Belgian Genetic Tests database

ANALYTE:
ABHD5

NAME:	abhydrolase domain containing 5, lysophosphatidic acid acyltransferase
SYMBOL:	ABHD5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CGI-58 NCIE2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Source URL: http://gentest.healthdata.be/analyte/3965

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Congenital malformation (gene panel - 1721 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Ichthyosis (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Neuromuscular disorders (232 genes (= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia)
- Neuromuscular disorders (548 genes)
- Neuromuscular disorders (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Neutral lipid storage disease with ichthyosis

Related Gene Panels

- Ataxia (348 genes) - ULB

- Congenital malformation (1721 genes) - ULB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ichthyosis and erythroderma (98 genes) - KUL
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Skin disorders - UGent
