

Na⁺ CP type IX α (H-17): sc-130096

BACKGROUND

Voltage-gated sodium channels are selective ion channels that regulate the permeability of sodium ions in excitable cells. During the propagation of an action potential, sodium channels allow an influx of sodium ions, which rapidly depolarizes the cell. Na⁺ CP type IX α , also known as SCN9A (sodium channel protein type 9 subunit α), NENA, PN1, Nav1.7 or ETHA, is a 1,988 amino acid multi-pass membrane protein that belongs to the voltage-gated sodium channel family. Expressed in dorsal root ganglion, smooth muscle cells and in the central and peripheral nervous system, Na⁺ CP type IX α functions to mediate the voltage-dependent sodium ion permeability of membranes, specifically forming a sodium-selective ion channel through which sodium may pass. Via its ability to control the flow of sodium in and out of excitable membranes, Na⁺ CP type IX α plays an important role in the inflammatory pain response. Defects in the gene encoding Na⁺ CP type IX α are the cause of primary erythralgia, autosomal recessive congenital indifference to pain and paroxysmal extreme pain disorder (PEPD), all of which are genetic pain disorders.

REFERENCES

1. Klugbauer, N., et al. 1995. Structure and functional expression of a new member of the tetrodotoxin-sensitive voltage-activated sodium channel family from human neuroendocrine cells. *EMBO J.* 14: 1084-1090.
2. Sangameswaran, L., et al. 1997. A novel tetrodotoxin-sensitive, voltage-gated sodium channel expressed in rat and human dorsal root ganglia. *J. Biol. Chem.* 272: 14805-14809.
3. Raymond, C.K., et al. 2004. Expression of alternatively spliced sodium channel α subunit genes. Unique splicing patterns are observed in dorsal root ganglia. *J. Biol. Chem.* 279: 46234-46241.
4. Yang, Y., et al. 2004. Mutations in SCN9A, encoding a sodium channel α subunit, in patients with primary erythralgia. *J. Med. Genet.* 41: 171-174.
5. Ahmad, S., et al. 2007. A stop codon mutation in SCN9A causes lack of pain sensation. *Hum. Mol. Genet.* 16: 2114-2121.
6. Drenth, J.P. and Waxman, S.G. 2007. Mutations in sodium-channel gene SCN9A cause a spectrum of human genetic pain disorders. *J. Clin. Invest.* 117: 3603-3609.
7. Sheets, P.L., et al. 2007. A Nav1.7 channel mutation associated with hereditary erythralgia contributes to neuronal hyperexcitability and displays reduced lidocaine sensitivity. *J. Physiol.* 581: 1019-1031.
8. Drenth, J.P., et al. 2008. Primary erythralgia as a sodium channelopathy: screening for SCN9A mutations: exclusion of a causal role of SCN10A and SCN11A. *Arch Dermatol.* 144: 320-324.
9. Tadc, A., Baskaya, O., Victor, A., Lieb, K., Höppner, W. and Dahmen, N. 2008. Association analysis of SCN9A gene variants with borderline personality disorder. *J Psychiatr Res.* PMID: 18439623

CHROMOSOMAL LOCATION

Genetic locus: SCN9A (human) mapping to 2q24.3; Scn9a (mouse) mapping to 2 C1.3.

SOURCE

Na⁺ CP type IX α (H-17) is a purified rabbit polyclonal antibody raised against Na⁺ CP type IX α of human origin.

PRODUCT

Each vial contains 50 μ g IgG in 0.5 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

Na⁺ CP type IX α (H-17) is recommended for detection of Na⁺ CP type IX α 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), immunohistochemistry (including paraffin-embedded sections) (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 Na⁺ CP type IX α siRNA (h): sc-94458 and Na⁺ CP type IX α siRNA (m): sc-149784; and as shRNA Plasmid control antibody for Na⁺ CP type IX α shRNA Plasmid (h): sc-94458-SH and Na⁺ CP type IX α shRNA Plasmid (m): sc-149784-SH.

Molecular Weight of Na⁺ CP type IX α : 226 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (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 goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941. 3) Immunohistochemistry: use ImmunoCruz™: sc-2051 or ABC: sc-2018 rabbit IgG Staining Systems.

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 **Na⁺ CP type IX α (5A11): sc-293298** our highly recommended monoclonal alternative to Na⁺ CP type IX α (H-17).