

Product Data Sheet

Anti-c-Jun (Phospho-T239) Antibody

Catalog #	Source	Reactivity	Applications			
CPA4696	Rabbit	H, M, R, B, C, P	WB, IH, IP			
Description	Rabb	Rabbit polyclonal antibody to c-Jun (Phospho-T239)				
Immunogen	KLH-	conjugated synthetic phosphop	eptide corresponding to residues surrounding			
	Т239	of human c-Jun protein. The e	kact sequence is proprietary.			
Purification	The a	antibody was purified by immu	nogen affinity chromatography.			
Specificity	Reco	gnizes endogenous levels of c-J	un protein only when phosphorylated at T239.			
Clonality	Polyc	Polyclonal				
Conjugation						
Form	Liqui	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), IP (1/10 - 1/100)			
Gene Symbol	JUN					
Alternative Na	ames Trans	scription factor AP-1; Activator	protein 1; AP1; Proto-oncogene c-Jun; V-jun			
	aviar	sarcoma virus 17 oncogene ho	omolog; p39			
Entrez Gene	3725	(Human); 16476 (Mouse); 245	16 (Rat)			
SwissProt	P054	12 (Human); P05627 (Mouse);	P17325 (Rat)			
Storage/Stabi	lity Shipp	oed at 4°C. Upon delivery aliqu	ot and store at -20°C for one year. Avoid			
	freez	e/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

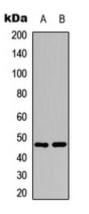
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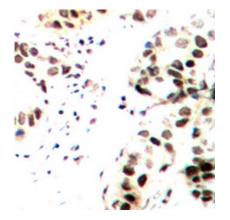
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For research purposes only, not for human use

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Western blot analysis of c-Jun (Phospho-T239) expression in HEK293T UV-treated (A), HeLa UV-treated (B) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 48 kD)



Immunohistochemical analysis of c-Jun (Phospho-T239) 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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