Technical Data Sheet

BV510 Mouse Anti-Human CD20

Product Information

Material Number: 563067

Alternate Name: MS4A1; B1; Bp35; LEU-16; S7

Immunogen:Human 6.16c1.3 B cell lineIsotype:Mouse (C57BL/6) IgG2b, κ Reactivity:QC Testing: Human

Tested in Development: Rhesus, Cynomolgus, Baboon

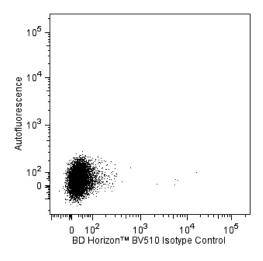
Workshop: II B22; III B739, NL382; IV B201

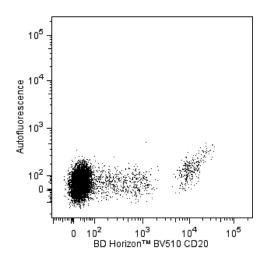
Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

Description

The 2H7 monoclonal antibody specifically binds to CD20 that is encoded by the MS4A1 (Membrane-spanning 4-domains, subfamily A, member 1) gene. CD20 is a 33-37 kDa unglycosylated four-transmembrane phosphoprotein. CD20 is expressed on pre-B-cells, resting and activated B cells and follicular dendritic cells but not on plasma cells. Low level CD20 expression is observed on a small subset of normal circulating T lymphocytes. The CD20 molecule is involved in the regulation of B-cell activation.

The antibody was conjugated to BD Horizon™ BV510 which is part of the BD Horizon™ Brilliant Violet™ family of dyes. With an Ex Max of 405-nm and Em Max at 510-nm, BD Horizon™ BV510 can be excited by the violet laser and detected in the BD Horizon™ V500 (525/50-nm) filter set. BD Horizon™ BV510 conjugates are useful for the detection of dim markers off the violet laser.





Two-color flow cytometric analysis of CD20 expression on human peripheral blood lymphocytes. Whole blood was stained with either BD Horizon™ BV510 Mouse IgG2b, k Isotype Control (Cat. No. 563025; Left Panel) or with BD Horizon™ BV510 Mouse Anti-Human CD20 antibody (Cat. No. 563067; Right Panel). Erythrocytes were lysed with BD Pharm Lyse™ Lysing Buffer (Cat. No. 555899). Two color flow cytometric dot plots show the correlated expression of CD20 (or Ig Isotype Control staining) versus cellular autofluorescence derived from gated events with the forward 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 HorizonTM BV510 under optimum conditions, and unconjugated antibody and free BD HorizonTM BV510 were removed.

Application Notes

Application

Flow cytometry Routinely Tested

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 United States
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 877.232.8995
 800.979.9408
 32.53.720.550
 0120.8555.90
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Suggested Companion Products

Catalog Number	<u>Name</u>	Size	Clone	
554656	Stain Buffer (FBS)	500 ml	(none)	
563025	BV510 Mouse IgG2b, κ Isotype Control	50 μg	27-35	
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.
- 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. Brilliant VioletTM 510 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

Clark EA, Yokochi T. Human B cell and B cell blast-associated surface molecules defined with monoclonal antibodies. In: Bernard A, Boumsell L, Dausset J, Milstein C, Schlossman SF, ed. *Leukocyte Typing*. Berlin: Springer-Verlag; 1984:339-346. (Clone-specific: Blocking, Flow cytometry, Immunoprecipitation) Hultin LE, Hausner MA, Hultin PM, Giorgi JV. CD20 (pan-B cell) antigen is expressed at a low level on a subpopulation of human T lymphocytes. *Cytometry*. 1993; 14(2):193-204. (Biology)

Knapp W, Dörken B, Gilks WR, et al, ed. Leucocyte Typing IV. New York, NY: Oxford University Press; 1989:1-1182. (Biology)

Ledbetter JA, Clark EA. Surface phenotype and function of tonsillar germinal center and mantle zone B cell subsets. *Hum Immunol.* 1986; 15:30-43. (Immunogen: Blocking, Flow cytometry)

Loken MR, Shah VO, Dattilio KL, Civin Cl. Flow cytometric analysis of human bone marrow. II. Normal B lymphocyte development. *Blood.* 1987; 70(5):1316-1324. (Biology)

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