Belgian Genetic Tests database

ANALYTE:
INVS

NAME:	inversin
SYMBOL:	INVS
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	nephrocystin 2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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Source URL: http://gentest.healthdata.be/analyte/2051

RELATED CONTENT

Related Genetic Tests

- Cholestasis (gene panel)
- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- End-stage renal disease, ESRD (gene panel)
- Hepatology (gene panel)
- Hepatorenal disorders (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Infantile nephronophthisis
- Senior-Loken syndrome

Related Gene Panels

- Cholestasis (40 genes) - UCL
- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB

- End-stage renal disease (106 genes) - IPG
- Hepatology panel - UGent
- Hepatorenal disorders (13 genes) - UCL
- Heterotaxie PCD - UGent
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
