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
-

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*Źródło

orphanet