GENETIC TEST:
alpha-globin hemoglobinopathies

FULL NAME:	alpha-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, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis MLPA based techniques
RIZIV CODE:	565471-565482

EQA:	• 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:	26 Jul 2019 - 09:49
CHANGED:	19 Jan 2024 - 13:23

RELATED CONTENT

Related Diseases

- Alpha-thalassemia

Related Laboratories

- Centre de Génétique Médicale UCL

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- hemoglobin subunit alpha 1