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

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*Źródło

orphanet