

Product Data Sheet

Recombinant Anti-CD45 Rabbit mAb

Catalog #	Source	Reactivity	Applications
CMA1004	Rabbit	Н	WB, IH, IF/IC
Description	Re	ecombinant rabbit mon	oclonal antibody to CD45
Immunogen	Re	ecombinant fusion prote	ein of human CD45. The exact sequence is proprietary.
Purification	Tł	ne antibody was purified	d by immunogen affinity chromatography.
Specificity	Re	ecognizes endogenous l	evels of CD45 protein.
Clonality	М	Ionoclonal	
Conjugation			
Form	Lie	quid in PBS, pH 7.3, 50%	6 glycerol, and 0.05% Proclin300.
Dilution	W	/B (1/500 - 1/1000), IH (1	/100 - 1/500), IF/IC (1/50 - 1/200)
Gene Symbol	PT	TPRC	
Alternative Na	ames CI	D45; Receptor-type tyro	sine-protein phosphatase C; Leukocyte common antigen;
	L-	CA; T200; CD45	
Entrez Gene	57	788 (Human)	
SwissProt	PC	08575 (Human)	
Storage/Stabi	lity Sł	nipped at 4°C. Upon deli	very aliquot and store at -20°C for one year. Avoid
	fre	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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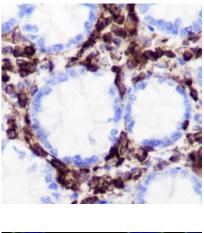
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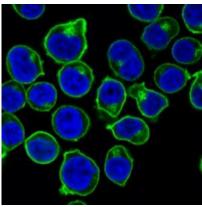
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Western blot analysis of CD45 expression in Jurkat (A) whole cell lysates. (Predicted band size: 147 kD; Observed band size: 250 kD)



Immunohistochemical analysis of CD45 staining in human colon 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 CD45 staining in Jurkat 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 humidified chamber. Cells were washed with PBST and incubated with an AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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