

## **Product datasheet for TP304007**

#### OriGene Technologies, Inc.

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# Tropomyosin 2 (TPM2) (NM\_213674) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human tropomyosin 2 (beta) (TPM2), transcript variant 2, 20 μg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC204007 protein sequence or AA Sequence: Red=Cloning site Green=Tags(s)

MDAIKKKMQMLKLDKENAIDRAEQAEADKKQAEDRCKQLEEEQQALQKKLKGTEDEVEKYSESVKEAQE

Κ

LEQAEKKATDAEADVASLNRRIQLVEEELDRAQERLATALQKLEEAEKAADESERGMKVIENRAMKDEEK MELQEMQLKEAKHIAEDSDRKYEEVARKLVILEGELERSEERAEVAESRARQLEEELRTMDQALKSLMAS EEEYSTKEDKYEEEIKLLEEKLKEAETRAEFAERSVAKLEKTIDDLEETLASAKEENVEIHQTLDQTLLE

LNNL

**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK
Predicted MW: 32.8 kDa

Concentration:

Predicted MW: 32.8 kDa

**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

>0.05 µg/µL as determined by microplate BCA method

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.

RefSeg: NP 998839

**Locus ID:** 7169





#### Tropomyosin 2 (TPM2) (NM\_213674) Human Recombinant Protein - TP304007

**UniProt ID:** <u>P07951</u>

RefSeq Size: 1182 Cytogenetics: 9p13.3 RefSeq ORF: 852

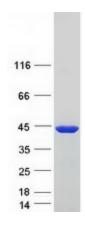
Synonyms: AMCD1; DA1; DA2B; DA2B4; HEL-S-273; NEM4; TMSB

Summary: This gene encodes beta-tropomyosin, a member of the actin filament binding protein family,

and mainly expressed in slow, type 1 muscle fibers. Mutations in this gene can alter the expression of other sarcomeric tropomyosin proteins, and cause cap disease, nemaline myopathy and distal arthrogryposis syndromes. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Mar 2009]

**Protein Pathways:** Cardiac muscle contraction, Dilated cardiomyopathy, Hypertrophic cardiomyopathy (HCM)

### **Product images:**



Coomassie blue staining of purified TPM2 protein (Cat# TP304007). The protein was produced from HEK293T cells transfected with TPM2 cDNA clone (Cat# [RC204007]) using MegaTran 2.0 (Cat# [TT210002]).