Congenital plasminogen activator inhibitor type 1 deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital plasminogen activator inhibitor type 1 deficiency. ORPHA:465

Congenital plasminogen activator inhibitor type 1 (PAI-1) deficiency is a rare genetic bleeding disorder characterized by premature lysis of hemostatic clots and a moderate bleeding tendency.