

Product Data Sheet

Anti-Cathepsin B Antibody

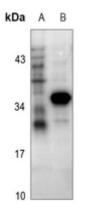
Catalog #	Source	Reactivity	Applications			
CPA4890	Rabbit	H, M, R, Mk	WB, IH			
Description	Rabb	Rabbit polyclonal antibody to Cathepsin B				
Immunogen	KLH-0	conjugated synthetic pep	tide encompassing a sequence within the center			
	regio	region of human Cathepsin B. The exact sequence is proprietary.				
Purification	The a	ntibody was purified by	immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous level	s of Cathepsin B protein.			
Clonality	Polyc	lonal				
Conjugation						
Form	Liqui	d in 0.42% Potassium ph	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	CTSB					
Alternative Na	ames CPSB	; Cathepsin B; APP secre	tase; APPS; Cathepsin B1			
Entrez Gene	1508	(Human); 13030 (Mouse	e)			
SwissProt	P078	58 (Human); P10605 (M	ouse); P00787 (Rat)			
Storage/Stabi	lity Shipp	ed at 4°C. Upon delivery	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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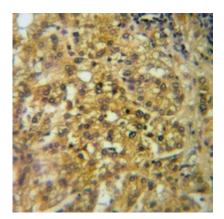
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For research purposes only, not for human use

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Western blot analysis of Cathepsin B expression in HCT116 (A), rat lung (B) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 25; 38 kD)



Immunohistochemical analysis of Cathepsin B 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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