Belgian Genetic Tests database

ANALYTE:
GSN

NAME:	gelsolin
SYMBOL:	GSN
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DKFZp313L0718 amyloidosis, Finnish type
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3829

RELATED CONTENT

Related Genetic Tests

- Corneal dystrophy (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- End-stage renal disease, ESRD (gene panel)
- Hereditary spastic paraplegia (gene panel - 249 genes)
- 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)
- Neuromuscular disorders (548 genes)
- Neuropathy (gene panel)
- Peripheral neuropathy (gene panel)

Related Diseases

- AGel amyloidosis

Related Gene Panels

- Corneal dystrophy - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Hereditary spastic paraplegia (188 genes) - ULB
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent

- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Neuromuscular disorders (548 genes) - ULB
- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
- Panel Nephro-ULG-V1
