ANALYTE:
DCDC2

NAME:	doublecortin domain containing 2
SYMBOL:	DCDC2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DCDC2A KIAA1154 NPHP19 RU2 nephronophthisis 19
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt Reactome
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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)
- End-stage renal disease, ESRD (gene panel)
- Hearing loss (deafness), (gene panel)
- Hepatology (gene panel)
- Hepatorenal disorders (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)

Related Diseases

- Isolated neonatal sclerosing cholangitis
- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB
- Senior-Boichis 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

- Hearing loss (deafness) (genepanel) - UZA
- Hepatology panel - UGent
- Hepatorenal disorders (13 genes) - UCL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
