

Product Data Sheet

Anti-DUSP6 Antibody

Catalog #	Source	Reactivity	Applications
CPA4390	Rabbit	H, M, R, B, D, P	WB, IH, IF/IC
Description	Ral	bit polyclonal antibody to I	DUSP6
Immunogen	KLł	I-conjugated synthetic pept	ide encompassing a sequence within the center
	reg	ion of human DUSP6. The e	xact sequence is proprietary.
Purification	The	e antibody was purified by i	mmunogen affinity chromatography.
Specificity	Red	cognizes endogenous levels	of DUSP6 protein.
Clonality	Pol	yclonal	
Form	Liq	uid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	d 0.01% sodium azide.	
Dilution	WE	8 (1/500 - 1/1000), IH (1/100 ·	- 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	DU	SP6	
Alternative Na	ames Mk	P3; PYST1; Dual specificity	protein phosphatase 6; Dual specificity protein
	pho	osphatase PYST1; Mitogen-a	activated protein kinase phosphatase 3; MAP kinase
	pho	osphatase 3; MKP-3	
Entrez Gene	184	18 (Human); 67603 (Mouse)	; 116663 (Rat)
SwissProt	Q1	6828 (Human); Q9DBB1 (M	ouse); Q64346 (Rat)
Storage/Stabi	lity Shi	pped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid
	fre	eze/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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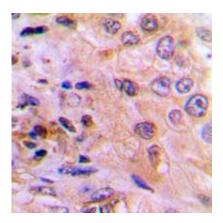
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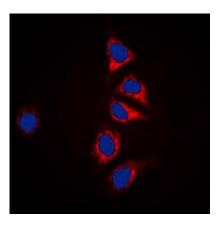
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Western blot analysis of DUSP6 expression in HepG2 (A), NIH3T3 (B) whole cell lysates.



Immunohistochemical analysis of DUSP6 staining in human lung 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of DUSP6 staining in HepG2 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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