ANALYTE:
IQCB1

NAME:	IQ motif containing B1
SYMBOL:	IQCB1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KIAA0036 NPHP5 SLSN5 nephrocystin-5
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- 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)
- Congenital malformation gene panel
- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Leber congenital amaurosis
- Senior-Loken syndrome

Related Gene Panels

- 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
- Congenital malformation gene panel - VUB
- End-stage renal disease (106 genes) - IPG
- Leber - Retinal, early onset
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1

- Retinal dystrophy - UGent
