

GENETIC TEST:**Beta-globin hemoglobinopathies, Sickle cell anemia (HbS) (HBB hot spot mutation - p.Glu6Val)**

FULL NAME:	Beta-globin hemoglobinopathies, Sickle cell anemia (HbS) (HBB hot spot mutation - p.Glu6Val)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis, Pre-implantation genetic diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi, Cell culture
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565390-565401

ACCREDITATION (ISO 15189):	2021-10-07 / 2026-06-14
EQA:	• DNA Sequencing - Sanger, • DNA Sequencing - Sanger, • DNA Sequencing - Sanger, • DNA Sequencing - Sanger, • DNA Sequencing - Sanger, • DNA Sequencing - Sanger, • DNA Sequencing - Sanger , • DNA Sequencing - Sanger
TURNAROUND TIME (MAXIMUM):	2 months (10 working days for prenatal diagnosis)
CREATED:	23 Aug 2019 - 16:28
CHANGED:	01 Mar 2023 - 16:29
URL:	https://laboguide.uzbrussel.be/laboguide#Analyses:Hemoglobinopathie%20S&&&&975...

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