Opis choroby *

Definicja

A rare inherited bleeding disorder due to reduced activity of factor II (FII, prothrombin) and characterized by mucocutaneous and soft tissue bleeding symptoms.

Dane

Klasyfikacja
Choroba

Synonimy
Dysprothrombinemia
Dysprotrombinemia
Hipoprothrombinemia
Niedobór protrombiny
Hypoprothrombinemia
Prothrombin deficiency

Kod ORPHA
325

Kod OMIM
613679

Kod ICD10
D68.2

Kod ICD11
3B14.Z

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*Źródło

orphanet