Store at -20C

Phospho-p70 S6 Kinase (Thr389) Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.

Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 70, 85	Source: Rabbit	UniProt ID: #P23443	Entrez-Gene Id: 6198
Ар	plication			Dilution	
We	estern Blotting			1:1000	
Sin	nple Western™			1:10 - 1:50	
	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
pho	Phospho-p70 S6 Kinase (Thr389) Antibody detects endogenous levels of p70 S6 kinase only when phosphorylated at threonine 389. This antibody also detects p85 S6 kinase when phosphorylated at the analogous site (Thr412), and possibly S6KII phosphorylated at Thr388.				
to re	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues around Thr389 of human p70 S6 kinase. Antibodies are purified by protein A and peptide affinity chromatography.				
	HMRMk Ap We Sin Sup 20°0 vity Pho pho ana n Poly to re	Application Western Blotting Simple Western™ Supplied in 10 mM sodi 20°C. Do not aliquot the vity Phospho-p70 S6 Kinase phosphorylated at three analogous site (Thr412) n Polyclonal antibodies at to residues around Thr3	Application Western Blotting Simple Western™ Supplied in 10 mM sodium HEPES (pH 7.5 20°C. Do not aliquot the antibody. Vity Phospho-p70 S6 Kinase (Thr389) Antibody phosphorylated at threonine 389. This antili analogous site (Thr412), and possibly S6K Polyclonal antibodies are produced by imm to residues around Thr389 of human p70 S	Application Western Blotting Simple Western™ Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 1 20°C. Do not aliquot the antibody. Vity Phospho-p70 S6 Kinase (Thr389) Antibody detects endogenor phosphorylated at threonine 389. This antibody also detects panalogous site (Thr412), and possibly S6KII phosphorylated at the produced by immunizing animals with to residues around Thr389 of human p70 S6 kinase. Antibodices	Application Western Blotting Simple Western™ Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% of 20°C. Do not aliquot the antibody. Phospho-p70 S6 Kinase (Thr389) Antibody detects endogenous levels of p70 S6 kinase phosphorylated at threonine 389. This antibody also detects p85 S6 kinase when phosphorylated at threonine 389. This antibody also detects p85 S6 kinase when phosphorylated at Thr388. Polyclonal antibodies are produced by immunizing animals with a synthetic phosphore to residues around Thr389 of human p70 S6 kinase. Antibodies are purified by protein

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

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- 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).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

WB: Western Blotting W-S: Simple Western™

Applications Key

Cross-Reactivity Key

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dq: dog Pq: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

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

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