

Opis choroby *

Definicja

Beta-thalassemias associated with hemoglobin (Hb) anomalies result in a variable clinical spectrum, ranging from asymptomatic to severe, depending on the severity of the thalassemia mutation and on the type of the Hb anomaly [hereditary persistence of fetal Hb, delta-beta-thalassemia, Hb C - beta-thalassemia, Hb E - beta-thalassemia and Hb S - beta-thalassemia (see these terms)].

Dane

Klasyfikacja

Kategoria	Synonimy
	Beta-thalassemia associated with another Hb anomaly
	Beta-talasemia z towarzyszącą inną anomalią Hb

Kod ORPHA

231230

Kod OMIM

-

Kod ICD10

D58.2

Kod ICD11

-

[*Źródło](#)

orphanet