Belgian Genetic Tests database

ANALYTE:
MYO1E

NAME:	myosin IE
SYMBOL:	MYO1E
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HuncM-IC MGC104638 MYO1C myosin-IC
XREF(S):	Orphanet Reactome SwissProt Ensembl Genatlas HGNC OMIM
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3647

RELATED CONTENT

Related Genetic Tests

- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel

Related Diseases

- Familial idiopathic steroid-resistant nephrotic syndrome with focal segmental hyalinosis
- Genetic steroid-resistant nephrotic syndrome

Related Gene Panels

- End-stage renal disease (106 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1

Source URL: http://gentest.healthdata.be/analyte/3647