

Product Data Sheet

Anti-CD134 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4281	Rabbit	H, M, R	WB, IF/IC		
Description	Ra	abbit polyclonal antibody	to CD134		
Immunogen	KL	_H-conjugated synthetic p	eptide encompassing a sequence within the center		
	re	gion of human CD134. Th	e exact sequence is proprietary.		
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of CD134 protein.		
Clonality	Pc	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.			
Dilution	W	/B (1/500 - 1/1000), IF/IC (1	/50 - 1/200)		
Gene Symbol	T	NFRSF4			
Alternative Na	ames TX	GP1L; Tumor necrosis fac	tor receptor superfamily member 4; ACT35 antigen;		
	O)	X40L receptor; TAX transc	riptionally-activated glycoprotein 1 receptor; CD134		
Entrez Gene	72	293 (Human); 22163 (Mou	se); 25572 (Rat)		
SwissProt	P2	43489 (Human); P47741 (I	Mouse); P15725 (Rat)		
Storage/Stabi	lity Sł	nipped at 4°C. Upon delive	ry 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

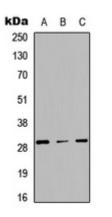
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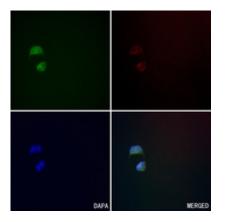


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Western blot analysis of CD134 expression in HEK293T (A), Raw264.7 (B), H9C2 (C) whole cell lysates. (Predicted band size: 29 kD; Observed band size: 30 kD)



Immunofluorescent analysis of CD134 staining in A549 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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