# ANALYTE: DYNC2H1

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| **NAME:** | dynein cytoplasmic 2 heavy chain 1 |
| **SYMBOL:** | DYNC2H1 |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | DHC1b<br>DHC2<br>DYH1B<br>hdhc11 |
| **XREF(S):** | Orphanet<br>Genatlas<br>HGNC<br>OMIM<br>Reactome<br>SwissProt<br>Ensembl |
| **CREATED:** | 13 May 2019 - 01:01 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# 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
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Retinal dystrophy / RETNET (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

- Jeune syndrome
- Short rib-polydactyly syndrome, Majewski type
- Short rib-polydactyly syndrome, Saldino-Noonan type
- 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
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent

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