GENETIC TEST:
Leber Congenital Amaurosis - Retinal dystrophy, early onset (gene panel)

FULL NAME:	Leber Congenital Amaurosis - Retinal dystrophy, early onset (gene panel)
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:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10
TURNAROUND TIME (MAXIMUM):	3 - 4 months
CREATED:	30 Jul 2019 - 13:22
CHANGED:	31 May 2022 - 13:58

RELATED CONTENT

Related Diseases

- Cone rod dystrophy
- Leber congenital amaurosis
- Retinitis pigmentosa
- Senior-Loken syndrome
- Severe early-childhood-onset retinal dystrophy

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- aryl hydrocarbon receptor interacting protein like 1
- centrosomal protein 290
- crumbs cell polarity complex component 1
- cone-rod homeobox
- guanylate cyclase 2D, retinal
- IQ motif containing E
- lebercilin LCA5
- lecithin retinol acyltransferase
- nicotinamide nucleotide adenylyltransferase 1
- retinol dehydrogenase 12
- retinoid isomerohydrolase RPE65

Related Gene Panels

- Leber Congenital Amaurosis - UGent
