

Product Data Sheet

Anti-Cytokeratin 8 Antibody

Catalog #	Source	Reactivity	Applications	
CPA3251	Rabbit	H, M, R	WB, IH, IF/IC	
Description	Rabbit polyclonal antibody to Cytokeratin 8			
Immunogen	К	LH-conjugated synthetic pe	ptide encompassing a sequence within the center	
	re	egion of human Cytokeratir	a 8. The exact sequence is proprietary.	
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.	
Specificity	R	ecognizes endogenous leve	els of Cytokeratin 8 protein.	
Clonality	P	olyclonal		
Form	Li	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	a	nd 0.01% sodium azide.		
Dilution	W	VB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	К	RT8		
Alternative Na	ames C	YK8; Keratin, type II cytosk	eletal 8; Cytokeratin-8; CK-8; Keratin-8; K8; Type-II	
	ke	eratin Kb8		
Entrez Gene	3	856 (Human); 16691 (Mou	se); 25626 (Rat)	
SwissProt	P	05787 (Human); P11679 (N	1ouse); Q10758 (Rat)	
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	fr	reeze/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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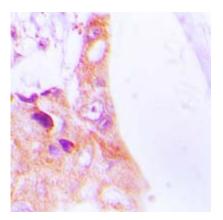
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kDa A B C

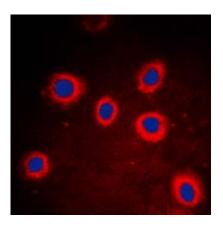
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Western blot analysis of Cytokeratin 8 expression in DLD (A), mouse muscle (B), rat muscle (C) whole cell lysates.



Immunohistochemical analysis of Cytokeratin 8 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.



Immunofluorescent analysis of Cytokeratin 8 staining in DLD 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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