

## Product datasheet for **SC329157**

### TRPV4 (NM\_001177428) Human Untagged Clone

#### Product data:

**Product Type:** Expression Plasmids  
**Product Name:** TRPV4 (NM\_001177428) Human Untagged Clone  
**Tag:** Tag Free  
**Symbol:** TRPV4  
**Synonyms:** BCYM3; CMT2C; HMSN2C; OTRPC4; SMAL; SPSMA; SSQTL1; TRP12; VRL2; VROAC  
**Mammalian Cell Selection:** None  
**Vector:** [pCMV6-XL4](#)  
**E. coli Selection:** Ampicillin (100 ug/mL)

**Fully Sequenced ORF:** >OriGene sequence for NM\_001177428 edited  
GCGGATCTCCCGCCCGCGGCCAGCCCGTCCCGGAGGCTGAGCAGTGCAGACGGGCC  
GGGGCAGGCATGGCGGATTCCAGCGAAGGCCCGCGCGGGGCCGGGGAGGTGGCTGAG  
CTCCCCGGGGATGAGAGTGGCACCCAGGTGGGAGGCTTTTCTCTCTCCTCCCTGGCC  
AATCTGTTTGAGGGGAGGATGGCTCCCTTTCGCCCTCACCGGCTGATGCCAGTCGCCCT  
GCTGGCCAGGCGATGGGCGACCAATCTGCGCATGAAGTTCCAGGGCGCCTTCCGCAAG  
GGGGTGCCAACCCCATCGATCTGCTGGAGTCCACCCTATATGAGTCCTCGGTGGTGCCT  
GGGCCAAGAAGCACCATGGACTCACTGTTTACTACGGCACCTATCGTCACCACTCC  
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CCTGCCCTCAGCCGCCCCCATCCTCAAAGTCTTCAACCGGCCATCCTCTTTGACATC  
GTGTCCCGGGGCTCCACTGCTGACCTGGACGGGCTGCTCCATTCTTGTGACCCACAAG  
AAACGCCTAACTGATGAGGAGTTTCGAGAGCCATCTACGGGAAGACCTGCCTGCCCAAG  
GCCTTGCTGAACCTGAGCAATGGCCGCAACGACACCATCCTGTGCTGCTGGACATCGCG  
GAGCGCACCGGCAACATGCGGGAGTTTCACTAAGTCCGCCCTTCCGTGACATCTACTATCGA  
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GTGCTCAACAACGACGGCCTCTCGCCCTCATGATGGCTGCCAAGACGGGCAAGATTGGG  
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GACACGTGTGGGAAGAGGCCTCCGTGCTGGAGATCCTGGTGTACAACAGCAAGATTGAG  
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ACGGTGGACTACCTGCGGCTGGCTGGCGAGGTCAATACGCTCTTACTGGGGTCTCTTTC  
TTCTTCCCAACATCAAAGACTTGTTTCATGAAGAAATGCCCTGGAGTGAATTCTCTCTTC



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ATTGATGGCTCCTTCCAGCTGCTCTACTTCATCTACTCTGCTCCTGGTGATCGTCTCAGCA
GCCCTCTACCTGGCAGGGATCGAGGCTACCTGGCCGTGATGGTCTTTGCCCTGGTCCTG
GGCTGGATGAATGCCCTTTACTTCACCCGTGGGCTGAAGCTGACGGGGACCTATAGCATC
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TTTCTAGTCCAGCCGATTTTCCAGAGTGCCTTCTGGGGTGTCCCCCACACCCTGCTTTG
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GGCTCTGCCTCCCCACCTGGGGTGGGGCTCCCGGCCACCTGTCTTGTCTCTATGGAGT
CACATAAGCCAACGCCAGAGCCCCTCCACCTCAGGCCCCAGCCCCTGCCTCTCCATTATT
TATTTGCTCTGCTCTCAGGAAGCGACGTGACCCCTGCCCCAGCTGGAACCTGGCAGAGGC
CTTAGCACCCCGTTCCAAGTGCCTGCCCCGCAAGCCCAGCCTCAGCCTGCGCCTGAG
CTGCATGCGCCACCAATTTTGGCAGCGTGGCAGCTTTGCAAGGGGCTGGGCCCCTCGGC
TGGGGCCATGCCTTCTGTGTGTTCTGTAGTGTCTGGGATTTGCCGGTGTCAATAAATGT
TTATTCATTGAAAAAAAAAAAAAAAAAAAAA
    
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- Restriction Sites:** Please inquire
- ACCN:** NM\_001177428
- Insert Size:** 3100 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).
- OTI Annotation:** This TrueClone is provided through our Custom Cloning Process that includes sub-cloning into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.
- 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_001177428.1, NP_001170899.1</u>
RefSeq Size:	3051 bp
RefSeq ORF:	2475 bp
Locus ID:	59341
UniProt ID:	<u>Q9HBA0</u>
Cytogenetics:	12q24.11
Protein Families:	Druggable Genome, Ion Channels: Transient receptor potential, Transmembrane
Gene Summary:	<p>This gene encodes a member of the OSM9-like transient receptor potential channel (OTRPC) subfamily in the transient receptor potential (TRP) superfamily of ion channels. The encoded protein is a Ca<sup>2+</sup>-permeable, nonselective cation channel that is thought to be involved in the regulation of systemic osmotic pressure. Mutations in this gene are the cause of spondylometaphyseal and metatropic dysplasia and hereditary motor and sensory neuropathy type IIC. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Apr 2010]</p> <p>Transcript Variant: This variant (5) lacks an in-frame exon in the coding region, compared to variant 1. The encoded isoform (c) is shorter, compared to isoform a.</p>