### **Technical Data Sheet**

# PE Mouse Anti-Human TNFR Related Protein (LTβR)

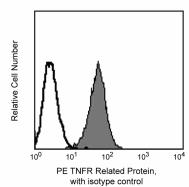
#### **Product Information**

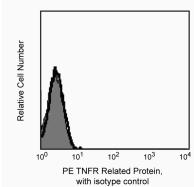
**Material Number:** 551503 Size: 0.2 mg 0.2 mg/mlConcentration: hTNFR-RP-M12 Clone: Human LTβR-Fc protein Immunogen: Mouse IgG1, κ **Isotype:** QC Testing: Human Reactivity:

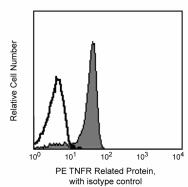
Aqueous buffered solution containing ≤0.09% sodium azide. Storage Buffer:

#### Description

The hTNFR-RP-M12 antibody reacts with the extracellular domain of the 61 kDa receptor for the human cytokines, LTα1β2 and LIGHT. This receptor is referred as the Lymphotoxin  $\beta$  Receptor (LT $\beta$ R). The gene encoding LT $\beta$ R has been designated as the TNFRSF3 by the Human Gene Nomenculture Committee (http://www.gene.ucl.ac.uk/users/hester/tnftop.html). LTBR was previously known as TNFRIII and TNF receptor-related protein (TNFRrp). LT $\beta$ R is a type I transmembrane glycoprotein and member of the TNF Receptor Superfamily. LT $\beta$ R are expressed on stromal cells in lymphoid tissue, normal dermal fibroblasts, bronchial airway epithelial cells and in a variety of adherent cell lines including FDC-1, U937, HT-29, HeLa and HEK 293 cells. LTBR are absent on human peripheral blood T and B cells and expressed at low levels by monocytes.







Overlapping Histograms Figure: Expression of cell surface LT\$\text{LT}\$\$ by HeLa, Jurkat and human monocytes. HeLa (left panel) and Jurkat (center panel) cells were stained with R-PE-conjugated hTNFR-RP-M12 (0.5 µg/10^6 cells, Cat No. 551503). Staining with the hTNFR-RP-M12 antibody (filled histograms) is compared to staining obtained using the isotype control antibody (open histograms). The histograms in the figure (left and center) were derived from gated events with the light scattering characteristics of viable cells. Similarly, whole human blood was first treated with PharmLyse™ (Cat No. 555899) to lyse erythrocytes prior to staining with hTNFR-RP-M12. The peripheral blood leucocytes were subsequently blocked with normal polyclonal human IgG (5 μg/10^6 cells) and stained with R-PE-conjugated hTNFR-RP-M12 (0.5 μg/10^6 cells, Cat No. 551503). Staining with the R-PE-conjugated hTNFR-RP-M12 antibody (filled histograms) is compared to staining obtained using the mouse IgG1 isotype control (open histograms). The histograms in the right panel were derived from gated events with the light scattering characteristics of monocytes. Note: Certain human cell lines or cell types (e.g., neutrophils, monocytes) can first be treated with reagents that block receptors for the Fc regions of immunoglobulin to avoid nonspecific immunofluorescent staining mediated by Fc receptors

#### **Preparation and Storage**

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.

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

### **Application Notes**

#### Application

Flow cytometry Routinely Tested

## **Suggested Companion Products**

| Catalog Number | Name                             | Size   | Clone   |
|----------------|----------------------------------|--------|---------|
| 554680         | PE Mouse IgG1, κ Isotype Control | 0.1 mg | MOPC-21 |
| 555899         | Lysing Buffer                    | 100 ml | (none)  |

#### **BD Biosciences**

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#### **Product Notices**

- 1. Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- 2. 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.

#### References

Browning JL, Dougas I, Ngam-ek A, et al. Characterization of surface lymphotoxin forms. Use of specific monoclonal antibodies and soluble receptors. *J Immunol.* 1995; 154(1):33-46. (Immunogen)

Crowe PD, VanArsdale TL, Walter BN, et al. A lymphotoxin-beta-specific receptor. Science. 1994; 264(5159):707-710. (Biology)

Murphy M, Walter BN, Pike-Nobile L,. Expression of the lymphotoxin beta receptor on follicular stromal cells in human lymphoid tissues. *Cell Death Differ.* 1998; 5(6):497-505. (Biology)

Rooney IA, Butrovich KD, Glass AA, et al. The lymphotoxin-beta receptor is necessary and sufficient for LIGHT-mediated apoptosis of tumor cells. *J Biol Chem.* 2000; 275(19):14307-14315. (Biology)

Tamada K, Shimozaki K, Chapoval AI, LIGHT, a TNF-like molecule, costimulates T cell proliferation and is required for dendritic cell-mediated allogeneic T cell response. *J Immunol.* 2000; 164(8):4105-4110. (Biology)

Zhai Y, Guo R, Hsu TL, et al. LIGHT, a novel ligand for lymphotoxin beta receptor and TR2/HVEM induces apoptosis and suppresses in vivo tumor formation via gene transfer.. J Clin Invest. 1998; 102(16):1142-1151. (Biology)

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