Phospho-eIF4B (Ser406) Antibody



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

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Applications: W, IP, IF-IC	Reactivity: H M	Sensitivity: Endogenous	MW (kDa): 80	Source/Isotype: Rabbit	UniProt ID: #P23588	Entrez-Gene Id: 1975	
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence	(Immunocytochem	iistry)		Dilution 1:1000 1:50 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.					
Specificity/Sen	sitivity	Phospho-eIF4B (Ser406) Antibody detects endogenous levels of eIF4B protein only when phosphorylated at Ser406.					
Species predic based on 100% homology		Rat					
Source / Purifi	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser406 of human eIF4B protein. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		Eukaryotic initiation factor 4B (eIF4B) is thought to assist the eIF4F complex in translation initiation. In plants, eIF4B is known to interact with the poly-(A) binding protein, increasing its poly-(A) binding activity (1). Heat shock and serum starvation cause dephosphorylation of eIF4B at multiple sites with kinetics similar to those of the corresponding inhibition of translation, while phosphorylation of eIF4B following insulin treatment correlates well with an observed increase in translation (2-5). Multiple kinases, including p70 S6 kinase, can phosphorylate eIF4B <i>in vitro</i> , and at least one serum-inducible eIF4B phosphorylation site is sensitive to rapamycin and LY294002 (6). Recently, Ser406 was identified as a novel phosphorylation site regulated by mitogens (7), and the phosphorylation of this site is dependent on MEK and mTOR activity (7). This phosphorylation is shown to be essential for the translational activity of eIF4B (7).					
Background Re	eferences	1. Le, H. et al. (1997) <i>J. Biol. Chem.</i> 272, 16247-16255. 2. Duncan, R.F. and Hershey, J.W. (1989) <i>J. Cell Biol.</i> 109, 1467-1481. 3. Duncan, R.F. and Hershey, J.W. (1984) <i>J. Biol. Chem.</i> 259, 11882-11889. 4. Duncan, R. and Hershey, J.W. (1985) <i>J. Biol. Chem.</i> 260, 5493-5497. 5. Manzella, J.M. et al. (1991) <i>J. Biol. Chem.</i> 266, 2383-2389. 6. Gingras, A.C. et al. (2001) <i>Genes Dev.</i> 15, 807-826. 7. van Gorp, A.G. et al. (2009) <i>Oncogene</i> 28, 95-106.					
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	, western blot).	
Western Blot E	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.					
Applications K	ey	W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)					
Cross-Reactivit	ty Key	H: Human M: Mouse					
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