ANALYTE:
MRPS22

NAME:	mitochondrial ribosomal protein S22
SYMBOL:	MRPS22
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	C3orf5 GIBT GK002 MRP-S22
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Mitochondrial disorders (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel

Related Diseases

- 46,XX gonadal dysgenesis
- Hypotonia with lactic acidemia and hyperammonemia

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- mitochondrial disease, nuclear based (343 genes) - VUB
