

# **Product Data Sheet**

# Anti-CD27 Antibody

Catalog #	Source	Reactivity	Applications		
CQA1876	Rabbit	H, M, R	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody t	o CD27		
Immunogen	I	Recombinant full length pro	tein of human CD27		
Purification	-	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	els of CD27 protein.		
Clonality	I	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	v	WB (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	(	CD27			
Alternative Na	ames	TNFRSF7; CD27 antigen; CD2	27L receptor; T-cell activation antigen CD27; T14; Tumor		
	I	necrosis factor receptor sup	erfamily member 7; CD27		
Entrez Gene	9	939 (Human); 21940 (Mouse	2)		
SwissProt	I	P26842 (Human); P41272 (N	1ouse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	t	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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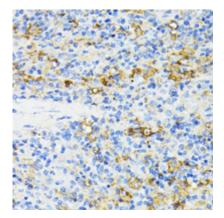
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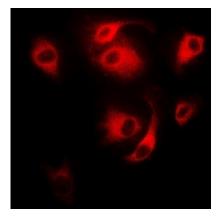
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Western blot analysis of CD27 expression in Raji (A), HL60 (B), mouse skeletal muscle (C) whole cell lysates. (Predicted band size: 29 kD; Observed band size: 55 kD)



Immunohistochemical analysis of CD27 staining in rat spleen 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 CD27 staining in MCF7 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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