Niedokrwistość sierpowata - beta-talasemia
Kod Orpha: 251359 Kod OMIM:

Opis choroby *

Definition
A rare, genetic hemoglobinopathy that affects red blood cells both in the production of abnormal hemoglobin, as well as the decreased synthesis of beta globin chains. Clinical manifestations depend on the amount of residual beta globin chains production, and are similar to sickle cell disease, including anemia, vascular occlusion and its complications, acute episodes of pain, acute chest syndrome, pulmonary hypertension, sepsis, ischemic brain injury, splenic sequestration crisis and splenomegaly.
Dane

Classification
Choroba

Synonyms
HbS-beta-thalassemia syndrome
HbS - beta-talasemia

ORPHA code
251359

OMIM code
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ICD10 code
D57.2

ICD11 code
3A51.4

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.