PE Anti-Human CD22 Monoclonal Antibody

 Catalog Number
 Vial Size

 H20221-09G
 25 tests

 H20221-09H
 100 tests



Market | 400-621-0003

marketing@sungenebiotech.com

Support | 022-66211636-8024

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Important Note: Centrifuge before opening to ensure complete recovery of vial contents. This product is guaranteed up to one year from purchase.

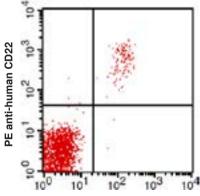
Purified Antibody Characterization

Clone	Isotype	Reactivity
HIB22	Mouse IgG1	Human

Description

CD22 is a 130 kD type I transmembrane glycoprotein also known as Siglec-2 and BL-CAM. It is a member of the immunoglobulin superfamily (sialoadhesion subgroup). CD22 is expressed in the cytoplasm of pro-B and pre-B cells, and on the surface of mature B and activated B cells, but not on plasma cells. CD22 is present in the B cell receptor complex and associates with SHP-1, Syk, Lck, Lyn, and phospholipase Cγ1. A primary function of CD22 is thought to be in limiting antigen receptor signaling by modulating B cell activation threshold. CD22 has been shown to bind to CD45RO and CD75, although the natural ligands for this molecule remain controversial

Illustration of Immunofluorescent Staining



FITC anti-human CD19

Human peripheral blood lymphocytes stained with FITC anti-human CD19 and PE anti-human CD22

Product Information

Conjugation: PE

Formulation: PBS pH 7.2, 0.09% NaN₃,

0.2% BSA

Storage: Keep as concentrated solution. Store at 4°C and protected from prolonged

exposure to light. Do not freeze.

Application: Recommended Application: FC

Usage: Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis (The amount of the reagent is suggested to be used from 20 μL to 5 μL per 100 μL of peripheral blood. Please check your vial). Since applications vary, the appropriate dilutions must be determined for individual use.

References

- [1] Schlossman, S., et al. Eds. 1995. Leukocyte Typing V:White Cell Differentiation Antigens. Oxford University Press. New York.
- [2] Clark, E., 1993. J. Immunol.. 150:4715.
- [3] Shan, D. and O. Press. 1995. J. Immunol.. 154:4466.

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