GENETIC TEST:
Fabry disease

FULL NAME:	Fabry disease
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
EQA:	• Fabry disease
TURNAROUND TIME (MAXIMUM):	3 months

CREATED:	30 Jul 2019 - 09:11
CHANGED:	19 May 2022 - 12:04

RELATED CONTENT

Related Diseases

- Fabry disease

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- galactosidase alpha