Choroba Von Willebranda
Kod Orpha: 903 Kod OMIM: 613554

Opis choroby *

Definicja
A rare, inherited bleeding disorder characterized by defective platelet adhesion and secondary coagulation defect that manifests as abnormal bleeding of variable severity occurring either spontaneously or in association with an invasive procedure. Three main subtypes are defined based on the type of von Willebrand factor defect: partial (type 1) or total (type 3) deficiency, and qualitative/functional anomalies (type 2).
Dane

Klasyfikacja
Choroba

Synonimy
Hereditary von Willebrand disease
Choroba Willebrand
Dziedziczna choroba Willebranda

Kod ORPHA
903

Kod OMIM
613554

Kod ICD10
D68.0

Kod ICD11
3B12

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.