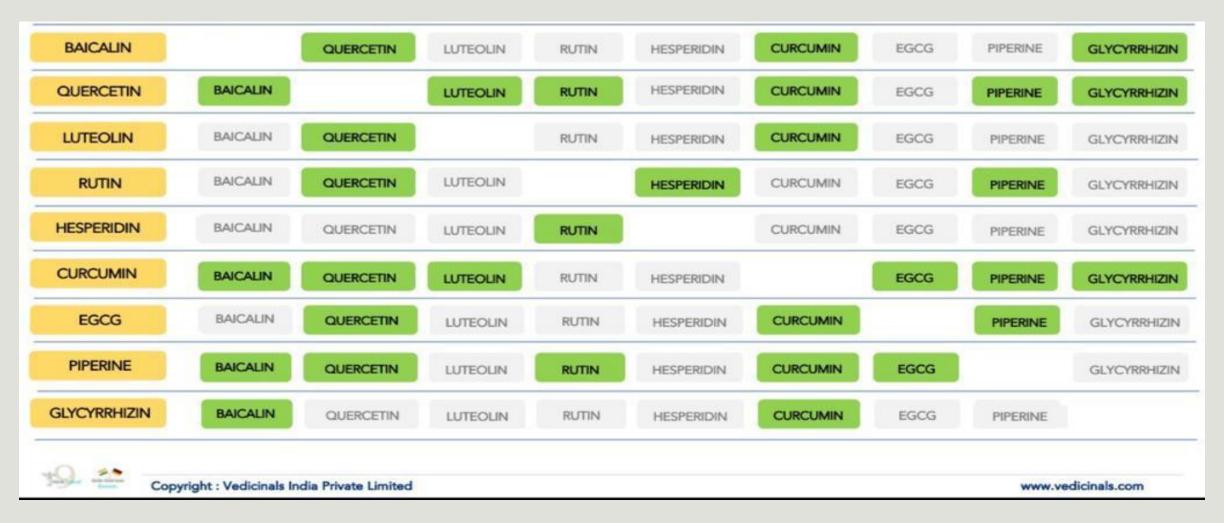








Synergies between Vedicinals compounds







Product Portfolio specialized for spike detox & persistent virus protocol





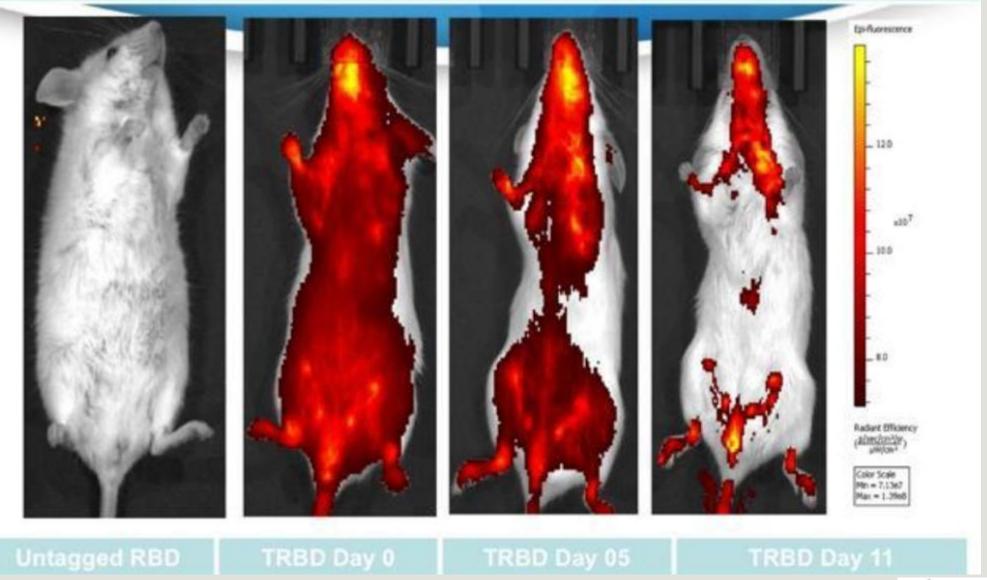








IR 780 Tagged RBD (TRDB) Spike Protein Injected in Animals (IP) Treated



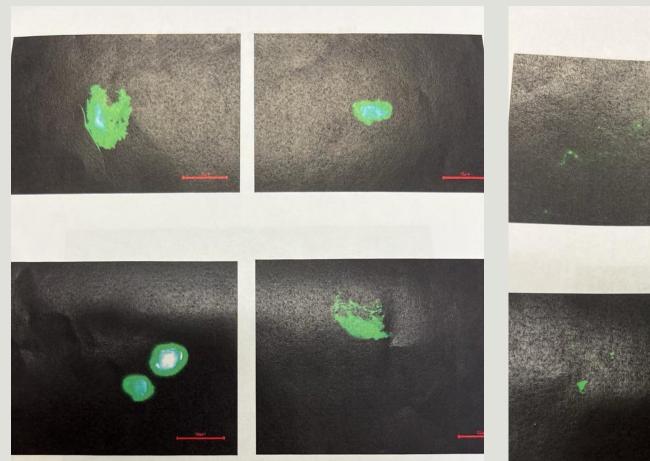




Biomarkers that are significantly elevated in our

- TGF-b1 patients
- D-dimer
- ABETA 42/40
- CV Biomarkers
- VEGF
- SARS Antibodies
- Histamine
- WBC
- Microthrombi (all our patients Grade 3/4-4/4)
- CRP
- ANA
- EBV

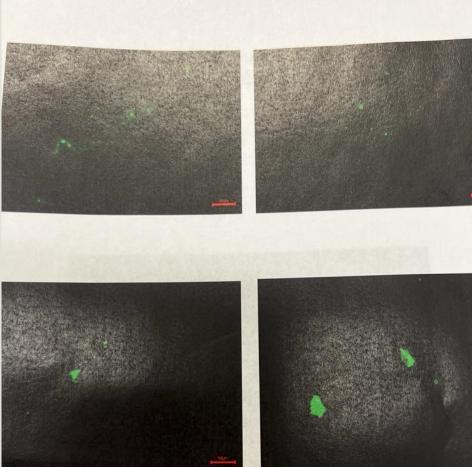




d staging of Amyloid Fibrin Microclots:

4 out of 4: Significant and Widespread

-clots come in all shapes and sizes. You may also see long, ing objects in your pictures. These are *Endothelial cast* and with *endothelial damage and inflammation*. This is a normal patients.

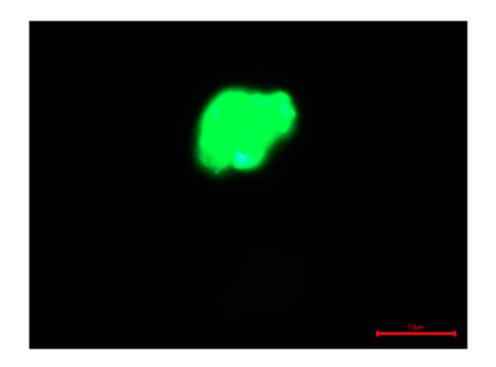


and staging of Amyloid Fibrin Microclots:

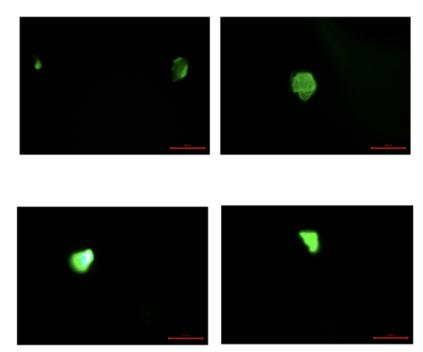
le 2 to 2.5 out of 4: Mild to Moderate Micro-clots.

ro-clots come in all shapes and sizes. You may also see loring objects in your pictures. These are **Endothelial cast** with **endothelial damage and inflammation**. This is a norm patients.

Microclot Results 3.5/4







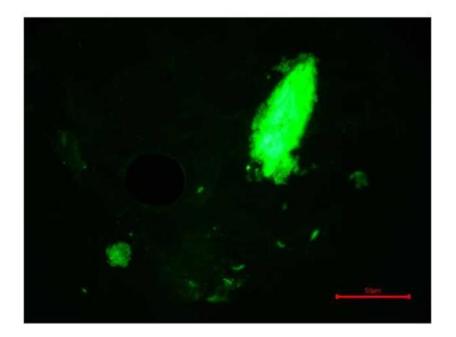
Comments and staging of Amyloid Fibrin Microclots:

Stage/Grade 3.5 out of 4: Moderate and Widespread

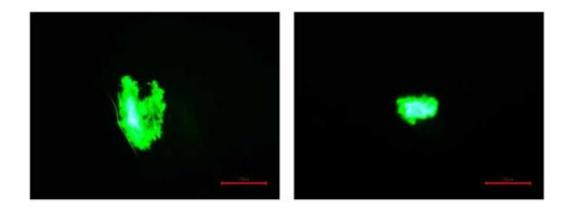
Note: Micro-clots come in all shapes and sizes. You may also see long, string-like appearing objects in your pictures. These are **Endothelial cast** and are associated with **endothelial damage and inflammation**. This is a normal finding for long-COVID patients.

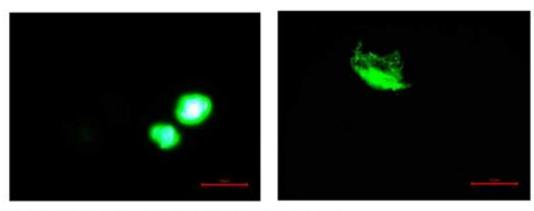


Microclot Results 4/4









Comments and staging of Amyloid Fibrin Microclots:

Stage/Grade 4 out of 4: Significant and Widespread

Note: Micro-clots come in all shapes and sizes. You may also see long, string-like appearing objects in your pictures. These are **Endothelial cast** and are associated with **endothelial damage and inflammation**. This is a normal finding for long-COVID patients.



Patient L History

- Male 66, Long Haul Covid, Suffering from:
 - Chronic fatigue a/w exercise intolerance
 - Dysautonomia/spontaneous tachycardia
 - Horrible brain fog



Patient L Before Treatment Biomarkers

- Out of Range:
 - o TGF-B1
 - SARS CoV2 Antibodies
 - o ABETA 42/40 Ratio
 - ANA Screen
 - EBV VCA IGM (U/mL)

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.50 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- TFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ABETA 42/40 Ratio: > OR = 0.160
- HS CRP: <1.0
- ANA: Negative
- EBV VCA IGM: <36 is a negative interpretation

Patient L		
Biomarker	Before Levels	
WBC (thous/uL)	5.4	
D-Dimer (mcg/mL)	0.43	
Histamine (ng/mL)	<1.5	
TGF-B1 (pg/mL)	9520 (High)	
VEGF (pg/mL)	83	
SARS CoV2 Antibodies	12.44 (High)	
ABETA 42/40 RATIO	0.126 (Low)	
HS CRP	0.8	
ANA	1:40 (High) Positive	
EBV VCA IGM (U/mL)	46 (High)	



Patient L Treatment Plan

- Long Hauler Protocol x 8 Weeks:
 - Vedicinals
 - Vitamin D
 - Vitamin C
 - O Zinc L-Carnosine
 - Lumbroxym
 - o Melatonin
 - o Rhizo Health
 - o Trans-Resveratrol
 - o EZ Trek
 - o Fisten



Patient L After Treatment Biomarkers

- Decreased Biomarkers
 - D-Dimer
 - o TGF-B1
 - o VEGF
 - SARS CoV2 Antibodies
 - HS CRP
 - o ANA
 - EBV VCA IGM (U/mL)

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.5 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- HTFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ABETA 42/40 Ratio: > OR = 0.160
- HS CRP: <1.0
- ANA: Negative
- EBV VCA IGM: <36 is a negative interpretation

Patient L		
Biomarker	Before Levels	After Levels
WBC (thous/uL)	5.4	5.5
D-Dimer (mcg/mL)	0.43	0.25
Histamine (ng/mL)	<1.5	<1.5
TGF-B1 (pg/mL)	9520 (High)	<mark>1820</mark>
VEGF (pg/mL)	83	38
SARS CoV2 Antibodies	12.44 (High)	11.34 (High)
ABETA 42/40 RATIO	0.126 (Low)	0.174
HS CRP	0.8	0.5
ANA	1:40 (Positive)	Negative
EBV VCA IGM (U/mL)	46 (High)	<36



Patient L Post Treatment

- Patient states he is feeling great:
 - Little to no brain fog
 - Fatigue has improved greatly
 - o Able to focus more
 - Able to exercise and play basketball
 - No more shortness of breath
 - No more tachycardia



Patient E History

- Female 60, Long Hauler Covid, suffering from:
 - o Dizziness
 - Exercise intolerance
 - Tachycardia
 - Shortness of breath
 - Chronic fatigue
 - Brain fog
 - Joint pain and muscle aches



Patient E Before Treatment Biomarkers

Out of Range:

- D-Dimer
- TGF-B1
- VEGF
- SAR CoV2 Antibodies
- EBV VCA IGG

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.5 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- TFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ANA: Negative
- EBV VCA IGM: <36 is a negative interpretation
- EBV VCA IGG: : <18 is a negative interpretation

Patient E		
Biomarker	Before Levels	
WBC (thous/uL)	4.7	
D-Dimer (mcg/mL)	0.67 (High)	
Histamine (ng/mL)	<1.5	
TGF-B1 (pg/mL)	4140 (High)	
VEGF (pg/mL)	<31 (Low)	
SARS CoV2 Antibodies	>150 (High)	
ANA	Negative	
EBV VCA IGM (U/mL)	<36	
EBV VCA IGG (U/mL)	248 (High)	



Patient E Treatment

- Long Haul Protocol
 - Vedicinals
 - Vitamin D
 - Vitamin C
 - ZInc L-Carnosine
 - Lumbroxym
 - o Melatonin
 - o Rhizo
 - Trans-Resveratrol
 - o EZ Trek



Patient After Treatment Biomarkers

- Decreased Biomarker
 - o D-Dimer
 - o TGF-B1
 - EBV VCA IGG

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.5 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- TFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ANA: Negative
- EBV VCA IGM: <36 is a negative interpretation
- EBV VCA IGG:: <18 is a negative interpretation

Patient E		
Biomarker	Before Levels	After Levels
WBC (thous/uL)	4.7	5.2
D-Dimer (mcg/mL)	0.67 (High)	0.27
Histamine (ng/mL)	<1.5	<1.5
TGF-B1 (pg/mL)	4140 (High)	2380
VEGF (pg/mL)	<31 (Low)	<31 (Low)
SARS CoV2 Antibodies	>150 (High)	>150 (High)
ANA	Negative	Negative
EBV VCA IGM (U/mL)	<36	<36
EBV VCA IGG (U/mL)	248 (High)	235 (High)



Patient E Post Treatment

- Patient states she is feeling much better:
 - Brain fog and chronic fatigue significantly improved
 - Joint pain improved
 - Slowly exercising again
 - Shortness of breath improved



Patient S Lab Work Prior to Treatment

- Out of Range
 - Histamine
 - o TGF-B1
 - SARS CoV2 Antibodies
 - o HS CRP
 - EBV VCA IGG

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.5 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- TFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ABETA 42/40 Ratio: > OR = 0.160
- HS CRP: <1.0
- ANA: Negative
- EBV VCA IGG: : <18 is a negative interpretation

Patient S		
Biomarker	Before Levels	
WBC (thous/uL)	10.6	
D-Dimer (mcg/mL)	0.26	
Histamine (ng/mL)	21.6 (High)	
TGF-B1 (pg/mL)	6500 (High)	
VEGF (pg/mL)	76	
SARS CoV2 Antibodies	>150.00 (High)	
ABETA 42/40 RATIO	0.163	
HS CRP	36.10 (High)	
ANA	Negative	
EBV VCA IGG (U/mL)	36.10 (High)	



Patient S Lab Work Prior to Treatment

- Out of Range
 - Histamine
 - o TGF-B1
 - SARS CoV2 Antibodies
 - o HS CRP
 - EBV VCA IGG

- White Blood Cell Count: 3.8-10.8 Thous/uL
- D-Dimer: <0.5 mcg/mL
- Histamine: < OR = 1.8 ng/mL
- TFG-B1: 344-2382 pg/mL
- VEGF: 31-86 pg/mL
- SARS CoV2 Antibodies: (<1 Index)
- ABETA 42/40 Ratio: > OR = 0.160
- HS CRP: <1.0
- ANA: Negative
- EBV VCA IGG: : <18 is a negative interpretation

Patient S		
Biomarker	Before Levels	
WBC (thous/uL)	10.6	
D-Dimer (mcg/mL)	0.26	
Histamine (ng/mL)	21.6 (High)	
TGF-B1 (pg/mL)	6500 (High)	
VEGF (pg/mL)	76	
SARS CoV2 Antibodies	>150.00 (High)	
ABETA 42/40 RATIO	0.163	
HS CRP	36.10 (High)	
ANA	Negative	
EBV VCA IGG (U/mL)	36.10 (High)	

