

PTP μ (C-20): sc-1115

BACKGROUND

Protein tyrosine phosphatases, or PTPs, are type I transmembrane proteins, membrane associated proteins or proteins localized in nuclei. Examples of transmembrane PTPs are LAR, PTP α , PTP β , PTP γ , PTP δ , PTP ϵ , PTP ζ , PTP κ and PTP μ . Transmembrane PTPs play diverse roles during development and in adult tissues. Immunodepletion studies have suggested LAR to be a regulator of Insulin receptor phosphorylation. PTP α activity is increased twofold in response to phorbol ester stimulation, resulting in serine phosphorylation either directly or indirectly by members of the PKC family. Overexpression of v-H-ras and Neu, but not Myc or Int2, in mammary tumors has been shown to induce PTP ϵ expression. An alternative splicing event leads to a nervous tissue-specific chondroitin sulfate proteoglycan called phosphacan, which represents the amino terminal portion of PTP ζ . PTP κ and PTP μ share a conserved amino terminal 160 amino acid MAM domain which facilitates homophilic binding. PTP μ localizes to points of cell contact and may be involved in regulating the assembly and disassembly of cadherin/catenin complexes *in vivo*.

REFERENCES

- Ahmad, F., et al. 1995. Increased abundance of the receptor-type protein-tyrosine phosphatase LAR accounts for the elevated Insulin receptor dephosphorylating activity in adipose tissue of obese human subjects. *J. Clin. Invest.* 95: 2806-2812.
- den Hertog, J., et al. 1995. Stimulation of receptor protein-tyrosine phosphatase α activity and phosphorylation by phorbol ester. *Cell Growth Differ.* 6: 303-307.
- Brady-Kalnay, S.M., et al. 1995. Receptor protein tyrosine phosphatase PTP μ associates with cadherins and catenins *in vivo*. *J. Cell. Biol.* 130: 977-986.
- Zondag, G.C., et al. 1995. Homophilic interactions mediated by receptor tyrosine phosphatases μ and κ . A critical role for the novel extracellular MAM domain. *J. Biol. Chem.* 270: 14247-14250.

CHROMOSOMAL LOCATION

Genetic locus: PTPRM (human) mapping to 18p11.2; Ptpm (mouse) mapping to 17 E1.1.

SOURCE

PTP μ (C-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of PTP μ of human origin.

PRODUCT

Each vial contains 200 μ g IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-1115 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

STORAGE

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

APPLICATIONS

PTP μ (C-20) is recommended for detection of PTP μ 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) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

PTP μ (C-20) is also recommended for detection of PTP μ in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for PTP μ siRNA (h): sc-44055, PTP μ siRNA (m): sc-45947, PTP μ shRNA Plasmid (h): sc-44055-SH, PTP μ shRNA Plasmid (m): sc-45947-SH, PTP μ shRNA (h) Lentiviral Particles: sc-44055-V and PTP μ shRNA (m) Lentiviral Particles: sc-45947-V.

Molecular Weight of PTP μ precursor: 200 kDa.

Molecular Weight of PTP μ subunits: 100 kDa

Positive Control: A549 whole cell lysate: sc-2413, T98G cell lysate: sc-2294 or SK-N-SH cell lysate: sc-2410

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (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 donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

SELECT PRODUCT CITATIONS

- Hiscox, S., et al. 1998. Association of PTP μ with catenins in cancer cells: a possible role for E-cadherin. *Int. J. Oncol.* 13: 1077-1080.
- Hiscox, S., et al. 1999. Association of the HGF/SF receptor, c-Met, with the cell-surface adhesion molecule, E-Cadherin, and Catenins in human tumor cells. *Biochem. Biophys. Res. Commun.* 261: 406-411.
- Basso, K., et al. 2004. Gene expression profiling of hairy cell leukemia reveals a phenotype related to memory B cells with altered expression of chemokine and adhesion receptors. *J. Exp. Med.* 199: 59-68.

RESEARCH USE

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


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Try PTP μ (2C10): sc-56957 or PTP μ (BK2): sc-33651, our highly recommended monoclonal alternatives to PTP μ (C-20).