

PTP μ (BK2): sc-33651

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*.

CHROMOSOMAL LOCATION

Genetic locus: PTPRM (human) mapping to 18p11.23; Ptpm (mouse) mapping to 17 E1.1.

SOURCE

PTP μ (BK2) is a mouse monoclonal antibody raised against a synthetic peptide from the extracellular segment of PTP μ .

PRODUCT

Each vial contains 200 μ g IgG_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

PTP μ (BK2) is recommended for detection of full length PTP μ (200 kDa) and extracellular, E-subunit (100 kDa) of mouse, rat, human and *Xenopus laevis* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)] and immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

PTP μ (BK2) is also recommended for detection of full length PTP μ (200 kDa) and extracellular, E-subunit (100 kDa) in additional species, including bovine.

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.

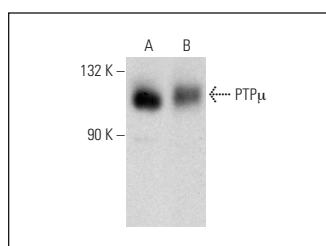
Molecular Weight of PTP μ subunits: 100 kDa.

Positive Controls: A549 cell lysate: sc-2413, HUVEC-C whole cell lysate: sc-364180 or T98G cell lysate: sc-2294.

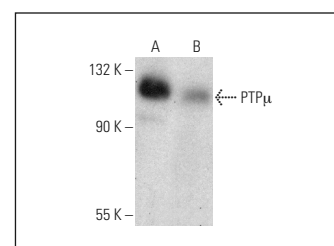
RECOMMENDED SUPPORT REAGENTS

To ensure optimal results, the following support reagents are recommended: 1) Western Blotting: use m-IgG κ BP-HRP: sc-516102 or m-IgG κ 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). 3) Immunofluorescence: use m-IgG κ BP-FITC: sc-516140 or m-IgG κ BP-PE: sc-516141 (dilution range: 1:50-1:200) with UltraCruz® Mounting Medium: sc-24941 or UltraCruz® Hard-set Mounting Medium: sc-359850.

DATA



PTP μ (BK2): sc-33651. Western blot analysis of PTP μ expression in T98G (A) and HUVEC-C (B) whole cell lysates.



PTP μ (BK2): sc-33651. Western blot analysis of PTP μ expression in A549 (A) and RAT2 (B) whole cell lysates.

SELECT PRODUCT CITATIONS

- Johnson, J.L., et al. 2009. Tetraspanin CD151 regulates RhoA activation and the dynamic stability of carcinoma cell-cell contacts. *J. Cell Sci.* 122: 2263-2273.
- Takahashi, K., et al. 2017. Expression of receptor-type protein tyrosine phosphatase in developing and adult renal vasculature. *PLoS ONE* 12: e0177192.

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.