## 4E-BP1 (53H11) Rabbit mAb



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

<b>Applications:</b> W, W-S, IP, IHC-P, IF-IC, FC-FP	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 15-20	Source/Isotype: Rabbit IgG	<b>UniProt ID:</b> #Q13541	Entrez-Gene Id 1978
Product Usage Information		Application Western Blotting			<b>Dilution</b> 1:1000	
		Simple Western™			1:50 -	1:250
		Immunoprecipitation			1:50	
		Immunohistochemist	ry (Paraffin)		1:1200	) - 1:4800
		Immunofluorescence	(Immunocytochem	istry)	1:800	- 1:3200
		Flow Cytometry (Fixed	l/Permeabilized)		1:1600	)
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/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) version of this product see product #42235.				
Specificity/Sensitivity		4E-BP1 (53H11) Rabbit mAb detects endogenous levels of total 4E-BP1 protein.				
Source / Purification		4E-BP1 (53H11) Rabbit mAb is produced by immunizing rabbits with a synthetic peptide corresponding to residues surrounding Ser112 of human 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 pathway and FRAP/mTOR kinase regulate 4E-BP1 activity (2,3). Multiple 4E-BP1 residues are phosphorylated <i>in vivo</i> (4). While phosphorylation by FRAP/mTOR at Thr37 and Thr46 does not prevent the binding of 4E-BP1 to eIF4E, it is thought to prime 4E-BP1 for subsequent phosphorylation at Ser65 and Thr70 (5).				
Background References		1. Pause, A. et al. (1994) <i>Nature</i> 371, 762-7. 2. Brunn, G.J. et al. (1997) <i>Science</i> 277, 99-101. 3. Gingras, A.C. et al. (1998) <i>Genes Dev</i> 12, 502-13. 4. Fadden, P. et al. (1997) <i>J Biol Chem</i> 272, 10240-7. 5. Gingras, A.C. et al. (1999) <i>Genes Dev</i> 13, 1422-37.				
Species Reactivi	ty	Species reactivity is de	etermined by testing	g in at least one approve	ed 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		<b>W:</b> Western Blotting <b>W-S:</b> Simple Western <sup>™</sup> <b>IP:</b> Immunoprecipitation <b>IHC-P:</b> Immunohistochemistry (Paraffin) <b>IF-IC:</b> Immunofluorescence (Immunocytochemistry) <b>FC-FP:</b> Flow Cytometry (Fixed/Permeabilized)				
Cross-Reactivity	Key	H: Human M: Mouse R: Rat Mk: Monkey				
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