Opis choroby *

Definicja

A rare genetic hematologic disease characterized by abnormal surface-mediated activation of fibrinolysis due to the deficiency of high-molecular-weight kininogen in plasma. Activated partial thromboplastin time (aPTT) may be prolonged. Clinically, patients are typically asymptomatic and do not show increased bleeding or thrombotic tendency.

Dane

Klasyfikacja
Choroba

Kod ORPHA
483

Kod OMIM
228960

Kod ICD10
D68.8

Kod ICD11
3B15

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*Źródło

orphanet