Brilliant Violet 650™ anti-mouse/human CD11b

Catalog # / Size: 1106295 / 50 μg

1106195 / 125 µl

Clone: M1/70

Isotype: Rat IgG2b, κ

Immunogen: C57BL/10 splenocytes

Reactivity: Human

Preparation: The antibody was purified by affinity

chromatography and conjugated with Brilliant Violet 650™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 650™ and

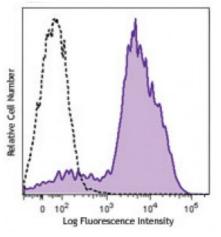
unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and BSA

(origin USA).

Concentration: test sizes: Lot-specific



C57BL/6 mouse bone marrow cells were stained with CD11b (clone M1/70) Brilliant Violet 650^{TM} . Data shown was gated on myeloid cell

population.

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 using the microg size, the suggested use of this reagent is ≤ 0.3 microg per million cells in 100 microL volume. For flow cytometric staining using the test size, the suggested use of this reagent is ≤ 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

Brilliant Violet 650™ excites at 405 nm and emits at 645 nm. The bandpass filter 660/20 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. **Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel.** Refer to your instrument manual or manufacturer for support. Brilliant Violet 650™ is a trademark of Sirigen Group Ltd.

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Application Notes:

Additional reported applications (for relevant formats of this clone) include: immunoprecipitation^{1,4}, *in vitro* blocking^{3,9,12}, depletion^{2,8}, immunofluorescence microscopy^{6,7,10}, and immunohistochemistry of acetone-fixed frozen sections^{5,11-13} and paraffin sections²⁸. The LEAF $^{\text{TM}}$ purified antibody (Endotoxin <0.1 EU/ μ g, Azide-Free, 0.2 μ m filtered) is recommended for functional assays (Cat. No. 101231). For *in vivo* studies or highly sensitive assays, we recommend Ultra-LEAF $^{\text{TM}}$ purified antibody (Cat. No. 101248) with a lower endotoxin limit than standard LEAF $^{\text{TM}}$ purified antibodies (Endotoxin <0.01 EU/microg).

Application References:

- 1. Springer T, et al. 1978. Eur. J. Immunol. 8:539. (IP)
- 2. Ault K and Springer T. 1981. *J. Immunol.* 126:359. (Deplete)
- 3. Springer TA, et al. 1982. Immunol. Rev. 68:171. (Block)
- 4. Ho MK and Springer TA. 1983. J. Biol. Chem. 258:2766. (IP)
- 5. Flotte TJ, et al. 1983. Am. J. Pathol. 111:112. (IHC)
- 6. Noel GJ, *et al.* 1990. J. Clin. Invest. 85:208. (IF)
- 7. Allen LA and Aderem A. 1996. J. Exp. Med. 184:627 (IF)
- 8. D'Amico A and Wu L. 2003. J. Exp. Med. 198:293. (Deplete)
- 9. Brickson SJ, et al. 2003. Appl Physiol. 95:969. (Block)
- 10. Clatworthy MR and Smith KG. 2004. J. Exp. Med. 199:717. (IF)
- 11. Hata H, et al. 2004. J. Clin. Invest. 114:582. (IHC)
- 12. Zhang Y, et al. 2002. J. Immunol. 168:3088. (IHC)
- 13. Iwasaki A and Kelsall BL. 2001. J. Immunol. 166:4884 (IHC, FC)
- 14. Tailleux L. 2003. J. Exp. Med. 197:121. (Block, FC)
- 15. Olver S, et al. 2006. Cancer Research 66:571. (FC)
- 16. Tan SL, et al. 2006. J. Immunol. 176:2872. (FC) PubMed
- 17. Ponomarev ED, et al. 2006. J. Immunol. 176:1402. (FC)
- 18. Dzhagalov I, et al. 2007. Blood 109:1620. (FC)
- 19. Fazilleau N, et al. 2007. Nature Immunol. 8:753.
- 20. Rasmussen JW, et al. 2006. Infect. Immun.74:6590. PubMed
- 21. Napimoga MH, et al. 2008. J. Immunol. 180:609. PubMed
- 22. Elqaraz-Carmon V, et al. 2008. J. Lipid. Res. 49:1894. PubMed
- 23. Kim DD, et al. 2008. Blood 112:1109. PubMed
- 24. Guo Y, et al. 2008. Blood 112:480. PubMed
- 25. Norian LA, et al. 2009. Cancer Res. 69:3086. (FC) PubMed
- 26. Baumgartner CK, et al. 2010. J. Immunol. 184:573. PubMed
- 27. Charles N, et al. 2010. Nat. Med. 16:701. (FC) PubMed
- 28. Whiteland J, et al. 1995. J. Histochem. Cytochem. 43:313. (IHC)
- 29. Wiesner DL, et al. 2015. PLoS Pathog. 11:1004701. PubMed

Description:

CD11b is a 170 kD glycoprotein also known as αM integrin, Mac-1 α subunit, Mol, CR3, and Ly-40. CD11b is a member of the integrin family, primarily expressed on granulocytes, monocytes/macrophages, dendritic cells, NK cells, and subsets of T and B cells. CD11b non-covalently associates with CD18 ($\beta 2$ integrin) to form Mac-1. Mac-1 plays an important role in cell-cell interaction by binding its ligands ICAM-1 (CD54), ICAM-2 (CD102), ICAM-4 (CD242), iC3b, and fibrinogen.

Antigen References:

- 1. Barclay A, et al. 1997. The Leukocyte Antigen FactsBook Academic Press.
- 2. Springer TA. 1994. Cell 76:301.
- 3. Coxon A, et al. 1996. Immunity 5:653.