

thrombotic events, and 19 patients (4.8%) had documented VTE despite routine anticoagulant thromboprophylaxis. Although many of the markers of inflammation and coagulation correlated with clinical outcomes, D-dimer emerged as the most useful. Thus, D-dimer levels over 2500 ng/mL, levels over 5 times higher than the upper limit of normal, were associated with sevenfold, twofold, and 15-fold increases in the risk of thrombosis, progression to critical illness, and mortality, respectively.

What are the new findings? First, the 7.6% rate of VTE in critically ill COVID-19 patients in this study is lower than what has previously been reported and is more in line with the rate of VTE found in critically ill patients without COVID-19.⁶ Second, with anticoagulant thromboprophylaxis, the authors report a rate of major bleeding of 5.6% in critically ill patients with COVID-19 and identify a baseline platelet count below $150 \times 10^9/L$ and D-dimer levels over 2500 ng/mL as independent predictors of a threefold increase in the risk of major bleeding.

The strengths of this study include the relatively large sample size and the comprehensive reporting of clinical outcomes. However, like many previous studies in patients with COVID-19, there are limitations, including the retrospective design as well as the potential for bias in case ascertainment and underestimation of VTE rates because of the inability to image all critically ill patients with suspected events.

The triggers responsible for COVID-19-associated coagulopathy remain elusive. Potential triggers include cytokine-induced overexpression of tissue factor, endothelial dysfunction with loss of its antithrombotic phenotype, stasis, and hypoxia (see figure). This study confirms the correlation between markers of inflammation and coagulation and supports the concept that inflammation is a major driver of the hypercoagulable state. An inflammation-driven hypercoagulable state has also been reported in critically ill patients with viral pneumonia caused by H1N1 or SARS-CoV-1.⁷ The VTE rate in such patients ranged from 5% to 25%, which is similar to the rates observed in patients with COVID-19.⁸⁻¹⁰ Although intensified anticoagulation regimens may reduce the risk of a thrombotic event, the results of this study raise the possibility that they

may increase major bleeding rates to unacceptable levels in critically ill patients. As the world waits for the second wave of COVID-19, randomized trials comparing anticoagulation dosing strategies are urgently needed. Fortunately, several such trials are under way.

Conflict-of-interest disclosure: The authors declare no competing financial interests. ■

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TRANSPLANTATION

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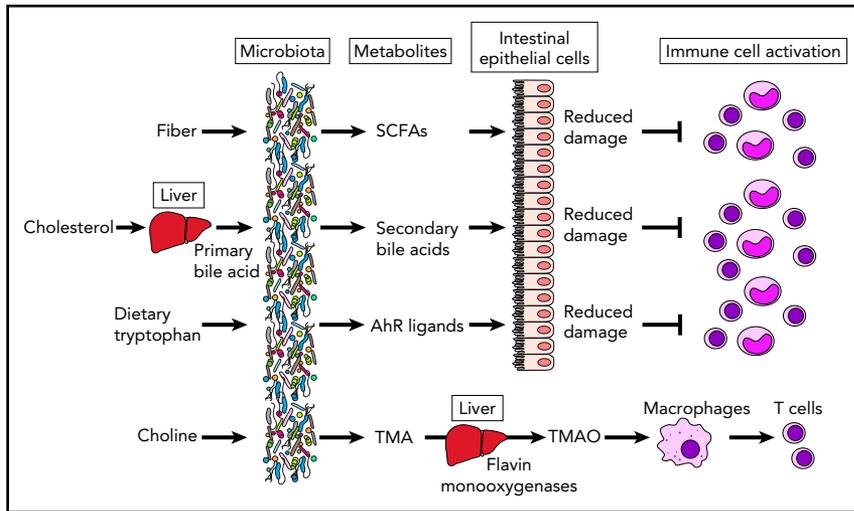
Too much TMAO and GVHD

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In this issue of *Blood*, Wu and colleagues¹ report that trimethylamine N-oxide (TMAO), an intestinal microbiome-dependent metabolite, worsens graft-versus-host disease (GVHD). They further found that TMAO induces M1 polarization of bone marrow-derived macrophages via the nucleotide-binding oligomerization domain-like receptor protein 3 (NLRP3). TMAO is produced by hepatic processing of intestinal bacteria-derived trimethylamine (TMA) following metabolism of certain dietary nutrients, including choline, lecithin, L-carnitine, and γ -butyrobetaine.

GVHD is an alloreactive, donor T-cell-mediated inflammatory disease that occurs often after allogeneic hematopoietic stem cell transplantation (allo-HSCT) and can involve the skin, liver, and gastrointestinal tract. The success of allo-HSCT continues to be significantly limited by GVHD, even though the field has seen improvements in both prevention and

clinical management. Targeting intestinal bacteria may be a promising novel approach to further improve allo-HSCT clinical outcomes. A disrupted intestinal microbiome, also known as dysbiosis, occurs frequently in allo-HSCT recipients and can manifest as loss of obligate anaerobic commensal bacteria, expansion of pathogenic bacteria, and reduced microbiome



Microbial metabolites modulate immune cell responses in GVHD. SCFAs derived from fiber and secondary bile acids protect epithelial cells from damage, leading to suppression of immune cell activation. AhR ligands derived from dietary tryptophan can also suppress immune cell activation through reduction of intestinal epithelial damage in GVHD. TMAO is produced by hepatic processing by flavin monooxygenases of TMA following bacterial metabolism of choline and other dietary components. TMAO augmented alloreactive T-cell proliferation via M1 polarization of bone marrow-derived macrophages.

diversity. Microbiome injury, in turn, has been strongly associated with acute GVHD and reduced overall survival.² However, the mechanisms by which GVHD severity is modulated by intestinal bacteria are not fully understood. Recent studies have uncovered relationships between the microbiome and microbiome-derived metabolites in both mouse models and the clinical setting. Metabolites such as short-chain fatty acids (SCFAs) can support tissue repair and regulate immune cell activation.^{3,4} Bile acids have also been identified as potent regulators of the immune system through the inhibition of the NLRP3-dependent inflammasome pathway.⁵ Aryl hydrocarbon receptor ligands (AhR) such as indole-3-aldehyde can directly regulate innate immunity and modulate T helper (Th)17 responses and promote tolerance via regulatory T cells and type 1 regulatory cells.⁶ In a metabolomic analysis of patients with acute GVHD, Michonneau and colleagues⁷ recently demonstrated that acute GVHD is associated with alterations in bile acids and decreased production of AhR ligands by the microbiome (see figure).

Wu and colleagues focused on a particularly well studied microbiome-derived metabolite, TMAO, which has been associated with cardiovascular complications arising from atherosclerosis and thrombosis, but has not been known to affect GVHD. TMAO can induce vascular inflammation and endothelial dysfunction

by formation and activation of NLRP3 inflammasomes in endothelial cells. The results show a novel link between TMAO and GVHD in mouse models and further demonstrate that TMAO induces secretion of M1-like cytokines from bone marrow-derived macrophages in an NLRP3-dependent fashion. TMAO augments alloreactive T-cell proliferation and Th1 subtype differentiation in mice with GVHD but not in an in vitro T-cell culture system, leading to an investigation to determine the role of TMAO in macrophage polarization.

NLRP3 has been shown to have conflicting contributions to GVHD. NLRP3-mediated signaling is essential for intestinal commensal bacterial products and the damage-associated molecular pattern that leads to interleukin-1 production.⁸ SCFAs, however, can also activate the NLRP3 inflammasome, and signaling in host nonhematopoietic cells has been found to be critical for reducing the severity of GVHD.⁴ The seemingly opposing effects of NLRP3 on GVHD may be related to differential activation in distinct host tissue compartments or immune cell populations, and further investigation is needed to better clarify the effects of microbiome-derived products in NLRP3-modulated GVHD.

High plasma levels of TMAO are associated with a variety of adverse outcomes, including cardiovascular diseases, glucose intolerance, kidney damage, obesity, and

diabetes, although other clinical factors, such as age, race, and cholic acid and bile salt levels, have also been associated. In patient who undergo allo-HSCT, plasma TMAO levels could be a novel biomarker for predicting GVHD, and an evaluation is warranted, given the preclinical findings by Wu and colleagues. One question is how TMAO levels are affected by anorexia and other dietary alterations, as well as antibiotic use, all of which are common in the setting of allo-HSCT.

If a clinical association is found, how could we target TMAO in allo-HSCT patients? Diet is known to play a key role in TMAO formation. TMAO can be found in fish or may be secondarily produced by bacterial metabolism of red meat or eggs, which are rich in choline and L-carnitine. Thus, strategies to reduce TMAO include limiting dietary intake of animal fat and protein, consuming a vegetarian diet, use of prebiotics/probiotics, and even supplementation with pistachios.⁹ Identification of bacterial subsets responsible for increased TMAO is another approach. The Firmicutes/Bacteroidetes ratio has been used to predict TMAO concentrations in plasma. Certain specific bacteria have also been identified as important in TMAO production, including *Anaerococcus hydrogenalis*, *Clostridium asparagiforme*, *Clostridium hathewayi*, *Clostridium sporogenes*, *Edwardsiella tarda*, *Escherichia fergusonii*, *Proteus penneri*, and *Providencia rettgeri*.¹⁰ Targeting those may be a challenging strategy, however, because they are taxonomically diverse and have not been reported to be associated with increased GVHD. Broad-spectrum antibiotics can almost totally suppress the production of TMAO by eliminating the gut microbiome. However, the levels of TMAO return to normal after withdrawal of antibiotics, and the long-term use of broad-spectrum antibiotics is not feasible given the likely selection for resistant bacteria. In the current study, the Wu et al demonstrated that administration of 3,3-dimethyl-1-butanol, an inhibitor of microbial TMA lyases, is effective in ameliorating aggravated GVHD. Blockade of hepatic flavin monooxygenases, which convert TMA to TMAO, is also being investigated in a clinical trial (www.clinicaltrials.gov #NCT03152097).

In summary, the findings in this study suggest that the microbial metabolite TMAO can be a potent contributor to the

severity of acute GVHD. Further studies in both animal models and human biospecimens could clarify the importance of the role played by TMAO with respect to the dynamics of the microbiome and microbial metabolites and the modulation of the immune system in the development of acute GVHD.

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