

Product datasheet for RC218941L1V

OriGene Technologies, Inc.

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TRPC4 (NM 016179) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: TRPC4 (NM_016179) Human Tagged ORF Clone Lentiviral Particle

Symbol: TRPC4

Synonyms: HTRP-4; HTRP4; TRP4

Mammalian Cell

Selection:

None

Vector: pLenti-C-Myc-DDK (PS100064)

Tag: Myc-DDK
ACCN: NM_016179

ORF Size: 2931 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC218941).

Sequence:

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This

naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

RefSeq: <u>NM 016179.1</u>

RefSeq Size: 3440 bp
RefSeq ORF: 2934 bp
Locus ID: 7223

UniProt ID: Q9UBN4

Cytogenetics: 13q13.3

Domains: ANK, ion_trans

Protein Families: Druggable Genome, Ion Channels: Transient receptor potential, Transmembrane





ORIGENE

MW: 111.9 kDa

Gene Summary:

This gene encodes a member of the canonical subfamily of transient receptor potential cation channels. The encoded protein forms a non-selective calcium-permeable cation channel that is activated by Gq-coupled receptors and tyrosine kinases, and plays a role in multiple processes including endothelial permeability, vasodilation, neurotransmitter release and cell proliferation. Single nucleotide polymorphisms in this gene may be associated with generalized epilepsy with photosensitivity. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene. [provided by RefSeq, Aug 2011]