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

---

*Źródło

orphanet