ANALYTE:
NPHP4

NAME:	nephrocystin 4
SYMBOL:	NPHP4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	KIAA0673 POC10 POC10 centriolar protein homolog (Chlamydomonas) SLSN4 nephroretinin
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/2054

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)
- Congenital structural heart defects (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)
- Retinal dystrophy / RETNET (gene panel)

Related Diseases

- Juvenile nephronophthisis
- Senior-Loken 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
- Congenital structural heart defects - UGent
- 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
- Retinal dystrophy - UGent
