# Niedokrwistość sierpowata - choroba hemoglobiny C

# Kod Orpha: 251365 Kod OMIM:

## 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

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*Źródło

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.