ANALYTE:
TERT

NAME:	telomerase reverse transcriptase
SYMBOL:	TERT
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SYNONYMS:	EST2 TCS1 TP2 TRT hEST2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Child Interstitial Lung Disease (child - gene panel)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Dyskeratosis Congenita (gene panel)
- Familial cancer predisposition (gene panel)
- Familial melanoma / Familial Atypical Multiple Mole Melanoma Syndrome, FAMMM (gene panel)
- Hereditary Melanoma Panel (7 genes)
- Intellectual disability (virtual gene panel)
- Maffucci syndrome (gene panel)
- Melanoma / Familial Atypical Multiple Mole Melanoma Syndrome (gene panel)
- Myeloid neoplasms with germline predisposition (Hereditary MDS/Acute Leukemia) (gene panel)
- Myeloid/lymphoid neoplasms with germline predisposition (gene panel)
- Overgrowth & vascular anomalies (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
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Adrenocortical carcinoma

- Clear cell sarcoma of kidney
- Differentiated thyroid carcinoma
- Dyskeratosis congenita
- Familial melanoma
- Hoyeraal-Hreidarsson syndrome
- Idiopathic aplastic anemia
- Idiopathic pulmonary fibrosis
- Meningioma

Related Gene Panels

- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Dyskeratosis Congenita (18 genes) - KUL
- Familial melanoma - UGent
- Hematologic Familiar Forms - ULG
- Hereditary Melanoma Panel (7 genes) - ULG
- Hereditary Myelodysplastic /Acute Leukemia Predisposition Syndromes (gene panel)
- Hereditary cancer predisposition - UGent
- Intellectual disability (gene panel)
- Maffucci syndrome (65 genes) - KUL
- Melanoma and Familial Atypical Multiple Mole Melanoma Syndrome (8 genes) - KUL
- Overgrowth & vascular anomalies (65 genes) - KUL
- 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
- Skeletal dysplasia - UGent
- Sturge-Weber syndrome (65 genes) - KUL
- chILD (35 genes) - KUL
