## Technical Data Sheet

# **BV711 Hamster Anti-Mouse CD11c**

#### **Product Information**

Material Number: 563048

Alternate Name: Cd11c; Itgax; Integrin alpha-X; Integrin αX; Cr4; Complement receptor 4

 Size:
 50 μg

 Concentration:
 0.2 mg/ml

 Clone:
 HL3

Immunogen: C57BL/6 Mouse Intestinal Intraepithelial Lymphocytes

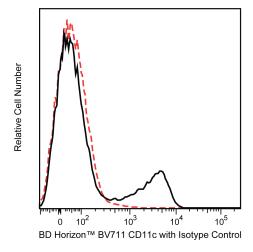
Isotype:Armenian Hamster IgG1,  $\lambda$ 2Reactivity:QC Testing: Mouse

Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

### Description

The HL3 monoclonal antibody specifically binds to the integrin  $\alpha x$  chain of gp150, 95 (CD11c/CD18). CD11c is expressed on dendritic cells, CD4- CD8+ intestinal intraepithelial lymphocytes (IEL) and some NK cells. It is upregulated on IEL and lymph-node T cells following *in vivo* activation. Cells of the monocyte/macrophage lineage have been reported to express low levels of CD11c. CD11c plays a role in binding of iC3b.

The antibody was conjugated to BD Horizon<sup>TM</sup> BV711 which is part of the BD Horizon<sup>TM</sup> Brilliant Violet<sup>TM</sup> family of dyes. This dye is a tandem fluorochrome of BD Horizon<sup>TM</sup> BV421 with an Ex Max of 405-nm and an acceptor dye with an Em Max at 711-nm. BD Horizon<sup>TM</sup> BV711 can be excited by the violet laser and detected in a filter used to detect Cy<sup>TM</sup>5.5 / Alexa Fluor® 700-like dyes (eg, 712/20-nm filter). Due to the excitation and emission characteristics of the acceptor dye, there may be moderate spillover into the Alexa Fluor® 700 and PerCP-Cy<sup>TM</sup>5.5 detectors. However, the spillover can be corrected through compensation as with any other dye combination.



Flow cytometric analysis of CD11c expression on mouse dendritic cells. C57BL/6 mouse splenocytes were cultured with recombinant mouse GM-CSF (Cat. No.554586; 5 ng/ml) overnight. The cells were then harvested and stained either with a BD Horizon™ BV711 Hamster IgG1, λ1 Isotype Control (Cat. No. 563049; dashed line histogram) or with the BD Horizon™ BV711 Hamster Anti-Mouse CD11c antibody (Cat. No. 563048; solid line histogram). The fluorescence histograms were derived from events with the forward and side light-scatter characteristics of viable dendritic cells. Flow cytometry was performed using a BD™ LSR II Flow Cytometer 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 BD Horizon™ BV711 under optimum conditions, and unconjugated antibody and free BD Horizon™ BV711 were removed.

## **Application Notes**

## Application

Flow cytometry

## **Suggested Companion Products**

Catalog Number	Name	Size	Clone	
554656	Stain Buffer (FBS)	500 ml	(none)	
563049	BV711 Hamster IgG1, λ1 Isotype Control	50 μg	G235-2356	
554586	Recombinant Mouse GM-CSF	10 ug	(none)	

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#### **Product Notices**

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 2. An isotype control should be used at the same concentration as the antibody of interest.
- 3. Brilliant Violet<sup>TM</sup> 711 is a trademark of Sirigen.
- 4. Cy is a trademark of Amersham Biosciences Limited.
- 5. Alexa Fluor® is a registered trademark of Molecular Probes, Inc., Eugene, OR.
- 6. Although hamster immunoglobulin isotypes have not been well defined, BD Biosciences Pharmingen has grouped Armenian and Syrian hamster IgG monoclonal antibodies according to their reactivity with a panel of mouse anti-hamster IgG mAbs. A table of the hamster IgG groups, Reactivity of Mouse Anti-Hamster Ig mAbs, may be viewed at http://www.bdbiosciences.com/documents/hamster\_chart\_11x17.pdf.
- Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 8. 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.
- 9. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- 10. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

#### References

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Burt BM, Plitas G, Stableford JA, Nguyen HM, Bamboat ZM, Pillarisetty VG, DeMatteo RP. CD11c identifies a subset of murine liver natural killer cells that responds to adenoviral hepatitis. *J Leukoc Biol.* 2008; 84(4):1039-1046. (Clone-specific: Flow cytometry)

Gao JX, Liu X, Wen J, et al. Differentiation of monocytic cell clones into CD8 alpha+ dendritic cells (DC) suggests that monocytes can be direct precursors for both CD8 alpha+ and CD8 alpha- DC in the mouse. *J Immunol.* 2003; 170(12):5927-5935. (Biology)

Huleatt JW, Lefrancois L. Antigen-driven induction of CD11c on intestinal intraepithelial lymphocytes and CD8+ T cells in vivo. *J Immunol.* 1995; 154(11):5684-5693. (Immunogen: Flow cytometry, Immunoprecipitation)

Maraskovsky E, Brasel K, Teepe M, et al. Dramatic increase in the numbers of functionally mature dendritic cells in Flt3 ligand-treated mice: multiple dendritic cell subpopulations identified. *J Exp Med.* 1996; 184(5):1953-1962. (Biology)

Pulendran B, Lingappa J, Kennedy MK, et al. Developmental pathways of dendritic cells in vivo: distinct function, phenotype, and localization of dendritic cell subsets in FLT3 ligand-treated mice. *J Immunol.* 1997; 159(5):2222-2231. (Biology)

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