

## Product datasheet for SC211691

## OriGene Technologies, Inc.

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## TM4SF2 (TSPAN7) (NM 004615) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

Product Name: TM4SF2 (TSPAN7) (NM\_004615) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: TSPAN7

**Synonyms:** A15; CCG-B7; CD231; DXS1692E; MRX58; MXS1; TALLA-1; TM4SF2; TM4SF2b

**ACCN:** NM\_004615

**Insert Size:** 1017 bp

Insert Sequence: >SC211691 3'UTR clone of NM\_004615

The sequence shown below is from the reference sequence of NM\_004615. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

 ${\tt TATTGTGAACATTTGTGATATATGTATTAATAAATAGAGCAATTACAAGCA}$ 

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

**Restriction Sites:** Sgfl-Mlul





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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 004615.4</u>

**Summary:** The protein encoded by this gene is a member of the transmembrane 4 superfamily, also

known as the tetraspanin family. Most of these members are cell-surface proteins that are characterized by the presence of four hydrophobic domains. The proteins mediate signal transduction events that play a role in the regulation of cell development, activation, growth and motility. This encoded protein is a cell surface glycoprotein and may have a role in the control of neurite outgrowth. It is known to complex with integrins. This gene is associated with X-linked cognitive disability and neuropsychiatric diseases such as Huntington's chorea,

fragile X syndrome and myotonic dystrophy. [provided by RefSeq, Jul 2008]

Locus ID: 7102 MW: 38.6