DISEASE:
Mitochondrial DNA-related progressive external ophthalmoplegia

NAME:	Mitochondrial DNA-related progressive external ophthalmoplegia
DESCRIPTION:	An Orphanet summary for this disease is currently under development. However, other data related to the disease are accessible from the Additional Information menu located on the right side of this page.
ORPHACODE:	663
SYNONYMS:	Maternally-inherited CPEO Maternally-inherited chronic progressive external ophthalmoplegia mtDNA-related progressive external ophthalmoplegia
XREF(S):	Orphanet ICD-10
ANALYTE(S):	MT-TS1 MT-TL2 MT-TN MT-TL1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Leigh / NARP Syndrome

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- mitochondrially encoded tRNA-Leu (UUA/G) 1
- mitochondrially encoded tRNA-Leu (CUN) 2
- mitochondrially encoded tRNA-Asn (AAU/C)
- mitochondrially encoded tRNA-Ser (UCN) 1

Related Gene Panels

- Leigh syndrome (mtDNA / 37 genes) - VUB