

# **Product Data Sheet**

## Anti-CD49e Antibody-PE labled

e Reactivity	Applications	
Μ	IF, FC	
Rat monoclonal antibody PE labled to CD49e		
Affinity-purified mouse VLA-5 protein ( $\alpha$ 5 $\beta$ 1, CD49e/CD29)		
The antibody was purified by affinity chromatography.		
Recognizes mouse, rat CD49e		
Monoclonal (clone: 5H10-27)		
PE		
Rat IgG2a kappa. Liquid in PBS, pH 7.3, 0.2% BSA, and 0.02% sodium azide.		
10 μl / assay		
ITGA5		
FNRA; Integrin alpha-5; CD49 antigen-like family member E; Fibronectin receptor		
subunit alpha; Integrin alpha-F; \	/LA-5; CD49e	
16402 (Mouse)		
P11688 (Mouse)		
1. Take 100 $\mu l$ peripheral blood anticoagulated by EDTA and add to the bottom of 5		
ml tube.		
2. Add 10 $\mu l$ labeled antibody to	the bottom of flow tube mixing with the whole	
blood, incubate for 20 minutes a	t room temperature away from light.	
3. Add 2 ml RBC lysis buffer, incu	bate for 10 minutes away from light after mixing,	
dissolve red blood cells.		
4. Sample tube is set to 1000 rpn	n centrifugation for 5 minutes, discard the	
supernatant.		
	M Rat monoclonal antibody PE lable Affinity-purified mouse VLA-5 pro The antibody was purified by affi Recognizes mouse, rat CD49e Monoclonal (clone: 5H10-27) PE Rat IgG2a kappa. Liquid in PBS, p 10 μl / assay ITGA5 FNRA; Integrin alpha-5; CD49 and subunit alpha; Integrin alpha-F; N 16402 (Mouse) P11688 (Mouse) 1. Take 100 μl peripheral blood a ml tube. 2. Add 10 μl labeled antibody to blood, incubate for 20 minutes a 3. Add 2 ml RBC lysis buffer, incu dissolve red blood cells. 4. Sample tube is set to 1000 rpm	

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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5. Add 2 ml PBS wash buffer to resuspend the cells, then 1000 rpm centrifugation for 5 minutes, discard the supernatant.

6. Add 0.5 ml PBS wash buffer to resuspend the cells and detect by flow cytometry (sample should be determined on the day on the machine and can also be added fixation overnight at 4 °C then measured).

**Storage/Stability** Shipped and store at 4°C for one year. Do not freeze.

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