

Product Data Sheet

Anti-MKK4 (Phospho-S80) Antibody

Catalo	g # Sourc	e Reactivity	Applications		
CPA20	54 Rabbi	t H, M, R, B, Z	WB, IH		
Description		Rabbit polyclonal antibody to MKK4 (Phospho-S80)			
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
		S80 of human MKK4 protein.	The exact sequence is proprietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of MKK4 protein only when phosphorylated at S80.			
Clonality		Polyclonal			
Conjug	ation				
Form		Liquid in 0.42% Potassium phe	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100)			
Gene Symbol		MAP2K4			
Alternative Names		JNKK1; MEK4; MKK4; PRKMK4; SEK1; SERK1; SKK1; Dual specificity			
		mitogen-activated protein kinase kinase 4; MAP kinase kinase 4; MAPKK 4;			
		JNK-activating kinase 1; MAP	C/ERK kinase 4; MEK 4; SAPK/ERK kinase 1; SEK1;		
		Stress-activated protein kinas	e kinase 1; S		
Entrez Gene		6416 (Human); 26398 (Mouse)			
SwissProt		P45985 (Human); P47809 (Mouse)			
Storage/Stability		Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
		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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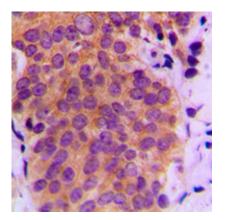
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For research purposes only, not for human use

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Western blot analysis of MKK4 (Phospho-S80) expression in zebrafish (A) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 44 kD)



Immunohistochemical analysis of MKK4 (Phospho-S80) 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.

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