p52 S6 kinase (K-16): sc-104570



The Power to Question

BACKGROUND

The phosphorylation and dephosphorylation of proteins on serine and threonine residues is an essential means of regulating a broad range of cellular functions in eukaryotes, including cell division, homeostasis and apoptosis. A group of proteins that are intimately involved in this process are the serine/threonine (Ser/Thr) protein kinases. p52 S6 kinase, also known as RPS6KC1 (ribosomal protein S6 kinase, 52 kDa, polypeptide 1) or RPK118, is a 1,066 amino acid member of the Ser/Thr kinase family that localizes to both the cytoplasm and the nucleus and contains one MIT domain, one PX domain and 2 protein kinase domains. Expressed at high levels in brain, placenta, heart, testis, kidney, liver and skeletal muscle, p52 S6 kinase catalyzes the ATP-dependent phosphorylation of target proteins and is thought to be involved in transmitting sphingosine-1 phosphate (SPP)-mediated signaling into the cell.

REFERENCES

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- Zhang, H., et al. 1999. Cloning, characterization, and chromosome mapping of RPS6KC1, a novel putative member of the ribosome protein S6 kinase family, to chromosome 12q12-q13.1. Genomics 61: 314-318.
- Hayashi, S., et al. 2002. Identification and characterization of RPK118, a novel sphingosine kinase-1-binding protein. J. Biol. Chem. 277: 33319-33324.
- 4. Ellson, C.D., et al. 2002. The PX domain: a new phosphoinositide-binding module. J. Cell Sci. 115: 1099-1105.
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- Ishida, S., et al. 2004. Differential modulation of PI 3-kinase/Akt pathway during all-trans retinoic acid- and Am80-induced HL-60 cell differentiation revealed by DNA microarray analysis. Biochem. Pharmacol. 68: 2177-2186.
- 7. Liu, L., et al. 2005. RPK118, a PX domain-containing protein, interacts with peroxiredoxin-3 through pseudo-kinase domains. Mol. Cells 19: 39-45.

CHROMOSOMAL LOCATION

Genetic locus: RPS6KC1 (human) mapping to 1q32.3; Rps6kc1 (mouse) mapping to 1 H6.

SOURCE

p52 S6 kinase (K-16) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of p52 S6 kinase of human origin.

STORAGE

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

PROTOCOLS

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

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-104570 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

p52 S6 kinase (K-16) is recommended for detection of p52 S6 kinase 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).

p52 S6 kinase (K-16) is also recommended for detection of p52 S6 kinase in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for p52 S6 kinase siRNA (h): sc-88609, p52 S6 kinase siRNA (m): sc-106342, p52 S6 kinase shRNA Plasmid (h): sc-88609-SH, p52 S6 kinase shRNA Plasmid (m): sc-106342-SH, p52 S6 kinase shRNA (h) Lentiviral Particles: sc-88609-V and p52 S6 kinase shRNA (m) Lentiviral Particles: sc-106342-V.

Molecular Weight of p52 S6 kinase: 118 kDa.

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

1. Zhou, X., et al. 2014. Rapamycin and everolimus facilitate hepatitis E virus replication: revealing a basal defense mechanism of PI3K-PKB-mTOR pathway. J. Hepatol. 61: 746-754.

RESEARCH USE

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

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