

## **Product datasheet for MC226789**

## Tpm2 (NM\_001277876) Mouse Untagged Clone

**Product data:** 

**Product Type:** Expression Plasmids

**Product Name:** Tpm2 (NM\_001277876) Mouse Untagged Clone

Tag: Tag Free Symbol: Tpm2

Synonyms:Tpm-; Tpm-2; Tro; Trop-2Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Cell Selection: Neomycin

Fully Sequenced ORF: >MC226789 representing NM\_001277876

Red=Cloning site Blue=ORF Orange=Stop codon

 ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT

ACAAGGATGACGACGATAAGGTTTAA

CTCAACAACCTGTAA

**Restriction Sites:** Sgfl-Mlul

**ACCN:** NM\_001277876

**Insert Size:** 855 bp



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## Tpm2 (NM\_001277876) Mouse Untagged Clone - MC226789

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

**OTI Annotation:** Clone contains native stop codon, and expresses the complete ORF without any c-terminal

tag.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

**Reconstitution Method:** 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 001277876.1</u>, <u>NP 001264805.1</u>

RefSeq Size: 2108 bp
RefSeq ORF: 855 bp
Locus ID: 22004
Cytogenetics: 4 A5

**Gene Summary:** This gene belongs to the tropomyosin family which encodes proteins that bind to actin

filaments and stabilize them by regulating access to actin modifying proteins. The encoded protein is a high molecular weight tropomyosin expressed in slow skeletal muscle. In humans, mutations in this gene are associated with nemaline myopathy, cap disease and distal arthrogryposis syndromes. Alternative splicing of this gene results in multiple transcript

variants encoding different isoforms. [provided by RefSeq, Apr 2013]

Transcript Variant: This variant (Tpm2.1, also known as variant 3) differs in the 3' UTR and has multiple differences in the 3' coding region, compared to variant Tpm2.2. It encodes isoform Tpm2.1sm/cy (also known as isoform 3), which is of the same size but has a different internal

segment and a distinct C-terminus, compared to isoform Tpm2.2st.