

Product Data Sheet

Anti-53BP1 (Phospho-S6) Antibody

Catalog #	Source	Reactivity	Applications		
CPA5864	Rabbit	H, M, R	WB, IH		
Description	Ra	Rabbit polyclonal antibody to 53BP1 (Phospho-S6)			
Immunogen	KL	H-conjugated synthetic pl.	hosphopeptide corresponding to residues surrounding S6		
	of	human 53BP1 protein. Th	ne exact sequence is proprietary.		
Purification	Th	ie antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of 53BP1 protein only when phosphorylated at S6.		
Clonality	Ро	blyclonal			
Conjugation					
Form	Lic	quid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/50) - 1/200)		
Gene Symbol	TP	253BP1			
Alternative Na	ames Tu	mor suppressor p53-bind	ing protein 1; 53BP1; p53-binding protein 1; p53BP1		
Entrez Gene	71	158 (Human)			
SwissProt	Q1	12888 (Human); P70399 (I	Mouse)		
Storage/Stabi	lity Sh	iipped at 4°C. Upon delive	ry 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

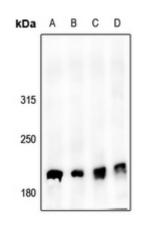
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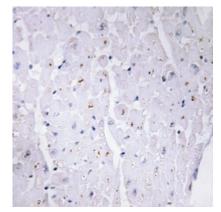


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Western blot analysis of 53BP1 (Phospho-S6) expression in Hela (A), U87MG (B), 3T3L1 (C), H9C2 (D) whole cell lysates. (Predicted band size: 213 kD; Observed band size: 220 kD)



Immunohistochemical analysis of 53BP1 (Phospho-S6) staining in human heart 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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