GENETIC TEST:
Maternally-inherited diabetes and deafness / Mitochondrial myopathy with reversible cytochrome C oxidase deficiency / mitochondrial tRNA glutamic acid

FULL NAME:	Maternally-inherited diabetes and deafness / Mitochondrial myopathy with reversible cytochrome C oxidase deficiency / mitochondrial tRNA glutamic acid
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi, Cell culture
METHOD CATEGORY:	Sequence analysis: entire coding region
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565456-565460

ACCREDITATION (ISO 15189):	2021-10-07 / 2026-06-14
TURNAROUND TIME (MAXIMUM):	3 months (10 working days for prenatal diagnosis)
CREATED:	26 Aug 2019 - 17:18
CHANGED:	29 Jul 2022 - 13:04

RELATED CONTENT

Related Diseases

- MELAS
- Maternally-inherited diabetes and deafness

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- mitochondrially encoded tRNA-Glu (GAA/G)