Technical Data Sheet

PE Rat anti-Mouse CD125

Product Information

Material Number: 558488

Alternate Name: IL-5 Receptor α-subunit

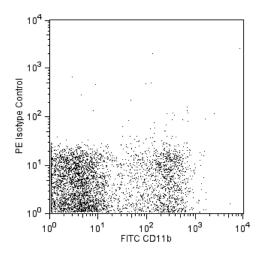
Size $0.1 \, \text{mg}$ 0.2 mg/ml Concentration: Clone: T21

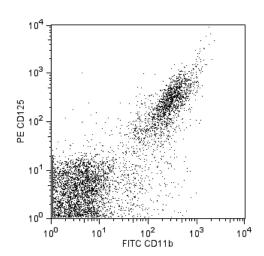
Mouse T88-M Immunogen: Isotype: Rat IgG1, λ Reactivity: QC Testing: Mouse

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

Description

The antibody clone T21 recognizes the mouse α-subunit (also known as CD125) of the IL-5 receptor. The IL-5 receptor is comprised of αand β -subunits where the α -subunit is specific to the binding of IL-5 ligand and the β -subunit is common among the IL-3, IL5 and granulocyte/macrophage colony-stimulating factor (GM-CSF) receptors. Murine IL-5 belongs to the hematopoietic growth factor family of cytokines and is known to enhance several functions of murine B-cells, including immunoglobulin production, growth and differentiation. Upon binding, T21 inhibits IL-5 induced B-cell proliferation and Ig-secretion.





Flow cytometric analysis of PE-conjugated anti-mouse CD125 on mouse bone marrow cells. Isolated murine bone marrow cells were stained with either PE anti-CD125 (clone T21, Cat. No. 558488, right panel) or a PE-conjugated rat IgG1 isotype control (Cat. No. 557078, left panel) and FITC anti-CD11b (clone M1/70, Cat. No. 553310). Flow cytometry was performed on a BD FACSCalibur™ System and the histograms were derived from the gated events based on light scattering characteristics of viable bone marrow cells

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 R-PE under optimum conditions, and unconjugated antibody and free PE were removed.

Application Notes

Application

Flow cytometry Routinely Tested

Neutralization Activity:

The T21 clone has been reported to inhibit IL-5 binding to it's receptor and to inhibit IL-5 induced B-cell proliferation and Ig-secretion.

Recommended Assay Procedure:

For flow cytometry of cell suspensions from peripheral lymphoid tissues, it is recommended that the cells be pre-incubated with Mouse BD Fc Block™ Purified anti-Mouse CD16/CD32 mAb, Clone 2.4G2 (Cat. No. 553141/553142).

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Suggested Companion Products

Catalog Number	Name	Size	Clone	
557078	PE Rat IgG1, λ Isotype Control	0.1 mg	A110-1	
553141	Purified Rat Anti-Mouse CD16/CD32 (Mouse BD Fc Block TM)	0.1 mg	2.4G2	
553310	FITC Rat Anti-Mouse CD11b	0.5 mg	M1/70	
554656	Stain Buffer (FBS)	500 ml	(none)	

Product Notices

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 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.
- 3. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
- 4. An isotype control should be used at the same concentration as the antibody of interest.
- 5. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

References

Adachi T, Alam R.. The mechanism of IL-5 signal transduction. Am J Physiol. 275(3 Pt 1):C623-C633. (Biology)

Hitoshi Y, Yamaguchi N, Mita S, et al. Distribution of IL-5 receptor-positive cells. Expression of IL-5 receptor in Ly-1 (CD5)+ B cells.. *J Immunol.* 1990; 144(11):4218-4225. (Biology)

Wetzel, GD. Interleukin 5 regulation of peritoneal Ly-1 B lymphocyte proliferation, differentiation, and autoantibody secretion. *Eur J Immunol.* 1989; 19(9):1701-1707. (Biology)

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