

Product Data Sheet

Anti-Myeloperoxidase 89k Antibody

Catalog #	Source	Reactivity	Applications			
CPA5685	Rabbit	Н, М	WB, IF/IC			
Description	Rabb	Rabbit polyclonal antibody to Myeloperoxidase 89k				
Immunogen	KLH-	conjugated synthetic p	eptide encompassing a sequence within the N-term			
	regio	region of human Myeloperoxidase 89k. The exact sequence is proprietary.				
Purification	Thea	antibody was purified l	oy immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous lev	els of Myeloperoxidase 89k protein.			
Clonality	Poly	clonal				
Conjugation						
Form	Liqui	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), IF/IC	(1/50 - 1/200)			
Gene Symbol	MPC	I				
Alternative Na	ames Mye	loperoxidase; MPO				
Entrez Gene	4353	(Human); 17523 (Mou	use)			
SwissProt	P051	.64 (Human); P11247 (Mouse)			
Storage/Stabi	lity Ship	oed at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid			
	freez	e/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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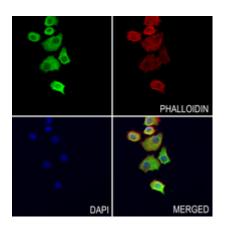
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Western blot analysis of Myeloperoxidase 89k expression in Jurkat (A), K562 (B), SP20 (C) whole cell lysates. (Predicted band size: 83 kD; Observed band size: 89 kD)



Immunofluorescent analysis of Myeloperoxidase 89k 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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