

**ANALYTE:**  
**CEP120**

<b>NAME:</b>	centrosomal protein 120
<b>SYMBOL:</b>	CEP120
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	FLJ36090
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</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

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## 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 \(gene panel - 1721 genes\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

### Related Diseases

- [Jeune syndrome](#)
- [Joubert syndrome](#)
- [Joubert syndrome with ocular defect](#)

### Related Gene Panels

- [Ciliopathy \(120 genes\) - UGent](#)
- [Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers \(146 genes\) - IPG](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Nephropathy panel - UGent](#)
- [Skeletal dysplasia \(394 genes\) - VUB](#)
- [Skeletal dysplasia \(genepanel\) - UZA](#)

- Skeletal dysplasia - UGent

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