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## eIF2 $\alpha$ Antibody



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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source:	UniProt ID:	Entrez-Gene Id:
WB	H M R Mk	Endogenous	38	Rabbit	#P05198	1965

<b>Product Usage Information</b>	<b>Application</b> Western Blotting	<b>Dilution</b> 1:1000
<b>Storage</b>	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.	
<b>Specificity / Sensitivity</b>	eIF2 $\alpha$ Antibody detects endogenous levels of total eIF2 $\alpha$ protein.	
<b>Source / Purification</b>	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy-terminal sequence of eIF2 $\alpha$ . Antibodies are purified by protein A and peptide affinity chromatography.	
<b>Background</b>	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).	
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Kimball, S.R. (1999) <i>Int. J. Biochem. Cell Biol.</i> 31, 25-29.</li> <li>2. de Haro, C. et al. (1996) <i>FASEB J.</i> 10, 1378-87.</li> <li>3. Kaufman, R.J. (1999) <i>Genes Dev.</i> 13, 1211-33.</li> <li>4. Sheikh, M.S. and Fornace Jr., A.J. (1999) <i>Oncogene</i> 18, 6121-8.</li> <li>5. Cheshire, J.L. et al. (1999) <i>J. Biol. Chem.</i> 274, 4801-6.</li> <li>6. Zamanian-Daryoush, M. et al. (2000) <i>Mol. Cell. Biol.</i> 20, 1278-90.</li> </ol>	

<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).
<b>Western Blot Buffer</b>	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.
<b>Applications Key</b>	<b>WB:</b> Western Blotting
<b>Cross-Reactivity Key</b>	<b>H:</b> human <b>M:</b> mouse <b>R:</b> rat <b>Hm:</b> hamster <b>Mk:</b> monkey <b>Vir:</b> virus <b>Mi:</b> mink <b>C:</b> chicken <b>Dm:</b> D. melanogaster <b>X:</b> Xenopus <b>Z:</b> zebrafish <b>B:</b> bovine <b>Dg:</b> dog <b>Pg:</b> pig <b>Sc:</b> S. cerevisiae <b>Ce:</b> C. elegans <b>Hr:</b> horse <b>GP:</b> Guinea Pig <b>Rab:</b> rabbit <b>All:</b> all species expected
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