Wrodzony Niedobór fibrynogenu
Kod Orpha: 335 Kod OMIM: 616004

Opis choroby *

Definicja
Congenital deficiencies of fibrinogen are coagulation disorders characterized by bleeding symptoms ranging from mild to severe resulting from reduced quantity and/or quality of circulating fibrinogen. Afibrinogenemia (complete absence of fibrinogen) and hypofibrinogenemia (reduced plasma fibrinogen concentration) (see these terms) correspond to quantitative anomalies of fibrinogen while dysfibrinogenemia (see this term) corresponds to a functional anomaly of fibrinogen. Hypo- and dysfibrinogenemia may be frequently combined (hypodysfibrinogenemia).
Dane

Klasyfikacja
Choroba

Kod ORPHA
335

Kod OMIM
616004

Kod ICD10
D68.2

Kod ICD11
3B14.0

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.