Belgian Genetic Tests database

ANALYTE:
COQ6

NAME:	coenzyme Q6, monooxygenase
SYMBOL:	COQ6
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CGI-10
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (gene panel)
- Mitochondrial disorders (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Diseases

- Familial steroid-resistant nephrotic syndrome with sensorineural deafness
- Full schwannomatosis

Related Gene Panels

- Ataxia (348 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG

- Panel Nephro-ULG-V1
- mitochondrial disease, nuclear based (343 genes) - VUB

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