Acquired hemophilia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Acquired hemophilia. ORPHA:73274

A rare hemorrhagic disorder due to an acquired coagulation factor defect characterized by sudden, spontaneous, and often severe bleeding, manifesting with skin, muscle and mucuous membrane hemorrhages, in persons without a previous bleeding tendency. Additional symptoms may include epistaxis, gastrointestinal and/or urogenital bleeding, spontaneous bruising, melena, and hematuria.