ANALYTE:
PODXL

NAME:	podocalyxin like
SYMBOL:	PODXL
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	Gp200 PC PCLP PDX PODXL1 gp135
XREF(S):	Orphanet HGNC OMIM Reactome Genatlas SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/1264

RELATED CONTENT

Related Genetic Tests

- Ataxia Spasticity (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Mitochondrial disorders (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Neurodegeneration (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- test

Related Diseases

- Atypical juvenile parkinsonism
- Young-onset Parkinson disease

Related Gene Panels

- Ataxia Spasticity - UGent

- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hepatology panel - UGent
- Intellectual disability & Epilepsy - UGent
- Movement Disorders - UGent
- Movement Disorders - ULG
- Nephropathies, hereditary (219 genes) - KUL
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neurodegeneration (99 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1
- mitochondrial disease, nuclear based (343 genes) - VUB
- test1
