Hemoglobina Lepore - beta-talasemia
Kod Orpha: 330032 Kod OMIM:

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

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.