Belgian Genetic Tests database

ANALYTE:
ABCD1

NAME:	ATP binding cassette subfamily D member 1
SYMBOL:	ABCD1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ALDP AMN adrenoleukodystrophy
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Adrenoleukodystrophy, X-linked
- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia Spasticity (gene panel)
- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hereditary Spastic Paraplegia (94 genes)
- Hereditary Spastic Paraplegia (gene panel)
- Hereditary spastic paraplegia (gene panel - 249 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodegeneration (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Neuromuscular disorders (548 genes)

Related Diseases

- Adrenomyeloneuropathy
- CADDS
- Hirschsprung disease
- X-linked cerebral adrenoleukodystrophy

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- 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)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Leukodystrophy - UGent
- Metabolic disorders (213 genes) - VUB
- Nephropathy panel - UGent
- Neurodegeneration (99 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB
