Belgian Genetic Tests database

ANALYTE:
RUNX1

NAME:	RUNX family transcription factor 1
SYMBOL:	RUNX1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	AMLCR1 PEBP2A2 aml1 oncogene
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Myeloid neoplasms with germline predisposition (Hereditary MDS/Acute Leukemia) (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders (gene panel)
- Pediatric oncopredisposition (gene panel)
- Trombosis - Hemostasis (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Acute myeloid leukemia with t(8;21)(q22;q22) translocation
- Aggressive systemic mastocytosis
- B-lymphoblastic leukemia/lymphoma with recurrent genetic abnormality
- Chronic myeloid leukemia
- Familial platelet disorder with associated myeloid malignancy
- Isolated delta-storage pool disease
- Precursor B-cell acute lymphoblastic leukemia

Related Gene Panels

- Hematologic Familiar Forms - ULG
- Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB

- Pediatric oncopredisposition - UGent
- Trombosis - Hemostasis (108 genes) - KUL
