

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

*Źródło

orphanet