

# **Product datasheet for TP300266L**

# OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

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

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human thioredoxin-like 4A (TXNL4A), 1 mg

Species: Human
Expression Host: HEK293T

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

MSYMLPHLHNGWQVDQAILSEEDRVVVIRFGHDWDPTCMKMDEVLYSIAEKVKNFAVIYLVDITEVPDFN KMYELYDPCTVMFFFRNKHIMIDLGTGNNNKINWAMEDKQEMVDIIETVYRGARKGRGLVVSPKDYSTK

Y RY

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

**Predicted MW:** 16.6 kDa

**Concentration:** >0.05 μg/μL 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



#### TXNL4A (NM\_006701) Human Recombinant Protein - TP300266L

RefSeq Size: 1415

Cytogenetics: 18q23 RefSeq 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

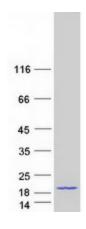
(snRNP), and is involved in pre-mRNA splicing. This protein contains a thioredoxin-like fold and it is expected to interact with multiple proteins. Protein-protein interactions have been observed with the polyglutamine tract-binding protein 1 (PQBP1). Mutations in both the coding region and promoter 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]).