

## Opis choroby \*

### Definicja

A rare inherited hemoglobinopathy characterized by impaired synthesis of two to all four alpha-globin chains leading to a variable clinical picture depending on the number of affected alleles.

Dane

### Klasyfikacja

Choroba

**Kod ORPHA**

846

**Kod OMIM**

604131

**Kod ICD10**

D56.0

**Kod ICD11**

3A50.0

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[\\*Źródło](#)

orphanet