GENETIC TEST:
Beta-globin hemoglobinopathies

FULL NAME:	Beta-globin hemoglobinopathies
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS) MLPA based techniques
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-03-25 / 2024-09-09

EQA:	• DNA for Haemoglobinopathies, • DNA for Haemoglobinopathies , • DNA for Haemoglobinopathies , • DNA for Haemoglobinopathies, • DNA for Haemoglobinopathies, • DNA for Haemoglobinopathies, • DNA for Haemoglobinopathies , • DNA for Haemoglobinopathies, • DNA for Haemoglobinopathies
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	24 Jul 2019 - 11:39
CHANGED:	19 Jan 2024 - 09:01
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM88

RELATED CONTENT

Related Diseases

- Beta-thalassemia intermedia
- Beta-thalassemia major
- Delta-beta-thalassemia
- Hemoglobin C disease
- Hemoglobin C-beta-thalassemia syndrome
- Hemoglobin D disease
- Hemoglobin E disease
- Hemoglobin E-beta-thalassemia syndrome
- Hemoglobin Lepore-beta-thalassemia syndrome
- Hemoglobin M disease
- Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome
- Hereditary persistence of fetal hemoglobin-sickle cell disease syndrome
- Sickle cell anemia
- Sickle cell-beta-thalassemia disease syndrome
- Sickle cell-hemoglobin C disease syndrome
- Sickle cell-hemoglobin D disease syndrome
- Sickle cell-hemoglobin E disease syndrome

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- hemoglobin subunit beta

Source URL: http://gentest.healthdata.be/genetic_test/583