ANALYTE:
PKHD1

NAME:	PKHD1 ciliary IPT domain containing fibrocystin/polyductin
SYMBOL:	PKHD1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ARPKD FCYT FPC fibrocystin fibrocystin/polyductin complex polyductin tigmin
XREF(S):	Orphanet OMIM SwissProt Ensembl Genatlas HGNC
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Caroli Disease
- 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)
- Congenital malformation gene panel
- End-stage renal disease, ESRD (gene panel)
- Hepatology (gene panel)
- Hepatorenal disorders (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)

Related Diseases

- Autosomal recessive polycystic kidney disease
- Caroli disease

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
- Congenital malformation gene panel - VUB
- End-stage renal disease (106 genes) - IPG
- Hepatology panel - UGent
- Hepatorenal disorders (13 genes) - UCL
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
