

**PerCP/Cy5.5 anti-mouse CD11a/CD18 (LFA-1)**

**Catalog # / Size:** 1305040 / 100 µg  
1305035 / 25 µg

**Clone:** H155-78

**Isotype:** Rat IgG1, κ

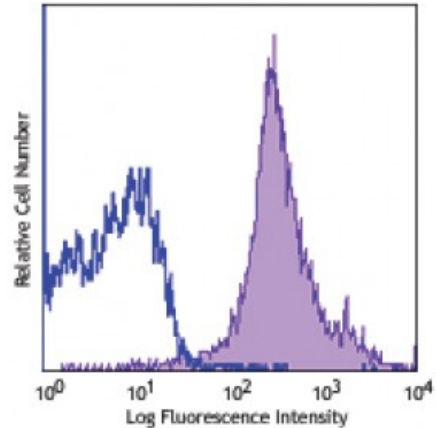
**Immunogen:** T<sub>H</sub>-derived anti-I-Ak CTL clone A1 5.1.17

**Reactivity:** Mouse

**Preparation:** The antibody was purified by affinity chromatography and conjugated with PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cy5.5 and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.

**Concentration:** 0.2



C57BL/6 splenocytes were stained with LFA-1 (clone H155-78) PerCP/Cy5.5 (filled histogram) or rat IgG1 PerCP/Cy5.5 isotype control (open histogram).

**Applications:**

**Applications:** Flow Cytometry

**Recommended Usage:** Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is ≤1.0 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

**Application Notes:** This clone is reported to have some effect on increasing LFA-1-mediated adhesion and does not block LFA-1 binding1.

**Application References:**

1. Pont S, *et al.* 1986. *J. Immunol.* 136:3750. (FC)
2. Dupont CD, *et al.* 2014. *PLoS Pathog.* 10:1004047. [PubMed](#)
3. Markey KA, *et al.* 2014. *J Immunol.* 192:5426. [PubMed](#)

**Description:** Lymphocyte function-associated antigen 1 or LFA-1 is a heterodimer composed of two members of the integrin family, α<sub>L</sub> (CD11a) and β<sub>2</sub> (CD18). LFA-1 is expressed primarily on lymphocytes, monocytes/macrophages, and granulocytes. It is involved in cell adhesion and costimulation processes. The ligands for LFA-1 are ICAM-1 (CD54), ICAM-2 (CD102), and ICAM-3 (CD50).

**Antigen References:**

1. Dugger KJ, *et al.* 2009. *J. Neuroimmunol.* 206:22.
2. Barclay A, *et al.* 1997. *The Leukocyte Antigen Facts Book* Academic Press.
3. Springer TA. 1994. *Cell* 76:301.
4. Lub M, *et al.*