Belgian Genetic Tests database

ANALYTE:
CP

NAME:	ceruloplasmin
SYMBOL:	CP
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ferroxidase
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3537

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia (gene panel)
- Ataxia Spasticity (gene panel)
- Dystonia (gene panel)
- Hemochromatosis (17 genes)
- Intellectual disability (virtual gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Neurodegeneration with Brain Iron Accumulation (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel

Related Diseases

- Aceruloplasminemia

Related Gene Panels

- Ataxia (141 genes) - KUL
- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Dystonia (68 genes) - KUL
- Hemochromatosis (17 genes) - ULB
- Intellectual disability (gene panel)

- Movement Disorders - UGent
- Movement Disorders - ULG
- Neurodegeneration with Brain Iron Accumulation (NBIA) - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB

Source URL: http://gentest.healthdata.be/analyte/3537