GENETIC TEST:
Neurofibromatosis type 1 / Legius syndrome

FULL NAME:	Neurofibromatosis type 1 / Legius syndrome
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis MLPA based techniques
RIZIV CODE:	565552-565563
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	3 - 4 months
CREATED:	31 Jul 2019 - 14:22
CHANGED:	20 May 2022 - 11:46

RELATED CONTENT

Related Diseases

- 17q11 microdeletion syndrome
- Legius syndrome
- Neurofibromatosis type 1
- Neurofibromatosis type 1 due to NF1 mutation or intragenic deletion

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- neurofibromin 1
- sprouty related EVH1 domain containing 1