Niedokrwistość sierpowata
Kod Orpha: 232 Kod OMIM: 603903

Opis choroby *

Definicja
A severe form of sickle cell disease (SCD) characterized by homozygosity for the sickle hemoglobin (HbS) gene and which acutely manifests with severe anemia, susceptibility to severe bacterial infections, and ischemic vasoocclusive accidents (VOA). It is a red cell disease of genetic origin which manifests with hemolytic disease and loss of red cell deformability leading to other occlusive events.

Dane

Klasyfikacja
Choroba

Synonimy
Anemia sierpowata

Kod ORPHA
232

Kod OMIM
603903

Kod ICD10
D57.2

Kod ICD11
3A51.2

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.