Opis choroby *

Definicja

A rare, genetic, congenital vitamin K-dependant coagulation factor deficiency disorder characterized by decreased levels or absence of coagulation factor VII (FVII), resulting in bleeding diathesis of variable severity.

Dane

Klasyfikacja
Choroba

Synonimy
Congenital proconvertin deficiency
Hipoprokonwertynemia
Wrodzony Niedobór prokonwertyny
Hypoproconvertinemia

Kod ORPHA
327

Kod OMIM
227500

Kod ICD10
D68.2

Kod ICD11
3B14.Z

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*Źródło

orphanet