

Product datasheet for MC209557

Tnnt1 (NM_011618) Mouse Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: Tnnt1 (NM_011618) Mouse Untagged Clone

Tag: Tag Free
Symbol: Tnnt1

Synonyms: AW146156; ss; ssTnT; sT; sTnT; Tn; Tnt

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Cell Selection: Neomycin

Fully Sequenced ORF: >MC209557 representing NM_011618

Red=Cloning site Blue=ORF Orange=Stop codon

TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC

GCCGCGATCGCC

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT

ACAAGGATGACGACGATAAGGTTTAA

Restriction Sites: Sgfl-Mlul ACCN: NM_011618

Insert Size: 786 bp



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Tnnt1 (NM_011618) Mouse Untagged Clone - MC209557

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).

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 011618.2</u>, <u>NP 035748.1</u>

 RefSeq Size:
 1027 bp

 RefSeq ORF:
 786 bp

 Locus ID:
 21955

 UniProt ID:
 088346

 Cytogenetics:
 7 2.6 cM

Gene Summary: This gene encodes the slow skeletal tropomyosin-binding subunit of the troponin complex

and plays an essential role in the regulation of striated muscle contraction. In humans, mutations in this gene are associated with nemaline myopathy type 5. Alternative splicing of this gene results in multiple transcript variants encoding different isoforms. [provided by

RefSeq, Apr 2013]

Transcript Variant: This variant (2) uses an alternate in-frame splice site in the 5' coding region, compared to variant 1. This results in a shorter protein (isoform 2, also known as high

Mr isoform 2), compared to isoform 1.