

Brilliant Violet 785™ anti-mouse/human CD11b

Catalog # / Size: 1106215 / 50 µg

Clone: M1/70

Isotype: Rat IgG2b, κ

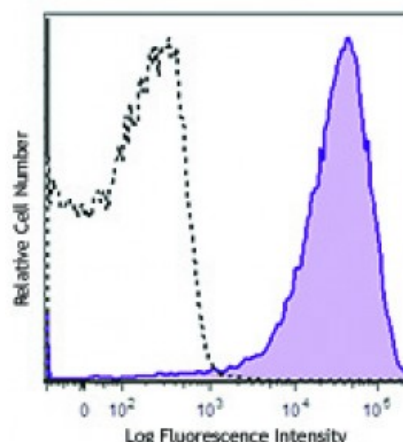
Immunogen: C57BL/10 splenocytes

Reactivity: Human

Preparation: The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 785™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 785™ and unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).

Concentration: 0.2



C57BL/6 mouse bone marrow cells were stained with CD11b (clone M1/70) Brilliant Violet 785™ (filled histogram) or rat IgG2b, κ Brilliant Violet 785™ isotype control (open histogram). Data shown was gated on the 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, the suggested use of this reagent is ≤0.25 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.

Brilliant Violet 785™ excites at 405 nm and emits at 785 nm. The bandpass filter 780/60 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 785™ 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™ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 101231). For *in vivo* studies or highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 101248) with a lower endotoxin limit than standard LEAF™ 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)
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Description: CD11b is a 170 kD glycoprotein also known as α M 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 (β 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.