ANALYTE:
SGPL1

NAME:	sphingosine-1-phosphate lyase 1
SYMBOL:	SGPL1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	SPL
XREF(S):	Orphanet SwissProt OMIM Genatlas Reactome HGNC Ensembl
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RELATED CONTENT

Related Genetic Tests

- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Congenital malformation (gene panel - 1721 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- End-stage renal disease, ESRD (gene panel)
- Ichthyosis (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Neuromuscular disorders (548 genes)
- Neuropathy (gene panel)
- Neuropathy (gene panel)
- Primary immune deficiencies (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Familial steroid-resistant nephrotic syndrome with adrenal insufficiency

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL

- Congenital malformation (1721 genes) - ULB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Ichthyosis and erythroderma (98 genes) - KUL
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Neuropathy (148 genes) - IPG
- Neuropathy panel - UGent
- Panel Nephro-ULG-V1
- Primary immune deficiencies - UGent
- Skin disorders - UGent
