

Product Data Sheet

Anti-Cathepsin L HC Antibody

Catalog #	Source	Reactivity	Applications			
CPA3545	Rabbit	H, M, R, C, Mk	WB, IF/IC			
Description	Ral	Rabbit polyclonal antibody to Cathepsin L HC				
Immunogen	KLł	KLH-conjugated synthetic peptide encompassing a sequence within the center				
	reg	region of human Cathepsin L HC. The exact sequence is proprietary.				
Purification	The	e antibody was purified by	immunogen affinity chromatography.			
Specificity	ecificity Recognizes endogenous levels of Cathepsin L HC protein.					
Clonality	Pol	lyclonal				
Conjugation						
Form	Liq	uid in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.				
Dilution	WE	3 (1/500 - 1/1000), IF/IC (1/5	0 - 1/200)			
Gene Symbol	CTS	SL1				
Alternative Na	ames CTS	SL1; Cathepsin L1; Catheps	in L; Major excreted protein; MEP			
Entrez Gene	15:	14 (Human); 13039 (Mouse	e); 25697 (Rat)			
SwissProt	PO	7711 (Human); P06797 (Mo	ouse); P07154 (Rat)			
Storage/Stabi	lity Shi	pped at 4°C. Upon delivery	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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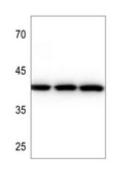
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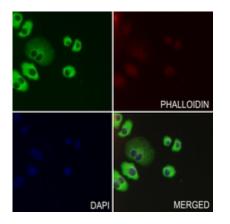
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kDa A B C



Western blot analysis of Cathepsin L HC expression in U87MG (A), A549 (B), HepG2 (C) whole cell lysates. (Predicted band size: 37 kD; Observed band size: 42 kD)



Immunofluorescent analysis of Cathepsin L HC staining in SGC7901 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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