Diabetes mellitus is associated with an increased risk for cardiovascular disease, but the link between hyperglycemia and atherothrombotic disease is not completely understood. Patients with diabetes often show hyporesponsiveness to antiplatelet therapies, and it has been suggested that hyperreactive reticulated platelets underlie this altered therapeutic response. In this issue of the *JCI*, Kraakman et al. uncover a previously unknown link between hyperglycemia and enhanced platelet production and reactivity. The authors demonstrate that high blood glucose levels trigger neutrophil release of S100 calcium-binding protein A8/A9 (S100A8/A9), which binds to the receptor for advanced glycation end products (RAGE) on Kupffer cells, ultimately leading to increased thrombopoietin (TPO) production in the liver. TPO causes megakaryocyte proliferation and increased platelet production. This study demonstrates the importance of glycemic control and identifies potential therapeutic targets in the normalization of platelet numbers and function in diabetes.
Sugar makes neutrophils RAGE: linking diabetes-associated hyperglycemia to thrombocytosis and platelet reactivity

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Diabetes mellitus is associated with an increased risk for cardiovascular disease, but the link between hyperglycemia and atherothrombotic disease is not completely understood. Patients with diabetes often show hypersensiveness to antiplatelet therapies, and it has been suggested that hyperreactive reticulated platelets underlie this altered therapeutic response. In this issue of the JCI, Kraakman et al. uncover a previously unknown link between hyperglycemia and enhanced platelet production and reactivity. The authors demonstrate that high blood glucose levels trigger neutrophil release of S100 calcium-binding protein A8/A9 (S100A8/A9), which binds to the receptor for advanced glycation end products (RAGE) on Kupffer cells, ultimately leading to increased thrombopoietin (TPO) production in the liver. TPO causes megakaryocyte proliferation and increased platelet production. This study demonstrates the importance of glycemic control and identifies potential therapeutic targets in the normalization of platelet numbers and function in diabetes.

Introduction
Diabetes mellitus is a disease characterized by an inability of the body to produce or respond to insulin, resulting in hyperglycemia. Type 1 diabetes (T1D) is caused by autoimmune destruction of insulin-producing pancreatic β cells (also known as juvenile diabetes). Type 2 diabetes (T2D) results from progressive resistance to insulin, often found in older individuals, and is associated with excessive body weight, poor diet, and lack of exercise (also known as adult-onset diabetes). T2D accounts for approximately 90% of all diabetes cases, affecting almost 10% of the adult population (1). Uncontrolled diabetes results in various pathologies, including thrombotic complications such as myocardial infarction and stroke. The risk of cardiovascular disease (CVD) can persist even in diabetic patients whose blood glucose levels are controlled (2).

But how is diabetes linked to an increased risk for CVD? The answer, at least to some extent, seems to lie in dysregulation of the number and function of platelets. The main physiological role of platelets is in the prevention of blood loss at sites of vascular injury (hemostasis). To guard the integrity of the vasculature, platelets are equipped with specialized surface-expressed adhesion and agonist receptors and a unique intracellular signaling machinery, all of which are critical for adhesion and hemostatic plug formation under high shear stress conditions such as are found in arterioles and arteries. However, these activating mechanisms are highly sensitive and pose a risk of platelet hyperreactivity, which can lead to the development of atherosclerotic plaques (atherosclerosis) and the thrombotic complications associated with plaque rupture (atherothrombosis). Under physiological conditions, platelet numbers are tightly controlled, and the peripheral platelet count reflects both the rate of platelet production and the clearance of platelets from the blood. Once released into the circulation by megakaryocytes (MKs) within the bone marrow, platelets circulate for approximately 10 days before getting cleared, primarily by phagocytes of the reticuloendothelial system. Platelet generation (thrombopoiesis) depends on the proliferation and differentiation of MKs, which are driven by thrombopoietin (TPO) produced by various cell types, including marrow stromal cells and hepatocytes (3). Platelets newly released into the blood are larger in size, contain more mRNA (“reticulated”), and are more reactive to agonist stimulation than are older cells (4). These highly reactive reticulated platelets may be the key to understanding the enhanced risk for CVD and thrombotic events in diabetes.

It is well established that diabetes leads to significant changes in blood platelets, including increased mean platelet volume (MPV) and greater numbers of reticulated platelets (5), a phenotype often documented in T2D patients. Consistently, circulating platelets in diabetic patients are hyperreactive to agonist stimulation and are more likely to form aggregates with circulating leukocytes, thereby increasing inflammation in these patients (6). Thus, it is proposed that a high reticulated platelet count contributes to the high risk for thrombotic events in diabetic patients (Figure 1) and that normalization of platelet function could positively affect outcomes in these patients (7). Current strategies to prevent atherothrombosis include the use of inhibitors of cyclooxygenase-1 (aspirin) and inhibitors of the platelet ADP receptor P2Y12 such as clopidogrel, prasugrel, or ticagrelor. While effective in most individuals, significant pharmacological hypersensiveness to these inhibitors has been documented for various disease states, including diabetes. Therefore, it is critical

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to identify the mechanisms underlying enhanced reticulated platelet counts and platelet reactivity in diabetes to better tailor antiplatelet therapies for these patients.

In this issue, Kraakman and colleagues describe a novel link between hyperglycemia and increased reticulated platelet counts and thrombocytosis (8). Their elegant mechanistic studies identify various approaches with the potential to normalize platelet function in diabetes. Moreover, these studies highlight the complexities of metabolic perturbations on various organ and cell systems throughout the body.

**Neutrophils drive hepatic TPO production and thrombopoiesis in diabetes**

In mice, T1D can be modeled using treatment with the pancreatic β cell toxin streptozotocin (STZ). Kraakman and colleagues demonstrate that STZ-treated diabetic mice exhibit robust thrombocytosis, a substantial increase in reticulated platelet numbers, upregulation of P-selectin on the platelet surface, and higher numbers of platelet-neutrophil and platelet-monocyte aggregates in the blood. In accordance with elevated platelet counts, diabetic mice have higher levels of plasma TPO. Under physiological conditions, hepatocytes are a major source for circulating TPO. Analysis of livers in STZ-treated mice uncovered greater numbers of resident macrophages (Kupffer cells). Additionally, Kupffer cells from STZ-treated animals produce more IL-6 than do Kupffer cells from normoglycemic mice. Kraakman et al. suggest that this local IL-6 production then induces TPO production in hepatocytes. Finally, the increase in Kupffer cell abundance and IL-6 expression is shown to be in response to neutrophil-derived S100 calcium-binding protein A8/A9 (S100A8/A9, also known as myeloid-related protein 8/14) binding to the receptor for advanced glycation end products (RAGE) (Figure 1). Kraakman and colleagues thus connect diabetes-induced hyperglycemia to thrombocytosis and an increase in reticulated platelets, independently of any direct action of glucose on MK function and/or platelet survival.

Neutrophils are glucose-sensitive inflammatory cells and have previously been shown to secrete the proinflammatory protein S100A8/A9 under conditions of hyperglycemia (9). Plasma S100A8/A9 is a member of the large family of calcium-binding proteins that is predominantly found in the cytoplasm of blood cells and in plasma. Elevated plasma levels of S100A8/A9 are considered a biomarker for chronic inflammatory disorders such as inflammatory bowel disease and rheumatoid arthritis (10). In addition, enhanced release of
S100A8/A9 from neutrophils has been shown to modulate myelopoiesis (9). Kraakman et al. use a variety of genetic and therapeutic approaches to demonstrate the connection between hyperglycemia, neutrophil-derived S100A8/A9, and thrombocytosis. Thrombocytosis is normalized in STZ-treated mice lacking S100A9 (which also lack S100A8) or treated with an inhibitor of S100A8/A9 (ABR-215757). Moreover, depletion of circulating neutrophils in STZ-treated animals normalizes thrombocytosis. Consistent with these findings in mice, plasma S100A8/A9 levels are elevated in human T2D patients (8) and in T1D patients with coronary artery disease (CAD) (9). Unfortunately, total and reticulated platelet numbers were not reported for the T1D patients. Finally, Kraakman and colleagues demonstrate that treatment with the S100A9 inhibitor ABR-215757 ameliorates diabetes-induced atheroprogession in STZ-treated mice. Altogether, these results confirm a role for S100A8/A9 in CVD and implicate these proteins in the regulation of hyperglycemia-induced reticulated thrombocytosis in mice.

Remaining questions and future directions

The studies by Kraakman et al. suggest an intriguing and novel mechanism for how hyperglycemia in diabetes patients is coupled to alterations in MK biology, leading to thrombocytosis and an increase in the reticulated platelet count. These studies provide important answers to questions regarding the molecular mechanisms that underlie altered platelet function in diabetes but also raise a number of important questions that should be addressed in the future. Arguably, the most important question is, how well do these studies in mice translate to humans with diabetes? Compared with the marked thrombocytosis observed in STZ-treated mice (~2-fold increase in platelet counts), patients with T1D do not typically present with thrombocytosis (11), and increases in platelet counts are variable in patients with T2D. Diabetic patients, however, do show a marked increase in reticulated platelet counts, even in the absence of thrombocytosis. For example, T2D patients in the Kraakman et al. study had approximately $3.2 \times 10^{10}$ reticulated platelets per liter of blood compared with approximately $1.4 \times 10^{10}$.
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