Immune thrombocytopenic purpura

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Immune thrombocytopenic purpura. ORPHA:3002

Immune thrombocytopenic purpura (or immune thrombocytopenia; ITP) is an autoimmune coagulation disorder characterized by isolated thrombocytopenia (a platelet count <100,000/microL), in the absence of any underlying disorder that may be associated with thrombocytopenia.