

Opis choroby *

Definicja

A rare, genetic hemoglobinopathy characterized by anemia, reticulocytosis and erythrocyte abnormalities including target cells, irreversibly sickled cells and crystal-containing cells. Clinical course is similar to sickle cell disease, but less severe and with less complications. Signs and symptoms may include acute episodes of pain, splenic infarction and splenic sequestration crisis, acute chest syndrome, focal segmental glomerulosclerosis, ischemic brain injury, peripheral retinopathy, and osteonecrosis.

Dane

Klasyfikacja

Choroba

Synonimy

HbSC disease

Choroba HbSC

Kod ORPHA

251365

Kod OMIM

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Kod ICD10

D57.2

Kod ICD11

3A51.3

*Źródło

orphanet