## eIF2α (D7D3) XP<sup>®</sup> Rabbit mAb (Biotinylated)



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

<b>Applications:</b> W	Reactivity: H M R Mk	<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		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 140 mM NaCl, 3 mM KCI, 10 mM sodium phosphate (pH 7.4) dibasic, 2 mM potassium phosphate monobasic, 2 mg/mL BSA, and 50% glycerol. Store at –20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		eIF2 $\alpha$ (D7D3) XP $^{\otimes}$ Rabbit mAb (Biotinylated) detects endogenous levels of total eIF2 $\alpha$ protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a purified recombinant human eIF2 $\alpha$ protein.				
Description		This Cell Signaling Technology antibody is conjugated to biotin under optimal conditions. The biotinylated antibody is expected to exhibit the same species cross-reactivity as the unconjugated eIF2 $\alpha$ (D7D3) XP $^{\otimes}$ Rabbit mAb #5324.				
Background		Phosphorylation of the eukaryotic initiation factor 2 (eIF2) α 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 α subunit of eIF2 (3,4). This phosphorylation stabilizes the eIF2-GDP-eIF2B complex and inhibits the turnover of eIF2B. Induction of PKR by IFN-γ and TNF-α induces potent phosphorylation of eIF2α at Ser51 (5,6).				
Background References		<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 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

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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