eIF4B Antibody	Cell Signaling TECHNOLOGY®		
Stor	Orders:	877-616-CELL (2355) orders@cellsignal.com	
	Support:	877-678-TECH (8324)	
359	Web:	info@cellsignal.com cellsignal.com	
#	3 Trask Lane Danvers Mas	sachusetts 01923 USA	
For Research Use Only. Not for Use in Diagnostic Procedures.			

Applications: W	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 80	Source/Isotype: Rabbit	UniProt ID: #P23588	Entrez-Gene Id: 1975		
Product Usage Information	e	Application Western Blotting			Dilution 1:1000			
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/Ser	nsitivity	eIF4B Antibody recognizes endogenous levels of eIF4B, independent of phosphorylation.						
Source / Purifi	e / Purification Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues at the amino terminus of human eIF4B. 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 R	eferences	1. Le, H. et al. (1997) <i>J</i> . 2. Duncan, R.F. and He 3. Duncan, R.F. and He 4. Duncan, R. and Her 5. Manzella, J.M. et al. 6. Gingras, A.C. et al. (7. van Gorp, A.G. et al	Biol. Chem. 272, 14 ershey, J.W. (1989) J ershey, J.W. (1984) J shey, J.W. (1985) J. L (1991) J. Biol. Chen 2001) Genes Dev. 1 . (2009) Oncogene 2	5247-16255. <i>Cell Biol.</i> 109, 1467-148 <i>Biol. Chem.</i> 259, 11882- <i>Biol. Chem.</i> 260, 5493-54 1. 266, 2383-2389. 5, 807-826. 28, 95-106.	1. -11889. 97.			
Species Reacti	ivity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot l	Buffer	IMPORTANT: For west TBS, 0.1% Tween® 20	or western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X n® 20 at 4°C with gentle shaking, overnight.					
Applications K	(ey	W: Western Blotting						
Cross-Reactivi	ity Key	H: Human M: Mouse R: Rat Mk: Monkey						
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