ANALYTE:
SLX4

NAME:	SLX4 structure-specific endonuclease subunit
SYMBOL:	SLX4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FANCP Fanconi anemia, complementation group P KIAA1784 KIAA1987
XREF(S):	Orphanet Ensembl HGNC Genatlas OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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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)
- 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
