Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome. ORPHA:46532

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.