

## 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