

ANALYTE:
AFG3L2

NAME:	AFG3 like matrix AAA peptidase subunit 2
SYMBOL:	AFG3L2
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SYNONYMS:	SPAX5
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
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Related Diseases

- [Early-onset spastic ataxia-myoclonic epilepsy-neuropathy syndrome](#)

- Spinocerebellar ataxia type 28

Related Gene Panels

- Ataxia (141 genes) - KUL
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- Ataxia Spasticity - UGent
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- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Hereditary Spastic Paraplegia (94 genes) - KUL
- Hereditary spastic paraplegia (188 genes) - ULB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Movement Disorders - UGent
- Movement Disorders - ULG
- Neurodegeneration (99 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Optic atrophy - UGent
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Retinal dystrophy - UGent
- Spastic Paraplegia (89 genes) - IPG
- mitochondrial disease, nuclear based (343 genes) - VUB
- test

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