

## **Product Data Sheet**

## **Anti-CD29 Antibody**

Catalog #	Source	Reactivity	Applications	
CPA7148	Rabbit	H, M, R, B, C, P, S	WB, IH, IF/IC	
Description	R	Rabbit polyclonal antibody to (	CD29	
Immunogen	K	(LH-conjugated synthetic pept	ide encompassing a sequence within the C-term	
	r	egion of human CD29. The exa	act sequence is proprietary.	
Purification	Т	The antibody was purified by in	nmunogen affinity chromatography.	
Specificity	R	Recognizes endogenous levels	of CD29 protein.	
Clonality	Р	Polyclonal		
Conjugation				
Form	L	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	а	and 0.01% sodium azide.		
Dilution	V	NB (1/500 - 1/1000), IH (1/100	) - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	LI.	TGB1		
Alternative Na	ames F	NRB; MDF2; MSK12; Integrin	beta-1; Fibronectin receptor subunit beta;	
	G	Glycoprotein IIa; GPIIA; VLA-4	subunit beta; CD29	
Entrez Gene	3	3688 (Human); 16412 (Mouse)	; 24511 (Rat)	
SwissProt	Р	205556 (Human); P09055 (Mo	use); P49134 (Rat)	
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid	
	fı	reeze/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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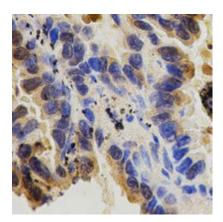
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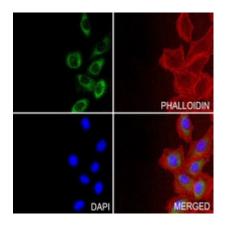
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KDA A B C 130 95 72

Western blot analysis of CD29 expression in U87MG (A), A375 (B), rat skin (C) whole cell lysates. (Predicted band size: 88 kD; Observed band size: 115 kD)



Immunohistochemical analysis of CD29 staining in human lung cancer 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 CD29 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 hidified chamber. Cells were washed with PBST and incubated with an 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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