## 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