Technical Data Sheet

APC-Cy™7 Rat Anti-Mouse CD11b

Product Information

Material Number:
Alternate Name:
Size:
Concentration:
Clone:
Immunogen:
Isotype:
Reactivity:

Storage Buffer:

Description

The M1/70 antibody reacts with the 170-kDa α [M] chain of Mac-1 (CD11b/CD18, α [M] β [2] integrin), also known as complement receptor 3 (CR3), which mediates adhesion to C3bi and ICAM-1 (CD54). Mac-1 is expressed at varying levels on granulocytes, macrophages, myeloid-derived dendritic cells, natural killer cells, microglia, and B-1 cells. Mac-1 expression is rapidly up-regulated on neutrophils after activation, in the same time period that CD62L (L-selectin) is shed from the cell surface. M1/70 antibody reportedly blocks cell adherence and C3bi binding, but it does not block cell-mediated lysis. Cross-reaction of mAb M1/70 with CD11b on human monocytes, polymorphonuclear leukocytes, and NK cells has been reported.

561039

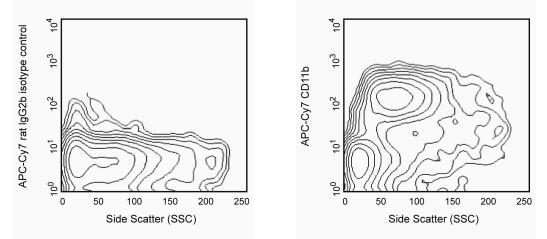
25 μg 0.2 mg/ml M1/70

Mouse Splenic Cells Rat (DA) IgG2b, κ QC Testing: Mouse

Tested in Development: Human

CR-3 alpha chain; Itgam; Integrin alpha M; Ly-40; Mac-1 alpha

Aqueous buffered solution containing $\leq 0.09\%$ sodium azide.



Expression of CD11b on bone-marrow myeloid cells. C57Bl/6 bone-marrow leukocytes were stained with either APC-Cy7-conjugated Rat (gG2b κ isotype control A95-1 (Cat. No. 552773, left panel) or APC-Cy7-conjugated M1/70 monoclonal antibodies (right panel). Please note that the population of cells having the lowest SSC (erythroid and lymphoid cells) show little expression of CD11b, while cells with moderate-to-high SSC (myeloid cells) are almost uniformly CD11b positive (right panel). Flow cytometry was performed on a BD FACSVantageTM SE flow cytometry system.

Preparation and Storage

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

The antibody was conjugated with APC-Cy7 under optimum conditions, and unconjugated antibody and free APC-Cy7 were removed.

Application Notes

Routinely Tested				
ion Products				
Name			Size	Clone
APC-Cy™7 Rat IgG2b κ Isotype Control			0.1 mg	A95-1
		· · ·		
Europe Japan	Asia Pacific	Latin America/Caribbean		e BI
nformation, visit bdbiosciences.c ed herein is not to be construed as a re not be held responsible for patent infri not include or carry any right to resell c her product. Any use of this product of	om/how_to_orde commendation to us ngement or other vie or transfer this produ ther than the permit	rr/ ie the above product in violation olations that may occur with the ict either as a stand-alone		
	APC-Cy TM 7 Rat IgG2 ary, each investigator should Europe Japan 32.53.720.550 0120.8555.90 iformation, visit bdbiosciences.c red herein is not to be construed as a re not be held responsible for patent infri not include or carry any right to resel	Name APC-Cy ^{TM7} Rat IgG2b κ Isotype Cor ary, each investigator should titrate the reag Europe Japan Asia Pacific 32.53.720.550 0120.8555.90 65.6861.0633 aformation, visit bdbiosciences.com/how_to_orde ed herein is not to be construed as a recommendation to us not include or carry any right to resell or transfer this production	ion Products <u>Name</u> APC-Cy™7 Rat IgG2b κ Isotype Control ary, each investigator should titrate the reagent to obtain optimal results. Europe Japan Asia Pacific Latin America/Caribbean	Europe Japan Asia Pacific Latin America/Caribbean 32.53.720.550 0120.8555.90 65.6861.0633 0800.771.7157 rformation, visit bdbiosciences.com/how_to_order/ ed herein is not to be construed as a recommendation to use the above product in violation not be held responsible for patent infringement or other violations that may occur with the not include or carry any right to resell or transfer this product either as a stand-alone

- 2. An isotype control should be used at the same concentration as the antibody of interest.
- 3. APC-Cy7 tandem fluorochrome emission is collected in a detector for fluorescence wavelengths of 750 nm and higher.
- 4. APC-Cy7 is a tandem fluorochrome composed of Allophycocyanin (APC), which is excited by laser lines between 595 and 647 nm and serves as an energy donor, coupled to the cyanine dye Cy7[™], which acts as an energy acceptor and fluoresces at 780 nm. BD Biosciences Pharmingen has maximized the fluorochrome energy transfer in APC-Cy7, thus maximizing its fluorescence emission intensity, minimizing residual emission from APC, and minimizing required electronic compensation in multilaser-laser flow cytometry systems. Note: Although every effort is made to minimize the lot-to-lot variation in residual emission from APC, it is strongly recommended that every lot be tested for differences in the amount of compensation required and that individual compensation controls are run for each APC-Cy7 conjugate.
- 5. Warning: Some APC-Cy7 and PE-Cy7 conjugates show changes in their emission spectrum with prolonged exposure to formaldehyde. If you are unable to analyze fixed samples within four hours, we recommend that you use BDTM Stabilizing Fixative (Cat. No. 338036).
- 6. Please observe the following precautions: Absorption of visible light can significantly alter the energy transfer occurring in any tandem fluorochrome conjugate; therefore, we recommend that special precautions be taken (such as wrapping vials, tubes, or racks in aluminum foil) to prevent exposure of conjugated reagents, including cells stained with those reagents, to room illumination.
- 7. This conjugated product is sold under license to the following patent: US Patent No. 5,714,386.
- 8. This conjugated product is sold under license to the following patents: US Patent No. 5,798,276.
- 9. Cy is a trademark of Amersham Biosciences Limited. This conjugated product is sold under license to the following patents: US Patent Nos. 5,486,616; 5,569,587; 5,569,766; 5,627,027.
- 10. This product is subject to proprietary rights of Amersham Biosciences Corp. and Carnegie Mellon University and made and sold under license from Amersham Biosciences Corp. This product is licensed for sale only for research. It is not licensed for any other use. If you require a commercial license to use this product and do not have one return this material, unopened to BD Biosciences, 10975 Torreyana Rd, San Diego, CA 92121 and any money paid for the material will be refunded.
- 11. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 12. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
- 13. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

References

Ault KA, Springer TA. Cross-reaction of a rat-anti-mouse phagocyte-specific monoclonal antibody (anti-Mac-1) with human monocytes and natural killer cells. J Immunol. 1981; 126(1):359-364. (Biology)

Beavis AJ, Pennline KJ. Allo-7: a new fluorescent tandem dye for use in flow cytometry. Cytometry. 1996; 24(4):390-395. (Biology)

Beller DI, Springer TA, Schreiber RD. Anti-Mac-1 selectively inhibits the mouse and human type three complement receptor. J Exp Med. 1982; 156(4):1000-1009. (Clone-specific: Blocking)

Kantor AB, Stall AM, Adams S, Herzenberg LA, Herzenberg LA. Differential development of progenitor activity for three B-cell lineages. Proc Natl Acad Sci U S A. 1992; 89(8):3320-3324. (Biology)

Kishimoto TK, Jutila MA, Berg EL, Butcher EC. Neutrophil Mac-1 and MEL-14 adhesion proteins inversely regulated by chemotactic factors. *Science*. 1989; 245(4923):1238-1241. (Biology)

Lagasse E, Weissman IL. Flow cytometric identification of murine neutrophils and monocytes. *J Immunol Methods*. 1996; 197(1-2):139-150. (Biology) Leenen PJ, de Bruijn MF, Voerman JS, Campbell PA, van Ewijk W. Markers of mouse macrophage development detected by monoclonal antibodies. *J Immunol Methods*. 1994; 174(1-2):5-19. (Biology)

Lodge PA, Sriram S. Regulation of microglial activation by TGF-beta, IL-10, and CSF-1. J Leukoc Biol. 1996; 60(4):502-508. (Biology) Lub M, van Kooyk Y, Figdor CG. Competition between lymphocyte function-associated antigen 1 (CD11a/CD18) and Mac-1 (CD11b/CD18) for binding to

intercellular adhesion molecule-1 (CD54). *J Leukoc Biol.* 1996; 59(5):648-655. (Biology) Roederer M, Kantor AB, Parks DR, Herzenberg LA. Cy7PE and Cy7APC: bright new probes for immunofluorescence. *Cytometry*. 1996; 24(3):191-197. (Methodology: Flow cytometry)

Sanchez-Madrid F, Simon P, Thompson S, Springer TA. 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. J Exp Med. 1983; 158(2):586-602. (Clone-specific: Blocking)

Springer T, Galfre G, Secher DS, Milstein C. Monoclonal xenogeneic antibodies to murine cell surface antigens: identification of novel leukocyte differentiation antigens. *Eur J Immunol.* 1978; 8(8):539-551. (Immunogen)

Springer T, Galfre G, Secher DS, Milstein C. Mac-1: a macrophage differentiation antigen identified by monoclonal antibody. *Eur J Immunol.* 1979; 9(4):301-306. (Clone-specific)

Springer TA, Davignon D, Ho MK, Kurzinger K, Martz E, Sanchez-Madrid F. LFA-1 and Lyt-2,3, molecules associated with T lymphocyte-mediated killing; and Mac-1, an LFA-1 homologue associated with complement receptor function. *Immunol Rev.* 1982; 68:171-195. (Clone-specific: Blocking) Vremec D, Zorbas M, Scollay R, et al. The surface phenotype of dendritic cells purified from mouse thymus and spleen: investigation of the CD8 expression by a

Vremec D, Zorbas M, Scollay R, et al. The surface phenotype of dendritic cells purified from mouse thymus and spleen: investigation of the CD8 expression by a subpopulation of dendritic cells. J Exp Med. 1992; 176(1):47-58. (Biology)