# **Technical Data Sheet**

# V500 Rat anti-Mouse CD11b

## **Product Information**

Material Number:
Alternate Name:
Entrez Gene ID:
Size:
<b>Concentration:</b>
Clone:
Immunogen:
Isotype:
Reactivity:
Storage Buffer:

# 562128

Itgam; Integrin alpha-M; Ly-40; Mac-1a; Mac-1 alpha; CR3A; CR-3 alpha chain 16409 25 µg 0.2 mg/ml M1/70 Mouse Splenic Cells Rat (DA) IgG2b, ĸ QC Testing: Mouse Aqueous buffered solution containing protein stabilizer, glycerol and ≤0.09% sodium azide.

# Description

The M1/70 monoclonal antibody specifically binds to CD11b, also known as Integrin alpha M (Itgam or  $\alpha$ M). CD11b is a 170-kDa type 1 transmembrane glycoprotein and belongs to the Integrin alpha chain family. CD11b serves as the alpha chain of the heterodimeric Mac-1 integrin (CD11b/CD18, αMβ2), also known as complement receptor 3 (CR3). Mac-1 mediates adhesion to ICAM-1 (CD54), ICAM-2 (CD102), fibrinogen and binding to C3bi. Mac-1 is expressed at varying levels on granulocytes, macrophages, myeloid-derived dendritic cells, natural killer cells, microglia, and B-1 B lymphocytes. 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. The M1/70 antibody reportedly blocks cell adherence and C3bi binding but does not block cell-mediated lysis. Cross-reaction of the M1/70 antibody with CD11b expressed on human monocytes, polymorphonuclear leukocytes, and NK cells has been reported.

The antibody is conjugated to BD Horizon™ V500, which has been developed for use in multicolor flow cytometry experiments and is available exclusively from BD Biosciences. It is excited by the Violet laser with an Ex max of 415 nm and Em Max at 500 nm. BD Horizon V500 conjugates emit at a similar wavelength to Amcyan yet exhibit reduced spillover into the FITC channel. For more information on BD Horizon V500, visit bdbiosciences.com/colors.

When compensating dyes in this spectral range (such as Horizon™ V500 and AmCyan), the most accurate compensation can be obtained using single stained cellular controls. Due to spectral differences between cells and beads in this channel, using BD CompBeads can result in spillover errors for V500 and AmCyan reagents. Therefore, the use of BD CompBeads or BD CompBeads Plus to determine spillover values for these reagents is not recommended. Different V500 reagents (e.g. CD4 vs. CD45) can have slightly different fluorescence spillover therefore, it may also be necessary to use clone specific compensation controls when using these reagents.

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Expression of CD11b on mouse bone-marrow cells. Mouse BALB/c bone marrow leukocytes were stained with either V500 Rat IgG2b, ĸ Isotype Control antibody (Cat. No. 560784; Left Panel) or V500 Rat Anti-Mouse CD11b (Cat. No. 562128; Right Panel). Flow cytometric dot plots show the expression of side-scattered light characteristics versus CD11b (or background Ig Isotype Control staining) for events with the forward and side light-scatter characteristics of viable bone marrow leukocytes. Flow cytometry was performed using a BD FACSCanto™ II Flow Cytometer System.

## **Preparation and Storage**

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

The antibody was conjugated with BD Horizon<sup>TM</sup> V500 under optimum conditions, and unreacted BD Horizon<sup>TM</sup> V500 was removed. The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography.

#### **Application Notes**

#### Application

Flow cytometry Routinely Tested
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Suggestee	l Compan	ion Pi	roducts
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Catalog Number	Name	Size	Clone
560784	V500 Rat IgG2b, κ Isotype Control	0.1 mg	A95-1
554656	Stain Buffer (FBS)	500 ml	(none)
562127	V500 Rat anti-Mouse CD11b	0.1 mg	M1/70

#### **Product Notices**

- Since applications vary, each investigator should titrate the reagent to obtain optimal results. 1.
- 2. An isotype control should be used at the same concentration as the antibody of interest.
- BD Horizon<sup>™</sup> V500 has a maximum absorption of 415 nm and maximum emission of 500 nm. Before staining with this reagent, please 3. confirm that your flow cytometer is capable of exciting the fluorochrome and discriminating the resulting fluorescence.
- For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at 4. www.bdbiosciences.com/colors.
- Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before 5. discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 6. 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. (Immunogen: Immunoprecipitation)

Greimers R, Trebak M, Moutschen M, Jacobs N, Boniver J. Improved four-color flow cytometry method using fluo-3 and triple immunofluorescence for analysis of intracellular calcium ion ([Ca2+]i) fluxes among mouse lymph node B- and T-lymphocyte subsets. Cytometry. 1996; 23(3):205-217. (Methodology: Flow cytometry) Lagasse E, Weissman IL. Flow cytometric identification of murine neutrophils and monocytes. J Immunol Methods. 1996; 197(1-2):139-150. (Methodology: Flow cytometry)

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: Immunoprecipitation)

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: Immunoprecipitation)

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. (Biology)

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