

Product Data Sheet

Anti-CD44 Antibody-PE/Cy5 labled

Source	Reactivity	Applications		
Mouse	Н	IF, FC		
Description Mouse monoclonal antibody PE/Cy5 labled to CD44				
	Native purified human CD4	4.		
	The antibody was purified by affinity chromatography.			
	Recognizes human CD44			
	Monoclonal (clone: F10-44-2)			
	PE/Cy5			
	Mouse IgG2a kappa. Liquid in PBS, pH 7.3, 0.2% BSA, and 0.02% sodium azide.			
	10 μl / assay			
	CD44			
ames	LHR; MDU2; MDU3; MIC4; CD44 antigen; CDw44; Epican; Extracellular ma			
	receptor III; ECMR-III; GP90) lymphocyte homing/adhesion receptor; HUTCH-I;		
	Heparan sulfate proteoglycan; Hermes antigen; Hyaluronate receptor; Phagocytic			
	glycoprotein 1; PGP-1; Pha	go		
	960 (Human)			
	P16070 (Human)			
Use	1. Take 100 μl peripheral blood anticoagulated by EDTA and add to the bottom of 5			
	ml tube.			
	2. Add 10 μl labeled antibo	dy to the bottom of flow tube mixing with the whole		
	blood, incubate for 20 min	utes at room temperature away from light.		
	3. Add 2 ml RBC lysis buffe	, incubate for 10 minutes away from light after mixing,		
	dissolve red blood cells.			
	Mouse	MouseΗMouse monoclonal antibodNative purified human CD4The antibody was purified IRecognizes human CD44Monoclonal (clone: F10-44PE/Cy5Mouse IgG2a kappa. Liquid10 μl / assayCD44CD44LHR; MDU2; MDU3; MIC4; receptor III; ECMR-III; GP90Heparan sulfate proteogly0glycoprotein 1; PGP-1; Phag960 (Human)P16070 (Human)Use1. Take 100 μl peripheral bl ml tube.2. Add 10 μl labeled antibo blood, incubate for 20 minu 3. Add 2 ml RBC lysis buffer		

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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4. Sample tube is set to 1000 rpm centrifugation for 5 minutes, discard the supernatant.

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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