

Product Data Sheet

Anti-CD234 Antibody

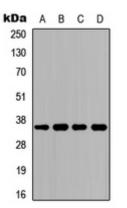
Catalog #	Source	Reactivity	Applications		
CPA3554	Rabbit	H, M, R	WB, IF/IC		
Description	Ra	Rabbit polyclonal antibody to CD234			
Immunogen	KL	-H-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	re	gion of human CD234. The	e exact sequence is proprietary.		
Purification	Th	ie antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of CD234 protein.		
Clonality	Рс	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/	100 - 1/500)		
Gene Symbol	DA	ARC			
Alternative Na	ames AC	CKR1; FY; GPD; Duffy antig	en/chemokine receptor; Atypical chemokine receptor 1;		
	Fy	glycoprotein; GpFy; Glyco	protein D; Plasmodium vivax receptor; CD234		
Entrez Gene	25	532 (Human); 13349 (Mou	se)		
SwissProt	Q	16570 (Human); Q9QUI6 (Mouse)		
Storage/Stabi	lity Sh	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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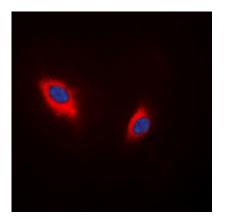




For research purposes only, not for human use

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Western blot analysis of CD234 expression in HeLa (A), K562 (B), Raw264.7 (C), PC12 (D) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 35 kD)



Immunofluorescent analysis of CD234 staining in HeLa 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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