

## **Product Data Sheet**

### **Anti-ADAMTS7 Antibody**

Catalog #	Source	e Reactivity	Applications		
CPA3754	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to ADAMTS7			
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
		region of human ADAMTS7.	The exact sequence is proprietary.		
Purification		The antibody was purified by	v immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of ADAMTS7 protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 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		ADAMTS7			
Alternative Na	ames	A disintegrin and metallopro	teinase with thrombospondin motifs 7; ADAM-TS 7;		
		ADAM-TS7; ADAMTS-7; COM	IPase		
Entrez Gene		11173 (Human); 108153 (Mouse); 315879 (Rat)			
SwissProt	Prot Q9UKP4 (Human); Q68SA9 (Mouse); Q1EHB3 (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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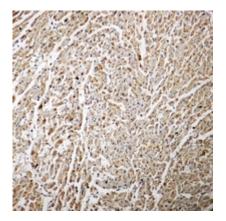
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Western blot analysis of ADAMTS7 expression in HEK293T (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 184 kD; Observed band size: 185 kD)



Immunohistochemical analysis of ADAMTS7 staining in human heart 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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