Technical Data Sheet

BV711 Mouse Anti-Human CD183

Product Information

Material Number: 563156

Alternate Name: CXCR3; C-X-C chemokine receptor type 3; GPR9; IP10R; MigR; CKRL2; CMKAR3

 Size:
 50 test

 Vol. per Test:
 5 μl

Clone: 1C6/CXCR3

 Immunogen:
 Human CXCR3 Peptide

 Isotype:
 Mouse (BALB/c) IgG1, κ

 Reactivity:
 QC Testing: Human

Tested in Development: Rhesus, Cynomolgus, Baboon

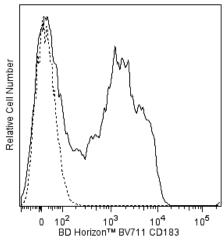
Workshop: VII 70500

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

Description

The 1C6/CXCR3 monoclonal antibody specifically binds to human CD183, also known as the CXCR3 chemokine receptor. CD183 is a 40-41 kDa seven-transmembrane protein and member of the G protein-coupled receptor family. CD183 is expressed primarily on activated T cells that infiltrate inflammatory sites. It has also been detected on some circulating T cells, B cells, and NK cells. Reports show that some CXCR3-positive T cells also express CCR5 and are mostly CD45RO-positive cells. Three ligands for CXCR3 have been identified. They are CXCL9 (Mig/monokine induced by interferon-γ), CXCL10 (IP-10/interferon-γ inducible 10-kD protein), and CXCL11 (I-TAC/interferon-inducible T-cell alpha chemoattractant). These chemokines are produced by a variety of cells upon stimulation by IFN-γ and interact with CXCR3 to mediate T-cell chemotaxis. This reagent has been reported to be suitable for immunohistochemical staining of acetone-fixed, frozen sections and/or formalin-fixed, paraffin-embedded tissue sections with citrate pretreatment. Clone 1C6/CXCR3 also cross reacts with a subset of peripheral blood lymphocytes of baboon, and both rhesus and cynomolgus macaque monkeys. The distribution of lymphocytes is similar to that observed with CD183-positive peripheral blood lymphocytes from normal human donors. CXCR3 has been clustered as CD183 in the VIIth HLDA workshop.

The antibody was conjugated to BD HorizonTM BV711 which is part of the BD HorizonTM Brilliant VioletTM family of dyes. This dye is a tandem fluorochrome of BD HorizonTM BV421 with an Ex Max of 405-nm and an acceptor dye with an Em Max at 711-nm. BD HorizonTM BV711 can be excited by the violet laser and detected in a filter used to detect CyTM5.5 / 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-CyTM5.5 detectors. However, the spillover can be corrected through compensation as with any other dye combination.



Flow cytometric analysis of CD183 (CXCR3) expression on human peripheral blood lymphocytes. Whole blood was stained with BD Horizon™ BV711 Mouse anti-Human CD183 antibody (Cat. No. 563156; solid line histogram) or with a BD Horizon™ BV711 Mouse IgG1, κ Isotype Control (Cat. No. 563044; dashed line histogram). The erythrocytes were lysed with BD Pharm Lyse™ Lysing Buffer (Cat. No. 555899). The fluorescence histograms were derived from events with the forward and side light-scatter characteristics of viable lymphocytes. Flow cytometric analysis 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.

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Application Notes

Application

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Flow cytometry	Routinely Tested

Suggested Companion Products

Catalog Number	Name Name	Size	Clone	
554656	Stain Buffer (FBS)	500 ml	(none)	
563044	BV711 Mouse IgG1, k Isotype Control	50 μg	X40	
555899	Lysing Buffer	100 ml	(none)	

Product Notices

- This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1 × 10⁶ cells in a 100-μl experimental sample (a test).
- 2. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 3. An isotype control should be used at the same concentration as the antibody of interest.
- 4. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- 5. 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.
- 6. For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- 7. Alexa Fluor® is a registered trademark of Molecular Probes, Inc., Eugene, OR.
- 8. Cy is a trademark of Amersham Biosciences Limited.
- 9. Brilliant VioletTM 711 is a trademark of Sirigen.
- Species testing during development may have been performed with a different format of the same clone. Selected applications have been tested for cross-reactivity.

References

Loetscher M, Gerber B, Loetscher P, et al. Chemokine receptor specific for IP10 and mig: structure, function, and expression in activated T-lymphocytes. *J Exp Med.* 1996; 184(3):963-969. (Biology)

Marcher C, Moller BK, Lillevang ST, Kristensen T. CXCR4 and IL17R are downregulated on cord-blood CD34-positive cells during short-term culture. In: Mason D, Simmons D, Buckley C, et al., ed. Leucocyte Typing VII: White Cell Differentiation Antigens. New York: Oxford University Press; 2002:629-632. (Clone-specific: Flow cytometry)

Piali L, Weber C, LaRosa G, et al. The chemokine receptor CXCR3 mediates rapid and shear-resistant adhesion-induction of effector T lymphocytes by the chemokines IP10 and Mig. Eur J Immunol. 1998; 28(3):961-972. (Biology)

Qin S, Rottman JB, Myers P, et al. The chemokine receptors CXCR3 and CCR5 mark subsets of T cells associated with certain inflammatory reactions. *J Clin Invest.* 1998; 101(4):746-754. (Immunogen: Blocking, Flow cytometry, Immunohistochemistry, Inhibition)

Uguccioni M, Willimann K. Cytokine/Chemokine Receptors: Section report. In: Mason D, Simmons D, Buckley C, et al, ed. *Leukocyte Typing VII*. New York: Oxford University Press; 2002:237-243. (Clone-specific: Flow cytometry)

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