Belgian Genetic Tests database

ANALYTE:
HNF1A

NAME:	HNF1 homeobox A
SYMBOL:	HNF1A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HNF1 HNF1a LFB1
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital structural heart defects (gene panel)
- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (gene panel)
- Hyperinsulinism (gene panel)
- Lipodystrophy and/or hyperinsulinism (gene panel)
- MODY : Maturity onset Diabete of the Young (gene panel)
- Maffucci syndrome (gene panel)
- Maturity onset Diabete of the Young (MODY), type 5 / Renal cysts and diabetes syndrome (gene panel)
- Nephropathies, hereditary (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Sturge-Weber syndrome (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Chromophobe renal cell carcinoma
- Clear cell papillary renal cell carcinoma
- Hyperinsulinism due to HNF1A deficiency
- MODY

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital structural heart defects - UGent

- Diabetes neonatal / Maturity onset Diabete of the Young (MODY) / Hyperinsulinism (genepanel) - UZA
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- MODY (7 genes) - UZA
- MODY - Maturity onset Diabete of the Young (21 genes) - IPG
- Maffucci syndrome (65 genes) - KUL
- Nephropathies, hereditary (219 genes) - KUL
- Overgrowth & vascular anomalies (65 genes) - KUL
- Sturge-Weber syndrome (65 genes) - KUL
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
