Opis choroby *

Definicja

An intermediate form of alpha-thalassemia characterized by increased hemolysis and mild to severe anemia with marked microcytosis and hypochromia. Hemoglobin H disease (HbH) disease belongs to the group of nontransfusion-dependent thalassemia.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Alpha-thalassemia intermedia
Alfa talasemia pośrednia
Choroba HbH
HbH disease

Kod ORPHA
93616

Kod OMIM
613978

Kod ICD10
D56.0

Kod ICD11
3A50.02

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*Źródło

orphanet