

ANALYTE:
IFT80

NAME:	intraflagellar transport 80
SYMBOL:	IFT80
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FAP167 KIAA1374
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
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RELATED CONTENT

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- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophthisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation gene panel](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
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- [Skeletal dysplasia \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)

Related Diseases

- [Jeune syndrome](#)
- [Short rib-polydactyly syndrome, Beemer-Langer type](#)
- [Short rib-polydactyly syndrome, Verma-Naumoff type](#)

Related Gene Panels

- [Ciliopathy \(120 genes\) - UGent](#)
- [Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophthisis, 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
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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