

Product datasheet for CL033B

OriGene Technologies, Inc.

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Cd72 (CD72.1 alloantigen) Mouse Monoclonal Antibody [Clone ID: CT-72.1]

Product data:

Product Type: Primary Antibodies

Clone Name: CT-72.1

Applications: FC

Recommended Dilution: Flow Cytometry.

Reactivity: Mouse
Host: Mouse
Isotype: IgG2a

Clonality: Monoclonal

Specificity: This monoclonal antibody reacts with the CD72 alloantigen CD72.1, a B-cell surface protein

that is encoded by the Cd72a allele. CD72.1 is expressed on cells of the B cell lineage, except plasma cells.1 Mouse strains expressing CD72.1 include C57L/-, C58/-, DBA/1, DBA/2, and

SWR/J.

Formulation: PBS containing 0.09% sodium azide (NaN3) as preservative and EIA grade BSA as a stabilizing

protein to bring total protein concentration to 4-5 mg/ml.

Label: Biotin

State: Liquid purified Ig fraction

Concentration: lot specific
Conjugation: Biotin

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: CD72 antigen

Database Link: Entrez Gene 12517 Mouse

P21855

Synonyms: Lyb-2, Ly-32, Ly32, B-Cell marker





Note:

Protocol: **FLOW CYTOMETRY ANALYSIS:**

Method:

- 1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population using Lympholyte-M cell separation medium.
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of 2x10e7 cells/ml in media A. Add $50 \mu l$ of this suspension to each tube (each tube will then contain 1x10e6 cells, representing 1 test).
- 4. To each tube, add \sim 0.25 µg of this antibody per 10e6 cells.
- 5. Vortex the tubes to ensure thorough mixing of antibody and cells.
- 6. Incubate the tubes for 30 minutes at 4°C.
- 7. Wash 2 times at 4°C.
- 8. Add 100 μ l of detection reagent (Streptavidin-FITC) at 1/50 dilution.
- 9. Incubate the tubes at 4°C for 30-60 minutes.
- (It is recommended that the tubes are protected from light since most fluorochromes are light sensitive).
- 10. Wash 2 times at 4°C in media B.
- 11. Resuspend the cell pellet in 50 µl ice cold media B.
- 12. Transfer to suitable tubes for flow cytometric analysis containing 15 μ l of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

Media:

- A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 μ l of 2M sodium azide in 100 mls).
- B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 μ l of 2M sodium azide in 100 mls).