# SANTA CRUZ BIOTECHNOLOGY, INC.

# TRPM5 (N-20): sc-27366



## BACKGROUND

Transient receptor potential ion channels (TRPCs) are a superfamily of six transmembrane segment-spanning, gated cation channels. TRPC subtypes mediate store-operated  $Ca^{2+}$  entry, a process involving  $Ca^{2+}$  influx and replenishment of  $Ca^{2+}$  stores formerly emptied through the action of inositol 1,4,5trisphospate production and other  $Ca^{2+}$  mobilizing agents. TRP ion channels influence calcium-depletion induced calcium influx processes in response to chemo-, mechano- and osmoregulatory events. TRPM5 forms a cation channel that is directly activated by micromolar concentrations of intracellular  $Ca^{2+}$ . Sustained exposure to  $Ca^{2+}$  desensitizes TRPM5 channels, but phosphatidylinositol-4,5-bisphosphate reverses desensitization, partially restoring channel activity. TRPM5 channels are nonselective among monovalent cations and not detectably permeable to divalent cations.

## REFERENCES

- 1. Liu, D. and Liman, E.R. 2003. Intracellular Ca<sup>2+</sup> and the phospholipid PIP2 regulate the taste transduction ion channel TRPM5. PNAS 100: 15160-15165.
- Prawitt, D., Monteilh-Zoller, M.K., Brixel, L., Spangenberg, C., Zabel, B., Fleig, A. and Penner, R. 2003. TRPM5 is a transient Ca<sup>2+</sup>-activated cation channel responding to rapid changes in [Ca<sup>2+</sup>]i. Proc. Natl. Acad. Sci. USA 100: 15166-15171.
- Hofmann, T., Chubanov, V., Gudermann, T. and Montell, C. 2003. TRPM5 is a voltage-modulated and Ca<sup>2+</sup>-activated monovalent selective cation channel. Curr. Biol. 13: 1153-1158.
- Pérez, C.A., Margolskee, R.F., Kinnamon, S.C. and Ogura, T. 2003. Making sense with TRP channels: store-operated calcium entry and the ion channel Trpm5 in taste receptor cells. Cell Calcium 33: 541-549.
- Perraud, A.L., Knowles, H.M. and Schmitz, C. 2004. Novel aspects of signaling and ion-homeostasis regulation in immunocytes. The TRPM ion channels and their potential role in modulating the immune response. Mol. Immunol. 41: 657-673.

## CHROMOSOMAL LOCATION

Genetic locus: TRPM5 (human) mapping to 11p15.5; Trpm5 (mouse) mapping to 7 F5.

#### SOURCE

TRPM5 (N-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the N-terminus of TRPM5 of human origin.

## PRODUCT

Each vial contains 200  $\mu g$  IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-27366 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

# **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

## **APPLICATIONS**

TRPM5 (N-20) is recommended for detection of TRPM5 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

TRPM5 (N-20) is also recommended for detection of TRPM5 in additional species, including equine, canine and bovine.

Suitable for use as control antibody for TRPM5 siRNA (h): sc-44169, TRPM5 siRNA (m): sc-154695, TRPM5 shRNA Plasmid (h): sc-44169-SH, TRPM5 shRNA Plasmid (m): sc-154695-SH, TRPM5 shRNA (h) Lentiviral Particles: sc-44169-V and TRPM5 shRNA (m) Lentiviral Particles: sc-154695-V.

Molecular Weight of TRPM5: 131 kDa.

## **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluo-rescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

## SELECT PRODUCT CITATIONS

 Teruyama, R., Sakuraba, M., Kurotaki, H. and Armstrong, W.E. 2011. Transient receptor potential channel m4 and m5 in magnocellular cells in rat supraoptic and paraventricular nuclei. J. Neuroendocrinol. 23: 1204-1213.

## **STORAGE**

Store at 4° C, \*\*D0 NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.