

Opis choroby *

Definicja

A rare hemorrhagic disorder due to a constitutional platelet anomaly characterized by moderate to severe deficiency in both platelet alpha-granules and dense bodies, resulting in impaired platelet function and decreased aggregation responses. Patients present increased bleeding tendency with symptoms like easy bruising, or menorrhagia.

Dane

Klasyfikacja

Choroba

Synonimy

Alpha dense granule deficiency

Niedobór ziarnistości alfa

Combined alpha-delta platelet storage pool deficiency

Kod ORPHA

734

Kod OMIM

185050

Kod ICD10

D69.1

Kod ICD11

3B62.4

*Źródło

orphanet