## Phospho-eIF2α (Ser51) (119A11) Rabbit



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

<b>Applications:</b> W, IP, IHC-P	<b>Reactivity:</b> H M R Mk Dm	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 38	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P05198	Entrez-Gene Id: 1965
Product Usage Information		Application Western Blotting			<b>Dilution</b> 1:1000	
		Immunoprecipitation			1:50	
		Immunohistochemistry (Paraffin)			1:75 - 1:300	
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) ver	sion of this product see	product #63795.	
Specificity/Sensitivity		Phospho-eIF2alpha (Ser51) RmAb detects endogenous eIF2alpha only when phosphorylated at Ser51. The antibody does not recognize elF2alpha phosphorylated at other sites. Human eIF2alpha residue Ser52 historically has been referenced as Ser51.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser51 of human eIF2alpha.				
Background		Phosphorylation of the eukaryotic initiation factor 2 (eIF2) $\alpha$ subunit is a well-documented mechanism to downregulate protein synthesis under a variety of stress conditions. eIF2 binds GTP and Met-tRNAi and transfers Met-tRNA to the 40S subunit to form the 43S preinitiation complex (1,2). eIF2 promotes a new round of translation initiation by exchanging GDP for GTP, a reaction catalyzed by eIF2B (1,2). Kinases that are activated by viral infection (PKR), endoplasmic reticulum stress (PERK/PEK), amino acid deprivation (GCN2), or heme deficiency (HRI) can phosphorylate the $\alpha$ subunit of eIF2 (3,4). This phosphorylation stabilizes the eIF2-GDP-eIF2B complex and inhibits the turnover of eIF2B. Induction of PKR by IFN- $\gamma$ and TNF- $\alpha$ induces potent phosphorylation of eIF2 $\alpha$ at Ser51 (5,6).				
Background Re	ferences	<ol> <li>Kimball, S.R. (1999) <i>Int. J. Biochem. Cell Biol.</i> 31, 25-29.</li> <li>de Haro, C. et al. (1996) <i>FASEB J.</i> 10, 1378-87.</li> <li>Kaufman, R.J. (1999) <i>Genes Dev.</i> 13, 1211-33.</li> <li>Sheikh, M.S. and Fornace Jr., A.J. (1999) <i>Oncogene</i> 18, 6121-8.</li> <li>Cheshire, J.L. et al. (1999) <i>J. Biol. Chem.</i> 274, 4801-6.</li> <li>Zamanian-Daryoush, M. et al. (2000) <i>Mol. Cell. Biol.</i> 20, 1278-90.</li> </ol>				
Species Peactiv	•.	6. Zamanian-Daryousl	. ,	<u> </u>		

**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 IP: Immunoprecipitation IHC-P: Immunohistochemistry (Paraffin)

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

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