p-PI 3-kinase p85α (Tyr 467): sc-293115



The Power to Question

BACKGROUND

Phosphatidylinositol 3-kinase (PI 3-kinase) phosphorylates the 3' OH position of the inositol ring of inositol lipids and is composed of p85 and p110 subunits. PI 3-kinase p85 lacks PI 3-kinase activity and acts as an adapter, coupling p110 to activated protein tyrosine kinase. Two forms of p85 have been described (p85 α and p85 β), each possessing one SH3 and two SH2 domains. PI 3-kinase p85 α , also known as GRB1, phosphatidylinositol 3-kinase regulatory 1 or p85, is a 724 amino acid protein that exists as 4 alternatively spliced isoforms. Involved in Insulin metabolism, defects in the PI 3-kinase p85 α gene have been linked to Insulin resistance. PI 3-kinase p85 α is polyubiquitinated in T-cells by CbI-b, and has multiple phosphorylated amino acid residues, including a phosphorylated tyrosine residue at position 467.

REFERENCES

- Skolnik, E.Y., et al. 1991. Cloning of PI3 kinase-associated p85 utilizing a novel method for expression/cloning of target proteins for receptor tyrosine kinases. Cell 65: 83-90.
- 2. Van Horn, D.J., et al. 1994. Direct activation of the phosphatidylinositol 3'-kinase by the Insulin receptor. J. Biol. Chem. 269: 29-32.
- Craparo, A., et al. 1995. Non-SH2 domains within Insulin receptor substrate-1 and SHC mediate their phosphotyrosine-dependent interaction with the NPEY motif of the Insulin-like growth factor I receptor. J. Biol. Chem. 270: 15639-15643.

CHROMOSOMAL LOCATION

Genetic locus: PIK3R1 (human) mapping to 5q13.1; Pik3r1 (mouse) mapping to 13 D1.

SOURCE

p-Pl 3-kinase p85 α (Tyr 467) is a rabbit polyclonal antibody raised against a short amino acid sequence containing phosphorylated Tyr 467 of Pl 3-kinase p85 α of human origin.

PRODUCT

Each vial contains 100 μg IgG in 1.0 ml PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

p-Pl 3-kinase p85 α (Tyr 467) is recommended for detection of Tyr 467 phosphorylated Pl 3-kinase p85 α of mouse, rat and human 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).

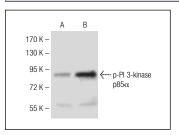
Suitable for use as control antibody for PI 3-kinase p85 α siRNA (h): sc-36217, PI 3-kinase p85 α siRNA (m): sc-36218, PI 3-kinase p85 α shRNA Plasmid (h): sc-36217-SH, PI 3-kinase p85 α shRNA Plasmid (m): sc-36218-SH, PI 3-kinase p85 α shRNA (h) Lentiviral Particles: sc-36217-V and PI 3-kinase p85 α shRNA (m) Lentiviral Particles: sc-36218-V.

Molecular Weight of p-Pl 3-kinase p85 α isoforms: 84/53/50/84 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (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) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



p-Pl 3-kinase p85 α (Tyr 467): sc-293115. Western blot analysis of Pl 3-kinase p85 α phosphorylation expression in untreated (**A**) and sorbitol treated (**B**) 293 whole cell lysafes

SELECT PRODUCT CITATIONS

- Chen, H.T., et al. 2011. Stromal cell-derived factor-1/CXCR4 promotes IL-6 production in human synovial fibroblasts. J. Cell. Biochem. 112: 1219-1227.
- Tsai, C.H., et al. 2013. High glucose induces vascular endothelial growth factor production in human synovial fibroblasts through reactive oxygen species generation. Biochim. Biophys. Acta 1830: 2649-2658.
- 3. Tsou, H.K., et al. 2013. HGF and c-Met interaction promotes migration in human chondrosarcoma cells. PLoS ONE 8: e53974.
- Yu, H.S., et al. 2013. Involvement of intercellular adhesion molecule-1 upregulation in bradykinin promotes cell motility in human prostate cancers. Int. J. Mol. Sci. 14: 13329-13345.

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 or our catalog for detailed protocols and support products.

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