• Hemophilia A, Classic hemophilia, Factor VIII (eight) deficiency, or Hemophilia A carrier
• Hemophilia B, Christmas disease, Factor IX (nine) deficiency, or Hemophilia B carrier
• von Willebrand (vWD) disease: Type 1, Type 2 (2A, 2B, 2M, and 2N), Type 3, and other
• Factor I (one) deficiency, including fibrinogen deficiency; afibrinogenemia; hypofibrinogenemia and dysfibrinogenemia
• Factor II (two) deficiency, including prothrombin deficiency; hypoprothrombinemia and dysprothrombinemia
• Factor V (five) deficiency
• Factor V-VIII (five and eight) deficiency
• Factor VII (seven) deficiency
• Factor X (ten) deficiency
• Factor XI (eleven) deficiency or Hemophilia C
• Factor XIII (thirteen) deficiency
• Platelet disorder
  o Glanzmann's thrombasthenia (GT)
  o Bernard Soulier syndrome (BSS)
  o Grey platelet syndrome (Alpha storage pool disease; Alpha SPD; GPS)
  o Hermansky-Pudlak syndrome (HPS)
  o Inherited thrombocytopenia
  o Platelet storage pool defect (SPD; platelet secretion disorder)
  o Platelet release defect
  o Other platelet disorder