Belgian Genetic Tests database

ANALYTE:
LAMB2

NAME:	laminin subunit beta 2
SYMBOL:	LAMB2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	NPHS5 laminin S
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (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)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- LAMB2-related infantile-onset nephrotic syndrome
- Pierson syndrome
- Synaptic congenital myasthenic syndromes

Related Gene Panels

- End-stage renal disease (106 genes) - IPG
- Intellectual disability (gene panel)
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (166 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
