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p70 S6 Kinase (49D7) Rabbit mAb



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Source/Isotype: Applications: Reactivity: Sensitivity: MW (kDa): **UniProt ID:** Entrez-Gene Id: WB Н Endogenous 70, 85 Rabbit IgG #P23443 6198 **Product Usage** Application Dilution Information Western Blotting 1:1000 Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than **Storage** 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody. p70 S6 Kinase (49D7) Rabbit mAb detects endogenous levels of total p70 S6 kinase protein. The antibody Specificity / Sensitivity also recognizes p85 S6 kinase. Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to Source / Purification residues surrounding the amino-terminus of human p70 S6 kinase.

Background

p70 S6 kinase is a mitogen activated Ser/Thr protein kinase that is required for cell growth and G1 cell cycle progression (1,2). p70 S6 kinase phosphorylates the S6 protein of the 40S ribosomal subunit and is involved in translational control of 5' oligopyrimidine tract mRNAs (1). A second isoform, p85 S6 kinase, is derived from the same gene and is identical to p70 S6 kinase except for 23 extra residues at the amino terminus, which encode a nuclear localizing signal (1). Both isoforms lie on a mitogen activated signaling pathway downstream of phosphoinositide-3 kinase (PI-3K) and the target of rapamycin, FRAP/mTOR, a pathway distinct from the Ras/MAP kinase cascade (1). The activity of p70 S6 kinase is controlled by multiple phosphorylation events located within the catalytic, linker and pseudosubstrate domains (1). Phosphorylation of Thr229 in the catalytic domain and Thr389 in the linker domain are most critical for kinase function (1). Phosphorylation of Thr389, however, most closely correlates with p70 kinase activity in vivo (3). Prior phosphorylation of Thr389 is required for the action of phosphoinositide 3-dependent protein kinase 1 (PDK1) on Thr229 (4,5). Phosphorylation of this site is stimulated by growth factors such as insulin, EGF and FGF, as well as by serum and some G-protein-coupled receptor ligands, and is blocked by wortmannin, LY294002 (PI-3K inhibitor) and rapamycin (FRAP/mTOR inhibitor) (1,6,7). Ser411, Thr421 and Ser424 lie within a Ser-Pro-rich region located in the pseudosubstrate region (1). Phosphorylation at these sites is thought to activate p70 S6 kinase via relief of pseudosubstrate suppression (1,2). Another LY294002 and rapamycin sensitive phosphorylation site, Ser371, is an in vitro substrate for mTOR and correlates well with the activity of a partially rapamycin resistant mutant p70 S6 kinase (8).

Background References

- 1. Pullen, N. and Thomas, G. (1997) FEBS Lett 410, 78-82.
- 2. Dufner, A. and Thomas, G. (1999) Exp Cell Res 253, 100-9.
- 3. Weng, Q.P. et al. (1998) J Biol Chem 273, 16621-9.
- 4. Pullen, N. et al. (1998) Science 279, 707-10.
- 5. Alessi, D.R. et al. (1998) Curr Biol 8, 69-81.
- 6. Polakiewicz, R.D. et al. (1998) J Biol Chem 273, 23534-41.
- 7. Fingar, D.C. et al. (2002) Genes Dev 16, 1472-87.
- 8. Saitoh, M. et al. (2002) J Biol Chem 277, 20104-12.

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, Western Blot Buffer

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

WB: Western Blotting

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster **Cross-Reactivity Key**

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

3/23/24. 1:10 PM

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Limited Uses

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