Belgian Genetic Tests database

ANALYTE:
BLM

NAME:	BLM RecQ like helicase
SYMBOL:	BLM
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SYNONYMS:	BS RECQ2 RECQL3
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Bloom syndrome
- Bloom syndrome
- Breast cancer, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Constitutional Mismatch Repair Deficiency Syndrome + Bloom syndrome (5 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Familial cancer predisposition (gene panel)
- Hereditary cancer (Breast, ovary, colon) (26 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Lipodystrophy and/or hyperinsulinism (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders (gene panel)
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Skin disorders (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Bloom syndrome

Related Gene Panels

- Extended Breast Cancer Panel (26 gene) - VUB
- Cancer (Breast, ovary, colon,…) (26 genes) - ULG
- Congenital malformation (1721 genes) - ULB
- Constitutional Mismatch Repair Deficiency Syndrome / Bloom syndrome - KUL
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Hematologic Familiar Forms - ULG
- Hereditary breast and ovarian cancer (26 genes) - CHULg
- Hereditary cancer predisposition - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability (gene panel) - UZA
- Intellectual disability/Epilepsy (859 genes) - ULG
- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Skin disorders - UGent
