

**ANALYTE:**  
**FANCL**

<b>NAME:</b>	FA complementation group L
<b>SYMBOL:</b>	FANCL
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	FAAP43 FLJ10335 Pog
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a> <a href="#">Ensembl</a> <a href="#">Genatlas</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

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Congenital malformation \(gene panel\)](#)
- [Congenital structural heart defects \(gene panel\)](#)
- [Familial cancer predisposition \(gene panel\)](#)
- [Fanconi anemia \(gene panel\)](#)
- [Intellectual disability \(gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [Pediatric oncopredisposition \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [cleft lip with/without cleft palate \(virtual gene panel\)](#)
- « [Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

### Related Diseases

- [Fanconi anemia](#)

### Related Gene Panels

- [Cleft lip and palate / dysmorphic facial features / craniofacial anomalies \(255 genes\) - UCL](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
- [Congenital structural heart defects - UGent](#)
- [Fanconi anemia - UGent](#)
- [Hematologic Familiar Forms - ULG](#)

- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel) - UZA
- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB

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