

Product Data Sheet

Anti-Cytokeratin 8 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1659	Rabbit	H, M, R, B, D, P, Rb	WB, IH, IF/IC		
Description	Rabb	Rabbit polyclonal antibody to Cytokeratin 8			
Immunogen	KLH-	conjugated synthetic peptide ei	ncompassing a sequence within the C-term		
	regio	n of human Cytokeratin 8. The	exact sequence is proprietary.		
Purification	The a	antibody was purified by immu	nogen affinity chromatography.		
Specificity	Reco	gnizes endogenous levels of Cy	cokeratin 8 protein.		
Clonality	Polyc	lonal			
Conjugation					
Form	Liqui	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/100 - 1/200)), IF/IC (1/100 - 1/500)		
Gene Symbol	KRT8				
Alternative Na	ames CYK8	; Keratin, type II cytoskeletal 8;	Cytokeratin-8; CK-8; Keratin-8; K8; Type-II		
	kerat	in Kb8			
Entrez Gene	3856	(Human); 16691 (Mouse); 256	26 (Rat)		
SwissProt P0578		37 (Human); P11679 (Mouse); Q10758 (Rat)			
Storage/Stabi	lity Ship	oed at 4°C. Upon delivery alique	ot 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

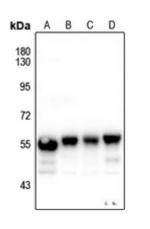
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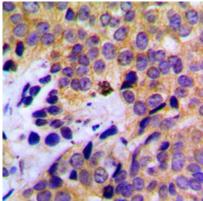


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Western blot analysis of Cytokeratin 8 expression in HCT116 (A), Hela (B), PC12 (C), AML12 (D) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 54 kD)



Immunohistochemical analysis of Cytokeratin 8 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 Cytokeratin 8 staining in NIH3T3 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.

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