

# **Product datasheet for TP300266**

### OriGene Technologies, Inc.

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### TXNL4A (NM 006701) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human thioredoxin-like 4A (TXNL4A), 20 μg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC200266 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MSYMLPHLHNGWQVDQAILSEEDRVVVIRFGHDWDPTCMKMDEVLYSIAEKVKNFAVIYLVDITEVPDFN KMYELYDPCTVMFFFRNKHIMIDLGTGNNNKINWAMEDKQEMVDIIETVYRGARKGRGLVVSPKDYSTKY

RY

**TRTRPL**EQKLISEEDLAANDILDYKDDDDK**V** 

Tag: C-Myc/DDK

**Predicted MW:** 16.6 kDa

Concentration:  $>0.05 \mu g/\mu L$  as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

**Preparation:** Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

**Note:** For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 006692

 Locus ID:
 10907

 UniProt ID:
 P83876

 RefSeq Size:
 1415



#### TXNL4A (NM\_006701) Human Recombinant Protein - TP300266

Cytogenetics: 18q23

RefSeg ORF: 426

Synonyms: BMKS; DIB1; DIM1; SNRNP15; TXNL4; U5-15kD

**Summary:** The protein encoded by this gene is a member of the U5 small ribonucleoprotein particle

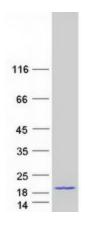
(snRNP), and is involved in pre-mRNA splicing. This protein contains a thioredoxin-like fold and it is expected to interact with multiple proteins. Protein-protein interactions have been observed with the polyglutamine tract-binding protein 1 (PQBP1). Mutations in both the coding region and promoter region of this gene have been associated with Burn-McKeown syndrome, which is a rare disorder characterized by craniofacial dysmorphisms, cardiac defects, hearing loss, and bilateral choanal atresia. A pseudogene of this gene is found on chromosome 2. Alternative splicing results in multiple transcript variants. [provided by

RefSeq, Mar 2015]

**Protein Families:** Druggable Genome

**Protein Pathways:** Spliceosome

# **Product images:**



Coomassie blue staining of purified TXNL4A protein (Cat# TP300266). The protein was produced from HEK293T cells transfected with TXNL4A cDNA clone (Cat# [RC200266]) using

MegaTran 2.0 (Cat# [TT210002]).