

Phospho-Mnk1 (Thr197/Thr202) Antibody

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Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W	H M R	Endogenous	55	Rabbit	#Q9BUB5	8569

Product Usage Information**Application**

Western Blotting

Dilution

1:1000

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

Phospho-Mnk1 (Thr197/Thr202) Antibody recognizes endogenous levels of Mnk1 protein only when phosphorylated at Thr197 and Thr202.

Source / Purification

Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr250 and Thr255 of human Mnk1 protein. These two residues correspond to residues Thr197 and Thr202 of a shorter mouse Mnk1 (Waskiewicz, A. J. et al. (1997) *EMBO J.* 16, 1909-1920). Antibodies are purified by protein A and peptide affinity chromatography.

Background

Eukaryotic initiation factor 4E (eIF4E) binds to the mRNA cap structure to mediate the initiation of translation (1,2). eIF4E interacts with eIF4G, a scaffold protein that promotes assembly of eIF4E and eIF4A into the eIF4F complex (2). eIF4B is thought to assist the eIF4F complex in translation initiation. Upon activation by mitogenic and/or stress stimuli mediated by Erk and p38 MAPK, Mnk1 phosphorylates eIF4E at Ser209 *in vivo* (3,4). Two Erk and p38 MAPK phosphorylation sites in mouse Mnk1 (Thr197 and Thr202) are essential for Mnk1 kinase activity (3). The carboxy-terminal region of eIF4G also contains serum-stimulated phosphorylation sites, including Ser1108, Ser1148, and Ser1192 (5). Phosphorylation at these sites is blocked by the PI3 kinase inhibitor LY294002 and by the FRAP/mTOR inhibitor rapamycin.

Background References

1. Sonenberg, N. et al. (1978) *Proc. Natl. Acad. Sci. USA* 75, 4843-47.
2. Gingras, A.C. et al. (1999) *Annu. Rev. Biochem.* 68, 913-63.
3. Waskiewicz, A. et al. (1999) *Mol. Cell. Biol.* 19, 1871-80.
4. Pyronnet, S. et al. (1999) *EMBO J.* 18, 270-9.
5. Raught, B. et al. (2000) *EMBO J.* 19, 434-44.

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 nonfat dry milk, 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

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