

## 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 | Synonimy  |
|--------------|---|
| Choroba      | Alpha dense granule deficiency<br>Niedobór ziarnistości alfa<br>Combined alpha-delta platelet storage pool deficiency |
| Kod ORPHA    | Kod OMIM  |
| 734          | 185050  |

| Kod ICD10 |
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| D69.1     |

| Kod ICD11 |
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| 3B62.4    |

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\*Źródło

orphanet