

Product Data Sheet

Anti-Mast Cell Tryptase Antibody

Catalog #	Source	Reactivity	Applications		
CPA5731	Rabbit	H, M, R	WB, IH		
Description	I	Rabbit polyclonal antibody to Mast Cell Tryptase			
Immunogen	I	KLH-conjugated synthetic pe	otide encompassing a sequence within the C-term		
	I	region of human Mast Cell Ti	yptase. The exact sequence is proprietary.		
Purification	-	The antibody was purified by	immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	ls of Mast Cell Tryptase protein.		
Clonality	I	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	i	and 0.01% sodium azide.			
Dilution	,	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	-	TPSAB1; TPSB2			
Alternative Na	ames	TPSAB1; TPS1; TPS2; TPSB1;	Tryptase alpha/beta-1; Tryptase-1; Tryptase I; Tryptase		
	i	alpha-1; TPSB2; TPS2; Trypta	se beta-2; Tryptase-2; Tryptase II		
Entrez Gene	•	7177 (Human); 100503895 (Mouse); 54271, 29268 (Rat)		
SwissProt		Q15661, P20231 (Human); C	02844, P21845 (Mouse); P27435, P50343 (Rat)		
Storage/Stabil	lity	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
	t	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

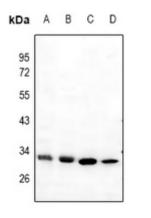
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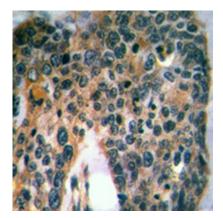


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Western blot analysis of Mast Cell Tryptase expression in MCF7 (A), mouse heart (B), PMVEC (C), A375 (D) whole cell lysates. (Predicted band size: 30 kD; Observed band size: 31 kD)



Immunohistochemical analysis of Mast Cell Tryptase staining in human lung 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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