Belgian Genetic Tests database

ANALYTE:
COL4A5

NAME:	collagen type IV alpha 5 chain
SYMBOL:	COL4A5
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XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Alport autosomal recessive and X-linked and hematuria (3 genes)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- End-stage renal disease, ESRD (gene panel)
- Hearing loss (deafness), (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)

Related Diseases

- NON RARE IN EUROPE: Benign familial hematuria
- X-linked Alport syndrome
- X-linked Alport syndrome-diffuse leiomyomatosis

Related Gene Panels

- Alport (X-linked and recessive) (3 genes) - IPG
- Alport (X-linked and recessive) (3 genes) - UZA
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- End-stage renal disease (106 genes) - IPG
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Nephropathies, hereditary (219 genes) - KUL

- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
