

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

*Źródło

orphanet