## Pacific Blue™ anti-human CD56 (NCAM)

Catalog # / Size: 2191625 / 25 µg

2191630 / 100 µg

Clone:

Isotype: Mouse IgG1, κ

Reactivity: Human

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

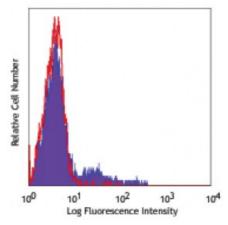
> chromatography, and conjugated with Pacific Blue<sup>™</sup> under optimal conditions. The solution is free of unconjugated

Pacific Blue™.

Formulation: Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.5



Human peripheral blood lymphocytes stained with HCD56 Pacific Blue™

## **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 ≤2.0 microg per million cells in 100 microL volume or 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

\* Pacific Blue™ has a maximum emission of 455 nm when it is excited at 405 nm. Prior to using Pacific Blue<sup>™</sup> conjugate for flow cytometric analysis, please verify your flow cytometer's capability of exciting and detecting the fluorochrome.

**Application** References: 1. Kishimoto T, et al. Eds. 1997. Leucocyte Typing VI. Garland Publishing Inc.

London

2. Correia DV, et al. 2011. Blood 118:992. (FC) PubMed

**Description:** CD56 is a single transmembrane glycoprotein also known as NCAM (Neural Cell

Adhesion Molecule), Leu-19, or NKH1. It is a member of the Ig superfamily. The 140 kD isoform is expressed on NK cells and NK-T cells. CD56 is also expressed in the brain (cerebellum and cortex) and at neuromuscular junctions. Certain large granular lymphocyte (LGL) leukemias, small-cell lung carcinomas, neuronal derived tumors, myelomas, and myeloid leukemias also express CD56. CD56 plays a role in homophilic and heterophilic adhesion via binding to itself or heparin sulfate.

**Antigen** References: 1. Lanier L, et al. 1991. J. Immunol. 146:4421. 2. Hemperly J, et al. 1990. J. Mol. Neurosci. 2:71.

3. Cremer H, et al. 1994. Nature 367:455.