

## TECHNICAL DATA SHEET

# PerCP-Cyanine5.5 Anti-Mouse CD11a (M17/4)

Catalog Number: 65-0111

## PRODUCT INFORMATION

**Contents:** PerCP-Cyanine5.5 Anti-Mouse CD11a (M17/4)

**Isotype:** Rat IgG2a, kappa

**Concentration:** 0.2 mg/mL

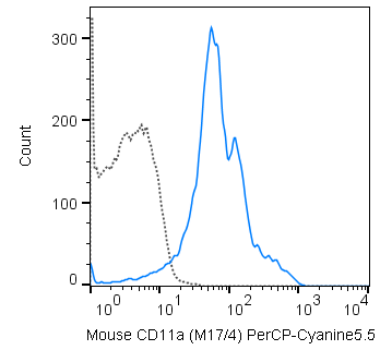
**Clone:** M17/4

**Reactivity:** Mouse

**Use By:** 6 months from date of receipt

**Storage Conditions:** 2-8°C protected from light

**Formulation:** 10 mM NaH<sub>2</sub>PO<sub>4</sub>, 150 mM NaCl, 0.09% NaN<sub>3</sub>,  
0.1% gelatin, pH7.2



C57Bl/6 splenocytes were stained with 0.25 ug PerCP-Cyanine5.5 Anti-Mouse CD11a (65-0111) (solid line) or 0.25 ug PerCP-Cyanine5.5 Rat IgG2a isotype control (dashed line).

## DESCRIPTION

The M17/4 monoclonal antibody reacts with mouse CD11a, an 180kD glycoprotein that is also known as integrin alpha L (ITGAL). CD11a associates non-covalently with CD18 (integrin beta 2) to form the lymphocyte function associated antigen-1 (LFA-1) heterodimer. CD11a is expressed on the cell surface of all leukocytes. LFA-1 plays a critical role in intercellular adhesion that is mediated through interactions with its ligands, ICAM-1 (CD54), ICAM-2 (CD102), and ICAM-3 (CD50). LFA-1 also functions in lymphocyte costimulatory signaling. On resting cells, LFA-1 exists in a low-affinity conformation that is rapidly induced to the high-affinity conformation following cell stimulation.

## PREPARATION & STORAGE

This monoclonal antibody was purified from tissue culture supernatant via affinity chromatography. The purified antibody was conjugated under optimal conditions, with unreacted dye removed from the preparation. It is recommended to store the product undiluted at 4°C, and protected from prolonged exposure to light. Do not freeze.

## APPLICATION NOTES

This antibody preparation has been quality-tested for flow cytometry using mouse spleen cells, or an appropriate cell type (where indicated). Please refer to the figure legend for the optimal concentration used to stain the tissue shown. We recommend titrating the antibody under your specific conditions to determine the optimal concentration of antibody needed in your experimental system.

## REFERENCES

Larson RS, Springer TA. 1990. Immunol Rev. 114:181-217. Sanders VM, Vitetta ES. 1991. Cell Immunol. 132(1):45-55. Kuhlman P, Moy VT, Lollo BA, Brian AA. 1991. J Immunol. 146(6):1773-1782. Springer TA. 1994. Cell. 76(2):301-314. van Kooyk Y, de Vries-van der Zwan A, de Waal LP, Figdor CG. 1994. Transplant Proc. 26(2):401-403.

Tonbo Biosciences tests all antibodies by flow cytometry. Citations are provided as a resource for additional applications that have not been validated by Tonbo Biosciences. Please choose the appropriate format for each application and consult Materials and Methods sections for additional details about the use of any product in these publications.

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