

**TRPV4 Antibody (N-term) Blocking Peptide**  
**Synthetic peptide**  
**Catalog # BP18990a****Specification**

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**TRPV4 Antibody (N-term) Blocking Peptide - Product Information**Primary Accession [Q9HBA0](#)**TRPV4 Antibody (N-term) Blocking Peptide - Additional Information****Gene ID** 59341**Other Names**

Transient receptor potential cation channel subfamily V member 4, TrpV4, Osm-9-like TRP channel 4, OTRPC4, Transient receptor potential protein 12, TRP12, Vanilloid receptor-like channel 2, Vanilloid receptor-like protein 2, VRL-2, Vanilloid receptor-related osmotically-activated channel, VR-OAC, TRPV4, VRL2, VROAC

**Format**

Peptides are lyophilized in a solid powder format. Peptides can be reconstituted in solution using the appropriate buffer as needed.

**Storage**

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C.

**Precautions**

This product is for research use only. Not for use in diagnostic or therapeutic procedures.

**TRPV4 Antibody (N-term) Blocking Peptide - Protein Information****Name** TRPV4**Synonyms** VRL2, VROAC**Function**

Non-selective calcium permeant cation channel involved in osmotic sensitivity and mechanosensitivity (PubMed:<a href="http://www.uniprot.org/citations/16293632" target="\_blank">16293632</a>, PubMed:<a href="http://www.uniprot.org/citations/18826956" target="\_blank">18826956</a>, PubMed:<a href="http://www.uniprot.org/citations/18695040" target="\_blank">18695040</a>, PubMed:<a href="http://www.uniprot.org/citations/29899501" target="\_blank">29899501</a>, PubMed:<a href="http://www.uniprot.org/citations/22526352" target="\_blank">22526352</a>, PubMed:<a href="http://www.uniprot.org/citations/23136043" target="\_blank">23136043</a>). Activation by exposure to hypotonicity within the physiological range exhibits an outward rectification (PubMed:<a href="http://www.uniprot.org/citations/18826956" target="\_blank">18826956</a>, PubMed:<a href="http://www.uniprot.org/citations/18695040" target="\_blank">18695040</a>, PubMed:<a href="http://www.uniprot.org/citations/29899501" target="\_blank">29899501</a>). Also activated by heat, low pH, citrate and phorbol esters (PubMed:<a

[16293632](http://www.uniprot.org/citations/16293632), PubMed: [18826956](http://www.uniprot.org/citations/18826956), PubMed: [18695040](http://www.uniprot.org/citations/18695040), PubMed: [25256292](http://www.uniprot.org/citations/25256292), PubMed: [20037586](http://www.uniprot.org/citations/20037586), PubMed: [21964574](http://www.uniprot.org/citations/21964574)). Increase of intracellular  $\text{Ca}^{2+}$  potentiates currents. Channel activity seems to be regulated by a calmodulin-dependent mechanism with a negative feedback mechanism (PubMed: [12724311](http://www.uniprot.org/citations/12724311), PubMed: [18826956](http://www.uniprot.org/citations/18826956)). Promotes cell-cell junction formation in skin keratinocytes and plays an important role in the formation and/or maintenance of functional intercellular barriers (By similarity). Acts as a regulator of intracellular  $\text{Ca}^{2+}$  in synoviocytes (PubMed: [19759329](http://www.uniprot.org/citations/19759329)). Plays an obligatory role as a molecular component in the nonselective cation channel activation induced by 4- $\alpha$ -phorbol 12,13-didecanoate and hypotonic stimulation in synoviocytes and also regulates production of IL-8 (PubMed: [19759329](http://www.uniprot.org/citations/19759329)). Together with PKD2, forms mechano- and thermosensitive channels in cilium (PubMed: [18695040](http://www.uniprot.org/citations/18695040)). Negatively regulates expression of PPARGC1A, UCP1, oxidative metabolism and respiration in adipocytes (By similarity). Regulates expression of chemokines and cytokines related to pro-inflammatory pathway in adipocytes (By similarity). Together with AQP5, controls regulatory volume decrease in salivary epithelial cells (By similarity). Required for normal development and maintenance of bone and cartilage (PubMed: [26249260](http://www.uniprot.org/citations/26249260)). In its inactive state, may sequester DDX3X at the plasma membrane. When activated, the interaction between both proteins is affected and DDX3X relocalizes to the nucleus (PubMed: [29899501](http://www.uniprot.org/citations/29899501)). In neurons of the central nervous system, could play a role in triggering voluntary water intake in response to increased sodium concentration in body fluid (By similarity).

### Cellular Location

Cell membrane. Apical cell membrane; Multi-pass membrane protein. Cell junction, adherens junction {ECO:0000250|UniProtKB:Q9EPK8}. Cell projection, cilium. Note=Assembly of the putative homotetramer occurs primarily in the endoplasmic reticulum (PubMed:16293632, PubMed:20037587, PubMed:20037588). Localization to the cell membrane is inhibited by WNK kinases (WNK1, WNK2, WNK3 or WNK4) in a kinase-independent mechanism (PubMed:16403833) [Isoform 5]: Cell membrane [Isoform 4]: Endoplasmic reticulum

### Tissue Location

Found in the synoviocytes from patients with (RA) and without (CTR) rheumatoid arthritis (at protein level)

## TRPV4 Antibody (N-term) Blocking Peptide - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Blocking Peptides](#)

## TRPV4 Antibody (N-term) Blocking Peptide - Images

## TRPV4 Antibody (N-term) Blocking Peptide - Background

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  $\text{Ca}^{2+}$ -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.

#### **TRPV4 Antibody (N-term) Blocking Peptide - References**

Shukla, A.K., et al. J. Biol. Chem. 285(39):30115-30125(2010)Cantero-Recasens, G., et al. J. Biol. Chem. 285(36):27532-27535(2010)Loukin, S., et al. J. Biol. Chem. 285(35):27176-27181(2010)Nishimura, G., et al. Am. J. Med. Genet. A 152A (6), 1443-1449 (2010):Zimon, M., et al. Brain 133 (PT 6), 1798-1809 (2010) :