Product Data Sheet

PE/Cy7 anti-human TCR α/β

Catalog # / 2133595 / 25 tests

Size: 2133600 / 100 tests

Clone: IP26

Isotype: Mouse IgG1, κ

Reactivity: Human

Preparation: The antibody was purified by affinity

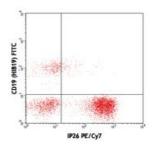
chromatography and conjugated with PE/Cy7 under optimal conditions. The solution is free of unconjugated PE/Cy7 and unconjugated antibody.

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 stained with CD19 (HIB19) FITC and IP26 PE/Cy7 (top) or mouse IgG1, κ PE/Cy7 isotype control (bottom)

IgG1.k PE/Cv7

Applications:

Applications: Flow Cytometry

Recommended Each lot of this

Usage:

Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis.

Test size products are

transitioning from 20 microL to 5 microL per test. Please check your vial or your CoA to find the suggested use of this reagent per million cells in 100 microL staining volume or per 100

microL of whole blood. It is

recommended that the reagent be titrated for optimal performance for

each application.

Application Notes:

Additional reported applications (for the relevant formats) include: T cell activation. When co-staining with anti-CD3, we recommend using clone UCHT1, since we have confirmed that IP26 does not compete with this clone. Other anti-CD3 clones may compete

out the binding of IP26.

Application References:

1. Schlossman S, et al. Eds. 1995. Leucocyte Typing V. Oxford University Press.

New York. (FC)

2. Joseph A, et al. 2008. J. Virol. 82:3078. (FC) <u>PubMed</u> 3. Pinto JP, et al. 2010. Immunology. 130:217. <u>PubMed</u>

Description: The IP26 antibody reacts with a monomorphic determinant of the α/β T-cell

receptor, which is expressed on greater than 95% of normal peripheral blood CD3⁺ T cells. The α/β TCR recognizes a peptide bound to MHC leading to T-cell

activation.

Antigen References:

1. Marchalonis J, et al. 2002. J. Mol. Recognit. 15:260.