

Product datasheet for **CL067R**

MHC Class II I-Abd Mouse Monoclonal Antibody [Clone ID: 28-16-8S]

Product data:

Product Type:	Primary Antibodies
Clone Name:	28-16-8S
Applications:	FC
Recommended Dilution:	Flow Cytometry (See Protocols).
Reactivity:	Mouse
Host:	Mouse
Isotype:	IgM
Clonality:	Monoclonal
Immunogen:	C3H.SW spleen Donor: C3H Fusion Partner: SP2/0Ag.14
Specificity:	Recognizes Mouse MHC class II I-Ab and cross reacts with I-Ad. This Monoclonal Antibody reacts with the I-Ab encoded MHC class II antigen expressed on Mouse strains of the H-2b haplotype. It also reacts with the I-Ad encoded MHC class II antigen expressed on Mouse strains of the H-2d haplotype. Class II antigens are most highly expressed on antigen presenting cells including B cells, macrophages, dendritic cells and certain epithelial cells.
Formulation:	PBS containing 0.02% Sodium Azide as a preservative as a and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml Label: PE State: Liquid purified IgM fraction
Concentration:	lot specific
Conjugation:	PE
Storage:	Store the antibody undiluted at 2-8°C. DO NOT FREEZE! This product is photosensitive and should be protected from light. Avoid repeated freezing and thawing.
Stability:	Shelf life: one year from despatch.



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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 with Lympholyte®-M cell separation medium.
2. Wash 2 times.
3. Resuspend the cells to a concentration of 2×10^7 cells/ml in media A. Add 50 μ l of this suspension to each tube (each tube will then contain 1×10^6 cells, representing 1 test).
4. To each tube, add ~ 0.2 μ g of CL067R 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.
(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 μ l 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 ml of 2M sodium azide in 100 mls).

B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 ml of 2M sodium azide in 100 mls).

Results:

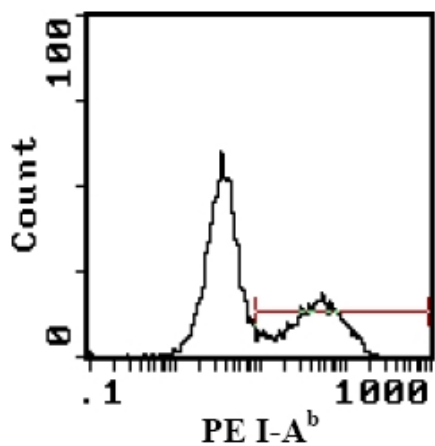
Tissue Distribution by Flow Cytometry Analysis:

Mouse strain: C57BL/6
Cell Concentration: 1×10^6 cells per test.
Antibody Concentration Used: 0.2 μ g/ 10^6 cells.
Isotypic Control: PE Mouse IgM

Cell Source: Percentage Stained Above Control:

Thymus: 28.2%
Spleen: 40.7%

Product images:



Cell Source: Spleen