

Product Data Sheet

Anti-MMP13 Antibody

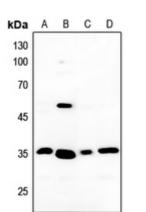
Catalog #	Source	Reactivity	Applications
CPA1756	Rabbit	H, M, R, Mk	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody to	0 MMP13
Immunogen	KLH-	conjugated synthetic pe	otide encompassing a sequence within the N-term
	regio	on of human MMP13. Th	e exact sequence is proprietary.
Purification	The	antibody was purified by	immunogen affinity chromatography.
Specificity	Recc	gnizes endogenous leve	ls of MMP13 protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium pł	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	1/500 - 1/1000), IH (1/10) - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	MM	213	
Alternative Na	ames Colla	genase 3; Matrix metall	oproteinase-13; MMP-13
Entrez Gene	4322	: (Human); 17386 (Mous	e)
SwissProt	P454	52 (Human); P33435 (N	ouse); P23097 (Rat)
Storage/Stabi	lity Ship	oed at 4°C. Upon deliver	y 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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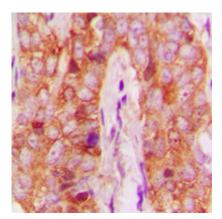
Coherion



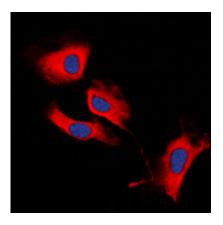
For research purposes only, not for human use

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Western blot analysis of MMP13 expression in HEK293T (A), Hela (B), mouse lung (C), rat lung (D) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 53; 34 kD)



Immunohistochemical analysis of MMP13 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 MMP13 staining in HeLa 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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