SANTA CRUZ BIOTECHNOLOGY, INC.

PP2Cα (7F12): sc-517264



BACKGROUND

The phosphorylation and dephosphorylation of proteins on serine and threonine residues is an essential means of regulating a broad range of cellular functions in eukaryotes, including division, homeostasis and apoptosis. A group of proteins that are intimately involved in this process are the serine/ threonine protein phosphatases. Protein phosphatase $2C\alpha$ (PP2C α) has broad specificity. It dephosphorylates and negatively regulates the activities of MAP kinases and MAP kinase-kinases while also inhibiting the activation of p38 and JNK kinase cascades, induced by environmental stresses. PP2C α also induces the expression of endogenous p53 and the p53-responsive gene p21, leading to cell cycle arrest and apoptosis. The PP2C α protein, which contains an active site containing a dinuclear metal ion center, shows highest expression in epithelial cells, as well as in the digestive tract, lung, kidney, breast, prostate, endocrine glands and brain.

REFERENCES

- Ueki, K., et al. 1992. Structure and expression of two isoforms of the murine calmodulin-dependent protein phosphatase regulatory subunit (calcineurin B). Biochem. Biophys. Res. Commun. 187: 537-543.
- 2. Cohen, P.T. 1993. Important roles for novel protein phosphatases dephosphorylating serine and threonine residues. Biochem. Soc. Trans. 21: 884-888.
- Yokoyama, N., et al. 1996. Purification and characterization of protein phosphatase 2C in rat parotid acinar cells: two forms of Mg²⁺-activated histone phosphatase and phosphorylation by cAMP-dependent protein kinase. Arch. Biochem. Biophys. 331: 1-8.
- 4. Takekawa, M., et al. 1998. Protein phosphatase $2C\alpha$ inhibits the human stress-responsive p38 and JNK MAPK pathways. EMBO J. 17: 4744-4452.
- 5. Lifschitz-Mercer, B., et al. 2001. Protein phosphatase $2C\alpha$ expression in normal human tissues: an immunohistochemical study. Histochem. Cell Biol. 116: 31-39.

CHROMOSOMAL LOCATION

Genetic locus: PPM1A (human) mapping to 14q23.1; Ppm1a (mouse) mapping to 12 C3.

SOURCE

PP2C α (7F12) is a mouse monoclonal antibody raised against a recombinant protein corresponding to amino acids 202-382 of PP2C α of human origin.

PRODUCT

Each vial contains 200 μg lgG_1 kappa light chain in 1.0 ml of PBS with <0.1% sodium azide and 0.1% gelatin.

PP2Cα (7F12) is available conjugated to agarose (sc-517264 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-517264 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-517264 PE), fluorescein (sc-517264 FITC), Alexa Fluor[®] 488 (sc-517264 AF488), Alexa Fluor[®] 546 (sc-517264 AF546), Alexa Fluor[®] 594 (sc-517264 AF594) or Alexa Fluor[®] 647 (sc-517264 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-517264 AF680) or Alexa Fluor[®] 790 (sc-517264 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

APPLICATIONS

PP2C α (7F12) is recommended for detection of PP2C α of mouse, rat and human 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)], flow cytometry (1 µg per 1 x 10⁶ cells) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for PP2C α siRNA (h): sc-45214, PP2C α siRNA (m): sc-45215, PP2C α shRNA Plasmid (h): sc-45214-SH, PP2C α shRNA Plasmid (m): sc-45215-SH, PP2C α shRNA (h) Lentiviral Particles: sc-45214-V and PP2C α shRNA (m) Lentiviral Particles: sc-45215-V.

Molecular Weight of PP2Ca: 46 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200, A-431 whole cell lysate: sc-2201 or CCRF-CEM cell lysate: sc-2225.

RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgGκ BP-HRP: sc-516102 or m-IgGκ BP-HRP (Cruz Marker): sc-516102-CM (dilution range: 1:1000-1:10000), Cruz Marker[™] Molecular Weight Standards: sc-2035, UltraCruz[®] 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).

DATA





 $PP2C\alpha$ (7F12): sc-517264. Western blot analysis of $PP2C\alpha$ expression in CCRF-CEM (**A**), Ramos (**B**) and PC-12 (**C**) whole cell lysates and mouse brain tissue extract (**D**).

 $PP2C\alpha$ (7F12): sc-517264. Western blot analysis of $PP2C\alpha$ expression in HeLa (A), A-431 (B), SH-SY5Y (C) and CCRF-CEM (D) whole cell lysates.

SELECT PRODUCT CITATIONS

- Flores, K., et al. 2020. Lower body weight in rats under hypobaric hypoxia exposure would lead to reduced right ventricular hypertrophy and increased AMPK activation. Front. Physiol. 11: 342.
- Yadav, Y., et al. 2022. PP2Cα positively regulates neuronal insulin signalling and aggravates neuronal insulin resistance. FEBS J. 289: 7561-7581.

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.

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