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Prolific Activation of a Network Immune-Inflammatory Crisis [PANIC]

Part II. High-Dose Methotrexate with Leucovorin Rescue for Severe COVID-19: An Immune Stabilization Strategy for SARS-CoV-2 Induced ‘PANIC’ Attack

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Abstract

Here, in Part II of a duology on the characterization and potential treatment for COVID-19, we characterize the application of an innovative treatment regimen for the prevention of the transition from mild to severe COVID-19, as well as detail an intensive immunotherapy intervention hypothesis.

We propose as a putative randomized controlled trial that high-dose methotrexate with leucovorin (HDMTX-LR) rescue can abolish ‘PANIC’, thereby ‘left-shifting’ severe COVID-19 patients to the group majority of those infected with the SARS-CoV-2 virus, who are designated as having mild, even asymptomatic, disease. HDMTX-LR is endowed with broadly pleiotropic properties and is a repurposed, generic, inexpensive, and widely available agent which can be administered early in the course of severe COVID-19 thus rescuing the critical and irreplaceable gas-exchange alveoli.

Further, we describe a preventative treatment intervention regimen for those designated as having mild to moderate COVID-19 disease, but who exhibit features which herald the
transition to the severe variant of this novel Coronavirus. Both of our proposed hypothesis-driven questions should be urgently subjected to rigorous assessment in the context of randomized controlled trials, in order to confirm or refute the contention that the approaches characterized herein, are in fact capable of exerting mitigating, if not abolishing, effects upon the SARS-CoV-2 triggering of the ‘PANIC Attack’. Confirmation of our immunotherapy hypothesis would have far-reaching ramifications for the current pandemic, along with yielding invaluable lessons from which we can more effectively prepare for the next challenge to global health.

**Hypothetical Strategies for Rescuing the Severely Affected COVID-19 Patient**

We submit that the best chance to rescue those afflicted with the severe variant of COVID-19 depends upon the decision to intervene with a corresponding assemblage of diverse immune modulatory agents, each directed toward a mechanistically distinctive component of the activated inflammatory network. Such a diverse combination of agents would require a synchrony of action in order to arrest the ‘irrationally exuberant’, poorly coordinated, and regulatory-deficient immune network in severe COVID-19 patients.

The prolific activation, and expansive constellation of coincidentally activated inflammatory cascades, confers potentially destructive consequences upon host tissue compartments which harbor the SARS-CoV-2 agent (i.e. bystander damage). In contrast to the application of a multi-component treatment strategy, the identification of a single agent endowed with pleiotropic mechanistic properties, and capable of reconstituting the normally coordinated immune-regulatory response characteristics to SARS-CoV-2, would represent a potential advance in our struggle to transition severely affected COVID-19 patients into the ‘mild’ phenotypic variant of this Coronavirus-mediated disorder.

**Survival of the ‘FOCUSED’**

The categorical designation of ‘mild COVID-19’ signifies a considerably more ‘focused’ activation of the immune network compared to severely affected patients. The majority of mildly infected patients demonstrate a short-lived flu-like illness with minimal long-lasting effects or deficits. Clinical manifestations, along with their magnitude and/or worsening, should
give pause to consider the prospect of acute and directed interventions, which may potentially serve to mitigate symptoms and reduce the risk of advancing to a more severe clinical state with an associated worsening prognosis. Such issues can involve recalcitrant cough, poor mucus clearance, perception of a diminished negative inspiratory force of breathing, and risk of secondary upper respiratory tract infection.

Treating a recalcitrant cough may be as simple as offering dextromethorphan, but the addition of quinidine can reduce the former’s metabolism and thereby prolong its action. A dose of 20mg/10mg respectively is the composition of Nuedexta, which is FDA-approved for pseudobulbar affect, but can be of great utility in the circumstances of recalcitrant cough.

An expectorant, such as guaifenesin, may aid in mucus clearance, whereas a bronchodilator may improve perception of a diminished negative inspiratory force of breathing. Smooth muscle relaxation at the terminal bronchioles and activation of the beta-2 adrenergic receptor on lung mononuclear cells may skew the immune milieu from a TH1 to a TH2 anti-inflammatory phenotype. The intracellular biomarker, secondary to transmembrane signaling at the beta-2 adrenergic receptor, is an escalation of cAMP an intracellular biomarker for this effect [1-4].

A bronchodilator in conjunction with inhalational steroids will likely reduce peripheral airway inflammation. Antibiotics, such as azithromycin, will mitigate risk of secondary upper respiratory tract infection.

Interventions that help detect worsening, such as home finger pulse oximetry, may be useful. This multi-faceted interventional approach may prevent the transformation of the mild phenotype into the severe disease course and its correspondingly worse prognosis [Vignette 1].

The recovery phase for mildly to moderately affected COVID-19 patients appears to be temporally linked with the emergence of detectable SARS-CoV-2 antibodies; although their relevance to recovery (i.e. are they endowed with viral-neutralization effects or even protection in the absence of ‘viral neutralization’ in vitro) and the establishment of a durable remission is currently a matter of vigorous debate [5]. Notwithstanding the controversy surrounding the therapeutic relevance of SARS-CoV-2 associated antibodies, the donation of plasma derived
from patients dichotomized to the mildly affected cohort, and administered to more severely affected patients, can exhibit impressive remission-exacting effects [5-7]. This salient observation, in conjunction with inadequate evidence to confirm the relevance of SARS-CoV-2 antibodies, makes it is entirely feasible to assume that one, or even a variety of plasma-derived soluble factors from recovered COVID-19 patients, may serve to promote recovery in severely affected patients.

Those dichotomized into the ‘severe COVID-19’ subgroup, because of their ‘PANIC Attack’ response phenotype, fail to organize the coordinated neutralization of the SARS-CoV-2 virus, in contradistinction to mildly to moderately affected patients who do in fact eventually orchestrate effective clearance of the COVID-19 pathoetiologic agent. Alternately, the severely affected cohort of patients will exhibit evidence of the coincident activation of both the innate and adaptive limbs of the inflammatory response network, which is in keeping with our definition of the ‘prolific activation of a network-immune-inflammatory crisis’ (i.e. PANIC), and provides an attractive explanation for why the most severely affected COVID-19 patients harbor stereotyped patterns of irreversible tissue destruction, including but not limited to, the obliteration of the life-sustaining process of gas exchange within the terminals of the bronchopulmonary tree.

**Anti-Cytokine Monoclonal Antibody Therapy**

The anti-IL-6 monoclonal antibody, tocilizumab, has been shown to provide a mitigating effect upon the CRS, in addition to reducing fevers, CRP levels, and improving the chest CT demonstrated lung abnormalities (e.g. the ground glass opacifications) in COVID-19 positive patients [8,9].

Tocilizumab is a humanized anti-interleukin-6-receptor (IL-6R) monoclonal antibody (mAb) that inhibits interleukin-6 (IL-6) signaling. However, morbidity and mortality are not markedly altered by the administration of this single-targeted immunotherapeutic intervention. Similarly, the IL-1 receptor antagonist anakinra, may improve the clinical disposition of septic patients with COVID, especially those who harbor evidence of the macrophage activation syndrome [10].
The observation of the CRS appears to be, at least in part, secondary to the accelerated viral replication characteristics within the peripheral lung alveolar network, ultimately culminating in the massive release of newly assembled virions during cellular apoptosis, a process which also results in the release of pro-inflammatory mediators. Apoptosis resulting in the exaggerated and paroxysmal release of inflammatory effector elements is now referred to as *pyroptosis* [11]. In fact, pyroptosis is the process most likely responsible for the pathobiologic underpinnings of the so-called ‘cytokine storm’ (representing an important albeit one mechanism among a constellation of pathobiological underpinnings, which collectively represent the new rubric, which we have defined as PANIC) which involves the rapid release of a broad repertoire of cytokine and chemokine effector elements including TNF-α, IFN-γ, IL-1β, IL-6, IL-12, IL-18, IL-33, TGFβ; CCL2, CCL3, CCL5, CXCL3, CXCL19, and CXCL10 [11].

In conjunction with the CRS, the SARS-CoV-2 virus is further able to provoke what we believe to represent a redistribution of immune effector cells into the lungs, with the consequent intensification of tissue damage secondary to such cells releasing their corresponding effector elements, free radicals and reactive oxygen species, such as super oxide (O’).

The viral spike glycoprotein which decorates the outside surface of SARS-CoV-2, and serves as the ligand for ACE-2 receptor, also happens to represent a strongly immunogenic portion of the viral architecture [12]. During the process of viral binding to the alveolar ACE-2 receptor, in addition to utilizing the endosomal pathway for its tropism, the SARS-CoV-2 can integrate its membranous envelope with that of the alveoli epithelium, leading to multinucleated cell formation [Figure 3 in Part I]. In virology, such cross-species fusion elements are designated as syncytia.

The specific SARS-CoV-2 agent fuses its membrane in such a way that the resulting syncytium includes the strategic externalization of the spike protein, which can then interact with anti-spike IgG (possibly also IgM) antibodies. The result is the establishment of antigen-antibody complexes capable of complement fixation, and ultimately complement-dependent cellular (alveoli epithelium representing the ‘kill-target’ in this case) cytotoxicity [see ‘H’ in Figure 3 of Part I].
Biomarkers Implicating SARS-CoV-2 Triggered ‘PANIC’ Attack

A primary aim of this 2-part series has been to advance and substantiate the hypothesis that the pathobiological underpinnings of severely affected COVID-19 patients is principally related to the ‘PANIC’ Attack we have described throughout. Specifically that such patients mount exaggerated and injurious inflammatory cascades spanning the broad dynamic range of the effector mechanisms which endow the human immune network. The unintended consequences of this process include bystander tissue injury, particularly destruction of the most distal microanatomy of the bronchopulmonary tree, given the proclivity of the SARS-CoV-2 spike glycoprotein binding to the robust expression of ACE2 receptors on the alveolar epithelium.

The pleiotropic mechanisms of tissue injury involved in severely affected COVID-19 patients mean that targeting a single regulatory cytokine (e.g. IL-1 or IL-6) is highly unlikely to adequately uncouple the confluence of convergent inflammatory cascades that appear to rapidly descend upon the bronchopulmonary tree, ultimately resulting in an irreversible and step-wise decrement of the alveolar gas-exchange apparatus.

When a critical corpus of lung alveoli are lost, there is an accelerated failure to adequately load oxygen sufficient to deliver the substrate required for aerobic respiration. Virtually every system of the body requires aerobic respiration in order to maintain the constant bioenergetic balance between supply and demand of energy substrate for purposes of homeostasis. The respiration demands of a multiorgan system that operates across a dynamic range of metabolic circumstances with corresponding and rapidly fluctuating energy production and delivery characteristics must be met or catastrophe ensues.

Our hypothesis is that in the ~20% of SARS-CoV-2 infected patients with severe COVID-19, activation of innate and adaptive immune cascades working in synchrony inflict a ‘PANIC Attack’ upon the terminal lung fields. The outcome inevitably involves a hyperacute, and potentially diffuse, malfunction of the incredibly delicate functional interface between the alveolar epithelium and the single cell thick endothelial tubes, which comprise the capillaries derived from the pulmonary arteries and those which subsequently become the oxygen-rich blood vessels which enter into the pulmonary veins. We can ill afford to permit such a PANIC Attack to proceed without interruption, knowing that a catastrophic loss of alveoli will ultimately
bring the severe COVID-19 patient to the pathophysiologic threshold beyond which the lungs will fail to both transfer and load oxygen onto red cell hemoglobin, and to expel carbon dioxide.

We propose a potentially innovative hypothesis, whereby the application of an available, intensive, and highly pleiotropic immunotherapy, is characterized by commensurately synchronous and diametrically opposed actions serving to attenuate the specific sequence of SARS-CoV-2 triggered PANIC Attack mechanisms. We have employed this strategy for a number of fulminant and treatment recalcitrant inflammatory disorders of the CNS, that upon careful analysis, suggests that the immune phenomenonology, whether triggered by a foreign microbe, an adjuvant-enriched vaccine, or the ‘ignition’ of a monumentally severe exacerbation of autoimmunity, all appear to exhibit common pathobiological mechanisms reminiscent of our definition for the SARS-CoV-2 triggered PANIC Attack.

Our proposed strategy may serve to abort the uncoordinated and poorly regulated host immune activities, which appear to be fundamentally germane to the development of severe COVID-19, and which determine its prognostic disposition if the PANIC Attack proceeds with impunity. Further, we present a vignette [Vignette 1] of a patient with initially mild COVID-19 disease, who began to exhibit what we suspected were transformational clinical features at high risk for advancing to severe COVID-19, and underscore our multi-modal intervention strategy for ‘left-shifting’ away from the transition to severe COVID-19.

We follow with a second vignette which underscores the utility of the application of the HDMTX-LR protocol (the details of which are provided as a supplement at the end of the paper, but also detailed in our flow diagram on dichotomizing mild/moderate COVID-19 from the severe variant, and their respective proposed treatment interventions [Flow Diagram].

HYPOTHESIS:

*Methotrexate & Leucovorin Rescue Will Abolish SARS-CoV-2-Induced PANIC Attack: A Mechanistically Pleiotropic Rescue Strategy For Severe COVID-19*

We suggest that high-dose methotrexate with leucovorin rescue (HDMTX-LR), an anti-inflammatory and chemotherapeutic agent, is endowed with impressive mechanistically pleiotropic properties [Table 1], which are capable of mitigating, if not abolishing, each of the
SARS-CoV-2 triggered mechanisms that compositionally make up the so-called ‘PANIC Attack’. Further, this array of HDMTX-mediated coordinate actions may provide for the development of active immunity towards the SARS-CoV-2 infection, reflecting the very real prospect that such an intervention may serve to ‘left-shift’ the severe COVID-19 patient to the milder phenotype, along with a correspondingly improved prognosis. Our prior experience with the application of this intensive treatment strategy, and the nearly stereotyped remission in our patients, who, upon presentation, were codified as harboring treatment refractory syndromes of fulminant inflammatory disorders of CNS (e.g. multiple sclerosis (MS), neuromyelitis optica spectrum disorder (NMOSD), Sjogren’s associated myelitis, post-vaccinal (dTap) encephalomyelitis, and a series of cases of monumentally severe post-viral encephalomyelitides).

**A Novel, Host-Mediated Syndrome of Deleterious Immune Network Activation: PANIC**

The observation of reminiscent ‘PANIC Attack’ mechanisms across our expanding set of fulminant inflammatory syndromes appears to be indistinguishable whether afflicting the CNS or the bronchopulmonary tree. These salient similarities have prompted us to extrapolate from our experience with treatment resistant, recalcitrant, and monumentally fulminant immune activation affecting our patients with MS, NMOSD, Sjogren’s syndrome, and in a host of other circumstances where such catastrophic immune-mediated inflammation, can, if not identified and treated accordingly, obliterate the triggering agent’s target tissue(s).

We have since utilized the same protocol in many patients, who we now believe in retrospect were suffering from a form of ‘PANIC’ Attack, except targeting the central nervous system. In Part I of this two-part publication, we reported a patient suffering from a severe adenovirus-triggered PANIC Attack, characterized by cytokine storm, and an encephalomyelitis, with a PRESS-like pattern distribution of lesions [ADD REFERENCE FOR PART I]. His protracted and treatment-recalcitrant course had come to the point where a decision was made to consider cessation of care. However, it was finally agreed to try the HDMTX-LR regimen, which ultimately led to the complete recovery of the patient and resolution of all of the brain lesions which characterized his syndrome [ADD REF FOR PART I HERE].
A more recent example considers the plight of a young accountant, who developed optic neuritis in August of 2017, approximately two weeks following a dTap vaccine and during the 3rd trimester of her first pregnancy. Declining steroids, for fear of early term delivery, she recovered in a self-limited fashion. However, by October she presented again, this time with optic neuritis in the fellow eye, in addition to some sensory and motor manifestations. Investigations revealed an elevated, blood-derived anti-MOG antibody at a titer of 1:1000, and consistent with a post-vaccinal, anti-MOG associated encephalomyelitis. Treated with corticosteroids, she failed to improve.

By December, our patient presented again with confusion, psychomotor slowing, and MRI demonstrating extensive and severe inflammatory lesions throughout much of the cerebral white matter. Coticosteroids once again were without effect, whereas the application of our HDMTX-LR protocol was tolerated without any adverse effects. This therapy rapidly attenuated the disease process, including normalization of her clinical manifestations, in conjunction with a remarkable disappearance of many of her white matter lesions [ADD REF FOR PART I]. A few months later, the appearance of new enhancing cerebral white matter lesions prompted us to advance our regimen to a full course of plasma exchange (5-full volume exchanges; which took 8 days), followed by another course of HDMTX-LR. From the end of that course of therapy in early 2018 to the writing of this manuscript, there has been no recurrence, clinically or radiographically. In fact, most of the lesional burden has disappeared [ADD REF FOR PART I].

**Calming ‘PANIC’ in Severely affected COVID-19 Patients:**

**A Pleiotropic Immune Stabilization Strategy for SARS-CoV-2 Induced ‘PANIC’ Attack**

The application of HDMTX for neoplastic and fulminant inflammatory syndromes of the CNS represents a critically important observation because, despite a steady-state blood to CSF ratio of MTX of about 30:1 [13], doses of 500mg/m² or more are necessary to achieve CNS concentrations of MTX to be sufficiently efficacious against such disorders. The ability to escalate dosing of MTX while rescuing peripheral folate with leucovorin (folinic acid)
represents a unique aspect of using this agent across a number of disease states, with more than compelling justification for the WHO to list MTX on the coveted list of ‘essential medications’ [13,14].

**Pleiotropic Mechanisms of Action for HDMTX**

Methotrexate (4-amino-10-methyl folic acid) is both an analog as well as an antagonist of folic acid, but is capable of exerting its action via a wide spectrum of pleiotropic mechanisms, such that it is considered by the WHO as an essential medicine [13,14] [Figure 1; Table 1].

**Cell Cycle S-Phase Inhibition via Folate-Dependent Enzyme Inhibition**

Cells that are rapidly dividing, such as those associated with malignancy, autoimmunity, and during our proposed PANIC syndrome associated with the SARS-CoV-2 infection, spend an increased time in the cell cycle synthesis (S) phase. However, MTX, in part, functions as an S phase inhibitor by depriving nucleoside precursors for both DNA and RNA synthesis and replication [15].

**DNA and RNA Base Synthesis Inhibition**

MTX accomplishes this via its potent inhibition of folate-dependent enzymes for both purine and pyrimidine base synthesis [Figure 1]. The vast majority of such rapidly dividing cells can manufacture base synthesis only by the de novo (i.e. from ‘scratch’) pathways involving inosine monophosphate dehydrogenase (IMDH) for purine, and dihydroorotate dehydrogenase (DHOD) for pyrimidine biosynthesis. Alternately, the vast majority of body cells can produce bases by either the de novo or via the salvage pathways, the latter characterized by the use of partially assembled or partially broken down bases [15].

Following its entry into the cellular cytoplasm, MTX is polyglutamated, which serves to both trap the drug within the intracellular compartment precluding its ability to efflux, and allow the drug to exert one of its principal actions: the inhibition of dihydrofolate reductase (DHFR), as well as other folate-dependent enzymes, such as thymidylate synthase, 5-amino-imidazole-4 carboxamide ribonucleotide (AICAR) transformylase (ATIC), and methylene tetrahydrofolate reductase (MTHFR) [Add REF FOR PART I].
Interestingly, folate supplementation does not abolish either the anti-inflammatory, the chemotherapeutic actions, or the efficacy of MTX. This suggests that other mechanisms, independent of folate antagonism, must figure prominently in the potent actions of MTX. For instance, MTX administration provokes the release of adenosine, a potent anti-inflammatory agent that exerts regulatory effects upon neutrophils, macrophages, and T cells [15].

**Adenosine Mediated Anti-Inflammatory Mechanisms**

Adenosine promotes its action through interaction with its receptor A2a and ADORA3, where transmembrane signaling activities result in the reduction of production and release of pro-inflammatory cytokines such as IL-1β, TNFα, IL-6, and reduction in the production of matrix metalloproteinases (MMPs) (the transcription of each requires NF-κB). An additional action of MTX is the inhibition of NFκB as well as the escalation in their corresponding tissue inhibitors (TIMPs). Methotrexate inhibits catabolism of adenosine and AMP, while increasing release of both ATP and ADP from inflammatory cells; an effect which modulates the bioenergetic activity, such that inflammatory cells are diminished in their ability to foment unchecked pro-inflammatory effector mechanisms [16,17].

**JAK/STAT Signaling Platform Suppression**

A significant pathway responsible for the transduction of multiple pro-inflammatory cytokines is the JAK/STAT signaling platform, which is potently suppressed by mechanisms independent of the folate inhibition effects of MTX, as such effects are not reversed by administration of folinic acid [Figure 1B].

**Methotrexate Modulation of Reactive Oxygen Species**

Methotrexate, through its modulation of reactive oxygen species (ROS), can induce T cell apoptosis, thereby reducing the cytokine production and elaboration; with a prominent effect upon the reduction of IL-6 [18-21]. Further, MTX activates JNK, thereby activating downstream targets such as c-JUN and cFOS, constitutive elements of the AP-1 complex
involved in the heightened expression of pro apoptotic genes which increase the propensity to apoptosis [Figure 1].

**Methotrexate Inhibits HMGB1 Binding to RAGE**

The high-mobility group box chromosomal protein 1 (HMGB1) is elaborated by activation or injured immune cells, including monocytes, macrophages, and dendritic cells, and mediates inflammation [Figure 1C]. One such example is called ‘alarmin’, which promotes acute inflammation followed by tissue repair [22-24]. The effects of HMGB1 is through its counterreceptor, a receptor for advanced glycan end products (RAGE), and is associated with cell maturation and migration to the site of injury, whereby chronic inflammation is perpetuated [25,26]. Methotrexate can inhibit HMGB1 by directly binding to RAGE or indirectly by inhibition of cytokine production of TNFα, IL-1, IL-6, IL-8.

**Treatment Protocol for the Application of High-Dose MTX with Leucovorin Rescue**

The proposed HDMTX-LR protocol has been the focus of prior investigation, the details of which have been published [27] [Table and Figure 1, Protocol, Flow Diagram]. Briefly, prior to therapy we ensure that the urine specific gravity is <1.1010, urinalysis and urine culture are without evidence of infection, the serum creatinine (Scr) is below 1.4mg/dl, and complete blood count (CBC), and the comprehensive metabolic panel (CMP) to include liver functions and electrolytes, are assessed prior to administration of HDMTX-LR. For purposes of reducing the risk of MTX crystalline deposition in the kidneys, D5W and sodium bicarbonate 100mEq are administered intravenously at a rate of 150ml/m²/hr for at least 8 hours prior to the administration of the HDMTX-LR protocol in order to achieve a urine pH of at least 8.0 before and during the entire HDMTX-LR infusion.

MTX at 2500mg/m² in 250ml of normal saline is administered intravenously over 2 hours. Precisely 12 hours after the inception of the MTX infusion, we began leucovorin (folinic acid) rescue with 80mg given intravenously. Thereafter, leucovorin at 35mg is administered intravenously precisely every 6 hours for a total of 12 doses [Figure 1 & Protocol]. Serum MTX concentration levels are assessed at 24hr, 36hr, 48hr, and 72hr after the administration of HDMTX-LR. Patient discharge cannot proceed until the MTX concentration falls below 0.05μmolar (typically requiring a period of 24-72 hours).
Nausea and vomiting tend to occur during or shortly after the HDMTX-LR administration, hence anti-emetics (e.g. odansetron) are given 30 minutes before the infusion and as needed thereafter. Dextromethorphan (5-10mg po prn every 2-4 hours; or 20mg with 10mg of quinidine; Nuedexta daily) is utilized to treat infusion-related headaches. CBC, Scr, and CMP are assessed at baseline and then at weeks 1 and 4 following treatment with HDMTX-LR.

Penicillin derivatives, probenecid, fluoroquinolones, sulfonamides, aspirin, and non-steroidal anti-inflammatory drugs are known to decrease renal excretion of MTX and are therefore avoided during HDMTX-LR treatment. Vancomycin and trimethoprim/sulfamethoxazole are avoided immediately prior to, and during HDMTX-LR administration to avoid potential additive or synergistic nephrotoxicity, and possible myelosuppression [13].

A range of potential toxicities can be associated with the HDMTX-LR protocol, each of which however, can be markedly mitigated, if not prevented or abolished, with careful attention to the precise proscriptions of our protocol, and the use of systematic and sequential stepwise transitions to each phase of the therapeutic plan [Protocol]. Most important is that the HDMTX-LR protocol can be safely employed, while reducing risk of toxicity by ensuring vigorous hydration (to avoid MTX crystalline deposition), urine alkalinization (crystal formation is more likely in acid pH of the genitourinary system), along with the precisely-timed administration of the leucovorin rescue (to prevent mucositis).

The addition of benfotiamine has been shown to reduce MTX mediated liver toxicity in those utilizing low dose weekly chronic dosing of the medication, and the addition of this agent to the high dose punctuated administration of MTX may also confer some tangible benefits, albeit a subject for further investigation. Further, adequate levels of vitamin D (to achieve 60-100ng/ml) are known to exert immunomodulatory effects [28].

**Management of Severe COVID-19: Practice Principles**

The principal thrust of this paper is to emphasize that the SARS-CoV-2 induced severe variant of COVID-19 is not mediated by a single renegade cytokine or immunological phenomenon (e.g. the cytokine storm). Instead, this third, and most ominous of the known
Coronaviruses, is characterized by its rapid and highly efficient transmissibility, the broad spectrum of clinical courses ranging from wholly asymptomatic (rendering such patients as a kind of ‘typhoid Mary’), to a mild/moderate flu-like illness, and ultimately to the severe variant. The latter is principally recognized by the requirement for respiratory support, culminating in pressure ventilation, at which point the prognosis is poor, with most succumbing to the disease via the paroxysmal cessation of gas exchange, and the pathophysiologic signature of irreversible demise, treatment refractory hypoxic-ischemia. At this point, further interventions become an unnecessary and psychologically devastating exercise in futility for the patient, their family, and for the front-line care team.

Herein we have underscored the serious nature of COVID-19, the manifestations and clinical and paraclinical factors which portend this ominous, and commonly irreversible, semiology. Further, we have dissected the manifold mechanisms, though not exhaustively, triggered by the SARS-CoV-2 virus, which has led us to forge new principles with respect to microbial associated immune activation in a naïve host. Mild to moderate COVID-19 is largely defined by the nature, magnitude, location, and whether such immune network responses are mechanistically coherent, and effective, while at the same time being sufficiently ‘focalized’. Focalized responses signify a coordinately regulated network immune response, one most capable of limiting the severity of end-organ tissue damage.

The primary goal of this communication is to help our front line workers recognize that despite the ‘prolific activation of a network-mediated inflammatory crisis’, or PANIC Attack, we do in fact have compositional immunotherapies, that when combined together (as in the era of discovery of highly active anti-retroviral therapy (HAART)), may serve to abort or at least mitigate, suffering while promoting an improved prognosis. Alternately, we herein hypothesize that a single agent endowed with a plethora of diverse and site selective targeting capabilities, each of which is sufficiently robust so as to uncouple the SARS-CoV-2 induced PANIC Attack, may be used to lead the counter-attack. It is precisely for this reason that the WHO has long considered MTX as a ‘necessary’ medicine.

CONCLUSIONS: TIME IS TISSUE

About 20% of COVID-19 patients will ultimately advance to severe disease, with assimilation of the corresponding risk escalation of both morbidity and mortality. Most COVID-
19 patients shall cycle through a number of clinical dispositions, each of which may be remediable, and potentially even mitigated with respect to ‘left-shifting’ the patient and their course, further away from those thresholds beyond which survival becomes increasingly more remote. To illustrate, we have prepared two scenarios as ‘call out boxes’ which contain an abbreviated vignette, followed by our specific plans of intervention [Box 1 and 2]. The goals for each scenario are to mitigate suffering, promote reconstitution of network immune coordinate regulation sufficient to prevent PANIC, but also to recognize and urgently treat PANIC when the features that define it are recognized by our front-line workers.

As a synopsis, the mildly affected COVID-19 patient is offered site-selected delivery of inhalational steroids followed by L-albuterol (R form of this agent, making it relatively lung selective). This promotes bronchial smooth muscle relaxation, as well as activates the beta-2 adrenergic receptor on immune cells, thereby rendering them diminished in their inflammatory activity via activation and escalation of intracellular cyclic AMP (essentially biasing immune cells from a TH1 to TH2 anti-inflammatory phenotype). Further, low-dose weekly MTX (7.5 to 25mg taken po weekly along with 2mg of folate daily) could be utilized in those patients with mild to moderate COVID-19, but who persist or exhibit what appears to represent a more protracted and slowly advancing clinical deterioration. When compared to our HDMTX-LR intervention, low dose weekly oral MTX is not as prolific, nor as rapidly acting to exact remission. It can, nevertheless, exert a constellation of attenuating influences in response to the SARS-CoV-2 triggering of the broad expanse of the host immune networks.

Pulmonary toilet (now referred to as pulmonary hygiene) can loosen and expectorate mucus in the distal bronchopulmonary tree, which is facilitated by the performance of postural percussion of the back and sides of the thoracic cage. For chronic cough, dextromethorphan + quinidine may be prescribed, the latter reducing the metabolism of dextromethorphan while also providing a derivative of quinine from the cinchona tree. Quinine, like chloroquine and hydroxychloroquine, has demonstrated interrupted viral synthesis in vitro, as well as the release of cytokines. Vitamin D3 taken daily, and at a dose commensurate to achieving a blood level of 60-100ng/ml, can promote immune regulatory networks, and improve energy (reduce fatigue) [29].
Ultimately, rather than targeting individual cytokines and their receptors, the HDMTX-LR protocol provides a broader spectrum of protection against the cytokine storm described herein, as well as the other mechanisms that collectively we now define as a PANIC Attack. Whether triggered by immune network dysregulation in autoimmune disorders, or secondary to a foreign microbe capable of triggering PANIC, the resultant process occur in a significant proportion of infected individuals. The pathobiological underpinnings which dichotomize those destined to experience mild vs severe COVID-19, shall, and already is, the subject of vigorous debate, and extensive investigations by the most gifted and talented physicians, scientists, and physician-scientists from around the globe. We truly are in this together, and by working together, we have our best prospects to conquer this pandemic, while preparing for the next ‘PANIC’ Attack.

Author Disclosures

Elliot Frohman: Has received speaker fees from Genzyme, Alexion, Novartis, and consulting fees from Biogen and Serono

Esther Melamed: served as a consultant and received honoraria from EMD Serono and Genentech

Roberto Alejandro Cruz: Has received speaker fees from Alexion

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Lawrence Steinman: Dr. Steinman is on the Editorial Boards of The Proceedings of the National Academy of Sciences, and the Journal of Neuroimmunology. He has served on the Editorial Board of the The Journal of Immunology and International Immunology. He has served as a member of grant review committees for the National Institutes of Health (NIH) and the National MS Society.

He has served, or serves, as a consultant and received honoraria from Atara Biotherapeutics, Atreca, Biogen-Idec, Celgene, Centocor, Coherus, EMD-Serono, Genzyme, Johnson and Johnson, Novartis, Roche/Genentech, Teva Pharmaceuticals, Inc., and TG Therapeutics. He
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Currently, Dr. Steinman receives research grant support from the NIH and Atara Biotherapeutics.

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Supplementary data

Supplementary material

References

1. Khoury SJ, Healy BC, Kivisäkk P, Viglietta V, Egrova S, Guttmann CR, Wedgwood JF, Hafler DA, Weiner HL, Buckle G, Cook S, Reddy J. A randomized controlled double-masked trial of albuterol add-on therapy in patients with multiple sclerosis. Arch Neurol. 2010 Sep;67(9):1055-61. doi: 10.1001/archneurol.2010.222. PubMed PMID: 20837847; PubMed Central PMCID: PMC2954052.

2. Makhlouf K, Weiner HL, Khoury SJ. Potential of beta2-adrenoceptor agonists as add-on therapy for multiple sclerosis: focus on salbutamol (albuterol). CNSDrugs. 2002;16(1):1-8. Review. PubMed PMID: 11772115.

3. Makhlouf K, Comabella M, Imitola J, Weiner HL, Khoury SJ. Oral salbutamol decreases IL-12 in patients with secondary progressive multiple sclerosis. J Neuroimmunol. 2001 Jul 2;117(1-2):156-65. PubMed PMID: 11431016.

4. Tsai CP, Lin FC, Lee CT. Beta2-adrenergic agonist use and the risk of multiple sclerosis: a total population-based case-control study. Mult Scler. 2014 Oct;20(12):1593-601. doi: 10.1177/1352458514528758. Epub 2014 Apr 14. PubMed PMID: 24732071.

5. Zhang L, Pang R, Xue X, Bao J, Ye S, Dai Y, Zheng Y, Fu Q, Hu Z, Yi Y. Anti-SARS-CoV-2 virus antibody levels in convalescent plasma of six donors who have recovered from COVID-19. Aging (Albany NY). 2020 Apr 22;12. doi:10.18632/aging.103102. [Epub ahead of print] PubMed PMID: 32320384.
6. Franchini M, Marano G, Velati C, Pati I, Pupella S, Liumbruno GM. Operational protocol for donation of anti-COVID-19 convalescent plasma in Italy. Vox Sang. 2020 Apr 23. doi: 10.1111/vox.12940. [Epub ahead of print] PubMed PMID: 32324899.

7. Epstein J, Burnouf T. Points to consider in the preparation and transfusion of COVID-19 convalescent plasma. Vox Sang. 2020 Apr 22. doi: 10.1111/vox.12939. [Epub ahead of print] PubMed PMID: 32319102.

8. DHOLARIA BR, BACHMEIER CA, LOCKE F: Mechanisms and management of chimeric antigen receptor T-cell therapy related toxicities. BioDrugs 2019; 33: 45-60.

9. LEE DW, GARDNER R, PORTER DL et al.: Current concepts in the diagnosis and management of cytokine release syndrome. Blood 2014; 124: 188-95.

10. SHAKOORY B, CARCILLO J, CHATHAM W et al.: Interleukin-1 receptor blockade is associated with reduced mortality in sepsis patients with features of macrophage activation syndrome: reanalysis of a prior phase iii trial. Crit Care Med 2016; 44: 275-81.

11. YANG M: Cell pyroptosis, a potential pathogenic mechanism of 2019-nCoV infection. SSRN Electron J 2020.

12. LIU L, WEI Q, LIN Q et al.: Anti-spike IgG causes severe acute lung injury by skewing macrophage responses during acute SARS-CoV infection. JCI Insight 2019; 4.

13. M. Joerger, A.D. Huitema, G. Illerhaus, A.J. Ferreri, Rational administration schedule for high-dose methotrexate in patients with primary central nervous system lymphoma, Leuk. Lymphoma 53 (2012) 1867–1875.

14. Conway, R.; Carey, J.J. Risk of liver disease in methotrexate treated patients. World J. Hepatol. 2017, 9, 1092–1100. [CrossRef] 9. WHO. WHO Model Lists of Essential Medicines 21th List. Available online: http://www.who.int/medicines/publications/essentialmedicines/en/ (accessed on 22 July 2019).

15. Bedoui Y, Guillot X, Sélambarom J, Guiraud P, Giry C, Jaffar-Bandjee MC, Ralandison S, Gasque P. Methotrexate: an Old Drug with New Tricks. Int J Mol Sci. 2019 Oct 10;20(20). pii: E5023. doi: 10.3390/ijms20205023. Review. PubMed PMID:31658782; PubMed Central PMCID: PMC6834162.
16. Brown, P.M.; Pratt, A.G.; Isaacs, J.D. Mechanism of action of methotrexate in rheumatoid arthritis, and the search for biomarkers. Nat. Rev. Rheumatol. 2016, 12, 731. [PubMed]

17. Montesinos, M.C.; Takedachi, M.; Thompson, L.F.; Wilder, T.F.; Fernández, P.; Cronstein, B.N. The antiinflammatory mechanism of methotrexate depends on extracellular conversion of adenine nucleotides to adenosine by ecto-5'-nucleotidase: Findings in a study of ecto-5'-nucleotidase gene-deficient mice. Arthritis Rheum. 2007, 56, 1440–1445.

18. Herman, S.; Zurgil, N.; Deutsch, M. Low dose methotrexate induces apoptosis with reactive oxygen species involvement in T lymphocytic cell lines to a greater extent than in monocytic lines. Inflamm. Res. 2005, 54, 273–280.

19. Phillips, D.C.; Woollard, K.J.; Griffiths, H.R. The anti-inflammatory actions of methotrexate are critically dependent upon the production of reactive oxygen species. Br. J. Pharm. 2003, 138, 501–511.

20. Chalupsky, K.; Cai, H. Endothelial dihydrofolate reductase: Critical for nitric oxide bioavailability and role in angiotensin I uncoupling of endothelial nitric oxide synthase. Proc. Natl. Acad. Sci. USA 2005, 102, 9056–9061. [PubMed]

21. Sung, J.Y.; Hong, J.H.; Kang, H.S.; Choi, I.; Lim, S.D.; Lee, J.K.; Seok, J.H.; Lee, J.H.; Hur, G.M. Methotrexate suppresses the interleukin-6 induced generation of reactive oxygen species in the synoviocytes of rheumatoid arthritis. Immunopharmacology 2000, 47, 35–44.

22. Klune, J.R.; Dhupar, R.; Cardinal, J.; Billiar, T.R.; Tsung, A. HMGB1: Endogenous Danger Signaling. Mol. Med. 2008, 14, 476–484.

23. Castiglioni, A.; Canti, V.; Rovere-Querini, P.; Manfredi, A.A. High-mobility group box 1 (HMGB1) as a master regulator of innate immunity. Cell Tissue Res. 2011, 343, 189–199.

24. Scaffidi, P.; Misteli, T.; Bianchi, M.E. Release of chromatin protein HMGB1 by necrotic cells triggers inflammation. Nature 2002, 418, 191–195. [PubMed]

25. Schmidt, A.M.; Yan, S.D.; Yan, S.F.; Stern, D.M. The multiligand receptor RAGE as a progression factor amplifying immune and inflammatory responses. J. Clin. Investig. 2001, 108, 949–955. [PubMed]
26. Bierhaus, A.; Humpert, P.M.; Stern, D.M.; Arnold, B.; Nawroth, P.P. Advanced glycation end product receptor-mediated cellular dysfunction. Ann. N. Y. Acad. Sci. 2005, 1043, 676–680. [PubMed]

27. Beh SC, Kildebeck E, Narayan R, Desena A, Schell D, Rowe ES, Rowe V, Burns D, Whitworth L, Frohman TC, Greenberg B, Frohman EM. High-dose methotrexate with leucovorin rescue: For monumentally severe CNS inflammatory syndromes. J Neurol Sci. 2017 Jan 15;372:187-195. doi: 10.1016/j.jns.2016.11.012. Epub 2016 Nov 15. PubMed PMID: 28017209.

28. Erdogan MA, Yalcin A. Protective effects of benfotiamine on irisin activity in methotrexate-induced liver injury in rats. Arch Med Sci. 2018 Nov 29; 14(6):205-211. doi: 10.5114/aoms.2018.80002. eCollection 2020. PubMed PMID: 32051725; PubMed Central PMCID: PMC6963132.

29. Jakovac H. COVID-19 and vitamin D-Is there a link and an opportunity for intervention? Am J Physiol Endocrinol Metab. 2020 May 1;318(5):E589. doi: 10.1152/ajpendo.00138.2020. PubMed PMID: 32297519.

**Figure 1:** Methotrexate has been recognized by the World Health Organization as one of the ‘essential medications’, principally due to its broad diversity of mechanisms of action, whether utilized at low doses for purposes of taking advantage of its dynamic range of anti-inflammatory mechanisms (particularly beneficial in the management of rheumatoid arthritis), or at high doses to eradicate a range of malignancies, or in circumstances of fulminant
inflammatory conditions where conventional approaches have been futile. In this figure we emphasize the pleiotropic and discretely targeted actions of high-dose methotrexate, as a hypothetical intervention for the SARS-CoV-2 triggered PANIC Attack. The viral infection will produce the severe variant of COVID-19, principally secondary to the simultaneous activation of the diverse limbs of the human immune network (Figure 1). As such, productive therapeutic strategies are likely those which either combine different agents which target distinctly different mechanisms of immune activation, or we need to identify novel or existing monotherapies which are endowed with a diversity of actions capable of counteracting the PANIC Attack triggered by the SARS-CoV-2 agent in those afflicted with the severe variant of the disease. In A) we showcase methotrexate as a potent inhibitor of the folate-dependent enzyme systems, especially those which are essential in rapidly dividing cells; such as those of the hyper-activated immune system, given that the cell cycle synthesis (S) phase requires nucleoside precursors for the synthesis of both DNA and RNA by the de novo pathways; and as such for RNA replication of the SARS-CoV-2 agent. Other body cells overcome this base synthesis blockade by utilizing the so-called salvage pathways, where partially assembled and partially broken down bases can be utilized to complete the synthesis; which are not available to rapidly dividing cells and microbes such as the SARS-CoV-2 virus. MTX is polyglutamated following entry into cells, a molecular modification which serves to prevent its efflux back out of the cell. Once inside this form of the molecule effectively inhibits dihydrofolate reductase among other folate dependent enzyme systems, which ultimately inhibits the de novo biosynthetic pathways for purines and pyrimidines (necessary for both DNA and RNA); and as such it reduces RNA replication of the virus. In B) at 1:00 we illustrate the inhibitory effects of MTX on multiple transmembrane signaling pathways involved in inflammation, including the JAK-STAT pathways that are activated by a host of inflammatory cytokines such as IL-2, IL-6, IL-12, IL-15, GM-CSF, and IFNγ. At 2:00 we illustrate the role of the NFκB signaling pathway for a series of inflammatory mediators such as IL-1β, IL-6, TNFα, and for matrix metalloproteases. MTX inhibits the production of these inflammatory mediators, but also liberates reactive oxygen species (ROS) which then engages JNK activation, which then can promote p53 activity which represents the final NFκB sequence inhibitor (thereby blocking the effector functions of proinflammatory cytokines). Further JNK activation also activates the c-Fos and c-Fun pathways which culminate in the gene expression of pro-apoptotic gene expression and
increased sensitivity to apoptosis. In C) we illustrate the effect of MTX in its ability to block HMGB1 to the RAGE receptor, and thereby aborting the multiple downstream immune effector injury mechanisms that converge to destroy the lung alveoli in severe COVID-19. The attenuation of immune cell maturation, migration and transmigration across the alveoli epithelia (utilizing matrix metalloproteinases which digest fibronectin, collagen, and other matrix elements that constitute the integrity of tissue elements), and the release of cytokines and other inflammatory mediators by either active release mechanisms, or via the passive release secondary to the expulsion from damaged or dying cells (a process referred to as pyroptosis; inflammatory apoptosis) are additional alveolar injury pathways that are blocked by MTX. In D) we emphasize a series of risk mitigation interventions that are crucial when using HD-MTX-LR (all of which are further discussed in the text).

Vignette Box 1

Treatment Algorithm for Mild-Moderate Severity of COVID-19

‘Left-Shifting’ for Prevention of PANIC

A telemedicine visit involved a 31-year-old woman with relapsing-remitting multiple sclerosis, treated with dimethyl fumarate at 240mg po bid, who was ‘sheltering in place’ and awoke one morning with new onset fever (Tmax = 101.4 deg F), non-productive cough, the perception of chest ‘tightness’, increased work of breathing, headache (long history of migraine and what appears to be consistent with transformed headache) extreme fatigue and malaise. With head flexion or hip flexion while laying supine and attempting to straighten the leg, there is no increased head pain to suggest meningeal root irritation (Brudsinski and Kernig signs respectively were absent). Conspicuously, and on an unsolicited basis, she describes the relatively sudden loss of smell followed by taste sensation.

Based upon the constellation of symptoms, we were suspicious of COVID-19 and I asked the
patient to be seen at her local ER for testing (either with RT-PCR or with CT chest imaging to reveal characteristic peripherally based ground-glass lesions; findings with 98% sensitivity compared to the ~70% sensitivity of the RT-PCR assay). While the ER physician’s diagnosis was COVID-19, no confirmatory testing was performed. However, laboratory testing did reveal lymphopenia (absolute number of 250), a CRP of 2.75, normal D-Dimer, and she was discharged to home.

Over subsequent days, and nearly daily telemedicine conference calls, it was evident that her flu-like symptoms persisted, if not intensifying (cough, perception of reduced negative inspiratory force (NIF), persistent fever, and the perception of chest tightness). A series of recommendations were made in an attempt to both mitigate symptoms, but also to potentially reduce the induction of the SARS-CoV-2 induced PANIC attack.

| Complaint | Intervention |
|-----------|--------------|
| Fever Infection | Azithromycin |
| Multiple Sclerosis | The patient was lymphopenic on dimethyl fumarate (a recognized association) and with suspected COVID-19. The significant lymphopenia places this patient at risk for PML, although the lymphopenia can also be secondary to COVID-19. |
| Disease Modifying Therapy Change | 2. A decision was made to discontinue the dimethyl fumarate, and to instead begin treatment with interferon beta 1a, taken SQ every two weeks (Plegridy). Further, given that the patient exhibited breakthrough on interferon in the past, it was also decided to treat concomitantly with teriflunomide; starting at 7mg po daily for two weeks; then escalating to 14mg daily thereafter. |
|  | 3. Type I interferons exert anti-viral actions, and teriflunomide is an inhibitor of pyrimidine base synthesis, via the inhibition of the de novo biosynthetic enzyme, dihydroorotate dehydrogenase. |
Rapidly dividing cells and microbes such as the SARS-CoV-2 agent must utilize the de novo pathway for DNA and RNA base synthesis, whereas other body cells can use either de novo or salvage pathways (using partially assembled or partially broken down bases for DNA and RNA synthesis).

1. Assess oxygen saturation with simple home finger pulse oximetry. Given parameters, patients can provide their managing physicians with crucial information as to whether the COVID-19 clinical course is deteriorating and hence the need for hospitalization.

2. Supplemental Oxygen via nasal canula

### Shortness of Breath

**Shortness of Breath**

(Chest Tightness)

1. Postural Percussion performed by husband to facilitate pulmonary hygiene (formerly pulmonary toilet to mobilize and promote mucus clearance)

2. Incentive Spirometry for deep breathing and exercise of respiratory muscles

### Cough: Non-Productive

Dextromethorphan

Dextromethorphan (DM) + Quinidine; Nuedexta; (the latter reduces the metabolism of DM). Quinidine is a derivative of quinine, from the Peruvian Cinchona tree, as are chloroquine and hydroxychloroquine; all of which have been observed to reduce viral replication and cytokine release in vitro.

### Chest Tightness:

- **A)** Liquify mucus
- **B)** Mucus mobilization
- **C)** Bronchopulmonary inflammation
- **D)** Deep terminal airway inflammation

- **A)** Guaifenesin bid for mucus liquification
- **B)** Consider Warm Steam Vaporizer
- **C)** Pulmicort Inhalational steroid
- **D)** Site-selective delivery of anti-inflammatory action into the terminal bronchopulmonary tree
1. **Bronchial constriction**
   - Levalbuterol; also known as L-albuterol (Xopenex) and is the active form of the drug called R-albuterol. Fewer side effects than with racemic albuterol (which is a 50:50 mixture of R and S form).
   - Pharmacologically acts via the beta-2 adrenergic receptor on bronchial smooth muscle thereby relaxing the distal extent of the bronchopulmonary tree. Can reduce the ‘tight lung’ perception, and via bronchodilation can facilitate mobilization of mucus and avoid inspissated mucus organization. Two puffs TID.

2. **Beta-2 adrenergic receptor mediated immune skewing**
   - Beta-2 adrenergic receptors are also expressed on circulating mononuclear cells. R albuterol binds to the beta-2 receptor and promotes conversion of cells of the TH1 proinflammatory phenotype to TH2 anti-inflammatory state [68-71]. Two puffs TID.

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### Elevated Blood Pressure

| 1. Bioenergetics | 1. Benfotiamine 300mg daily: Improved bioenergetics |
|------------------|--------------------------------------------------|
| 2. Reduce MTX tissue injury | 2. Reduced tissue damage from MTX and other inciting agents [94]. |

**Blood pressure monitoring and correlate with headache.**

### Vitamin D: 10,000 units of D3 daily for promoting immune modulation/Tregs

| 1. Vitamin D: 10,000 units of D3 daily for promoting immune modulation/Tregs | Vitamin D may play an immunoregulatory role in COVID-19 patients [95]. |

### Low Dose Weekly MTX

| Daily Folic Acid | 7.5-25mg po q weekly |
|------------------|----------------------|
| 2mg of Daily Folate |                       |

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### Vignette Box 2

**Prevention, Mitigation, or Abolishment of the SARS-CoV-2 Triggered PANIC**

A 55-year-old University School of Medicine hospital physician intensivist, and unit Director,
has been carefully and systematically managing mild, moderate, and severely affected COVID-19 patients since March 15, 2020. During this time he has consistently utilized PPE consisting of gloves, disposable gown, N95 masks with facial shields, and head and shoe covers. On April 22, his unit is visited by a senior member of the COVID-19 Federal Safety Task Force, refuses to wear a mask of any type, so that he is certain to express his abundant respect by not covering his face while he offers his congratulations to the unit Director, as the visit was orchestrated to recognize the heroic efforts of this senior faculty member, in the management of the most severe cases of COVID-19 in the region. The conversation between the two men is at a distance of about 3-4 feet from each other. In a lapse in judgment the unit Director removes his mask in order to exhibit his respect for the senior task force representative.

Within 24 hours, the unit Director has developed a fever of 101.4 F, a dry, non-productive cough, chest discomfort with inspiration, profound fatigue and malaise, increased work of breathing, myalgias and exercise intolerance. Laboratory assessment reveals elevated D-Dimer, a CRP of 3.75, CK of 1500, lymphopenia (absolute number = 400), ferritin of 700, and an O₂ saturation of 88%. Chest CT reveals multilobar ground glass lesions consistent with COVID-19; while the RT-PCR for SARS-CoV-2 is negative. There is a history of hypertension, currently being treated with Losartin.

| Complaint                  | Intervention                      |
|----------------------------|-----------------------------------|
| Fever Infection            | Azithromycin                      |
|                            | 1. Assess oxygen saturation (80%)|
|                            | 2. Supplemental Oxygen 4L via nasal canula |
| Shortness of breath        |                                   |
| Shortness of Breath (Chest Tightness) | Postural Percussion to facilitate pulmonary hygiene (formerly pulmonary toilet to mobilize and promote mucus clearance) |
|                            | 4. Incentive Spirometry for deep breathing and exercise of respiratory muscles |
| Cough:Non-Productive       | Dextromethorphan                  |
|                            | Dextromethorphan (DM) + Quinidine; Nuedexta; (the latter reduces the metabolism of DM). Quinidine is a derivative of quinine, from the Peruvian Cinchona tree, as are chloroquine and |
hydroxychloroquine; all of which have been observed to reduce viral replication and cytokine release in vitro.

Chest Tightness:

E) Liquify mucus  E) Guaifenesin bid for mucus liquification
F) Mucus mobilization  F) Consider Warm Steam Vaporizer
G) Bronchopulmonary inflammation  G) Pulmicort Inhalational steroid
H) Deep terminal airway inflammation  H) Site-selective delivery of anti-inflammatory action into the terminal bronchopulmonary tree

Reactive Airway

3. Bronchial constriction  3. Levalbuterol also known as L-albuterol (Xopenex) and is the active form of the drug called R-albuterol. Fewer side effects than with racemic albuterol (which is a 50:50 mixture of R and S form). Pharmacologically acts via the beta-2 adrenergic receptor on bronchial smooth muscle thereby relaxing the distal extent of the bronchopulmonary tree. Can reduce the ‘tight lung’ perception, and via bronchodilation can facilitate mobilization of mucus and avoid inspisated mucus organization. Two puffs TID.
4. Beta-2 adrenergic receptor mediated immune skewing  4. Beta-2 adrenergic receptors are also expressed on circulating mononuclear cells. R albuterol binds to the beta-2 receptor and promotes conversion of cells of the TH1 proinflammatory phenotype to TH2 anti-inflammatory state [Ref; Weiner Harvard]. Two puffs TID.

Elevated Blood Pressure

3. Bioenergetics  3. Benfotiamine 300mg daily : Improved bioenergetics
4. Reduce MTX tissue injury  4. reduced tissue damage from MTX and other inciting agents
2. Vitamin D: 10,000

Blood pressure monitoring and correlate with headache.
units of D3 daily for promoting immune modulation/Tregs

Prepare for HDMTX-LR Protocol

Flow chart
