

Leonard H. Calabrese, DO

Department of Rheumatic and Immunologic Diseases,
Orthopedic & Rheumatologic Institute,
Cleveland Clinic

Tiphaine Lenfant, MD

Department of Rheumatic and Immunologic Diseases,
Orthopedic & Rheumatologic Institute Department
of Infectious Disease, Cleveland Clinic; Assistance
Publique des Hôpitaux de Paris, Université de Paris;
Hôpital européen Georges Pompidou, Service de
médecine interne, Paris, France

Cassandra Calabrese, DO

Department of Rheumatic and Immunologic Diseases,
Orthopedic & Rheumatologic Institute, and
Department of Infectious Disease, Cleveland Clinic

Cytokine storm release syndrome and the prospects for immunotherapy with COVID-19, part 3: The role of GM-CSF

Posted July 30, 2020

■ ABSTRACT

Granulocyte-macrophage colony-stimulating factor (GM-CSF) has been used experimentally in patients with acute respiratory distress syndrome. Recombinant GM-CSF administered by direct inhalation is currently being studied in a cohort of patients with advanced COVID-19.

■ INTRODUCTION

In the first two parts of this series, we focused on the basic immunobiology of severe COVID-19 disease and the role of inflammatory cytokines in driving respiratory damage, coagulopathy, end-organ failure, and death, which is idealized in **Figure 1**.^{1,2} We know that 90% of patients with COVID-19 recover, but we are also aware that in about 10% of patients the disease is progressive, and that this is heavily influenced by a growing number of risk factors, including attendant cardiovascular disease, obesity, hypertension, and age. This stage 3 of COVID-19, when the disease may rapidly progress to fatality, has been continuously elucidated and referred to as “cytokine storm,” given the presence of elevated levels of inflammatory cytokines and chemokines including interleukin 1 (IL-1), IL-6, tumor necrosis factor (TNF), granulocyte-macrophage colony-stimulating factor (GM-CSF), gamma interferon, and monocyte chemoattractant protein-1 (MCP-1).² Furthermore, quantitative studies of these biomarkers have demonstrated that some may discriminate mild self-limiting forms of COVID-19 from severe progressive forms of the disease.³ But

still lacking is an understanding of the precise choreography of cytokine storm and of which cytokine or cytokines are upstream drivers vs those that may be late-stage effectors or amplifiers. While IL-6 and IL-1 are both logical therapeutic targets and the preliminary data from uncontrolled studies have shown promise, inhibition has not been curative and is not without real safety concerns. This Curbside Consult will focus on another cytokine of growing interest, GM-CSF.

■ BASIC IMMUNOLOGY OF GM-CSF

GM-CSF is a complex cytokine and a member of the colony-stimulating superfamily. While it does have a role as a growth factor for myeloid cells, this is considered a lesser function compared with other cytokines such as granulocyte-stimulating colony factor (G-CSF) and macrophage-stimulating colony factor (M-CSF), and GM-CSF is now considered a central player in the integrated immune response and a central mediator of tissue inflammation.^{4,5}

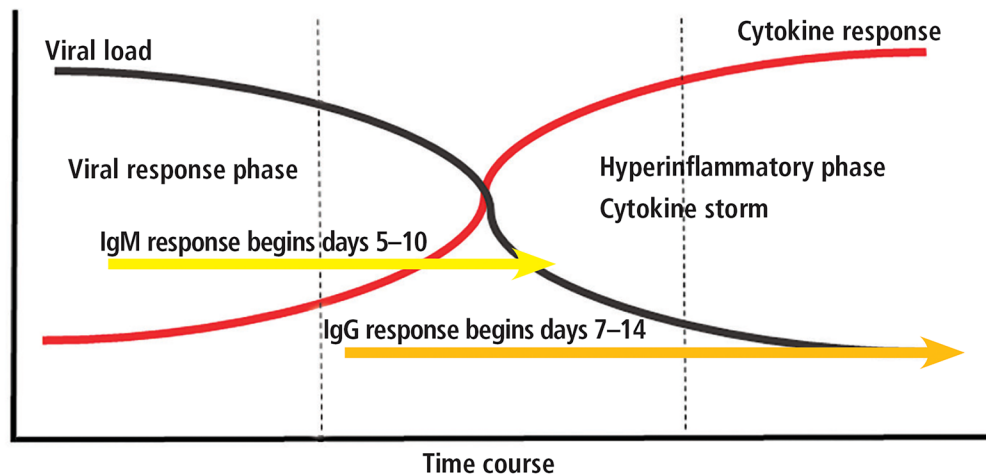
GM-CSF is produced by both hematopoietic (eg, T cells, B cells, macrophages, monocytes) and some viscerosomatic cells (type II alveolar epithelial cells, endothelial cells, fibroblasts) and can activate cells through the dimeric GM-CSF receptor expressed primarily on monocytes and macrophages but also on dendritic cells and other cells of the innate immune system.^{4,6} GM-CSF mediates its effects by signaling through the JAK-STAT pathway,⁷ a pathway inhibited by currently available therapeutics including tofacitinib and baricitinib. During inflammation GM-CSF serves two important roles. First, it functions to polarize mature myeloid cells into a pro-inflammatory phenotype capable of secreting other inflammatory cytokines such as IL-1, IL-6, and TNF, as well as a variety of chemokines governing trafficking of hema-

The statements and opinions expressed in COVID-19 Curbside Consults are based on experience and the available literature as of the date posted. While we try to regularly update this content, any offered recommendations cannot be substituted for the clinical judgment of clinicians caring for individual patients.

doi:10.3949/ccjm.87a.ccc057



	Stage 1	Stage 2	Stage 3
	Asymptomatic	Nonsevere symptomatic	Severe respiratory-inflammatory
Immune response over time:	Innate immune activation	Adaptive immune activation	Cytokine release syndrome
Self-limiting in 80%	Viral engagement of PAMPs	Generation of specific antibodies and T-cell response	IL-1, IL-6, TNF, GM-CSF, IFN-gamma, others
Severe in 15%–20%	Low type 1 IFN	Release of DAMPs	Coagulopathy
Fatal in 1%–2%			Complement



DAMPs= damage-associated molecular patterns; GM-CSF = granulocyte macrophage colony-stimulating factor; IFN = interferon; IgM = immunoglobulin M; IL-1 = interleukin 1; IL-6 = interleukin 6; PAMPs = pathogen-associated molecular patterns; TNF = tumor necrosis factor

Figure 1. Three stages of COVID-19 disease.

topoietic cells to areas of inflammation. GM-CSF also serves to activate dendritic cells to prime T cells during antigen-specific responses and thus links the myeloid compartment and pathogenic T cells (TH-1 and TH-17) in a positive feedback loop, capable of propagating inflammation and tissue injury.

Of particular relevance and importance regarding the immunobiology of GM-CSF in COVID-19 disease is its complex role in lung homeostasis and inflammation. In the healthy lung, GM-CSF has a critical role for maintaining the maturation and function of alveolar macrophages and surfactant metabolism⁵ and is required to maintain pulmonary function, as well as contributing to lung sentinel cell-mediated immunity. GM-CSF also appears central in driving inflammation locally and systemically,⁵ and experimental models of acute lung injury support this hypothesis, demonstrating that resident alveolar macrophages secrete a variety of inflammatory cytokines that lead to the influx of innate cells including neutrophils, further amplifying the activation of alveolar epithelial cells and tissue damage.⁸

The capacity of GM-CSF to amplify inflammatory response within the lung and its systemic effects, mediated by linking the release of upstream inflammatory cytokines such as IL-1, IL-6, and TNF across monocytes and macrophages and activated T cells in a positive feedback loop, highlight its potential importance in driving systemic inflammation and disease. However, as noted above, adding to the complexity is that GM-CSF is also a critical cytokine for healthy pulmonary function and is necessary for the maturation and maintenance of alveolar macrophages; and in some experimental models, it confers resistance to viral respiratory balance, underscoring that all putative inflammatory cytokines also play roles in integrated host defense.⁴

The central role of GM-CSF in the inflammatory response and because GM-CSF appears to be upstream of other key inflammatory cytokines invites targeting strategies against it in effort to down-modulate states of hypercytokinemia. GM-CSF levels are generally extremely low or undetectable in healthy individuals and are detectable in blood of patients

TABLE 1

Overview of anti-GM-CSF drugs, mechanisms, and studies

Mechanism of action	Drug	Study population	Phase
Humanized immunoglobulin G1 (IgG1) monoclonal antibody targeting granulocyte-macrophage colony-stimulating factor (GM-CSF)	TJ003234	Healthy individuals COVID-19	1 1b/2
	Gimsilumab	Ankylosing spondylitis COVID-19	1 2
	Lenzilumab	Chronic myelomonocytic leukemia Relapsed or refractory large B-cell lymphoma COVID-19	1 1/2 3
	Otilimab	Rheumatoid arthritis Inflammatory arthritis COVID-19	2b 2 2
	Namilumab	Rheumatoid arthritis Chronic plaque psoriasis Spondyloarthritis	2 2 2
	Mavrilimumab	Rheumatoid arthritis	2b
		Giant cell arthritis	2
		COVID-19	2
Humanized IgG4 monoclonal antibody targeting GM-CSF receptor alpha			
Betac-receptor-specific, fully human IgG4 monoclonal antibody (inhibitor of IL-3-, GM-CSF-, and IL-5-mediated functions)	CSL311	Asthma	1

with inflammatory manifestations of COVID-19,⁹ and CD14+CD16+ monocytes, a rich source of GM-CSF, are expanded in such patients as well.¹⁰ As noted in **Table 1**, GM-CSF targeting is actively being investigated in a variety of autoimmune diseases and has been successfully studied in a mouse model of cytokine release from CAR-T cells.¹¹ Based on these data, investigation of targeting GM-CSF in COVID-19 has commenced.

TARGETING GM-CSF IN AUTOIMMUNE AND INFLAMMATORY DISEASES AND COVID-19

There is a rich pipeline of biologic therapeutics that target GM-CSF directly or that targeting the GM-CSF receptor, and none are currently approved for any indication, though they are under investigation in many conditions, including rheumatoid arthritis, spondyloarthritis, giant cell arteritis, psoriasis, and certain malignancies (**Table 1**). To date, there are 8 clinical trials of varying design and size (**Table 2**) focused mainly on patients with advancing COVID-19 that does not require mechanical ventilation.

In the most detailed clinical report to date, De

Luca et al¹² described a single-center, nonrandomized, prospective cohort study in which 13 non-mechanically ventilated patients with pneumonia, hypoxia, and signs of systemic inflammation were treated with the anti-GM-CSF receptor mavrimumab given as single intravenous dose. They compared this group to a cohort of 26 patients treated simultaneously with similar baseline characteristics receiving the same standard of care without mavrimumab. At 28 days, no patients in the anti-GM-CSF group died, whereas 7 (27%) of 26 in the control group died. The agent was well tolerated with no infusion reactions.

This study, while encouraging, is cautionary on the basis of its design and the inherent risks of unobserved confounders. Robust trials are under way and will hopefully provide more definitive evidence (**Table 2**). A second single-center study from Mayo Clinic released but not yet peer-reviewed also described impressive results.¹³ In this observational noncontrolled study, 12 patients with COVID-19 pneumonia and risk factors for poor outcome were treated with 3 doses of lenzilumab, and clinical improvement was noted in 92% with a median time-to-discharge of

TABLE 2

Studies of anti-GM-CSF in COVID-19 (as of July 5, 2020)

Mechanism of action	Molecule	Trial	Trial name	Drugs	Phase	Primary end point
Immunoglobulin G1 (IgG1) monoclonal antibody targeting granulocyte-macrophage colony-stimulating factor (GM-CSF)	TJ003234	NCT04341116	A Phase 1b/2, Randomized, Double-Blind, Placebo-Controlled, Multi-Center Study to Evaluate the Safety and Efficacy of TJ003234 in Subjects With Severe Coronavirus Disease 2019 (COVID-19)	TJ003234 (3 or 6 mg/kg, single infusion) vs placebo	1b/2	Proportion (%) of patients experiencing deterioration in clinical status (changes from baseline on day 14)
	Gimsilumab	NCT04351243	A Multi-Center, Adaptive, Randomized, Double-blind, Placebo-controlled Study to Assess the Efficacy and Safety of Gimsilumab in Subjects With Lung Injury or Acute Respiratory Distress Syndrome Secondary to COVID-19 (BREATHE)	Gimsilumab (high dose day 1 + gimsilumab low dose day 8) vs placebo	2	Mortality at day 43
	Lenzilumab	NCT04351152	A Phase 3 Randomized, Placebo-Controlled Study of Lenzilumab in Hospitalized Patients With COVID-19 Pneumonia	Lenzilumab vs standard of care	3	Incidence of invasive mechanical ventilation and/or mortality (up to 28 days)
	Namilumab	EudraCT 2020-001684-89; ISRCTN40580903	CATALYST - A randomised phase II proof of principle multi-arm multi-stage trial designed to guide the selection of interventions for phase III trials in hospitalised patients with COVID-19 infection	Namilumab vs gentuzumab vs infliximab	2	SpO ₂ /FIO ₂ measured from randomization to day 14, hospital discharge, death
	Otilimab	NCT04376684	A Randomized, Double-blind, Placebo-controlled, Study Evaluating the Efficacy and Safety of Otilimab IV in Patients With Severe Pulmonary COVID-19-Related Disease	Otilimab as an IV infusion + standard of care vs placebo as IV infusion + standard of care	2	Proportion of participants alive and free of respiratory failure at day 28
IgG4 monoclonal antibody targeting GM-CSF receptor alpha	Mavrilimumab	NCT04447469	Study of Mavrilimumab (KPL-301) in Participants Hospitalized With Severe Corona Virus Disease 2019 (COVID-19) Pneumonia and Hyper-inflammation	Mavrilimumab vs placebo	2/3	Proportion of participants alive and without respiratory failure at day 15
		NCT04399980	Mavrilimumab to Reduce Progression of Acute Respiratory Failure in Patients With Severe COVID-19 Pneumonia and Systemic Hyper-inflammation (RCT)	Mavrilimumab vs placebo	2	Proportion of patients alive and off of oxygen at day 14
		NCT04397497	A Randomized, Double Blind, Placebo-controlled Trial of Mavrilimumab for Acute Respiratory Failure Due to COVID-19 Pneumonia With Hyper-inflammation (the COMBAT-19 Trial)	Mavrilimumab (single-dose IV) vs placebo	2	Reduction in the dependency on oxygen supplementation (within day 14 of treatment)

5 days. Detailed analysis of inflammatory biomarkers and cytokines also improved, and there were no treatment-related adverse events. In summary, while encouraging, these data make completion of randomized trials even more urgent.

■ SAFETY CONCERNS

Targeting GM-CSF carries with it a number of safety concerns derived from both preclinical modeling and human disease. GM-CSF is an integral part of the integrated immune response and, as noted above, is important in maintaining defense against infections. Thus, any use in COVID-19 must weigh this balance. Fortunately, in the collective clinical experience reported across many trials, targeting GM-CSF has been well tolerated with no major safety signals.⁵ Of high concern in all GM-CSF trials is the specter of pulmonary alveolar proteinosis, a rare disease characterized by accumulation of proteinaceous material in the lungs; some patients with this disease have autoantibodies directed against GM-CSF.¹⁴ This toxicity has been monitored as an event of special interest in clinical trials for autoimmune disease, and has not been observed to date.

Finally, a recurrent theme with all clinical trials in COVID-19 inflammatory disease is timing. Giving an immunomodulator too early in the course of the infection will compromise an evolving adaptive immune response and may theoretically prolong the disease and compromise the outcome.^{2,8} At the moment there is no definitive single biomarker or set of biomarkers capable of identifying the optimal timing of such interventions. How this therapy will fare used singly or in combination with antiviral therapy or glucocorticoids, which are at the moment in ascendency for treatment of COVID-19, will require larger and more carefully conducted trials.

■ RATIONALE FOR GIVING GM-CSF IN COVID-19

Consistent with the complex immunobiology of GM-CSF is a seemingly contradictory strategy of administering GM-CSF to patients with late-stage COVID-19 disease. This theory is based on the role of GM-CSF previously noted including its centrality in maintaining critical lung function and local anti-microbial defense.⁵ Numerous preclinical models appear to support this concept, and GM-CSF, an approved biologic agent, has been utilized experimentally in patients with acute respiratory distress syndrome. A trial of recombinant GM-CSF administered by direct inhalation in a cohort of patients with advanced COVID-

19 (NCT04326920) is under way.

These divergent strategies underscores our incomplete understanding of the immunobiology of COVID-19 and its most sinister clinical outcome, acute respiratory distress syndrome, and the need to carefully study such strategies in controlled settings.

■ REFERENCES

1. Calabrese C, Rajendram P, Sacha G, Calabrese L. Practical aspects of targeting IL-6 in COVID-19 disease. *Cleve Clin J Med* 2020. [Epub ahead of print] doi:10.3949/ccjm.87a.ccc018
2. Calabrese LH. Cytokine storm and the prospects for immunotherapy with COVID-19. *Cleve Clin J Med* 2020; 87(7):389–393. doi:10.3949/ccjm.87a.ccc008
3. Wang F, Hou H, Luo Y, et al. The laboratory tests and host immunity of COVID-19 patients with different severity of illness. *JCI Insight* 2020; 5(10):e137799. doi:10.1172/jci.insight.137799
4. Becher B, Tugues S, Greter M. GM-CSF: from growth factor to central mediator of tissue inflammation. *Immunity* 2016; 45(5):963–973. doi:10.1016/j.immuni.2016.10.026
5. Lotfi N, Thome R, Rezaei N, et al. Roles of GM-CSF in the pathogenesis of autoimmune diseases: an update. *Front Immunol* 2019; 10:1265. doi:10.3389/fimmu.2019.01265
6. Wicks IP, Roberts AW. Targeting GM-CSF in inflammatory diseases. *Nat Rev Rheumatol* 2016; 12(1):37–48. doi:10.1038/nrrheum.2015.161
7. Mehta P, Porter JC, Manson JJ, et al. Therapeutic blockade of granulocyte macrophage colony-stimulating factor in COVID-19-associated hyperinflammation: challenges and opportunities. *Lancet Respir Med* 2020. [Epub ahead of print] doi:10.1016/S2213-2600(20)30267-8
8. Huang C, Wang Y, Li X, et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet* 2020; 395(10223):497–506. doi:10.1016/S0140-6736(20)30183-5
9. Zhou Y, Fu B, Zheng X, et al. Aberrant pathogenic GM-CSF+ T cells and inflammatory CD14+CD16+ monocytes in severe pulmonary syndrome patients of a new coronavirus. *bioRxiv* 2020. doi:10.1101/2020.02.12.945576
10. Sterner RM, Sakemura R, Cox MJ, et al. GM-CSF inhibition reduces cytokine release syndrome and neuroinflammation but enhances CAR-T cell function in xenografts. *Blood* 2019; 133(7):697–709. doi:10.1182/blood-2018-10-881722
11. DeLuca G, Cavalli G, Caampochiaro C, et al. GM-CSF blockade with mavrilimumab in severe COVID-19 pneumonia and systemic hyperinflammation: a single center, prospective cohort study. *Lancet Rheumatol* 2020. [Epub ahead of print] doi:10.1016/S2665-9913(20)30170-3
12. Temesgen Z, Assi M, Vergidis P, et al. First clinical use of lenzilumab to neutralize GM-CSF in patients with severe COVID-19 pneumonia. *medRxiv* 2020. [Epub ahead of print] doi:10.1101/2020.06.08.20125369
13. Trapnell BC, Nakata K, Bonella F, et al. Pulmonary alveolar proteinosis. *Nat Rev Dis Primers* 2019; 5(1):16. doi:10.1038/s41572-019-0066-3

Correspondence: Leonard H. Calabrese, DO, Department of Rheumatic and Immunologic Diseases, A50, Cleveland Clinic, 9500 Euclid Avenue, Cleveland, OH 44195; calabrl@ccf.org