Coagulopathy

• Specify type:
  - Disseminated intravascular coagulation
  - Hereditary factor VIII deficiency
  - Hereditary factor IX deficiency
  - Von Willebrand’s disease
  - Hereditary factor XI deficiency
  - Hereditary deficiency of other clotting factors
  - Acquired coagulation factor deficiency
  - Primary thrombophilia
    ° Activated protein C resistance
    ° Prothrombin gene mutation
    ° Other primary thrombophilia
  - Other thrombophilia
    ° Antiphospholipid syndrome
    ° Lupus anticoagulant syndrome
  - Other Specified coagulation defects

• Document any associated diagnoses/conditions

• Hemorrhagic disorder due to circulating anticoagulants
  - Due to intrinsic circulating anticoagulants, antibodies, or inhibitors
    ° Acquired hemophilia
    ° Antiphospholipid antibody with hemorrhagic disorder
    ° Other hemorrhagic disorder due to intrinsic circulating anticoagulants, antibodies, or inhibitors
  - Hemorrhagic disorder due to extrinsic circulating anticoagulants
    ° Document specific drug, if drug-induced