

Product datasheet for CL008F

Cd8a Rat Monoclonal Antibody [Clone ID: CT-CD8a]

Product data:

Immunogen:

Product Type: Primary Antibodies

Clone Name: CT-CD8a

Applications: FC

Recommended Dilution: Flow Cytometry Analysis (see Protocols).

Murine thymocytes

Reactivity: Mouse

Host: Rat

Isotype: IgG2a

Clonality: Monoclonal

Specificity: This anti-mouse CD8a antigen monoclonal antibody recognizes the mouse CD8α chain. The α

chain of CD8 associates with the CD8 β chain to form a CD8 α / β heterodimer that is expressed by the majority of thymocytes and by the MHC class I restricted subset of mature T cells1. Mouse CD8 α can also form a CD8 α / α chain homodimer on subsets of CD8 positive cells. For this reason antibodies specific for CD8a rather than CD8b are recommended for a rigorous

delineation of CD8 positive cells.

Formulation: PBS, 0.09% NaN3 and EIA grade BSA as a stabilizing protein to bring total protein

concentration to 4-5 mg/ml

Label: FITC

State: Liquid purified IgG

Concentration: lot specific

Purification: Protein G Chromatography

Conjugation: FITC

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.

This product is photosensitive and should be proteced from light.

Stability: Shelf life: one year from despatch.

Gene Name: CD8 antigen, alpha chain



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Database Link: Entrez Gene 12525 Mouse

P01731

Background: The CD8 antigen is a cell surface glycoprotein found on most cytotoxic T lymphocytes that

mediates efficient cell to cell interactions within the immune system. The CD8 antigen, acting as a coreceptor, and the T cell receptor on the T lymphocyte recognize antigen displayed by an antigen presenting cell (APC) in the context of class I MHC molecules. The functional coreceptor is either a homodimer composed of two alpha chains, or a heterodimer composed of one alpha and one beta chain. Both alpha and beta chains share significant

homology to immunoglobulin variable light chains.

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 2x10e7 cells/ml in media A. Add $50 \mu l$ of this suspension to each tube (each tube will then contain $1 \times 10e6$ cells, representing 1 test).
- 4. To each tube, add \sim 1.0 µg* of this Ab per 1x10e6 cells.
- 5. Vortex the tubes to ensure thorough mixing of antibody and cells.
- 6. Incubate the tubes for 30 minutes at 4°C. (It is recommended that the tubes are protected from light, since most fluorochromes are light sensitive.)
- 7. Wash 2 times at 4°C.
- 8. Resuspend the cell pellet in 50 µl ice cold media B.
- 9. 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).

Results:

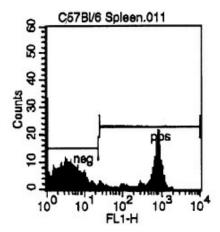
Mouse Strain: C57BL/6

<u>Cell Concentration</u>: 1x10e6 cells per test <u>Antibody Concentration Used</u>: 1.0 µg/10e6 cells

Isotypic Control: FITC Rat IgG2a



Product images:



Representative Histogram - Cell Source: CD3e Positive Spleen Cells Percentage of cells stained above control: 34.7%