

Product Data Sheet

Anti-MKK2 (Phospho-T394) Antibody

Catalog #	Source	Reactivity	Applications			
CPA3301	Rabbit	H, Mk	WB, IH, IF/IC, IP			
Description	F	Rabbit polyclonal antibody to MKK2 (Phospho-T394)				
Immunogen	k	KLH-conjugated synthetic pho	osphopeptide corresponding to residues surrounding			
	٦	T394 of human MKK2 proteir	. The exact sequence is proprietary.			
Purification	T	The antibody was purified by	immunogen affinity chromatography.			
Specificity	F	Recognizes endogenous level	s of MKK2 protein only when phosphorylated at T394.			
Clonality	F	Polyclonal				
Conjugation						
Form	L	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	ä	and 0.01% sodium azide.				
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200), IP (1/10 - 1/100)				
Gene Symbol	٦	MAP2K2				
Alternative N	ames N	MEK2; MKK2; PRKMK2; Dual	specificity mitogen-activated protein kinase kinase 2;			
	٦	MAP kinase kinase 2; MAPKK	2; ERK activator kinase 2; MAPK/ERK kinase 2; MEK 2			
Entrez Gene	5	5605 (Human)				
SwissProt	F	P36507 (Human)				
Storage/Stabi	ility S	Shipped at 4°C. Upon deliver	aliquot and store at -20°C for one year. Avoid			
	f	freeze/thaw cycles.				

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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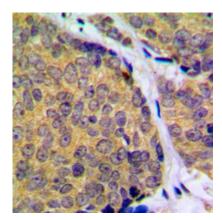
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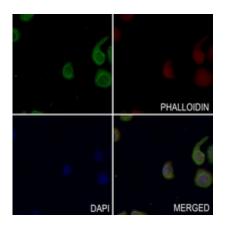
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kDa A B C 70 45 35 25 Western blot analysis of MKK2 (Phospho-T394) expression in HEK293T (A), Hela (B), DLD (C) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 44 kD)



Immunohistochemical analysis of MKK2 (Phospho-T394) staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of MKK2 (Phospho-T394) staining in SGC7901 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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