## **Product Data Sheet**

## PE anti-mouse IL-17RB

**Catalog # / Size:**  $1331525 / 25 \mu g$ 

1331530 / 100 µg

Clone: 9B10

**Isotype:** Rat IgG2a, κ

Immunogen: Recombinant mouse IL-17RB - Fc

chimera (human IgG1)

Reactivity: Mouse

**Preparation:** The antibody was purified by affinity

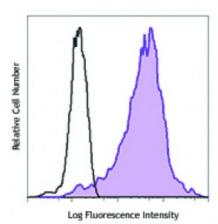
chromatography and conjugated with PE under optimal conditions. The solution is free of unconjugated PE and

unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.2



Mouse IL-17RB transfected 300.19 cells were stained with IL-17RB (clone 9B10) PE (filled histogram) or rat IgG2a, κ PE isotype control

(open histogram).

## **Applications:**

**Applications:** Flow Cytometry

Recommended

Usage:

Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is <0.25 microg per million cells in 100 microl, volume. It is

this reagent is ≤0.25 microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each

application.

Application References:

1. Wiesner DL, et al. 2015. PLoS Pathog. 11:1004701. PubMed

Description:

IL-17RB, also known as IL17RH1, belongs to the cytokine receptor family. IL-17RB possesses a unique intracellular signaling molecule called SEFIR, and is expressed similarily to fibroblast growth factor genes and IL-17R. It is reported to be expressed on iNKT cells, innate lymphoid cells (ILC), and Th2 cells. It binds IL-17B and IL-17E (IL-25) but not IL-17A or C. Its interaction with IL-25 has a higher affinity than that with IL-17B. Binding of IL-17RB and IL-25/IL-17E induces NF-kB mediated IL-8 production through interaction with TRAF6 and Act1. IL-17RB has been reported to play a role in autoimmune diseases such as rheumatoid arthritis and asthma.

Antigen References: 1. Hwang SY, et al. 2004. Arthritis Res. Ther. 6:R120. 2. Rickel EA, et al. 2008. J. Immunol. 181:4299.

3. Terashima A, et al. 2008. J. Exp. Med. 205:2727.