## Opis choroby *

Definicja

A rare beta-thalassemia associated with another hemoglobin anomaly characterized by the presence of the hemoglobin Lepore variant in association with beta-thalassemia. Clinical presentation is highly variable, depending on the type of beta-thalassemia, and ranges from severe hypochromic microcytic anemia and complete transfusion dependency to moderate, compensated anemia without a need for regular blood transfusions.

Dane

**Klasyfikacja**
Choroba

Synonimy
HbLepore-beta-thalassemia syndrome
HbLepore - beta-talasemia
Lepore - beta-talasemia
Lepore-beta-thalassemia syndrome

**Kod ORPHA**
330032

**Kod OMIM**
-

**Kod ICD10**
D56.8

**Kod ICD11**
-

---

*Źródło

orphanet