ANALYTE:
FANCG

NAME:	FA complementation group G
SYMBOL:	FANCG
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DNA repair protein XRCC9 FAG X-ray repair complementing defective repair in Chinese hamster cells 9 X-ray repair, complementing defective, in Chinese hamster, 9
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

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)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders (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)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB
