PerCP/Cy5.5 anti-human TCR Vα24-Jα18 (iNKT cell)

Catalog # / Size: 2314565 / 25 tests

2314570 / 100 tests

Clone: 6B11

Isotype: Mouse IgG1, κ

Reactivity: Human

Preparation: The antibody was purified by affinity

chromatography and conjugated with PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated PerCP/Cy5.5 and unconjugated

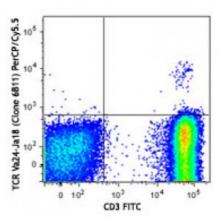
antibody.

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 $V\alpha24$ -J $\alpha18$ (clone 6B11, top) PerCP/Cy5.5 or mouse lgG1, κ PerCP/Cy5.5 isotype control (bottom).

10

104

0 102

10³ CD3 FITC

Mouse leg1, x PerCP/Cy5.5

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.

* PerCP/Cy5.5 has a maximum absorption of 482 nm and a maximum

emission of 690 nm.

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)

Description:

Encoded by the TCR V α 24-J α 18 germline configuration, V α 24-J α Q is expressed on a subset of NKT cells, namely invariant NKT (iNKT). V α 24-J α Q TCR interacts with the glycolipid loaded MHC class 1b molecule CD1d, inducing activation and 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

tissue homing potential.

Antigen 1. T

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

References: 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.