

# **Product Data Sheet**

### **Anti-Cathepsin L Antibody**

Catalog #	Source	Reactivity	Applications		
CPA5676	Rabbit	H, M, R	WB, IH		
Description	Rab	Rabbit polyclonal antibody to Cathepsin L			
Immunogen	KLH	-conjugated synthetic pe	ptide encompassing a sequence within the center		
	regi	on of human Cathepsin	. The exact sequence is proprietary.		
Purification	The	antibody was purified b	y immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous leve	els of Cathepsin L protein.		
Clonality	Poly	/clonal			
Conjugation					
Form	Liqu	uid in 0.42% Potassium p	hosphate, 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	CTS	L1			
Alternative Na	ames CTS	L1; Cathepsin L1; Cather	sin L; Major excreted protein; MEP		
Entrez Gene	151	1514 (Human); 13039 (Mouse); 25697 (Rat)			
SwissProt	P07	711 (Human); P06797 (N	1ouse); P07154 (Rat)		
Storage/Stabi	<b>lity</b> Ship	oped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	free	ze/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

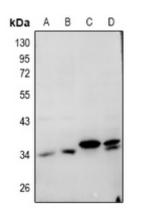
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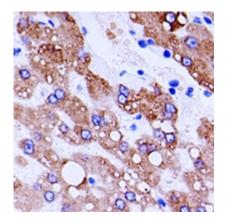


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Western blot analysis of Cathepsin L expression in C6 (A), CT26 (B), U87MG (C), A549 (D) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 35 kD)



Immunohistochemical analysis of Cathepsin L staining in human liver 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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