

Anti-Human/Mouse CD11b PE-Cyanine7

Catalog Number :03221-77

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

Product Information

Clone: M1/70

Format/Conjugate: PE-Cyanine7

Concentration: 0.2 mg/mL

Reactivity: Human, Mouse

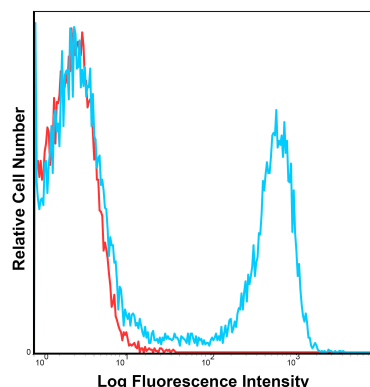
Laser: Blue (488nm), Yellow/Green (532-561nm)

Peak Emission: Not Applicable

Peak Excitation: Not Applicable

Filter: Not Applicable

Brightness (1=dim,5=brightest): Not Applicable



C57Bl/6 bone marrow cells were stained with PE-Cy7 M1/70 with relevant isotype control in Red.

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

Description

The M1/70 monoclonal antibody specifically reacts with the 170 kDa α M 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 ≤0.125 μ g per million cells in 100 μ 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.