

ANALYTE:
PARN

NAME:	poly(A)-specific ribonuclease
SYMBOL:	PARN
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SYNONYMS:	DAN deadenylation nuclease
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- [Child Interstitial Lung Disease \(child - gene panel\)](#)
- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophthisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [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 \(virtual gene panel\)](#)
- [Myeloid/lymphoid neoplasms with germline predisposition \(gene panel\)](#)
- [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

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (148genes) - IPG
- 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
- 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

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