PE Rabbit anti-Human CD79b/Igβ mAb

ABclonal

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Catalog No.: A24251

Basic Information

Observed MW

Calculated MW

14kDa/26kDa

Category

Primary antibody

Applications

Cross-Reactivity

Human

CloneNo number

ARC61823

Conjugate

PE. Ex:565nm. Em:574nm.

Background

The B lymphocyte antigen receptor is a multimeric complex that includes the antigenspecific component, surface immunoglobulin (Ig). Surface Ig non-covalently associates with two other proteins, Iq-alpha and Iq-beta, which are necessary for expression and function of the B-cell antigen receptor. This gene encodes the Ig-beta protein of the B-cell antigen component. Alternatively spliced transcript variants encoding different isoforms have been described.

Recommended Dilutions

FC

5 µl per 10^6 cells in 100 μl volume

Immunogen Information

Gene ID 974

Swiss Prot

P40259

Immunogen

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

Synonyms

CD79B; AGM6; B29; IGB; CD79b molecule

Contact

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Product Information

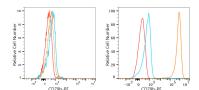
Source Isotype **Purification** Rabbit IgG Affinity purification

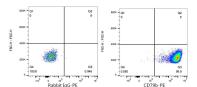
Storage

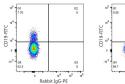
Store at 2-8°C. Avoid freeze.

Buffer: PBS containing 0.2% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

Validation Data









Flow cytometry: 1X10^6 HAP1 cells(negative control,left) and Daudi cells(right) were surface-stained with PE Rabbit anti-Human CD79b/lg β mAb(A24251,5 μ l/Test,orange line) or PE Rabbit IgG isotype control (5 μ l/Test,blue line).Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry:1X10^6 Daudi cells were surface-stained with PE Rabbit IgG isotype control (5 μ l/Test,left) or PE Rabbit anti-Human CD79b/lg β mAb(A24251,5 μ l/Test,right).

Flow cytometry:1X10^6 Human PBMC were surface-stained with FITC Rabbit anti-Human/Monkey CD19 mAb (A27989,5 μ I/Test) and PE Rabbit IgG isotype control (A24172,5 μ I/Test,left) or PE Rabbit anti-Human CD79b/Ig β mAb (A24251,5 μ I/Test,right). Cells in the lymphocyte gate were used for analysis.