

Product Data Sheet

Anti-Granzyme A Antibody

Catalog #	Source	Reactivity	Applications		
CPA5332	Rabbit	H, M, R	WB, IH, IF/IC		
Description	R	Rabbit polyclonal antibody to Granzyme A			
Immunogen	К	LH-conjugated synthetic p	eptide encompassing a sequence within the center		
	re	egion of human Granzyme	A. The exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous lev	els of Granzyme A protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form	Li	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/2000), IH (1/50) - 1/200), IF/IC (1/50 - 1/100)		
Gene Symbol	G	ZMA			
Alternative Na	a <mark>mes</mark> C	TLA3; HFSP; Granzyme A;	CTL tryptase; Cytotoxic T-lymphocyte proteinase 1;		
	F	ragmentin-1; Granzyme-1;	Hanukkah factor; H factor; HF		
Entrez Gene	3	001 (Human); 14938 (Μοι	ise)		
SwissProt	Р	12544 (Human); P11032 (I	Mouse)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fr	eeze/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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Product Data Sheet

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 A
 B

 250
 130

 130

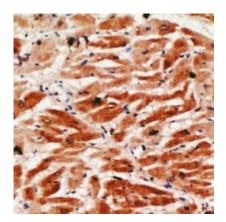
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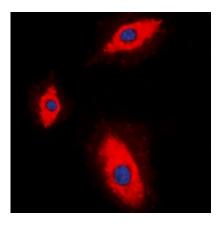
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Western blot analysis of Granzyme A expression in mouse blood (A), rat blood (B) whole cell lysates. (Predicted band size: 27; 28 kD; Observed band size: 28 kD)



Immunohistochemical analysis of Granzyme A 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.



Immunofluorescent analysis of Granzyme A staining in HepG2 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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