

ANALYTE:
FANCI

NAME:	FA complementation group I
SYMBOL:	FANCI
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ10719
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest-acc.healthdata.be/analyte/855>

RELATED CONTENT

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 \(virtual gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [Pediatric oncopredisposition \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [« Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

Related Diseases

- [Fanconi anemia](#)

Related Gene Panels

- [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 \(gene panel\)](#)

- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB

Source URL: <http://gentest-acc.healthdata.be/analyte/855>