

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	Synonimy
Choroba	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.