

Product datasheet for SC335377

TSSC4 (NM 001297659) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: TSSC4 (NM_001297659) Human Untagged Clone

Tag: Tag Free Symbol: TSSC4

Mammalian Cell Neomycin

Selection:

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

Fully Sequenced ORF: >SC335377 representing NM_001297659.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ACGCGTACGCGCCCCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

GAGGACCCAGGTGCTGAGGTCTGA

Restriction Sites: Sgfl-Mlul



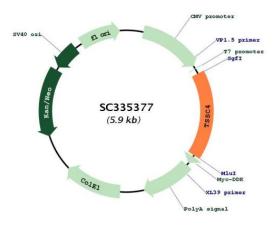
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Plasmid Map:



ACCN: NM_001297659

Insert Size: 990 bp

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.

Q9Y5U2

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: NM 001297659.1

RefSeq Size:1906 bpRefSeq ORF:990 bpLocus ID:10078

UniProt ID:



TSSC4 (NM_001297659) Human Untagged Clone - SC335377

Cytogenetics: 11p15.5

Protein Families: Druggable Genome

MW: 34.3 kDa

Gene Summary: This gene is one of several tumor-suppressing subtransferable fragments located in the

imprinted gene domain of 11p15.5, an important tumor-suppressor gene region. Alterations in this region have been associated with the Beckwith-Wiedemann syndrome, Wilms tumor, rhabdomyosarcoma, adrenocortical carcinoma, and lung, ovarian, and breast cancer. This gene is located among several imprinted genes; however, this gene, as well as the panhematopoietic expression gene (PHEMX), escapes imprinting. This gene may play a role in malignancies and disease that involve this region. Several transcript variants encoding two

different isoforms have been found for this gene. [provided by RefSeq, Jul 2014]

Transcript Variant: This variant (2) differs in the 5' UTR compared to variant 1. Variants 1, 2, 3,

and 4 all encode the same isoform (a).