Technical Data Sheet

BV510 Rat Anti-Mouse CD21/CD35

Product Information

 Material Number:
 563175

 Alternate Name:
 CR2/CR1

 Size:
 50 μg

 Concentration:
 0.2 mg/ml

 Clone:
 7G6

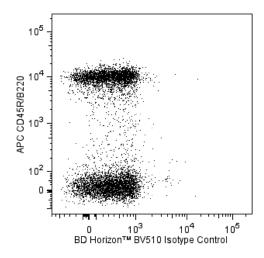
Immunogen:Purified Mouse CR1Isotype:Rat (SD) IgG2b, κ Reactivity:QC Testing: Mouse

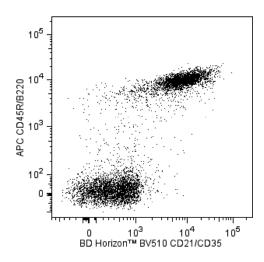
Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

Description

The 7G6 antibody recognizes an epitope shared by 145-150-kDa and 190-kDa complement receptor proteins, originally designated CR2 (CD21) and CR1 (CD35), respectively. In the mouse, CD21 and CD35 are expressed on the majority of peripheral B lymphocytes, on the majority of resident peritoneal macrophages and mast cells, on peripheral blood granulocytes after treatment with N-formyl-Met-Leu-Phe, and on follicular dendritic cells, but not on thymocytes, T cells, erythrocytes, or platelets. CD21 is a ligand-binding component of the CD19/CD21/CD81 signal-transduction complex associated with the antigen receptor on B lymphocytes. CD21/CD35 also co-localizes with CD19 on the surface of peritoneal mast cells. Cr2null mice display impaired inflammatory and humoral immune responses in vivo. The 7G6 mAb has been reported to inhibit rosette formation by C3d-bearing sheep erythrocytes, to block the complement dependent trapping of immune complexes by follicular dendritic cells, and to down-regulate mouse CD21/CD35 expression upon in vivo application, thus inhibiting primary antibody responses to immunization. Co-stimulation of B-cell differentiation via Sepharose-coupled 7G6 antibody has also been observed. The 7G6 mAb recognizes an epitope on CD35 distinct from the epitope recognized by anti-mouse CD35, clone 8C12 (Cat. No. 558768, for the purified antibody), and it does not block binding of 8C12 mAb to mouse CD35.

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





Two-color flow cytometric analysis of CD21/CD35 expression on BALB/c mouse splenocytes. BALB/c splenic leukocytes were stained with APC Rat Anti-Mouse CD45R/B220 antibody (Cat. No. 553092/561880) and either BD Horizon™ BV510 Rat IgG2b, κ Isotype Control (Cat. No. 562951, Left Panel) or BD Horizon™ BV510 Rat Anti-Mouse CD21/CD35 antibody (Cat. No. 563175, Right Panel) in the presence of Purified Rat Anti-Mouse CD16/CD32 antibody (Mouse BD Fc Block™) (Cat. No. 553141/553142). Two-color flow cytometric dot plots showing the correlated expression of CD21/CD35 (or Ig Isotype control staining) versus CD45R/B220 were derived from gated events with the forward and side light-scatter characteristics of viable splenocytes. Flow cytometric analysis was performed using a BD LSRFortessa™ Cell Analyzer System.

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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

Suggested Companion Products

Catalog Number	Name	Size	<u>Clone</u>	
554656	Stain Buffer (FBS)	500 ml	(none)	
562951	BV510 Rat IgG2b, κ Isotype Control	50 μg	R35-38	
555899	Lysing Buffer	100 ml	(none)	
553092	APC Rat Anti-Mouse CD45R/B220	0.1 mg	RA3-6B2	
561880	APC Rat Anti-Mouse CD45R/B220	25 μg	RA3-6B2	
553141	Purified Rat Anti-Mouse CD16/CD32 (Mouse BD Fc Block™)	0.1 mg	2.4G2	
553142	Purified Rat Anti-Mouse CD16/CD32 (Mouse BD Fc Block™)	0.5 mg	2.4G2	

Product Notices

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Source of all serum proteins is from USDA inspected abattoirs located in the United States. 2.
- An isotype control should be used at the same concentration as the antibody of interest. 3.
- 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.
- For fluorochrome spectra and suitable instrument settings, please refer to our Multicolor Flow Cytometry web page at www.bdbiosciences.com/colors.
- Brilliant VioletTM 510 is a trademark of Sirigen. 7.

References

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Axcrona K, Gray D, Leanderson T. Regulation of B cell growth and differentiation via CD21 and CD40. Eur J Immunol. 1996; 26(9):2203-2207. (Clone-specific: (Co)-stimulation, Flow cytometry)

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Fagarasan S, Muramatsu M, Suzuki K, Nagaoka H, Hiai H, Honio T, Critical roles of activation-induced cytidine deaminase in the homeostasis of gut flora. Science, 2002; 298(5597):1424-1427, (Clone-specific; Fluorescence microscopy, Immunofluorescence)

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Gommerman JL, Oh DY, Zhou X, et al. A role for CD21/CD35 and CD19 in responses to acute septic peritonitis: a potential mechanism for mast cell activation. J Immunol. 2000; 165(12):6915-6921. (Clone-specific: Flow cytometry, Fluorescence microscopy, Immunofluorescence)

Heyman B, Wiersma EJ, Kinoshita T. In vivo inhibition of the antibody response by a complement receptor-specific monoclonal antibody. J Exp Med. 1990; 172(2):665-668. (Clone-specific: Inhibition, In vivo exacerbation)

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Yoshida K, van den Berg TK, Dijkstra CD. Two functionally different follicular dendritic cells in secondary lymphoid follicles of mouse spleen, as revealed by CR1/2 and FcR gamma II-mediated immune-complex trapping. Immunology. 1993; 80(1):34-39. (Clone-specific: Inhibition)

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