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# Product datasheet for AM31884FC-L

# CD11a / ITGAL Rat Monoclonal Antibody [Clone ID: IBL-6/2]

## **Product data:**

Product Type:	Primary Antibodies
Clone Name:	IBL-6/2
Applications:	FC
Recommended Dilution:	This clone has been reported to work in immunohistochemistry (acetone-fixed frozen sections), flow cytometry and immunoprecipitation. The FITC conjugated format is especially useful for direct flow cytometry.
Reactivity:	Mouse
Host:	Rat
lsotype:	lgG1
Clonality:	Monoclonal
Immunogen:	EL4 (mouse thymoma cell line) <u>Donor:</u> Wistar spleen <u>Fusion Partner:</u> Sp-2/0Ag14
Specificity:	Anti-mouse CD11a monoclonal antibody (Clone: IBL-6/2) recognizes the LFA-1 (CD11a) antigen in mice. The LFA-1 is expressed on leukocytes and it mediates cell-cell and cell-adhesion by binding to CD54 and ICAM-2).
Formulation:	PBS containing 0.02% sodium azide (NaN3) as preservative and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml. Label: FITC State: Liquid purified Ig fraction Label: Fluorescein isothiocyanate isomer 1
Concentration:	lot specific
Purification:	Affinity chromatography on Protein G
Conjugation:	FITC
Storage:	Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.
Stability:	Shelf life: one year from despatch.
Database Link:	P24063



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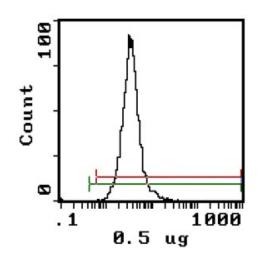
	CD11a / ITGAL Rat Monoclonal Antibody [Clone ID: IBL-6/2] – AM31884FC-L
Synonyms:	Integrin alpha-L, LFA1, LFA-1
Note:	Protocol: FLOW CYTOMETRY ANALYSIS:
	Method:
	1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell
	population with Lympholyte®-M cell separation medium.
	2. Wash 2 times.
	3. Resuspend the cells to a concentration of $2x10e7$ cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain 1 x 10e6 cells, representing 1 test).
	4. To each tube, add 0.5 – 0.2 $\mu$ g of this antibody per 10e6 cells.
	5. Vortex the tubes to ensure thorough mixing of antibody and cells.
	6. Incubate the tubes for 30 minutes at 4°C. (It is recommended that the tubes are protected
	from light, since most fluorochromes are light sensitive.)
	7. Wash 2 times at 4°C.
	8. Resuspend the cell pellet in 50 μl ice cold media B.
	9. Transfer to suitable tubes for flow cytometric analysis containing 15 µl of propidium iodide
	at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.
	Media: A Discussion to buffer and active $(n + 7, 2) + 50$ (non-model common of boot and size to active control (100)
	A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 μl of 2M sodium azide in 100 mls).
	B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 µl of
	2M sodium azide in 100 mls).
	Results:
	Tissue Distribution by Flow Cytometry Analysis:
	Mouse Strain: BALB/c
	<u>Cell Concentration:</u> 1x10e6 cells per test
	Antibody Concentration Used: 0.2 µg/10e6 cells
	Antibuly Concentration Osed. 0.2 µg/ 1000 Cells

Isotypic Control: FITC Rat IgG1

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## **Product images:**



Cell Source: Lymph Node Percentage of cells stained above control: 99.6%

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