ANALYTE:
COQ9

NAME:	coenzyme Q9
SYMBOL:	COQ9
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DKFZP434K046
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/2578

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia Spasticity (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Mitochondrial disorders (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Myopathy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neuromuscular disorders (548 genes)

Related Diseases

- Encephalopathy-hypertrophic cardiomyopathy-renal tubular disease syndrome

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Congenital malformation (1721 genes) - ULB

- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Movement Disorders - UGent
- Movement Disorders - ULG
- Myopathy (332 genes) - IPG
- Nephropathy panel - UGent
- Neuromuscular disorders (548 genes) - ULB
- mitochondrial disease, nuclear based (343 genes) - VUB
