

Product Data Sheet

Anti-Cathepsin D HC Antibody

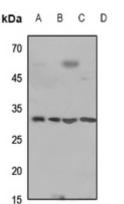
Catalog #	Source	Reactivity	Applications		
CPA1298	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to Cathepsin D HC			
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
		region of human Cathepsin	D HC. The exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of Cathepsin D HC protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid 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		CTSD			
Alternative N	ames	CPSD; Cathepsin D			
Entrez Gene		1509 (Human); 13033 (Mou	se)		
SwissProt		P07339 (Human); P18242 (N	1ouse); P24268 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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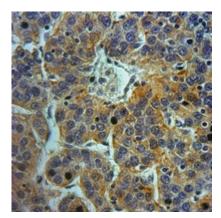




For research purposes only, not for human use

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Western blot analysis of Cathepsin D HC expression in mouse kidney (A), mouse heart (B), rat kidney (C), rat heart (D) whole cell lysates. (Predicted band size: 44 kD; Observed band size: 27 kD)



Immunohistochemical analysis of Cathepsin D HC 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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