

Product Data Sheet

Recombinant Anti-PSMC5 Rabbit mAb

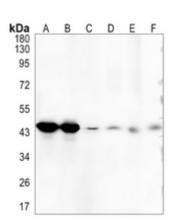
Catalog #	Source	e Reactivity	Applications		
CMA4566	Rabbit	H, M, R	WB, IH		
Description		Recombinant rabbit monoclonal antibody to PSMC5			
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within human PSMC5		
		protein. The exact sequence	is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	ls of PSMC5 protein		
Clonality		Monoclonal			
Conjugation					
Form		Liquid in PBS containing 50%	glycerol, 0.2% BSA and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol		PSMC5			
Alternative Names		SUG1; 26S protease regulatory subunit 8; 26S proteasome AAA-ATPase subunit			
		RPT6; Proteasome 26S subu	nit ATPase 5; Proteasome subunit p45; Thyroid hormone		
		receptor-interacting protein	1; TRIP1; p45/SUG		
Entrez Gene		5705 (Human); 19184 (Mouse); 81827 (Rat)			
SwissProt		P62195 (Human); P62196 (M	louse); P62198 (Rat)		
Storage/Stabi	lity	Shipped at 4 $^\circ$ C. Upon delive	ery aliquot and store at -20 $^\circ$ 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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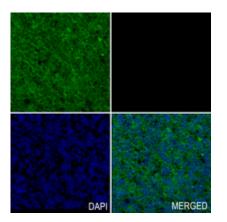
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For research purposes only, not for human use

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Western blot analysis of PSMC5 expression in Jurkat (A), K562 (B), mouse kidney (C), mouse muscle (D), rat kidney (E), rat muscle (F) whole cell lysates. (Predicted band size: 45 kD; Observed band size: 46 kD)



Immunohistochemical analysis of PSMC5 staining in human tonsil 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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