Opis choroby *

Definicja

Hereditary persistence of fetal hemoglobin (HPFH) associated with beta-thalassemia (see this term) is characterized by high hemoglobin (Hb) F levels and an increased number of fetal-Hb-containing-cells.

Dane

Klasyfikacja
Choroba

Synonimy
HPFH-beta-thalassemia syndrome
HPFH - beta-talasemia

Kod ORPHA
46532

Kod OMIM
613566

Kod ICD10
D56.4

Kod ICD11
3A50.4

---

*Źródło

orphanet