Opis choroby *

Definicja

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.

Dane

Klasyfikacja
Choroba

Synonimy
Fibrin-stabilizing factor deficiency
Niedobór czynnika stabilizującego włóknik

Kod ORPHA
331

Kod OMIM
613235

Kod ICD10
D68.2

Kod ICD11
3B14.Z

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*Źródło

orphanet