Bernard-Soulier syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Bernard-Soulier syndrome. ORPHA:274

Bernard Soulier syndrome (BSS) is an inherited platelet disorder characterized by mild to severe bleeding tendency , macrothrombocytopenia and absent ristocetin-induced platelet agglutination.