ANALYTE:
TTC21B

NAME:	tetratricopeptide repeat domain 21B
SYMBOL:	TTC21B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FAP60 FLA17 FLJ11457 IFT139B JBTS11 NPHP12 THM1
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/index.php/index.php/analyte/461

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)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- End-stage renal disease, ESRD (gene panel)
- Epilepsy gene panel
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Infantile nephronophthisis
- Jeune 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
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- End-stage renal disease (106 genes) - IPG
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
