

## **Product datasheet for TP304771**

## OriGene Technologies, Inc.

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## TIMM8A (NM\_004085) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human translocase of inner mitochondrial membrane 8 homolog A

(yeast) (TIMM8A), nuclear gene encoding mitochondrial protein, transcript variant 1, 20 µg

Species: Human
Expression Host: HEK293T

Expression riose.

**Expression cDNA Clone** >RC204771 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MDSSSSSSAAGLGAVDPQLQHFIEVETQKQRFQQLVHQMTELCWEKCMDKPGPKLDSRAEACFVNCVE

RF

IDTSQFILNRLEQTQKSKPVFSESLSD

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK

Predicted MW: 10.8 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 004076

Locus ID: 1678

UniProt ID: 060220





RefSeq Size: 1459

Cytogenetics: Xq22.1 RefSeq ORF: 291

Synonyms: DDP; DDP1; DFN1; MTS; TIM8

**Summary:** This translocase is involved in the import and insertion of hydrophobic membrane proteins

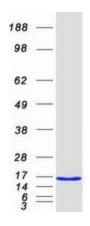
> from the cytoplasm into the mitochondrial inner membrane. The gene is mutated in Mohr-Tranebjaerg syndrome/Deafness Dystonia Syndrome (MTS/DDS) and it is postulated that MTS/DDS is a mitochondrial disease caused by a defective mitochondrial protein import system. Defects in this gene also cause Jensen syndrome; an X-linked disease with

opticoacoustic nerve atrophy and muscle weakness. This protein, along with TIMM13, forms a 70 kDa heterohexamer. Alternative splicing results in multiple transcript variants encoding

distinct isoforms.[provided by RefSeq, Mar 2009]

**Protein Families:** Druggable Genome

## **Product images:**



Coomassie blue staining of purified TIMM8A protein (Cat# TP304771). The protein was produced from HEK293T cells transfected with TIMM8A cDNA clone (Cat# [RC204771]) using MegaTran 2.0 (Cat# [TT210002]).