

eIF2α (D7D3) XP[®] Rabbit mAb (Biotinylated)



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Applications: W	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 38	Source/Isotype: Rabbit IgG	UniProt ID: #P05198	Entrez-Gene Id: 1965
Product Usage Information	2	Application Western Blotting			Dilution 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 α (D7D3) XP $^{\! 8}$ Rabbit mAb (Biotinylated) detects endogenous levels of total eIF2 α protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a purified recombinant human eIF2 α 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 eIF2α (D7D3) XP [®] 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		 Kimball, S.R. (1999) Int. J. Biochem. Cell Biol. 31, 25-29. de Haro, C. et al. (1996) FASEB J. 10, 1378-87. Kaufman, R.J. (1999) Genes Dev. 13, 1211-33. Sheikh, M.S. and Fornace Jr., A.J. (1999) Oncogene 18, 6121-8. Cheshire, J.L. et al. (1999) J. Biol. Chem. 274, 4801-6. Zamanian-Daryoush, M. et al. (2000) Mol. Cell. Biol. 20, 1278-90. 				

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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