

### **Product Data Sheet**

## Anti-Carbonic Anhydrase 2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1120	Rabbit	H, R, Mk	WB, IH		
Description	F	Rabbit polyclonal antibody to Carbonic Anhydrase 2			
Immunogen	k	KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	r	region of human Carbonic A	hydrase 2. The exact sequence is proprietary.		
Purification	Т	The antibody was purified by	r immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of Carbonic Anhydrase 2 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	WB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	C	CA2			
Alternative Na	ames (	Carbonic anhydrase 2; Carbo	nate dehydratase II; Carbonic anhydrase C; CAC;		
	C	Carbonic anhydrase II; CA-II			
Entrez Gene	7	760 (Human); 54231 (Rat)			
SwissProt	F	200918 (Human); P27139 (R	at)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	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

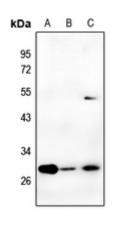
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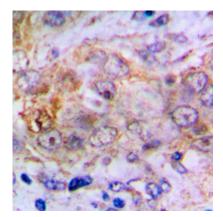
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Western blot analysis of Carbonic Anhydrase 2 expression in HEK293T (A), HCT116 (B), rat brain (C) whole cell lysates. (Predicted band size: 29 kD; Observed band size: 29 kD)



Immunohistochemical analysis of Carbonic Anhydrase 2 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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