

Product Data Sheet

Anti-USP50 Antibody

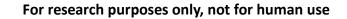
Catalog #	Source	Reactivity	Applications		
CPA5583	Rabbit	H, M, R	WB, IH		
Description	I	Rabbit polyclonal antibody to	0 USP50		
Immunogen	I	KLH-conjugated synthetic pe	otide encompassing a sequence within the C-term		
	ı	region of human USP50. The	exact sequence is proprietary.		
Purification	-	The antibody was purified by	immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	ls of USP50 protein.		
Clonality	I	Polyclonal			
Conjugation					
Form	l	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	ι	USP50			
Alternative Na	ames l	Inactive ubiquitin carboxyl-te	erminal hydrolase 50; Inactive ubiquitin-specific		
	i	peptidase 50			
Entrez Gene	ŝ	373509 (Human); 75083 (Mo	use)		
SwissProt	(Q70EL3 (Human); Q6P8X6 (N	1ouse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	freeze/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

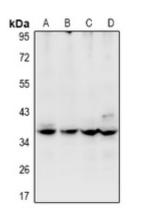
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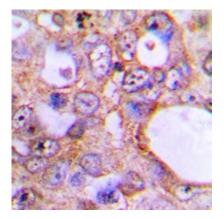




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Western blot analysis of USP50 expression in MCF7 (A), Hela (B), PC12 (C), CT26 (D) whole cell lysates. (Predicted band size: 38 kD; Observed band size: 38 kD)



Immunohistochemical analysis of USP50 staining in human lung 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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