DISEASE:
Maternally-inherited diabetes and deafness

NAME:	Maternally-inherited diabetes and deafness
DESCRIPTION:	A rare mitochondrial disease characterized by maternally transmitted diabetes and sensorineural deafness.
ORPHACODE:	225
SYNONYMS:	MIDD Maternally-inherited diabetes and hearing loss Mitochondrial diabetes
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	MT-TE MT-TL1 MT-TK
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Leigh / NARP Syndrome
- Maternally-inherited diabetes and deafness / Mitochondrial myopathy with reversible cytochrome C oxidase deficiency / mitochondrial tRNA glutamic acid
- Mitochondrial disorders, mitchondrial DNA based (Full sequencing of mtDNA genome)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- mitochondrially encoded tRNA-Glu (GAA/G)
- mitochondrially encoded tRNA-Lys (AAA/G)
- mitochondrially encoded tRNA-Leu (UUA/G) 1

Related Gene Panels

- Leigh syndrome (mtDNA / 37 genes) - VUB