Pacific Blue™ anti-human CD56 (NCAM)

Catalog # / Size: $2191630 / 100 \mu g$

2191625 / 25 µg

Clone: HCD56

Isotype: Mouse IgG1, κ

Reactivity: Human

Preparation: The antibody was purified by affinity

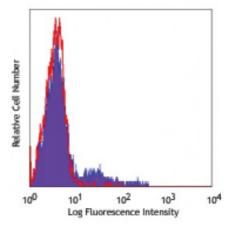
chromatography, and conjugated with Pacific Blue™ under optimal conditions. The solution is free of unconjugated

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™ 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.

Correia DV, et al. 2011. Blood 118:992. (FC) <u>PubMed</u>
Goodridge JP, et al. 2013. J Immunol. 191:3553. <u>PubMed</u>

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:

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

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