ANALYTE:
HNF1B

NAME:	HNF1 homeobox B
SYMBOL:	HNF1B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HNF1beta HNF1ß LFB3 MODY5 VHNF1 hepatocyte nuclear factor 1 beta
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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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)
- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (gene panel)
- End-stage renal disease, ESRD (gene panel)
- Familial cancer predisposition (gene panel)
- Hepatology (gene panel)
- Hepatorenal disorders (gene panel)
- Hyperinsulinism (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- MODY : Maturity onset Diabete of the Young (gene panel)
- Maturity onset Diabete of the Young (MODY), type 5 / Renal cysts and diabetes syndrome (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Renal cell carcinoma (kidney cancer) (gene panel)
- Renal cysts and diabetes syndrome
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- 17q12 microdeletion syndrome

- Autosomal dominant primary hypomagnesemia with hypocalciuria
- Bilateral multicystic dysplastic kidney
- Familial prostate cancer
- HNF1B-related autosomal dominant tubulointerstitial kidney disease
- Mayer-Rokitansky-Küster-Hauser syndrome type 2
- Medullary sponge kidney
- Renal dysplasia, bilateral
- Renal dysplasia, unilateral
- Unilateral multicystic dysplastic kidney

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- 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
- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (genepanel) - UZA
- End-stage renal disease (106 genes) - IPG
- Hepatology panel - UGent
- Hepatorenal disorders (13 genes) - UCL
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- MODY (7 genes) - UZA
- MODY - Maturity onset Diabete of the Young (21 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Renal cell carcinoma - UGent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
