

# **Product Data Sheet**

### Anti-Chromogranin A Antibody

Catalog #	Source	Reactivity	Applications		
CPA1229	Rabbit	H, M, R, B, Mk	WB, IH		
Description		Rabbit polyclonal antibody to Chromogranin A			
Immunogen		KLH-conjugated synthetic pep	tide encompassing a sequence within the center		
		region of human Chromogran	in A. The exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels	of Chromogranin A protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium pho	osphate, 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		CHGA			
Alternative Na	ames	Chromogranin-A; CgA; Pituita	ry secretory protein I; SP-I		
Entrez Gene		1113 (Human); 12652 (Mouse	)		
SwissProt		P10645 (Human); P26339 (Mo	ouse); P10354 (Rat)		
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

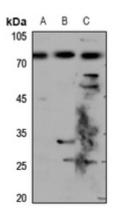
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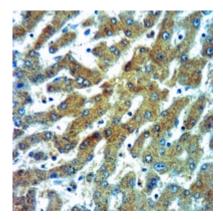


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Western blot analysis of Chromogranin A expression in H1688 (A), mouse lung (B), rat lung (C) whole cell lysates. (Predicted band size: 50 kD; Observed band size: 86 kD)



Immunohistochemical analysis of Chromogranin A 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. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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