

## **Product datasheet for SC202092**

## OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com
techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

## Troponin C1 (TNNC1) (NM\_003280) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: Troponin C1 (TNNC1) (NM\_003280) Human 3' UTR Clone

Symbol: TNNC1

Synonyms: CMD1Z; CMH13; TN-C; TNC; TNNC

**Mammalian Cell** 

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_003280

**Insert Size:** 205 bp

Insert Sequence: >SC202092 3'UTR clone of NM\_003280

The sequence shown below is from the reference sequence of NM\_003280. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul

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

point of reference. Note that the complete sequence of this clone is largely the same as the

reference sequence but may contain minor differences, e.g., single nucleotide

polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

**RefSeq:** <u>NM 003280.3</u>





## Troponin C1 (TNNC1) (NM\_003280) Human 3' UTR Clone - SC202092

**Summary:** Troponin is a central regulatory protein of striated muscle contraction, and together with

tropomyosin, is located on the actin filament. Troponin consists of 3 subunits: TnI, which is the inhibitor of actomyosin ATPase; TnT, which contains the binding site for tropomyosin; and TnC, the protein encoded by this gene. The binding of calcium to TnC abolishes the inhibitory action of TnI, thus allowing the interaction of actin with myosin, the hydrolysis of ATP, and the generation of tension. Mutations in this gene are associated with cardiomyopathy dilated

type 1Z. [provided by RefSeq, Oct 2008]

**Locus ID:** 7134

MW: 7.2