ANALYTE:
WDR35

NAME:	WD repeat domain 35
SYMBOL:	WDR35
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FAP118 IFT121 IFTA1 KIAA1336 MGC33196
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/1600

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
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Cranioectodermal dysplasia
- Short rib-polydactyly syndrome type 5
- Short rib-polydactyly syndrome, Verma-Naumoff type

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
- Congenital malformation gene panel - VUB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Heterotaxie PCD - UGent
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
