ANALYTE:
RTEL1

NAME:	regulator of telomere elongation helicase 1
SYMBOL:	RTEL1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	DKFZP434C013 KIAA1088 NHL RTEL bK3184A7.3
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest-acc.healthdata.be/analyte/719

RELATED CONTENT

Related Genetic Tests

- Child Interstitial Lung Disease (child - gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Dyskeratosis Congenita (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 immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Pulmonary Fibrosis (gene panel) + rs35705950 of MUC5B gene
- Respiratory disorders (gene panel): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Dyskeratosis congenita
- Hoyeraal-Hreidarsson syndrome
- Idiopathic pulmonary fibrosis

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Dyskeratosis Congenita (18 genes) - KUL
- Hematologic Familiar Forms - ULG
- 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 (444 genes) - KUL
- Primary immune deficiencies - UGent
- Pulmonary Fibrosis (21 genes) + rs35705950 (MUC5B gene) - KUL
- Respiratory Disorders panel (137 genes) - Ugent
- chILD (35 genes) - KUL
