Belgian Genetic Tests database

ANALYTE:
STAT3

NAME:	signal transducer and activator of transcription 3
SYMBOL:	STAT3
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	APRF
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Autoimmune disease, multisystem, infantile-onset (ADMIO) / Hyper-IgE recurrent infection syndrome
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation (gene panel)
- Congenital structural heart defects (gene panel)
- Erythrocytoses, polycythémies, thrombocytoses et neutropénies congénitales (gene panel)
- Myeloid neoplasms with germline predisposition (Hereditary MDS/Acute Leukemia) (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Respiratory disorders (gene panel): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)

Related Diseases

- Acute promyelocytic leukemia
- Autosomal dominant hyper-IgE syndrome
- Chronic lymphoproliferative disorder of natural killer cells
- Isolated permanent neonatal diabetes mellitus
- STAT3-related early-onset multisystem autoimmune disease
- T-cell large granular lymphocyte leukemia

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Erythocyoses, polycythémies, thrombocytoses congénitales (gene panel) - ULG
- Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes (gene panel)
- Immunogenetics (21 genes)
- Primary immune deficiencies (444 genes) - KUL
- Primary immune deficiencies - UGent
- Respiratory Disorders panel (137 genes) - Ugent
- Skeletal dysplasia (394 genes) - VUB
- Skin disorders - UGent
