ANALYTE:
SGSH

NAME:	N-sulfoglucosamine sulfohydrolase
SYMBOL:	SGSH
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HSS MPS3A SFMD mucopolysaccharidosis type IIIA sulfamidase
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- 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)
- Lysosomal Storage Disease (gene panel)
- Mucopolysaccharidosis (MPS) type IIIA / Sanfilippo A
- Mucopolysacharidosis type IIIa (N-sulfoglucosamine sulfohydrolase)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Sanfilippo syndrome type A

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- 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
- Lysomal Storage (64 genes) - VUB
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
