## PE/Cy7 anti-human TCR Vα24-Jα18 (iNKT cell)

Catalog # / Size: 2314560 / 100 tests

2314555 / 25 tests

Clone: 6B11

**Isotype:** Mouse IgG1, κ

Reactivity: Human

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

chromatography and conjugated with PE/Cy7 under optimal conditions. The solution is free of unconjugated PE/Cy7

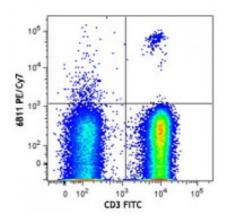
and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide and

0.2% (w/v) BSA (origin USA).

Concentration: Lot-specific



Human peripheral blood lymphocytes were stained with CD3 FITC and TCR V $\alpha$ 24-J $\alpha$ 18 (clone 6B11, top) PE/Cy7 or mouse IgG1,  $\kappa$  PE/Cy7 isotype control (bottom).

CD3 FITC

## **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 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for

each application.

Application Notes:

The 6B11 antibody recognizes the invariant CDR3 region of TCR  $V\alpha$ 24- $J\alpha$ Q.

Application References:

1. Rout N, et al. 2010. PLoS One 5:e9787. (FC)

Keterences:

**Description:** Encoded by the TCR Vα24-Jα18 germline configuration, Vα24-JαQ is expressed on a subset of NKT cells, namely invariant NKT (iNKT). Vα24-JαQ TCR interacts with the glycolipid loaded MHC class 1b molecule CD1d, inducing activation and subsequent cytoking production iNKT cells have been implicated in impune

subsequent cytokine production. iNKT cells have been implicated in immune regulation, tumor surveillance, and host response to pathogens. While iNKT cells occur at low frequency in the blood, assorted chemokines contribute to their

IgG1, K PE/Cy7

tissue homing potential.

Antigen References:

1. Thomas SY, et al. 2003. J. Immunol. 171:2571.

2. Exley MA, et al. 2008. Eur. J. Immunol. 38:1756.

3. Montoya CJ, et al. 2007. Immunology. 122:1.

4. Gansuvd B, et al. 2003.