Congenital factor XIII deficiency

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Congenital factor XIII deficiency. ORPHA:331

Congenital factor XIII deficiency is an inherited bleeding disorder due to reduced levels and activity of factor XIII (FXIII) and characterized by hemorrhagic diathesis frequently associated with spontaneous abortions and defective wound healing. Factor XIII deficiency is one of the most rare coagulation factor deficiencies.