

ANALYTE:
TINF2

NAME:	TERF1 interacting nuclear factor 2
SYMBOL:	TINF2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	TIN2
XREF(S):	<u>Orphanet</u> <u>Ensembl</u> <u>Genatlas</u> <u>HGNC</u> <u>OMIM</u> <u>Reactome</u> <u>SwissProt</u>
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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 \(gene panel\)](#)
- [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](#)
- [Skin disorders \(gene panel\)](#)
- [« Inherited bone marrow failures syndromes » with or without organ dysfunction](#)

Related Diseases

- [Dyskeratosis congenita](#)
- [Hoyeraal-Hreidarsson syndrome](#)
- [Revesz syndrome](#)

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
- Skin disorders - UGent

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