## SANTA CRUZ BIOTECHNOLOGY, INC.

# L-type Ca<sup>++</sup> CP α1S (IIC12D4): sc-21781



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 neuro-transmitter 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. L-type Ca<sup>2+</sup> currents initiate muscle contraction, endocrine secretion, and gene transcription, and can be regulated through second-messenger activated protein phosphorylation pathways. L-type calcium channels may form macromolecular signaling complexes with G protein-coupled receptors, thereby enhancing the selectivity of regulating specific targets. Calcium channels containing the  $\alpha$ -1S subunit play an important role in excitation-contraction coupling in skeletal muscle.

#### REFERENCES

- 1. Gregg, R.G., et al. 1993. Assignment of the human gene for the  $\alpha$  1 subunit of the skeletal muscle DHP-sensitive Ca<sup>2+</sup> channel (CACNL1A3) to chromosome 1q31-q32. Genomics 15: 107-112.
- 2. Perez-Reyes, E. and Schneider, T. 1995. Molecular biology of calcium channels. Kidney Int. 48: 1111-1124.

#### CHROMOSOMAL LOCATION

Genetic locus: CACNA1S (human) mapping to 1q32.1; Cacna1s (mouse) mapping to 1 E4.

## SOURCE

L-type Ca<sup>++</sup> CP  $\alpha$ 1S (IIC12D4) is a mouse monoclonal antibody raised against skeletal muscle triads of rabbit origin, recognizes skeletal DHPR  $\alpha$ 1S subunit.

#### PRODUCT

Each vial contains 200  $\mu g$  lgG\_1 kappa light chain in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

### **APPLICATIONS**

L-type Ca<sup>++</sup> CP  $\alpha$ 1S (IIC12D4) is recommended for detection of 170 kDa, L-type calcium channel  $\alpha$ 1S of mouse, rat, human and rabbit origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)] and immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

Suitable for use as control antibody for L-type Ca<sup>++</sup> CP  $\alpha$ 1S siRNA (h): sc-35772, L-type Ca<sup>++</sup> CP  $\alpha$ 1S siRNA (m): sc-35773, L-type Ca<sup>++</sup> CP  $\alpha$ 1S shRNA Plasmid (h): sc-35772-SH, L-type Ca<sup>++</sup> CP  $\alpha$ 1S shRNA Plasmid (m): sc-35773-SH, L-type Ca<sup>++</sup> CP  $\alpha$ 1S shRNA (h) Lentiviral Particles: sc-35772-V and L-type Ca<sup>++</sup> CP  $\alpha$ 1S shRNA (m) Lentiviral Particles: sc-35773-V.

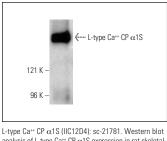
Molecular Weight of L-type Ca++ CP a1S: 170 kDa.

Positive Controls: rat skeletal muscle extract: sc-364810, mouse skeletal muscle extract: sc-364250 or Sol8 cell lysate: sc-2249.

#### **RECOMMENDED SUPPORT REAGENTS**

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG $\kappa$  BP-HRP: sc-516102 or m-IgG $\kappa$  BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker<sup>TM</sup> Molecular Weight Standards: sc-2035, UltraCruz<sup>®</sup> Blocking Reagent: sc-516214 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use m-IgG $\kappa$  BP-FITC: sc-516140 or m-IgG $\kappa$  BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz<sup>®</sup> Mounting Medium: sc-24941 or UltraCruz<sup>®</sup> Hard-set Mounting Medium: sc-359850.

#### DATA



L-type Ca<sup>++</sup> CP αTS (IICT2D4): SC-21781. Western blot analysis of L-type Ca<sup>++</sup> CP αTS expression in rat skeletal muscle tissue extract.

#### SELECT PRODUCT CITATIONS

- 1. Ren, X., et al. 2009. Cellular effect evaluation of micropollutants using transporter functions of renal proximal tubule cells. Chemosphere 77: 968-974.
- Wang, Z., et al. 2020. A temporal examination of cytoplasmic Ca<sup>2+</sup> levels, sarcoplasmic reticulum Ca<sup>2+</sup> levels, and Ca<sup>2+</sup>-handling-related proteins in different skeletal muscles of hibernating daurian ground squirrels. Front. Physiol. 11: 562080.

#### **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 for detailed protocols and support products.