

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

*Źródło

orphanet