

Wrodzony Niedobór prekalikreiny

Kod Orpha: 749 Kod OMIM: 612423

Opis choroby *

Definicja

A rare genetic coagulation disorder characterized by the usually incidental laboratory finding of a prolonged activated partial thromboplastin time (aPTT) but normal prothrombin time, due to a deficiency of normal prekallikrein or the presence of nonfunctional prekallikrein. Most patients remain clinically asymptomatic, although an association with cardiovascular conditions (hypertension, myocardial infarction, other coronary artery diseases, and ischemic strokes) and venous thrombosis, as well as rare cases with increased bleeding tendency have been reported.

Dane

Klasyfikacja

Choroba

Kod ORPHA

749

Kod OMIM

612423

Kod ICD10

D68.8

Kod ICD11

3B15

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.