Belgian Genetic Tests database

ANALYTE:
INF2

NAME:	inverted formin, FH2 and WH2 domain containing
SYMBOL:	INF2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	MGC13251 inverted formin 2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
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Source URL: http://gentest.healthdata.be/analyte/3640

RELATED CONTENT

Related Genetic Tests

- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (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)
- Peripheral neuropathy (gene panel)

Related Diseases

- Autosomal dominant intermediate Charcot-Marie-Tooth disease type E
- Familial idiopathic steroid-resistant nephrotic syndrome with focal segmental hyalinosis
- Genetic steroid-resistant nephrotic syndrome

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- End-stage renal disease (106 genes) - IPG
- 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 (genepanel) - UZA
- Neuropathy panel - UGent
- Panel Nephro-ULG-V1
