

Product Data Sheet

Anti-ASK1 (Phospho-T838) Antibody

Catalog #	Source	e Reactivity	Applications		
CPA6042	Rabbit	H, M	WB, IH		
Description	Description Rabbit polyclonal antibody to ASK1 (Phospho-T838)				
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
		T838 of human ASK1 protein. The exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous leve	ls of ASK1 protein only when phosphorylated at T838.		
Clonality	Clonality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 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		MAP3K5			
Alternative N	ames	ASK1; MAPKKK5; MEKK5; M	togen-activated protein kinase kinase kinase 5;		
		Apoptosis signal-regulating l	cinase 1; ASK-1; MAPK/ERK kinase kinase 5; MEK kinase		
		5; MEKK 5			
Entrez Gene		4217 (Human); 26408 (Mou	se)		
SwissProt		Q99683 (Human); O35099 (I	Mouse)		
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y 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

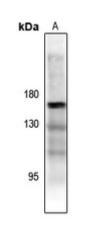
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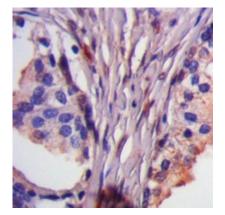
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Western blot analysis of ASK1 (Phospho-T838) expression in K562 (A) whole cell lysates. (Predicted band size: 154 kD; Observed band size: 155 kD)



Immunohistochemical analysis of ASK1 (Phospho-T838) staining in human prostate 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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