DISEASE:
Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome

NAME:	Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome
DESCRIPTION:	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.
ORPHACODE:	46532
SYNONYMS:	HPFH-beta-thalassemia syndrome
XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM ICD-10
ANALYTE(S):	HBB HBG1 HBG2 KLF1
CREATED:	13 May 2019 - 01:02

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Source URL: http://gentest.healthdata.be/disease/3032

RELATED CONTENT

Related Genetic Tests

- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies, phenotype modifiers (hot spot mutations - rs7482144 (Xmn1) at promoter 158 bp 5? upstream of HBG2 / 32C-T in the 5' UTR of the HBS1L)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique Médicale UCL

Related Analytes

- hemoglobin subunit beta
- hemoglobin subunit gamma 1
- hemoglobin subunit gamma 2
- KLF transcription factor 1