Familial platelet disorder with associated myeloid malignancy

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial platelet disorder with associated myeloid malignancy. ORPHA:71290

A rare, genetic, constitutional thrombocytopenia disease characterized by mild to moderate thrombocytopenia, abnormal platelet function and a propensity to develop hematological malignancies, mainly of myeloid origin.