

**ANALYTE:
ABHD5**

NAME:	abhydrolase domain containing 5, lysophosphatidic acid acyltransferase
SYMBOL:	ABHD5
VERSION OF ORPHANET:	2022-05-01 04:55:23
SYNONYMS:	CGI-58 NCIE2
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
CREATED:	13 May 2019 - 01:01
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- [Neuromuscular disorders \(232 genes \(= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia\)\)](#)
- [Neuromuscular disorders \(548 genes\)](#)
- [Neuromuscular disorders \(gene panel\)](#)
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Related Gene Panels

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- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Ichthyosis and erythroderma (95 genes) - KUL
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Myopathy (179 genes) - UZA
- Myopathy (332 genes) - IPG
- 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

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