

Product Data Sheet

Anti-SP1 (Phospho-T453) Antibody

Catalog #	Source	Reactivity	Applications	
CPA4253	Rabbit	H, M, R, B, C, Mk	WB, IH, IF/IC, ChIP	
Description	Ra	Rabbit polyclonal antibody to SP1 (Phospho-T453)		
Immunogen	KL	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding		
	T4	53 of human SP1 protein. The exact sequ	uence is proprietary.	
Purification	Th	e antibody was purified by immunogen	affinity chromatography.	
Specificity	Re	cognizes endogenous levels of SP1 prote	in only when phosphorylated at T453.	
Clonality	Ро	Polyclonal		
Conjugation				
Form	Lic	quid in 0.42% Potassium phosphate, 0.87	% Sodium chloride, pH 7.3, 30% glycerol,	
	an	d 0.01% sodium azide.		
Dilution	W	B (1/500 - 1/1000), IH (1/100 - 1/200), IF/IO	C (1/100 - 1/500), ChIP (Use at an assay	
	de	pendent concentration)		
Gene Symbol	SP	1		
Alternative Na	ames TS	FP1; Transcription factor Sp1		
Entrez Gene	66	67 (Human); 20683 (Mouse); 24790 (Rat	.)	
SwissProt	PO	8047 (Human); 089090 (Mouse); Q0171	.4 (Rat)	
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery aliquot and	store at -20°C for one year. Avoid	
	fre	eeze/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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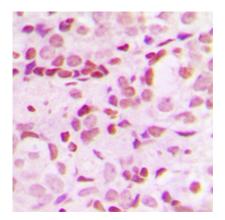
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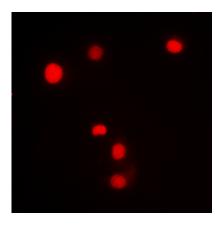
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Western blot analysis of SP1 (Phospho-T453) expression in H1688 (A) whole cell lysates. (Predicted band size: 80 kD; Observed band size: 80 kD)



Immunohistochemical analysis of SP1 (Phospho-T453) 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.



Immunofluorescent analysis of SP1 (Phospho-T453) staining in Jurkat 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.

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