Belgian Genetic Tests database

ANALYTE:
NOTCH2

NAME:	notch receptor 2
SYMBOL:	NOTCH2
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Alagille syndrome (2 genes)
- Cholestasis (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Alagille syndrome due to a NOTCH2 point mutation
- Hajdu-Cheney syndrome

Related Gene Panels

- Alagille (2 genes) -UCL
- Alagille syndrome (2 genes) - UCL
- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Cholestasis (40 genes) - UCL
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Hepatology panel - UGent
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
