

# **Product Data Sheet**

### **Anti-CD49c Antibody**

Catalog #	Source	e Reactivity	Applications		
CPA1611	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody t	o CD49c		
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
		region of human CD49c. The	exact sequence is proprietary.		
Purification		The antibody was purified by	/ immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of CD49c protein.			
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol		ITGA3			
Alternative N	ames	MSK18; Integrin alpha-3; CD	49 antigen-like family member C; FRP-2; Galactoprotein		
		B3; GAPB3; VLA-3 subunit al	pha; CD49c		
Entrez Gene		3675 (Human); 16400 (Mouse)			
SwissProt P26006 (Human); Q62470 (Mouse)		P26006 (Human); Q62470 (N	Nouse)		
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
		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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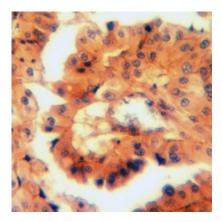
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**KDa** <u>A</u> <u>B</u> <u>C</u> <u>D</u> 180 130 95 Western blot analysis of CD49c expression in HepG2 (A), AML12 (B), PC12 (C), A549 (D) whole cell lysates. (Predicted band size: 116 kD; Observed band size: 125 kD)



Immunohistochemical analysis of CD49c 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.

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