

Product Data Sheet

Anti-CD39 Antibody

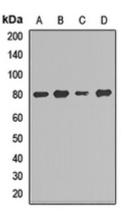
Catalog #	Source	Reactivity	Applications		
CQA2933	Rabbit	H <i>,</i> M <i>,</i> R	WB, IH, IF/IC		
Description	F	Rabbit polyclonal antibody t	o CD39		
Immunogen	ł	KLH-conjugated synthetic pe	ptide of human CD39		
Purification	٦	The antibody was purified b	<i>immunogen affinity chromatography.</i>		
Specificity	F	Recognizes endogenous leve	ls of CD39 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form	l	Liquid in 0.42% Potassium p	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	E	ENTPD1			
Alternative Names		CD39; Ectonucleoside triphosphate diphosphohydrolase 1; NTPDase 1; Ecto-ATP			
	C	diphosphohydrolase 1; Ecto-	ATPDase 1; Ecto-ATPase 1; Ecto-apyrase; Lymphoid cell		
	ä	activation antigen; CD39			
Entrez Gene	g	953 (Human); 12495 (Mouse	e); 64519 (Rat)		
SwissProt	F	P49961 (Human); P55772 (N	louse); P97687 (Rat)		
Storage/Stabi	ility S	Shipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	f	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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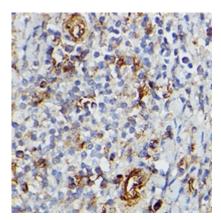




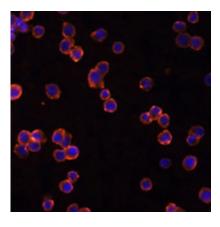
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Western blot analysis of CD39 expression in A549 (A), MCF7 (B), mouse lung (C), rat kidney (D) whole cell lysates. (Predicted band size: 34; 42; 46; 57; 58; 59 kD; Observed band size: 80 kD)



Immunohistochemical analysis of CD39 staining in human lung 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 CD39 staining in Raw264.7 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 a AF594-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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