Belgian Genetic Tests database

ANALYTE:
GATA2

NAME:	GATA binding protein 2
SYMBOL:	GATA2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	NFE1B
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Arteriovenous malformation (gene panel)
- Capillary malformation - arteriovenous malformation (2 genes)
- Cerebral cavernous malformation (gene panel)
- Child Interstitial Lung Disease (child - gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Emberger syndrome / Immunodeficiency 21
- Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)
- Hereditary cancer (gene panel)
- Intellectual disability (virtual gene panel)
- Myeloid neoplasms with germline predisposition (Hereditary MDS/Acute Leukemia) (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Primary lymphedema / fetal hydrops (gene panel)
- Pulmonary Fibrosis (gene panel) + rs35705950 of MUC5B gene
- Rendu-Osler-Weber disease (4 genes)
- Respiratory disorders (gene panel): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease
- Venous malformation (2 genes)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Deafness-lymphedema-leukemia syndrome
- Inherited acute myeloid leukemia
- Monocytopenia with susceptibility to infections
- Myelodysplastic syndrome
- Unclassified myelodysplastic syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Erythocyoses, polycythémies, thrombocytoses congénitales (gene panel) - ULG
- Hematologic Familiar Forms - ULG
- Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes (gene panel)
- Hereditary predisposition to cancer (47 genes) - IPG
- Immunogenetics (21 genes)
- Intellectual disability (gene panel)
- Lymphedema / fetal hydrops (27 genes) - UCL
- Pediatric oncopredisposition - UGent
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Pulmonary Fibrosis (21 genes) + rs35705950 (MUC5B gene) - KUL
- Respiratory Disorders panel (137 genes) - Ugent
- Vascular malformations (germline) (38 genes) - UCL
- chILD (35 genes) - KUL
- test-test
