

Product Data Sheet

Anti-Carbonic Anhydrase 1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6208	Rabbit	Η, Μ	WB, IH		
Description		Rabbit polyclonal antibody to Carbonic Anhydrase 1			
Immunogen		KLH-conjugated synthetic pe	eptide encompassing a sequence within the center		
		region of human Carbonic A	nhydrase 1. The exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of Carbonic Anhydrase 1 protein.		
Clonality	Polyclonal				
Conjugation					
Form		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 (1/50	- 1/100)		
Gene Symbol		CA1			
Alternative Na	ames	Carbonic anhydrase 1; Carbo	onate dehydratase I; Carbonic anhydrase B; CAB;		
		Carbonic anhydrase I; CA-I			
Entrez Gene		759 (Human); 12346 (Mous	e)		
SwissProt		P00915 (Human); P13634 (N	/louse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
		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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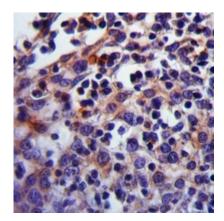
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Western blot analysis of Carbonic Anhydrase 1 expression in Myla2059 (A), Raw264.7 (B) whole cell lysates. (Predicted band size: 28 kD; Observed band size: 30 kD)



Immunohistochemical analysis of Carbonic Anhydrase 1 staining in human tonsil 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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