

Product Data Sheet

Anti-Pepsinogen C Antibody

Catalog #	Source	Reactivity	Applications		
CPA1881	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	Rabbit polyclonal antibody to Pepsinogen C			
Immunogen	KLI	H-conjugated synthetic p	eptide encompassing a sequence within the center		
	reg	region of human Pepsinogen C. The exact sequence is proprietary.			
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous lev	els of Pepsinogen C protein.		
Clonality	Po	Polyclonal			
Conjugation					
Form	Liq	juid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	d 0.01% sodium azide.			
Dilution	WE	B (1/500 - 1/1000), IH (1/50) - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	PG	C			
Alternative Na	ames Ga	stricsin; Pepsinogen C			
Entrez Gene	52	25 (Human); 109820 (Mo	use); 24864 (Rat)		
SwissProt	P20	0142 (Human); Q9D7R7 (Mouse); P04073 (Rat)		
Storage/Stabi	lity Shi	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eze/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

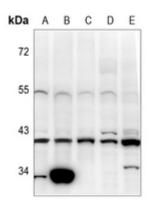
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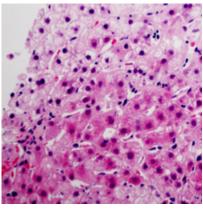
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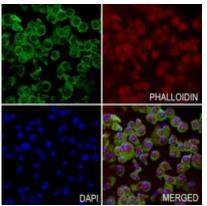


Western blot analysis of Pepsinogen C expression in HepG2 (A), LO2 (B), SGC7901 (C), AML12 (D), PC12 (E) whole cell lysates. (Predicted band size: 42 kD; Observed band size: 42; 34 kD)

Immunohistochemical analysis of Pepsinogen C staining in human liver 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. AEC was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX. Immunofluorescent analysis of Pepsinogen C staining in PANC1 cells. Formalin-fixed cells were permeabilized with 0.1% Triton



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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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