Belgian Genetic Tests database

ANALYTE:
FANCA

NAME:	FA complementation group A
SYMBOL:	FANCA
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SYNONYMS:	FA-H FAA FAH
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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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)
- Hepatology (gene panel)
- Intellectual disability (virtual gene panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrogenetics / Nephropathy (gene panel)
- Neuropathy (gene panel)
- Pediatric oncopredisposition (gene panel)
- Skeletal dysplasia (gene panel)
- Tubulopathy (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
- Hepatology panel - UGent
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- Metabolic disorders (213 genes) - VUB
- Nephropathy panel - UGent
- Neuropathy panel - UGent
- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
