

## **Product datasheet for TP710220**

## OriGene Technologies, Inc.

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## Troponin T1 (TNNT1) (NM\_003283) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human troponin T type 1 (skeletal, slow) (TNNT1), transcript

variant 1, full length, with C-terminal DDK tag, expressed in sf9, 20ug

Species: Human

**Expression Host:** Sf9

Expression cDNA Clone

or AA Sequence:

A DNA sequence from TrueORF clone, RC221318, encoding human full-length TNNT1

Tag: C-DDK

Predicted MW: 32.8 kDa

**Concentration:** >0.05 μg/μL as determined by microplate BCA method

**Purity:** > 80% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** 50 mM Tris-HCl, 100 mM glycine, pH 8.0, 10% glycerol

**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 003274

**Locus ID:** 7138

UniProt ID: P13805

RefSeq Size: 980

Cytogenetics: 19q13.42

RefSeq ORF: 834

Synonyms: ANM; NEM5; STNT; TNT; TNTS





**Summary:** 

This gene encodes a protein that is a subunit of troponin, which is a regulatory complex located on the thin filament of the sarcomere. This complex regulates striated muscle contraction in response to fluctuations in intracellular calcium concentration. This complex is composed of three subunits: troponin C, which binds calcium, troponin T, which binds tropomyosin, and troponin I, which is an inhibitory subunit. This protein is the slow skeletal troponin T subunit. Mutations in this gene cause nemaline myopathy type 5, also known as Amish nemaline myopathy, a neuromuscular disorder characterized by muscle weakness and rod-shaped, or nemaline, inclusions in skeletal muscle fibers which affects infants, resulting in death due to respiratory insufficiency, usually in the second year. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

Protein Families:

Druggable Genome

## **Product images:**

