Belgian Genetic Tests database

ANALYTE:
RECQL4

NAME:	RecQ like helicase 4
SYMBOL:	RECQL4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	RecQ4
XREF(S):	Orphanet HGNC OMIM SwissProt Ensembl Genatlas
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Craniosynostosis (gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Baller-Gerold syndrome
- RAPADILINO syndrome
- Rothmund-Thomson syndrome type 2

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB

- Craniosynostosis (32 genes) - KUL
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
