

PhosphoPlus ® GCN2 (Thr899) Antibody



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UniProt ID:	Entrez-Gene Id:	
#Q9P2K8	440275	

Product Includes	Product #	Quantity	Mol. Wt	Isotype/Source
Phospho-GCN2 (Thr899) (E1V9M) Rabbit mAb	94668	100 µl	220 kDa	Rabbit IgG
GCN2 (E7G7E) Rabbit mAb	65981	100 µl	220 kDa	Rabbit IgG

Please visit cellsignal.com for individual component applications, species cross-reactivity, dilutions, protocols, and additional product information.

Description

PhosphoPlus[®] Duets from Cell Signaling Technology (CST) provide a means to assess protein activation status. Each Duet contains an activation-state and total protein antibody to your target of interest. These antibodies have been selected from CST's product offering based upon superior performance in specified applications.

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

Background

Phosphorylation of the eukaryotic initiation factor 2 (eIF2) alpha subunit is a well-documented mechanism of downregulating protein synthesis under a variety of stress conditions. Kinases activated by viral infection (PKR), endoplasmic reticulum stress (PERK/PEK), amino acid deprivation (GCN2), and hemin deficiency (HRI) can phosphorylate the eIF2 alpha subunit (1,2). GCN2 is also required for UV light-induced translation inhibition, and *in vivo* phosphorylation of murine GCN2 at Thr898 is induced by both UV irradiation and by leucine deprivation (3). UV-induced activation of NF-kB also requires GCN2, which may act simply by preventing translation of IkB-alpha to replace pools that have been ubiquitinated and degraded (4). Interestingly, proteasome inhibitors (MG132 and ALLN) activate the GCN2/eIF2alpha pathway, suggesting a pivotal role for this kinase in stress response and ubiquitin-mediated signaling (5). *In vitro* autophosphorylation of yeast GCN2 within its activation loop at Thr882 and Thr887 (Thr898 and Thr903 in mouse) has also been reported (6).

Background References

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- 2. Sheikh, M.S. and Fornace, A.J. (1999) Oncogene 18, 6121-8.
- 3. Deng, J. et al. (2002) Curr Biol 12, 1279-86.
- 4. Jiang, H.Y. and Wek, R.C. (2005) *Biochem J* 385, 371-80.
- 5. Jiang, H.Y. and Wek, R.C. (2005) J Biol Chem 280, 14189-202.
- 6. Garcia-Barrio, M. et al. (2002) J Biol Chem 277, 30675-83.

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