

## Product datasheet for **CL033BX**

### 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 (NaN <sub>3</sub> ) 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:	<a href="#">Entrez Gene 12517 Mouse P21855</a>
Synonyms:	Lyb-2, Ly-32, Ly32, B-Cell marker



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- 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  $2 \times 10^7$  cells/ml in media A. Add 50  $\mu$ l of this suspension to each tube (each tube will then contain  $1 \times 10^6$  cells, representing 1 test).
  4. To each tube, add  $\sim 0.25$   $\mu$ g of this antibody per  $10^6$  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  $\mu$ 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  $\mu$ l ice cold media B.
  12. Transfer to suitable tubes for flow cytometric analysis containing 15  $\mu$ 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  $\mu$ l of 2M sodium azide in 100 mls).
  - B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).