

#2855

Phospho-4E-BP1 (Thr37/46) (236B4) Rabbit



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Applications: W, W-S, IHC-P, IF-IC, FC-FP	Reactivity: H M R Mk Dm	Sensitivity: Endogenous	MW (kDa): 15 to 20	Source/Isotype: Rabbit IgG	UniProt ID: #Q13541	Entrez-Gene Id: 1978
Product Usage Information		Application			Dilution	
Illioilliation		Western Blotting Simple Western™			1:10	
		•	ry (Paraffin)			- 1:50 0 - 1:3200
		Immunohistochemistry (Paraffin) Immunofluorescence (Immunocytochemistry)			1:200 - 1:800	
		Flow Cytometry (Fixed	•	iisti y)		- 1:200
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier free (BSA	and azide free) ver	sion of this product see	product #39788.	
Specificity/Sensitivity		Phospho-4E-BP1 (Thr37/46) (236B4) Rabbit mAb detects endogenous levels of 4E-BP1 only when phosphorylated at Thr37 and/or Thr46. This antibody may cross-react with 4E-BP2 and 4E-BP3 when phosphorylated at equivalent sites. Non-specific staining has been observed in mitotic cells by immunofluorescence.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr37 and Thr46 of mouse 4E-BP1.				
Background		Translation repressor protein 4E-BP1 (also known as PHAS-1) inhibits cap-dependent translation by binding to the translation initiation factor eIF4E. Hyperphosphorylation of 4E-BP1 disrupts this interaction and results in activation of cap-dependent translation (1). Both the PI3 kinase/Akt pathwa and FRAP/mTOR kinase regulate 4E-BP1 activity (2,3). Multiple 4E-BP1 residues are phosphorylated in vivo (4). While phosphorylation by FRAP/mTOR at Thr37 and Thr46 does not prevent the binding of 4BP1 to eIF4E, it is thought to prime 4E-BP1 for subsequent phosphorylation at Ser65 and Thr70 (5).				srupts this nase/Akt pathway hosphorylated <i>in</i> the binding of 4E-
Background References		 Pause, A. et al. (1994) Nature 371, 762-7. Brunn, G.J. et al. (1997) Science 277, 99-101. Gingras, A.C. et al. (1998) Genes Dev 12, 502-13. Fadden, P. et al. (1997) J Biol Chem 272, 10240-7. Gingras, A.C. et al. (1999) Genes Dev 13, 1422-37. 				
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.

Applications Key W: Western Blotting W-S: Simple Western™ IHC-P: Immunohistochemistry (Paraffin) IF-IC:

Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey Dm: D. melanogaster

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