

**ANALYTE:**  
**IFT52**

<b>NAME:</b>	intraflagellar transport 52
<b>SYMBOL:</b>	IFT52
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	CGI-53 NGD2 NGD5 dJ1028D15.1
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">Reactome</a> <a href="#">Ensembl</a> <a href="#">OMIM</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

- [Ciliopathy \(gene panel\)](#)
- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [cleft lip with/whitout cleft palate \(virtual gene panel\)](#)

### Related Diseases

- [Cranioectodermal dysplasia](#)

### 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](#)
- [Skeletal dysplasia \(genepanel\) - UZA](#)
- [Skeletal dysplasia - UGent](#)