

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Podtyp kliniczny	Alpha-thalassemia intermedia Alfa talasemia pośrednia Choroba HbH HbH disease

<b>Kod ORPHA</b> 93616	<b>Kod OMIM</b> 613978	<b>Kod ICD10</b> D56.0
<b>Kod ICD11</b> 3A50.02		

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### \*Źródło

orphanet