**COVID-Coag** 4-28-21



# COVID-Coag What is "Immuno-thrombo-inflammation?"

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The Fritsma Factor, Your Interactive Hemostasis Resource www.fritsmafactor.com ~ george@fritsmafactor.com

David L. McGlasson, MS MLS

Clot Club, by DiaPharma, Inc. [In absentia, with regrets]

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Bottom Line at the Start [BLAST] Keep it Short and Simple [KISS—no hope] Immuno-thrombo-inflammation

pathophysiology with disease Discusses possible COVID vaccine

Aligns COVID laboratory results and

laboratory assays results.

adverse events.

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## 32 YO Biker With Shortness of Breath



A motorcyclist was at the Sturgis, SD 8-8-2020 rally. He appears at his local ED experiencing a sore throat, cough, fever, and shortness of breath, O<sub>2</sub> sat. 88%. The physician suspects COVID-19. A lab scientist collects a nasopharyngeal specimen and employs a Cepheid GeneXpert® POC RT-PCR, which detects the SARS-Cov-2 virus in less than one hour. The patient is placed in isolation.

Smithgal, MC, Dowlatshahi M, Spitalnik SL, Hod EA, Rai AJ. Types of assays for SARS-CoV-2 testing: a review. Lab Med 2020 DOI: 10.1093/labmed/lmaa039

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The participant...

## COVID-19 Symptoms

- · Coronaviridae varieties: respiratory infections; the common cold
- COVID-19: the third Coronaviridae epidemic since 2000
  - Severe acute respiratory syndrome [SARS] 2002–3.
  - Middle East respiratory syndrome [MERS] 2012
- SARS and MERS: Interstitial pneumonia with progression to acute respiratory distress syndrome
- COVID: multisystem disorder: hyperimmune, inflammatory, progressing to profound hemostatic disturbance
  - High rates of pulmonary embolism and deep venous thrombosis
  - Smaller component of stroke, myocardial infarction
  - Mortality greatest over 60, males, hypertension, diabetes, obesity, cancer, pulmonary, renal, cardiovascular, liver, and neurological disorders

Lippi G, Sanchis-Gomar F, Favaloro EJ, Lavie CJ, Henry BM. Coronavirus disease 2019-associated coagulopathy. Mayo Clin Proc 2021; 96:203-17

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### Symptoms Specified in Vaccine Clinical Trials to Define COVID Infection

- Pfizer BioNTec. Moderna. Astra Zeneca. J&J/Janssen.
- No minimum duration: fever, dyspnea, shortness of breath
- · Minimum 2 days' duration: Chills, cough, fatigue, muscle or body aches, headache, loss of taste and/or smell, sore throat, congestion, runny nose, nausea, vomiting, diarrhea
- Confirm by reverse transcriptase polymerase chain reaction [RT-PCR] SARS-Cov2

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## Meta-analysis of 17,052 Patients

 Severe [3664]: respiratory frequency >30/min O<sub>2</sub> sat 93% at rest; artery PP of O<sub>2</sub>/inspired O<sub>2</sub> [PaO<sub>2</sub>/PiO<sub>2</sub>] 300 mmHg

· Mild [13,388]: Hx of exposure, fever, pneumonia

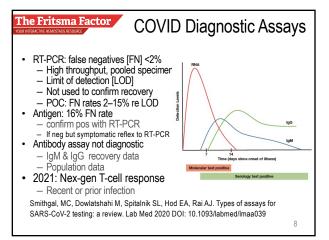


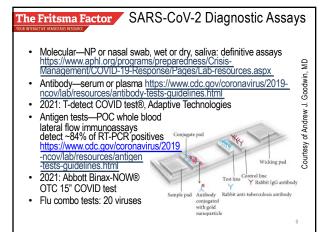
Chaudhary R, Garg J, Houghton DE. Thrombo-inflammatory biomarkers in COVID-19: systematic review and meta-analysis of 17,052 patients. Mayo Clin Proc pre-press 4-21

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Condition	Severe	Mild
Death	32%	1%
Age	64 Y	53 Y
Male sex	65%	54%
Hypertension	45%	23%
Diabetes	28%	16%
Cardiac/stroke	23%	8%
Chronic kidney disease	9%	3%
Chronic liver disease	5%	4% [NS]
Malignancy	10%	7%
COPD	9%	3%





COVID Progression: What is

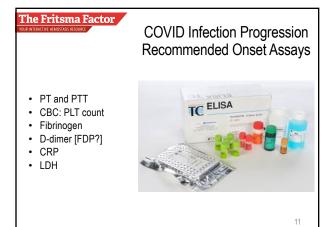
# COVID Progression: What is it?

- A hemostasis abnormality associates with COVID infection progression, but what is the abnormality? VTE? DIC? LAC? HIT? TMA? TTP? ITP? PLT activation? Sepsis? Hypofibrinolysis?
- It has earned a name, "immunothrombo-inflammation.
- We apply routine and specialized laboratory assay results to attempt a model and to develop antithrombotic therapy.



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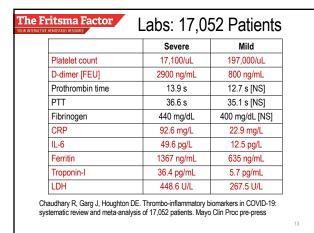
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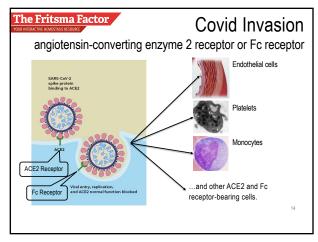


The Fritsma Factor 32 YO Biker with SOB Mean coagulation labs of 94 COVID patients Vs 40 healthy controls **Patients** Assay Control p-value Patient Onset Labs 29.01 s 28.65 s 0.518 PTT D-dimer: 854 ng/mL PT 12.43 s 12.08 s 0.678 FIB: 405 mg/dL П 18.00 s 18.34 s 0.137 PT: 12.5 s PTT: 31 s D-dimer 1036 ng/mL 260 ng/mL <0.001 PLT: 170,000/uL Fibrinogen 502 mg/dL 290 mg/dL < 0.001 AT 85.46% 98.83% <0.001 Han H et al. Clin Chem Lab Med. 2020;58:1116-20. Courtesy of Andrew J. Goodwin, MD.

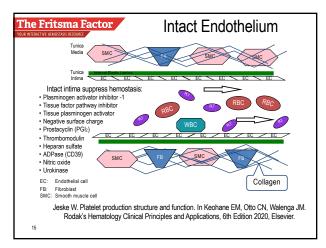
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DAMPs, PAMPs,
NETs, & APLs

• Damage-associated molecular patterns [DAMPs] or pathogenassociated molecular patterns [PAMPs]

— Array of pro-oxidant cytosolic proteins & nucleic acids released from
damaged or pathogen-invaded cells.

— Activate inflammatory cells such as T-cells and macrophages

• PMNs secrete neutrophil extracellular traps [NETs]

— Nucleic acids and cytoplasmic granule enzymes

— "Lasso" and destroy pathogens and foreign materials extracellularly

• Anti-phospholipid antibodies [APLs, lupus anticoagulants, LACs]
that interact with endothelium [some studies claim 60% of
patients]

Libby P, Luscher T. COVID-19 is, in the end, an endothelial disease. Euro Heart J
2020; 41: 3038–44.

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# The Fritsma Factor Activated Endothelium Properties Impaired barrier function exposes TF and pro-

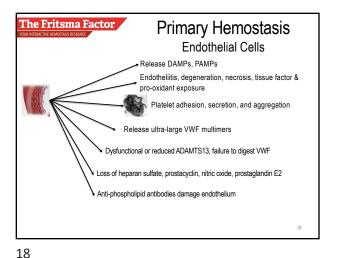
- oxidants

   Release ultra-large von Willebrand factor
- [ULVWF]
- Depleted ADAMTS13 [VWF-cleaving protease]
- Activates lymphocyte nuclear factor  $\kappa B$  [NFKB]
- Induces T cells and monocytes to secrete CD 40 ligand [B-cell activation], PAI-1, PMN adhesion molecules and NETs, inflammatory cytokines TNF, IL-1, IL-6, IL-8, IL-12, transforming factor β, IFN-γ, and more inflammatory cytokines



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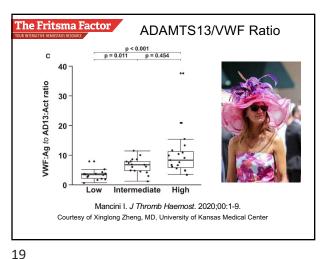
 $Libby\ P,\ Luscher\ T.\ COVID-19\ is,\ in\ the\ end,\ an\ endothelial\ disease.\ Euro\ Heart\ J\ 2020;\ 41:\ 3038-44.$ 

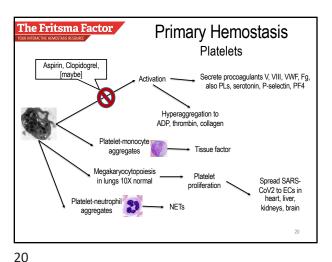


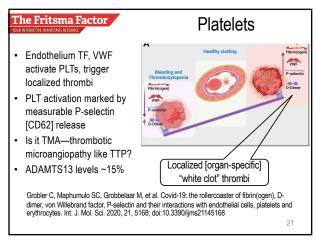
george@fritsmafactor.com

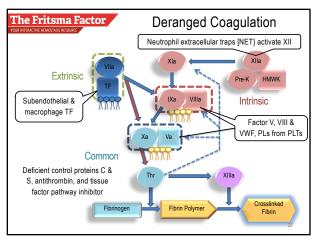
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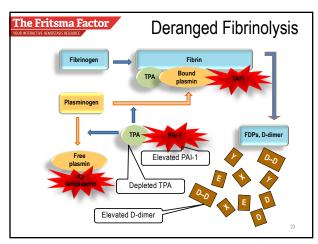


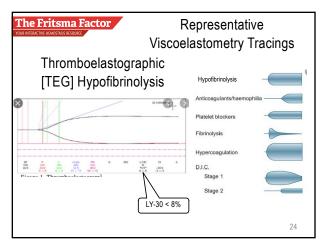






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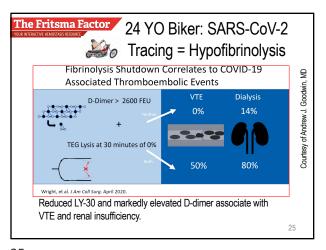




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The Fritsma Factor **D-dimer Correlated to Severity** Mean Onset Intermediate Severe Assay

19,110 ng D-dimer [ng/mL] 260 ng/mL 2140 ng 20,040 ng FDP [mg/L] 1.55 mg/L 7.93 mg 60.01 mg 69.15 mg

Courtesy of Andrew J. Goodwin, MD

- · FDP used in China-automated, not in North America
- · D-dimer units [DDUs]
  - Normal limit per manufacturer <240 ng/mL or 0.24 mg/L or ug/mL
- Fibrinogen equivalent units [FEUs]
  - Normal limit per manufacturer <500 ng/mL or 0.5 mg/L or ug/mL</li>
- · Most research reports fail to specify FEUs or DDUs; and fail to specify units
- Favaloro EJ, Thachil J. Reporting of D-dimer data in COVID-19: some confusion and potential for misinformation. Clin Chem Lab Med 2020; 58: 1191-9.
- Han H et al. Clin Chem Lab Med. 2020;58:1116-20.

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# The D-dimer Assay

- · "Promiscuous"
- Screening is ineffective
  - Baseline necessary when there are indications
  - R/O DVT or PE when there is low pre-test probability
- · Normal limits are age-adjusted
  - If over 50, multiply age by 10 ng/mL FEUs
  - For example, age 60 = <600 ng/mL FEUs, age 70 = <700 ng/mL FEUs</li>
- · Marked elevation implies DIC
- Tang: "Recent studies described that severe COVID-19 is commonly complicated with coagulopathy, DIC may exist in the majority of deaths."

Tang N, Bai H, Chen X, et al. Anticoagulant treatment is associated with decreased mortality in severe coronavirus disease. JTH 2020 as doi: 10.1111/JTH.14817

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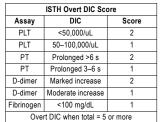


- Mild thrombocytopenia
- No schistocytes
- PT prolonged to 16.2 s
- PTT unaffected Coag factors normal No bleeding

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- Thrombosis
- D-dimer: 4300 ng/uL

#### 32 YO Biker: Is it DIC?



lha et al. J. Thromb Haemost, 2019:17:1989-94

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- Tpenia: 90,000/uL
- PT prolonged—16.2s
- PTT unaffected
- Coag factors normal
- Thrombosis D-dimer 4300 ng/uL
- Is it sepsis?

lha et al .l Thromb Haemost. 2019;17:1989-94

#### 32 YO Biker, 5 Days: Is it SIC?

Assay	Score	ISTH DIC	SIC
PLT	2	<50,000/uL	<100,000/uL
PLT	1	50-100,000/uL	100-150,000/uL
PT/PT Ratio	2	> 6 s	> 1.4
PT/PT Ratio	1	3–6 s	1.2-1.4
D-dimer	2	Marked increase	-
D-dimer	1	Moderate increase	-
Fibrinogen	1	< 100 mg/dL	-
SOFA Score	2	-	2 or more
SOFA Score	1	-	1
Overt DIC:	when tot	al = 5 or more	4 or more

- SIC = sepsis-induced intravascular coagulation SOFA = sequential organ failure assessment;
- respiratory, cardiovascular, renal, hepatic

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#### Sepsis: Infection Response



Endothelium normally releases cytokines to fight infection. Sepsis occurs when cytokine response is out of balance, damaging multiple organs. If sepsis progresses to shock, BP drops.

Organ failure accompanies severe sepsis. Livedo reticularis, anuria, thrombocytopenia, dyspnea, abnormal heart rhythm, chills, weakness, loss of consciousness. Treat with fluids, antibiotics, norepinephrine and vasopressin. Target a mean arterial pressure of 65 mm Hg.

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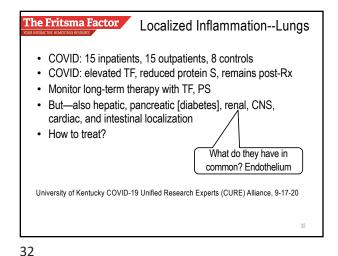
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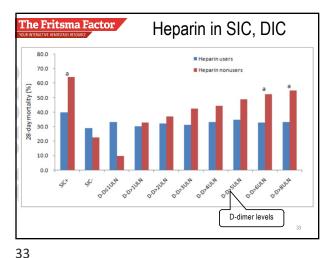
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	STASIS RESOURCE	Patient	Reference Interval
	Assay		
	PMNs	750/uL	17–7500/uL
<b>30</b>	Pelgeroid	10%	
_	Lymphs	900/uL	1000-11500/uL
Courtesy of Dr. Susan LeClair and Dave McGlasson	Reactive	15%	
san	PLTs	65,000/uL	150-450,000/uL
Sin Sin	CRP	1425 ug/dL	< 820 ug/dL
ř.	Serum ferritin	850 ng/mL	40-400 ng/mL
sy of	IL-6	28.2 pg/L	12.5 pg/L
anc	Procalcitonin	1.6 ug/L	< 0.5 ug/L
Š	Anti β-2-glycoprotein 1	Positive	Negative
	Antithrombin, PC, PS	< 60%	> 60%

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LMWH Anti-inflammatory The Fritsma Factor Animal studies: Heparin reduces pulmonary microvascular EC barrier dysfunction via microtubule stabilization in a sepsis mouse model and in-vitro human ECs Retrospective study in COVID-19 patients: LMWH reduced nuclear factor κB reduces IL-6 & increases lymphocyte % Heparin reduces expression of VWF and fibrinogen in animal lung injury models Tang et al. J Thromb Haemost. 2020 Mu et al. Respir Res. 2008 Li et al. Zhonghua 2019. 34

The Fritsma Factor Vaccine Adverse Events CDC reporting site: <u>VAERS</u> accessed May 30

— Self-reported or physician-reported adverse events No attempt to establish causation, collect all data
 Breakthrough Covid infection: 2/100,000 in 46 states, all vaccines
 Anaphylaxis 2–5/m; all vaccines, 30 m after injection Transient myo- or peri-carditis in teens, incidence not established Moderna and Pfizer 4863 deaths/285 m doses, 1.7/100,000 causation not established

Vaccine-induced thrombocytopenia with thrombosis [VITT] Cerebral venous sinus thrombosis [CVST]
Abdominal splanchnic vein thrombosis
In US, only J&J vaccine In Europe, AZ vaccine Thrombotic thrombocytopenic purpura: 1/million Same incidence as unselected baseline 35

The Fritsma Factor Vaccine-Induced Thrombosis with Thrombocytopenia [VITT] · Events documented after "replication-defective" adenovirus vector vaccines from J&J, AZ Adverse events documented 7–14 days after exposure · Autoantibody with PF4-specificity resembles heparininduced thrombocytopenia with thrombosis [HIT] Ab Muir KL, Kallam A, Koepsell SA, Gundabolu K. Thrombotic thrombocytopenia after Ad26.COV2.S vaccination [Letter]. NEJM 2021, DOI: 10.1056/NEJMc2105869

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J&J VITT



- CDC: 10.2 m doses as of 5-24-21
  - 32 females got VITT, median age 37, report rate 3.7/million
    - 7-14 days after injection
  - Females VITT 50+ YO, report rate 1/million, no males
  - Risk factors: obese, OC, hypothyroid, hypertension
- CVST, DVT/PE, abdominal vein, acute myocardial infarction
- · Strongly positive in EIA PF4/polyvinyl sulfonate fixed target, but not in functional HIT assays such as serotonin release
- Theoretic adverse response to heparin Rx as though it is HIT, use alternate such as argatroban, fondaparinux, DOACs

Strieff MB. Pathogenesis and management of thrombosis with thrombocytopenia syndrome. Presented 4-23-21 at CDC conference

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Astra-Zeneca VITT Incidence April 2021

- · EU: 18 fatalities
  - 62 cases of cerebral venous sinus thrombosis [CVST] with thrombocytopenia
  - 24 cases of abdominal vein thrombosis with thrombocytopenia
  - Most in females <60 years of age
- Variant data collection, cannot exclude age/gender as risks
- UK: 20.2 million doses, 19 fatalities
- Incidence ~4/million, slightly higher in younger age
- 79 cases of TTS
- 44 cases of CVST [14 fatalities] 35 cases of other clots [DVT/PE, 5 fatalities]
- 51 [65%] of cases were female



Schultz NH, Sorvall IH, Michelson AE, et al. Thrombosis and thrombocytopenia after CHAdOx1 nCoV19 vaccination [brief report, Norway, NEJM 2021; DOI 10.1056 NEJM0a2104882. Greinecher A, Thiele T, Warkentin TEet al. Thrombotic thrombocytopenia after CHAdOx1 nCoV19 vaccination [Germany, NEJM 2021; DOI 10.1056NEJM0a2104840.

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The Fritsma Factor J&J: 3.7/million

**Risk of Blood Clots COVID** AstraZeneca **Birth Control Pill** Infection Vaccine 500 - 1200 cases in 1,000,000 1,000,000 1,000,000 1.000.000 **Smokers** Cases 0.0004% 0.05% to 0.12% 0.18% 16.5%

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ISTH Vaccine-induced Immune Thrombocytopenic Purpura [VITT] Guidance

- Watch for fever, chills, nausea, headache, vision change, seizures, leg pain, chest pain, SOB, abdominal pain
- Clots on imaging and PLT count <150,000/uL - Mean nadir 27,000/uL
- PT, PTT, D-dimer, Fg, immunoassay for anti-PF4 AB
- Ivlg infusion, steroids, PLEX, Fg concentrate, no PLT concentrate
- Non-heparin AC: fondaparinux, argatroban, DOACs - Heparin OK if anti-PF4-negative

ISTH interim guidance for the diagnosis and treatment on vaccine-induced immune thrombotic thrombocytopenia. 4-21-21

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# Ongoing COVID-Coag Challenges

- · Is COVID-coagulopathy DIC? Is it SIC?
- Is it platelet-triggered thrombotic microangiopathy
- Control platelets with antiplatelet drugs such as aspirin or clopidogrel?
- Is monitoring UFH & LMWH required in COVID-coagulopathy? What is the target therapeutic level of
- anticoagulation? Are vaccines the cause of VITT?
- Report adverse events to Vaccine Adverse Event Reporting System [VAERS] at vars.hhs.gov, 800-822-7967, info@vaers.org



<u> The Fritsma Factor</u>

Bottom Line at the End [BLEAT] Kept it Short and Simple [KISS—nope]

The participant...

- Proposed the pathophysiology and comorbidities associated with COVID.
- Applied relevant COVID progression laboratory assays results.
- Aligned COVID laboratory results and pathophysiology with disease
- Discussed possible COVID vaccine adverse events.





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