

## PP2A-A $\beta$ (C-20): sc-6113

### BACKGROUND

In eukaryotes, the phosphorylation and dephosphorylation of proteins on serine and threonine residues is an essential means of regulating a broad range of cellular functions, including division, homeostasis and apoptosis. A group of proteins that are intimately involved in this process are the protein phosphatases. In general, the protein phosphatase (PP) holoenzyme is a trimeric complex composed of a regulatory subunit, a variable subunit and a catalytic subunit. Four major families of protein phosphatase catalytic subunits have been identified, designated PP1, PP2A, PP2B (calcineurin) and PP2C. The PP2A family comprises subfamily members PP2A $\alpha$  and PP2A $\beta$ . An additional protein phosphatase catalytic subunit, PPX (also known as PP4) is a putative member of a novel PP family. The PP2A catalytic subunit associates with a variety of regulatory subunits. Regulatory subunits include PP2A-A $\alpha$  and -A $\beta$ , PP2A-B $\alpha$  and -B $\beta$ , PP2A-C $\alpha$  and -C $\beta$ , and PP2A-B56 $\alpha$  and -B56 $\beta$ .

### CHROMOSOMAL LOCATION

Genetic locus: PPP2R1B (human) mapping to 11q23.1.

### SOURCE

PP2A-A $\beta$  (C-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of PP2A-A $\beta$  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-6113 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

### APPLICATIONS

PP2A-A $\beta$  (C-20) is recommended for detection of PP2A-A $\beta$  of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], 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); may cross react with PP2A-A $\alpha$ .

PP2A-A $\beta$  (C-20) is also recommended for detection of PP2A-A $\beta$  in additional species, including canine and porcine.

Suitable for use as control antibody for PP2A-A $\beta$  siRNA (h): sc-39179, PP2A-A $\beta$  shRNA Plasmid (h): sc-39179-SH and PP2A-A $\beta$  shRNA (h) Lentiviral Particles: sc-39179-V.

Molecular Weight of PP2A-A $\alpha$ : 55 kDa.

Molecular Weight of PP2A-A $\beta$ : 65 kDa.

Positive Controls: H4 cell lysate: sc-2408, SW480 cell lysate: sc-2219 or Jurkat whole cell lysate: sc-2204.

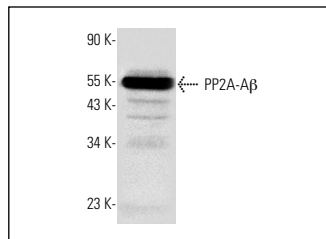
### RESEARCH USE

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

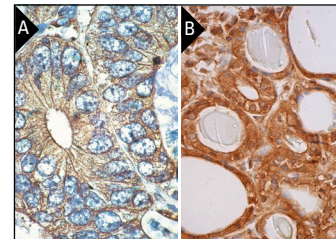
### STORAGE

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

### DATA



PP2A-A $\beta$  (C-20): sc-6113. Western blot analysis of PP2A-A $\beta$  expression in Jurkat whole cell lysate.



PP2A-A $\beta$  (C-20): sc-6113. Immunoperoxidase staining of formalin-fixed, paraffin-embedded human colon carcinoma tissue showing cytoplasmic localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human thyroid gland tissue showing cytoplasmic, membrane and nuclear staining of glandular cells (B).

### SELECT PRODUCT CITATIONS

- Wang, S.S., et al. 1998. Alterations of the PPP2R1B gene in human lung and colon cancer. *Science* 282: 284-287.
- Goodarzi, A.A., et al. 2004. Autophosphorylation of ataxia-telangiectasia mutated is regulated by protein phosphatase 2A. *EMBO J.* 23: 4451-4461.
- Jayadeva, G., et al. 2010. B55 $\alpha$  PP2A holoenzymes modulate the phosphorylation status of the retinoblastoma-related protein p107 and its activation. *J. Biol. Chem.* 285: 29863-29873.
- Clarke, C.J., et al. 2011. Neutral sphingomyelinase-2 mediates growth arrest by retinoic acid through modulation of ribosomal S6 kinase. *J. Biol. Chem.* 286: 21565-21576.
- Kurimchak, A., et al. 2013. Activation of p107 by FGF, which is essential for chondrocyte cell cycle exit, is mediated by the PP2A/B55 $\alpha$  holoenzyme. *Mol. Cell. Biol.* 33: 3330-3342.

### PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) or our catalog for detailed protocols and support products.



Try **PP2A-A $\alpha$ / $\beta$  (4G7): sc-13600** or **PP2A-A $\alpha$ / $\beta$  (A-5): sc-74580**, our highly recommended monoclonal alternatives to PP2A-A $\beta$  (C-20).