Belgian Genetic Tests database

DISEASE:
Hemoglobin C disease

NAME:	Hemoglobin C disease
DESCRIPTION:	Hemoglobin C disease (HbC) is a hemoglobinopathy characterized by production of abnormal variant hemoglobin known as hemoglobin C, with no or mild clinical manifestations (hemolytic anemia).
ORPHACODE:	2132
XREF(S):	Orphanet MedDRA MeSH ICD-10
ANALYTE(S):	HBB
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/1546

RELATED CONTENT

Related Genetic Tests

- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies
- Beta-globin hemoglobinopathies (Hot-spot mutations : Sickle cell disease (HBS), hemoglobin C, hemoglobin E or Hemoglobin D)
- Beta-globin hemoglobinopathies (full sequencing)
- Beta-globin hemoglobinopathies, Hemoglobinopathy C (HbC) (HBB hot spot mutation - p.Glu6Lys)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique Humaine - Erasme ULB
- Centre de Génétique Médicale UCL
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- hemoglobin subunit beta