Technical Data Sheet

BV786 Mouse Anti-Human CD25

Product Information

Material Number: 563701

Alternate Name: IL-2R; IL2RA; IL-2Rα; TCGFR; TAC antigen; p55

 $\begin{array}{lll} \textbf{Size:} & 100 \text{ tests} \\ \textbf{Vol. per Test:} & 5 \text{ } \mu \text{I} \\ \textbf{Clone:} & \text{M-A251} \\ \textbf{Isotype:} & \text{Mouse IgG1, } \kappa \\ \textbf{Reactivity:} & \text{QC Testing: Human} \end{array}$

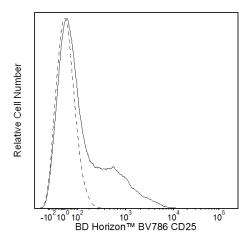
Workshop: IV A053

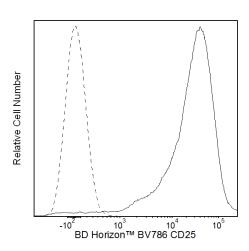
Storage Buffer: Aqueous buffered solution containing BSA and ≤0.09% sodium azide.

Description

The M-A251 monoclonal antibody specifically binds to the 55 kDa type I transmembrane glycoprotein known as the low-affinity interleukin-2 receptor alpha chain subunit (IL-2R α). CD25 is expressed on regulatory T cells and on activated lymphocytes (T and B) and monocytes. It associates with the IL-2R β /CD122 and the IL-2R γ /CD132 receptor chains to form the high-affinity IL-2R complex. CD25 expression on T and B lymphocytes is upregulated by antigenic or mitogenic stimulation. Soluble CD25/IL-2R α is produced as a consequence of lymphocyte stimulation and is found in biological fluids following inflammatory responses.

The antibody was conjugated to BD Horizon™ BV786 which is part of the BD Horizon™ Brilliant Violet™ family of dyes. This dye is a tandem fluorochrome of BD Horizon™ BV421 with an Ex Max of 405-nm and an acceptor dye with an Em Max at 786-nm. BD Horizon™ BV786 can be excited by the violet laser and detected in a filter used to detect Cy7™-like dyes (eg, 780/60-nm filter).





Flow cytometric analysis of CD25 expression on unstimulated and stimulated human peripheral blood lymphocytes.

Left Panel: Whole blood was stained with either BD Horizon™ BV786 Mouse IgG1, κ Isotype Control (Cat. No. 563330; dashed line histogram) or BD Horizon™ BV786 Mouse Anti-Human CD25 antibody (Cat. No. 563700/563701; solid line histogram). The erythrocytes were subsequently lysed with BD Pharm Lyse™ Lysing Buffer (Cat. No. 555899).

Right Panel: Phytohemagglutinin-stimulated (3 days) human peripheral blood mononuclear cells were stained with either BD Horizon™ BV786 Mouse Anti-Human CD25 antibody (solid line histogram) or with BD Horizon™ BV786 Mouse IgG1, κ Isotype Control (dashed line histogram).

The fluorescence histograms showing CD25 expression (or Ig Isotype control staining) were derived from gated events with the forward and side light-scatter characteristics of viable lymphocytes (Left Panel) or lymphoblasts (Right Panel). 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™ BV786 under optimum conditions, and unconjugated antibody and free BD Horizon™ BV786 were removed.

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Application Notes

Application

Flow cytometry	Routinely Tested

Suggested Companion Products

Catalog Number	Name Name	Size	Clone	
554656	Stain Buffer (FBS)	500 ml	(none)	
563330	BV786 Mouse IgG1, k Isotype Control	50 μg	X40	
563700	BV786 Mouse Anti-Human CD25	25 tests	M-A251	
555899	Lysing Buffer	100 ml	(none)	
349202	BD FACS TM Lysing Solution	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. Cy is a trademark of Amersham Biosciences Limited.
- Brilliant Violet™ 421 is a trademark of Sirigen.
- Brilliant VioletTM 786 is a trademark of Sirigen.

References

Janszen M, Buck D, Maino VC. Functional and molecular properties of CD25 monoclonal antibodies. In: Knapp W, Dörken B, Gilks WR, et al, ed. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:403-406. (Clone-specific: Blocking, Flow cytometry, Immunoprecipitation)

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

Nakamura K, Kitani A, Fuss I, Pedersen A, Harada N, Nawata H, Strober W. TGF-β1 plays an important role in the mechanism of CD4+CD25+ regulatory T cell activity in both humans and mice. *J Immunol.* 2004; 172(2):834-842. (Clone-specific: Flow cytometry, Fluorescence activated cell sorting)

Ravoet AM, Latinne D, Couvreur B, Zenebergh A, De Bruyère, Ninane J, Sokal G. CD25 mab: Epitopes recognized, effect on lymphocyte activation, mediation of ADCC. In: Knapp W, Dörken B, Gilks WR, et al, ed. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:408-411. (Clone-specific: Functional assay)

Schlossman SF, Bournsell L, Gilks W, et al, ed. *Leukocyte Typing V: White Cell Differentiation Antigens*. New York: Oxford University Press; 1995. (Biology) Schwarting R, Stein H. Cluster report: CD25. In: Knapp W, Dörken B, Gilks WR, et al, ed. *Leucocyte Typing IV: White Cell Differentiation Antigens*. New York, NY: Oxford University Press; 1989:399-403. (Clone-specific: Flow cytometry, Immunoprecipitation)

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