ANALYTE:
ACD

NAME:	ACD shelterin complex subunit and telomerase recruitment factor
SYMBOL:	ACD
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	POT1 and TIN2 organizing protein Pip1 Ptop TIN2 interacting protein 1 Tint1 Tpp1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia Spasticity (gene panel)
- 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)
- Epilepsy gene panel
- Familial cancer predisposition (gene panel)
- Familial melanoma / Familial Atypical Multiple Mole Melanoma Syndrome, FAMMM (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Leukodystrophy (gene panel)
- Lysosomal Storage Disease (gene panel)
- Melanoma / Familial Atypical Multiple Mole Melanoma Syndrome (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Pediatric oncopredisposition (gene panel)
- Primary immune deficiencies (gene panel)
- Primary immune deficiencies (gene panel)
- Progressive Myoclonic Epilepsy (PME) (gene panel)
- Pulmonary Fibrosis (gene panel) + rs35705950 of MUC5B gene
- test

Related Diseases

- Familial melanoma
- Hereditary isolated aplastic anemia
- Hoyeraal-Hreidarsson syndrome

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- 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
- Familial melanoma - UGent
- Hereditary cancer predisposition - UGent
- Intellectual disability & Epilepsy - UGent
- Leukodystrophy - UGent
- Lysomal Storage (64 genes) - VUB
- Melanoma and Familial Atypical Multiple Mole Melanoma Syndrome (8 genes) - KUL
- 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
- Progressive Myoclonic Epilepsy - UGent
- Pulmonary Fibrosis (21 genes) + rs35705950 (MUC5B gene) - KUL
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- test
- test1
