

# Anti-Human/Mouse CD11b PerCP-Cy5.5

Catalogue Number: 03221-70

RUO: For Research Use Only. Not for use in diagnostic procedures.

#### **Product Information**

**Clone:** M1/70

Format/Conjugate: PerCP-Cy5.5 Concentration: 0.2 mg/mL Reactivity: Human, Mouse Laser: Blue (488nm)

**Peak Emission:** 695nm **Peak Excitation:** 482nm

Filter: 695/40

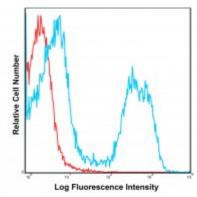
Brightness (1=dim,5=brightest): 3

Isotype: Rat IgG2b, kappa

Formulation: Phosphate-buffered aqueous solution, ≤0.09% Sodium azide, may contain carrier protein/stabilizer, ph7.2.

Storage: Product should be kept at 2-8°C and protected from prolonged exposure to light.

**Applications: FC** 



C57BI/6 bone marrow cells were stained with PerCP-Cy5.5 M1/70 with relevant isotype control

in Red.

## **Description**

The M1/70 monoclonal antibody specifically reacts with the 170 kDa αM integrin chain of mouse CD11b from the Mac-1 integrin (CD11b/CD18). Mac-1 binds to C3bi, CD54 (ICAM-1), and fibrinogen, and it is expressed by granulocytes, macrophages, NK cells, myeloid-derived dendritic cells, microglia, activated lymphocytes, and mouse B-1 cells. The expression is up-regulated on activated neutrophils at the same time that L-selectin is shed from the cell surface. The M1/70 antibody is used for the detection of monocytes, granulocytes, and a subset of natural killer cells in human peripheral blood.

M1/70 blocks C3bi binding and cell adherence, but not cell-mediated lysis and it cross-reacts with human CD11b.

### **Preparation & Storage**

The product should be stored undiluted at 4°C and should be protected from prolonged exposure to light. Do not freeze. The monoclonal antibody was purified utilizing affinity chromatography and unreacted dye was removed from the product.

## **Application Notes**

The antibody has been analyzed for quality through the flow cytometric analysis of the relevant cell type. For flow cytometric staining, the suggested use of this reagent is  $\leq 0.25$  ug per million cells in 100  $\mu$ l volume. It is recommended that the reagent be titrated for optimal performance for each application.

## References

- 1.Sanchez-Madrid, F., Simon, P., Thompson, S., ; Springer, T. A. (1983). Mapping of antigenic and functional epitopes on the alpha-and beta-subunits of two related mouse glycoproteins involved in cell interactions, LFA-1 and Mac-1.The Journal of experimental medicine,;158(2), 586-602.
- 2. Ault, K. A., ; Springer, T. A. (1981). Cross-reaction of a rat-anti-mouse phagocyte-specific monoclonal antibody (anti-Mac-1) with human monocytes and natural killer cells.; The Journal of Immunology,; 126(1), 359-364.
- 3. Springer, T., Galfre, G., Secher, D. S., ; Milstein, C. (1978). Monoclonal xenogeneic antibodies to murine cell surface antigens: identification of novel leukocyte differentiation antigens.; European journal of immunology,; 8(8), 539-551.