GENETIC TEST:
Achromatopsia

FULL NAME:	Achromatopsia
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
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
TURNAROUND TIME (MAXIMUM):	3 - 4 months
CREATED:	26 Jul 2019 - 12:50
CHANGED:	10 May 2022 - 08:03

RELATED CONTENT

Related Diseases

- Achromatopsia
- Cone rod dystrophy
- Progressive cone dystrophy
- Stargardt disease

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- cyclic nucleotide gated channel subunit alpha 3
- cyclic nucleotide gated channel subunit beta 3

Related Gene Panels

- Achromatopsia (2 genes) - UGent