Belgian Genetic Tests database

DISEASE:
Familial idiopathic steroid-resistant nephrotic syndrome with minimal changes

NAME:	Familial idiopathic steroid-resistant nephrotic syndrome with minimal changes
ORPHACODE:	93216
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PTPRO NPHS1 NPHS2 EMP2
CREATED:	13 May 2019 - 01:02
CHANGED:	01 Jul 2019 - 06:57

Source URL: http://gentest.healthdata.be/disease/3116

RELATED CONTENT

Related Genetic Tests

- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- epithelial membrane protein 2
- NPHS1 adhesion molecule, nephrin
- NPHS2 stomatin family member, podocin
- protein tyrosine phosphatase receptor type O

Related Gene Panels

- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG