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PP2C β (m): 293T Lysate: sc-122721

BACKGROUND

Eukaryotic protein phosphorylation and dephosphorylation on serine and threonine residues regulates numerous cell functions, including division, homeostasis and apoptosis. A group of proteins that play a major role in this process are the serine/threonine protein phosphatases. Protein phosphatase (PP) holoenzyme is a trimeric complex that contains a regulatory subunit, a variable subunit and a catalytic subunit. Families of PP catalytic subunits include PP1 (PP1 α , β and γ), PP2A (α and β), PP2B (calcineurin, PP2B α , β and γ), PP2C (α , β , γ , η and Wip1), PP4 (PPX) and PP5 (PPT). PP2C family members are negative regulators of cell stress response pathways. The PP2C β enzyme has broad specificity and is highly expressed in the heart and skeletal muscle. It may be involved in cell cycle control as it dephosphorylates the cyclin-dependent kinases (CDKs), CDK2 and CDK6, *in vitro*. Overexpression of PP2C β can cause cell growth arrest or cell death.

REFERENCES

1. Marley, A.E., Kline, A., Crabtree, G., Sullivan, J.E. and Beri, R.K. 1998. The cloning expression and tissue distribution of human PP2C β . FEBS Lett. 431: 121-124.
2. Cheng, A., Kaldis, P. and Solomon, M.J. 2000. Dephosphorylation of human cyclin-dependent kinases by protein phosphatase type 2C α and β 2 isoforms. J. Biol. Chem. 275: 34744-34749.
3. Online Mendelian Inheritance in Man, OMIM™. 2002. Johns Hopkins University, Baltimore, MD. MIM Number: 603770. World Wide Web URL: <http://www.ncbi.nlm.nih.gov/omim/>
4. Brautigan, D.L., Brown, M., Grindrod, S., Chinigo, G., Kruszewski, A., Lukasik, S.M., Bushweller, J.H., Horal, M., Keller, S., Tamura, S., Heimark, D.B., Price, J., Larner, A.N. and Larner, J. 2005. Allosteric activation of protein phosphatase 2C by D-chiro-inositol-galactosamine, a putative mediator mimetic of Insulin action. Biochemistry 44: 11067-11073.
5. Hufnagel, B., Dworak, M., Soufi, M., Mester, Z., Zhu, Y., Schaefer, J.R., Klumpp, S. and Kriegstein, J. 2005. Unsaturated fatty acids isolated from human lipoproteins activate protein phosphatase type 2C β and induce apoptosis in endothelial cells. Atherosclerosis 180: 245-254.

CHROMOSOMAL LOCATION

Genetic locus: Ppm1b (mouse) mapping to 17 E4.

PRODUCT

PP2C β (m): 293T Lysate represents a lysate of mouse PP2C β transfected 293T cells and is provided as 100 μ g protein in 200 μ l SDS-PAGE buffer.

STORAGE

Store at -20° C. Repeated freezing and thawing should be minimized. Sample vial should be boiled once prior to use. Non-hazardous. No MSDS required.

PROTOCOLS

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

APPLICATIONS

PP2C β (m): 293T Lysate is suitable as a Western Blotting positive control for mouse reactive PP2C β antibodies. Recommended use: 10-20 μ l per lane.

Control 293T Lysate: sc-117752 is available as a Western Blotting negative control lysate derived from non-transfected 293T cells.

RESEARCH USE

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