

Product datasheet for CL010P

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Cd8a Mouse Monoclonal Antibody [Clone ID: AD4(15)]

Product data:

Product Type: Primary Antibodies

Clone Name: AD4(15)
Applications: CT, FC

Recommended Dilution: Cytotoxicity assays.

Flow Cytometry.

Results:

Tissue Distribution by Flow Cytometry Analysis:

Mouse Strain: BALB/c

Cell Concentration: 1x106 cells per test.

Antibody Concentration Used: 2.0 µg /106 cells.

Isotypic Control: Mouse IgM.

Cell Source (Percentage of cells stained above control): Spleen (10.2%), Thymus (66.0%).

Reactivity: Mouse
Host: Mouse
Isotype: IgM

Clonality: Monoclonal Immunogen: C57BL/6

Donor: B6-Ly-2a spleen

Fusion Partner: Myeloma P3/X63-Ag8

Specificity: Anti-mouse CD8a (Ly 2.2) monoclonal antibody reacts with a sub-population of T-

lymphocytes from Mouse strains expressing the Ly-2.2 phenotype but does not react with

lymphocytes from strains expressing the Ly-2.1 phenotype.

Formulation: PBS

State: Purified

State: Liquid purified IgG fraction Preservative: 0.02% Soidium Azide

Concentration: lot specific

Purification: Euglobin precipitation

Conjugation: Unconjugated



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Storage: Store 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: CD8 antigen, alpha chain

Database Link: Entrez Gene 12525 Mouse

P01731

Synonyms: CD8 alpha chain, CD8A, MAL

Note: Protocol: FLOW CYTOMETRY ANALYSIS:

Method:

- 1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte®-M cell separation medium.
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of $2x10^{\circ}$ cells/ml in media A. Add 50 μ l of this suspension to each tube (each tube will then contain $1x10^{6}$ cells, representing 1 test).
- 4. To each tube, add 2.0 μg of this antibody*Cat.-No* CL010P.
- 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 μl of secondary antibody (FITC Goat anti-mouse IgM-3 (H+L)) at 1/500 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.

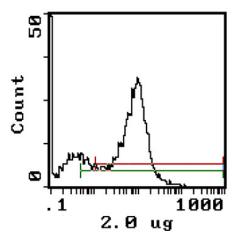
Media:

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).



Product images:



Cell Source: Thymus. Percentage of cells stained above control: 66.0%