GENETIC TEST:
Retinitis pigmentosa, X-Linked

FULL NAME:	Retinitis pigmentosa, X-Linked
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:	Next Generation Sequencing (NGS) MLPA based techniques
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	3 - 4 months
CREATED:	01 Aug 2019 - 14:42
CHANGED:	20 May 2022 - 10:50

RELATED CONTENT

Related Diseases

- Cone rod dystrophy
- Retinitis pigmentosa

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- OFD1 centriole and centriolar satellite protein
- RP2 activator of ARL3 GTPase
- retinitis pigmentosa GTPase regulator

Related Gene Panels

- Retinitis pigmentosa, X-linked - UGent