

Product Data Sheet

Anti-c-Jun (Phospho-S63) Antibody

Catalog #	Source	Reactivity	Applications		
CPA4695	Rabbit	H, M, R, B, Rb, S	WB, IH		
Description	Rab	bit polyclonal antibody to	c-Jun (Phospho-S63)		
Immunogen	KLH	-conjugated synthetic pho	sphopeptide corresponding to residues surrounding		
	S63	of human c-Jun protein. T	he exact sequence is proprietary.		
Purification	The	antibody was purified by i	immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous levels	of c-Jun protein only when phosphorylated at S63.		
Clonality	Poly	/clonal			
Conjugation					
Form	Liqu	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 (1/100	- 1/200)		
Gene Symbol	JUN	I			
Alternative Na	ames Trar	scription factor AP-1; Acti	vator protein 1; AP1; Proto-oncogene c-Jun; V-jun		
	avia	in sarcoma virus 17 oncoge	ene homolog; p39		
Entrez Gene	372	5 (Human); 16476 (Mouse); 24516 (Rat)		
SwissProt	P05	412 (Human); P05627 (Mc	ouse); P17325 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	free	ze/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

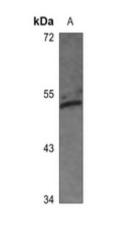
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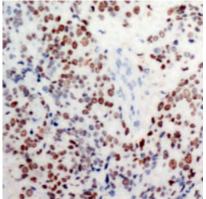
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For research purposes only, not for human use

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Western blot analysis of c-Jun (Phospho-S63) expression in A549 (A) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 48 kD)



Immunohistochemical analysis of c-Jun (Phospho-S63) 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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