Belgian Genetic Tests database

ANALYTE:
ERCC4

NAME:	ERCC excision repair 4, endonuclease catalytic subunit
SYMBOL:	ERCC4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FANCQ RAD1 xeroderma pigmentosum, complementation group F
XREF(S):	Orphanet HGNC OMIM Reactome SwissProt Ensembl Genatlas
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel)
- Ataxia Spasticity (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation (gene panel)
- Congenital structural heart defects (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Familial cancer predisposition (gene panel)
- Fanconi anemia (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Pediatric oncopredisposition (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Cockayne syndrome type 1
- Fanconi anemia
- Xeroderma pigmentosum
- Xeroderma pigmentosum-Cockayne syndrome complex

Related Gene Panels

- Ataxia - ULB
- Ataxia Spasticity - UGent
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Fanconi anemia - UGent
- Hematologic Familiar Forms - ULG
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (859 genes) - ULG
- Pediatric oncopredisposition - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia - UGent
- Skin disorders - UGent
