GENETIC TEST:
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)

FULL NAME:	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)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Real-time PCR
RIZIV CODE:	565390-565401
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	22 Aug 2019 - 09:45
CHANGED:	02 Mar 2022 - 11:58

Source URL: http://gentest.healthdata.be/genetic_test/584

RELATED CONTENT

Related Diseases

- Hemoglobinopathy Toms River
- Hereditary persistence of fetal hemoglobin-beta-thalassemia syndrome
- Hereditary persistence of fetal hemoglobin-sickle cell disease syndrome

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- BCL11 transcription factor A
- hemoglobin subunit gamma 2
- HBS1 like translational GTPase

Related Gene Panels

- Beta-globin hemoglobinopathies, phenotype modifiers (3 genes) - ULB