# T-type Ca<sup>++</sup> CP $\alpha$ 1H (N-18): sc-16261



The Power to Question

## **BACKGROUND**

Voltage-dependent Ca<sup>2+</sup> channels mediate Ca<sup>2+</sup> entry into excitable cells in response to membrane depolarization, and they are involved in a variety of Ca<sup>2+</sup>-dependent processes, including muscle contraction, hormone or neurotransmitter release and gene expression. Calcium channels are highly diverse, multimeric complexes composed of an  $\alpha 1$  subunit, an intracellular  $\beta$  subunit, a disulfide linked  $\alpha 2/\delta$  subunit and a transmembrane  $\gamma$  subunit. Ca<sup>2+</sup> currents are characterized on the basis of their biophysical and pharmacologic properties and include L-, N-, T-, P-, Q-, and R- types. T-type Ca<sup>++</sup> currents are activated and inactivated more rapidly and at more negative membrane potentials than other Ca<sup>2+</sup> current types. T-type Ca<sup>2+</sup> channels enhance odor sensitivity by lowering the threshold of spike generation in olfactory receptor cells (ORCs).

## **REFERENCES**

- 1. Perez-Reyes, E., et al. 1995. Molecular biology of calcium channels. Kidney Int. 48: 1111-1124.
- 2. Randall, A.D. 1998. The molecular basis of voltage-gated Ca<sup>2+</sup> channel diversity: is it time for T. J. Membr. Biol. 161: 207-213.

#### **CHROMOSOMAL LOCATION**

Genetic locus: CACNA1H (human) mapping to 16p13.3; Cacna1h (mouse) mapping to 17 A3.3.

## **SOURCE**

T-type Ca<sup>++</sup> CP  $\alpha$ 1H (N-18) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the N-terminus of T-type Ca<sup>++</sup> CP  $\alpha$ 1H of human origin.

## **PRODUCT**

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

Blocking peptide available for competition studies, sc-16261 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

# **APPLICATIONS**

T-type Ca<sup>++</sup> CP  $\alpha$ 1H (N-18) is recommended for detection of T-type calcium channel  $\alpha$ 1H 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).

Suitable for use as control antibody for T-type Ca++ CP  $\alpha$ 1H siRNA (h): sc-42706, T-type Ca++ CP  $\alpha$ 1H siRNA (m): sc-42707, T-type Ca++ CP  $\alpha$ 1H shRNA Plasmid (h): sc-42706-SH, T-type Ca++ CP  $\alpha$ 1H shRNA Plasmid (m): sc-42707-SH, T-type Ca++ CP  $\alpha$ 1H shRNA (h) Lentiviral Particles: sc-42706-V and T-type Ca++ CP  $\alpha$ 1H shRNA (m) Lentiviral Particles: sc-42707-V.

Molecular Weight (predicted) of T-type Ca++ CP α1H: 259 kDa.

Molecular Weight (observed) of T-type Ca<sup>++</sup> CP  $\alpha$ 1H: 247-257 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) Immunofluorescence: 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**

- Trevino, C.L., et al. 2004. Expression and differential cell distribution of low-threshold Ca<sup>2+</sup> channels in mammalian male germ cells and sperm. FEBS Lett. 563: 87-92.
- De Proost, I., et al. 2007. Pulmonary expression of voltage-gated calcium channels: special reference to sensory airway receptors. Histochem. Cell Biol. 128: 301-316.
- Escoffier, J., et al. 2007. Expression, localization and functions in acrosome reaction and sperm motility of CaV3.1 and CaV3.2 channels in sperm cells: an evaluation from CaV3.1 and CaV3.2 deficient mice. J. Cell. Physiol. 212: 753-763.
- 4. Morikawa, K., et al. 2010. Identification, isolation and characterization of HCN4-positive pacemaking cells derived from murine embryonic stem cells during cardiac differentiation. Pacing Clin. Electrophysiol. 33: 290-303.
- 5. Zhang, Y., et al. 2014. Peripheral pain is enhanced by Insulin-like growth factor 1 through a G protein-mediated stimulation of T-type calcium channels. Sci. Signal. 7: ra94.

## **STORAGE**

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

## **RESEARCH USE**

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

## **PROTOCOLS**

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



Try T-type Ca++ CP  $\alpha$ 1H (G-10): sc-377510, our highly recommended monoclonal alternative to T-type Ca++ CP  $\alpha$ 1H (N-18).

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