# Belgian Genetic Tests database

## ANALYTE:
## FANCM

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|---|---|
| **NAME:** | FA complementation group M |
| **SYMBOL:** | FANCM |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | FAAP250 |
| **XREF(S):** | Orphanet<br>Ensembl<br>Genatlas<br>HGNC<br>OMIM<br>Reactome<br>SwissProt |
| **CREATED:** | 13 May 2019 - 01:01 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

Source URL: http://gentest-acc.healthdata.be/analyte/854

# RELATED CONTENT

## Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation (gene panel)
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - Primary Adrenal Insuffiency (gene panel)
- Familial cancer predisposition (gene panel)
- Fanconi anemia (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

## Related Diseases

- Fanconi anemia
- Male infertility with azoospermia or oligozoospermia due to single gene mutation

## Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - Primary Adrenal Insufficiency - UGent
- Fanconi anemia - UGent
- Hematologic Familiar Forms - ULG
- Hereditary cancer predisposition - UGent