Belgian Genetic Tests database

ANALYTE:
CBL

NAME:	Cbl proto-oncogene
SYMBOL:	CBL
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	RNF55 c-Cbl oncogene CBL2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (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)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Intellectual Disability (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (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)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Primary immune deficiencies (gene panel)
- RASopathy (gene panel)
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)
- Skin disorders (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Aggressive systemic mastocytosis
- Juvenile myelomonocytic leukemia
- Noonan syndrome
- Noonan syndrome-like disorder with juvenile myelomonocytic leukemia

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Growth retardation/short stature (genepanel) - UZA
- Hematologic Familiar Forms - ULG
- Heterotaxie PCD - UGent
- Intellectual Disability (104 genes) - IPG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability (gene panel) - UZA
- Intellectual disability/Epilepsy (859 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies - UGent
- RASopathy - KUL
- Short Stature (46 genes) - IPG
- Skin disorders - UGent

- cardiopathy panel - UGent
