

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

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.