ANALYTE:
CTC1

NAME:	CST telomere replication complex component 1
SYMBOL:	CTC1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	AAF132 Conserved telomere capping protein 1 Conserved telomere maintenance component 1 FLJ22170 alpha accessory factor 132 conserved telomere capping protein 1 conserved telomere maintenance component 1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Dyskeratosis Congenita (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Movement Disorders (gene panel)
- Movement Disorders (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)
- « Inherited bone marrow failures syndromes » with or without organ dysfunction

Related Diseases

- Coats plus syndrome
- Dyskeratosis congenita

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Dyskeratosis Congenita (18 genes) - KUL
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hematologic Familiar Forms - ULG
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability (gene panel) - UZA
- Intellectual disability/Epilepsy (859 genes) - ULG
- Leukodystrophy - UGent
- Movement Disorders - UGent
- Movement Disorders - 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
