ANALYTE:
PSAP

NAME:	prosaposin
SYMBOL:	PSAP
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	saposin-A saposin-B saposin-C saposin-D variant Gaucher disease and variant metachromatic leukodystrophy
XREF(S):	Orphanet SwissProt Ensembl Genatlas HGNC OMIM Reactome
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RELATED CONTENT

Related Genetic Tests

- Ataxia Spasticity (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Lysosomal Storage Disease (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel

Related Diseases

- Atypical Gaucher disease due to saposin C deficiency
- Encephalopathy due to prosaposin deficiency
- Infantile Krabbe disease
- Metachromatic leukodystrophy, adult form
- Metachromatic leukodystrophy, juvenile form
- Metachromatic leukodystrophy, late infantile form

Related Gene Panels

- Ataxia Spasticity - UGent
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Leukodystrophy - UGent
- Lysomal Storage (64 genes) - VUB
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
