

Product Data Sheet

Anti-53BP1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3348	Rabbit	H, M, R	WB, IH		
Description	Rabb	Rabbit polyclonal antibody to 53BP1			
Immunogen	KLH-	conjugated synthetic p	eptide encompassing a sequence within the N-term		
	regio	on of human 53BP1. Th	e exact sequence is proprietary.		
Purification	The	antibody was purified l	by immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous lev	els of 53BP1 protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium ı	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/5	0 - 1/200)		
Gene Symbol	TP53	BP1			
Alternative Na	ames Tum	or suppressor p53-bind	ling protein 1; 53BP1; p53-binding protein 1; p53BP1		
Entrez Gene	7158	3 (Human)			
SwissProt	Q128	888 (Human); P70399 (Mouse)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ery aliquot 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

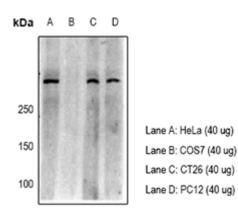
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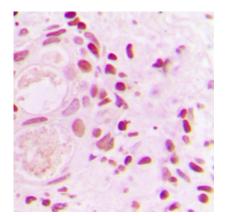


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Western blot analysis of 53BP1 expression in HeLa (A), COS7 (B), CT26 (C), PC12 (D) whole cell lysates. (Predicted band size: 213; 214 kD; Observed band size: 255 kD)



Immunohistochemical analysis of 53BP1 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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