### **Product Data Sheet**

#### **APC/Fire™ 750 anti-mouse/human CD11b**

Catalog # / 1106310 / 100 µg

Size: 1106305 / 25 µg

Clone: M1/70

Isotype: Rat IgG2b, ĸ

Immunogen: C57BL/10 splenocytes

Reactivity: Human, Mouse, Non-human primate,

Other

The antibody was purified by affinity Preparation:

chromatography and conjugated with

APC/Fire&trade

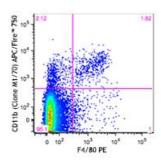
Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Workshop **Number:** 

750 under optimal conditions.

Concentration: 0.2 mg/ml



C57BL/6 splenocytes were blocked with TruStain fcX™ (antimouse CD16/32) Antibody then stained with F4/80 PE and CD11b (clone M1/80) APC/Fire™ 750 (top) or Rat IgG2b, κ APC/Fire™ 750 isotype control (bottom).

#### **Applications:**

Applications: Flow Cytometry

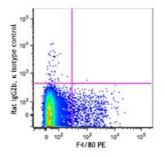
Recommended

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 μg per million cells in 100 µl volume. It is recommended that the reagent be titrated for optimal performance for each

application.

\* APC/Fire™ 750 has a maximum excitation of 650 nm and a maximum emission of 787 nm.



Human paraffin-embedded intestine tissue slices were prepared with a standard protocol of deparaffinization and rehydration. Antigen retrieval was done with Sodium Citrate H.I.E.R. 1X at 95°C for 40 minutes. Tissue was washed with PBS/0.05% Tween 20 twice for five minutes, permeabilized with 0.5% Triton-X 100 for ten minutes and blocked with 5% FBS and 0.2% gelatin for 30 minutes. Then, the tissue was stained with 10 μg/mL of anti-human CD44 (clone IM7) Spark YG<sup>™</sup> 570 (red) at 4°C overnight. Nuclei were counterstained with DAPI (blue). The image was captured with a 10X objective.

## Application Notes:

Clone M1/70 has been verified for immunocytochemistry (ICC) and frozen immunohistochemistry (IHC-F).

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  $^{5,11-13}$ . For in vivo studies or highly sensitive assays, we recommend Ultra-LEAF  $^{\text{TM}}$  purified antibody (Endotoxin < 0.01 EU/ $\mu$ g, Azide-Free, 0.2  $\mu$ m filtered) (Cat. No. 101248).

# Application References:

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  Noel GJ, et al. 1990. J. Clin. Invest. 85:208. (IF)
- 7. Allen LA and Aderem A. 1996. *J. Exp. Med.* 184:627 (IF)
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- 13. Iwasaki A and Kelsall BL. 2001. *J. Immunol.* 166:4884 (IHC, FC)
- 14. Tailleux L. 2003. J. Exp. Med. 197:121. (Block, FC)
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- 27. Charles N, et al. 2010. Nat. Med. 16:701. (FC) PubMed
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- 32. Doni A, et al. 2015. J Exp Med. 212:905. PubMed
- 33. Ferreira R, et al. 2016. J Infect Dis. 213: 669 673. PubMed
- 34. Peterson VM, et al. 2017. Nat. Biotechnol. 35:936. (PG)

#### **Description:**

CD11b is a 170 kD glycoprotein also known as  $\alpha M$  integrin, Mac-1  $\alpha$  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.