Belgian Genetic Tests database

ANALYTE:
WFS1

NAME:	wolframin ER transmembrane glycoprotein
SYMBOL:	WFS1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DIDMOAD WFS
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)
- Ataxia (gene panel)
- Cataract (gene panel)
- Deafness, autosomal dominant 6/14 / Wolfram syndrome
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hearing loss (deafness), (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephropathies, hereditary (gene panel)
- Optic atrophy (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Retinal dystrophy / RETNET (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Early-onset nuclear cataract
- Rare autosomal dominant non-syndromic sensorineural deafness type DFNA
- Wolfram syndrome
- Wolfram-like syndrome

Related Gene Panels

- Ataxia (141 genes) - KUL
- Ataxia (348 genes) - ULB
- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Cataract - UGent
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Optic atrophy - UGent
- Retinal dystrophy - UGent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
