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

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