

Young NS, Bacigalupo A, Marsh JC: Aplastic anemia: pathophysiology and treatment. *Biol Blood Marrow Transplant* 16:S119, 2010. [A discussion of the role of the immune system and telomerase mutations in aplastic anemia.]

Bleeding Disorders

Arepally GM, Ortel TL: Heparin-induced thrombocytopenia. *Annu Rev Med* 61:77, 2010. [A discussion of pathogenesis, clinical features, diagnostic criteria, and therapeutic approaches in HIT.]

De Meyer SF, Deckmyn H, Vanhoorelbeke K: von Willebrand factor to the rescue. *Blood* 113:5049, 2009. [An update on the molecular pathogenesis and treatment of vWD.]

Noris M, Remuzzi G: Atypical hemolytic uremic syndrome. *N Engl J Med* 361:1676, 2009. [An article focused on the role of excessive activation of the alternative complement pathway in some forms of HUS.]

Pawlinski R, Mackman N: Cellular sources of tissue factor in endotoxemia and sepsis. *Thromb Res* 125(S1):S70, 2010. [An overview of the role of cellular procoagulants in DIC associated with bacterial infection.]

Stasi R: Immune thrombocytopenia: pathophysiologic and clinical update. *Semin Thromb Hemost* 38:454, 2012. [Discussion of the role of T cells in immune thrombocytopenia and treatment with immunomodulatory agents and growth factors.]

Zhou Z, Nguyen TC, Guchhait P, et al: Von Willebrand factor, ADAMTS-13, and thrombotic thrombocytopenia purpura. *Semin Thromb Hemost* 36:71, 2010. [A review focused on the role of vWF deregulation and ADAMTS 13 deficiency in TTP.]