

Store at -20C  
#2332**eEF2 Antibody**
**Orders:** 877-616-CELL (2355)  
orders@cellsignal.com

**Support:** 877-678-TECH (8324)

**Web:** info@cellsignal.com  
cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W, IF-IC	H M R Mk Dm	Endogenous	95	Rabbit	#P13639	1938

**Product Usage Information****Application**Western Blotting  
Immunofluorescence (Immunocytochemistry)**Dilution**1:1000  
1:50**Storage**

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.

**Specificity/Sensitivity**

eEF2 Antibody detects endogenous levels of total eEF2 independent of phosphorylation.

**Species predicted to react based on 100% sequence homology**

Hamster, Chicken

**Source / Purification**

Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues at the amino-terminus of human eEF2. Antibodies are purified by protein A and peptide affinity chromatography.

**Background**

Eukaryotic elongation factor 2 (eEF2) catalyzes the translocation of peptidyl-tRNA from the A site to the P site on the ribosome. It has been shown that phosphorylation of eEF2 at threonine 56 by eEF2 kinase inhibits its activity (1-4). eEF2 kinase is normally dependent on Ca<sup>2+</sup> ions and calmodulin (5,6). eEF2 kinase can also be activated by PKA in response to elevated cAMP levels (7-9), which are generally increased in stress- or starvation-related conditions. A variety of treatments known to raise intracellular Ca<sup>2+</sup> or cAMP levels have been shown to result in increased phosphorylation of eEF2, and thus to inhibit peptide-chain elongation. The inactive phosphorylated eEF2 can be converted to its active nonphosphorylated form by a protein phosphatase, most likely a form of protein phosphatase-2A (PP-2A) (10). Insulin, which activates protein synthesis in a wide range of cell types, induces rapid dephosphorylation of eEF2 through mTOR signaling and may involve modulation of the activity of the PP-2A or the eEF2 kinase or both (11).

**Background References**

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4. Redpath, N.T. et al. (1993) *Eur. J. Biochem.* 213, 689-699.
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6. Palfrey, H.C. et al. (1987) *J. Biol. Chem.* 262, 9785-9792.
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8. Diggle, T. et al. (1998) *Biochem. J.* 336, 525-529.
9. Hovland, R. et al. (1999) *FEBS Lett.* 444, 97-101.
10. Redpath, N.T. et al. (1996) *EMBO J* 15, 2291-7.
11. Nilsson, A. and Nygård, O. (1995) *Biochim Biophys Acta* 1268, 263-8.

**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 **IF-IC:** Immunofluorescence (Immunocytochemistry)**Cross-Reactivity Key****H:** Human **M:** Mouse **R:** Rat **Mk:** Monkey **Dm:** D. melanogaster**Trademarks and Patents**

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