

Product datasheet for **SM083FS**

MHC Class II I-Ak Mouse Monoclonal Antibody [Clone ID: 14V.18]

Product data:

Product Type:	Primary Antibodies
Clone Name:	14V.18
Applications:	FC
Recommended Dilution:	Flow Cytometry (See Protocols).
Reactivity:	Mouse
Host:	Mouse
Isotype:	IgG2a
Clonality:	Monoclonal
Immunogen:	A.TL spleen. Donor: A.TH spleen. Fusion Partner: P3-X63-Ag8
Specificity:	This Monoclonal antibody is a cytotoxic antibody specific for cells expressing the Ia antigen coded for by the A subregion of the k haplotype. The reaction pattern of this antibody with a panel of inbred and recombinant haplotypes demonstrates that the antibody reacts with Ia.m2, a private specificity of the H-2k haplotype. This antibody can be used to quantitate or to eliminate cells bearing the I-Ak (Ia.m2) antigen.
Formulation:	PBS containing 0.02% Sodium Azide as preservative and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml Label: FITC State: Liquid purified IgG fraction Label: Fluorescein
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 protected from light.
Stability:	Shelf life: one year from despatch.



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Database Link: [P01910](#)

Synonyms: H2-Aa

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 1.0 μ g* of SM083FS 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 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:

Tissue Distribution by Flow Cytometry Analysis:

Mouse Strain: A.TL

Cell Concentration : 1×10^6 cells per test.

Antibody Concentration Used: 1.0 μ g/ 10^6 cells.

Isotypic Control: FITC Mouse IgG2a

Cell Source Percentage of cells stained above control:

Thymus: 32.5%

Spleen: 75.4%

Lymph Node: 32.1%

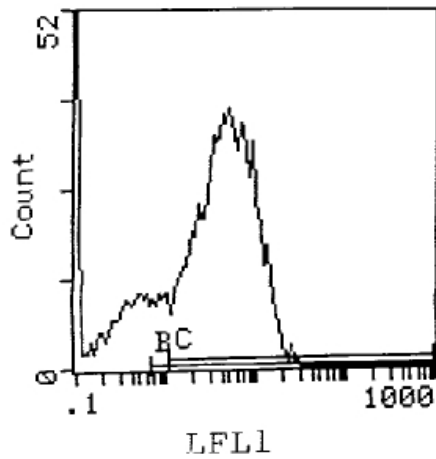
Bone Marrow: 21.6%

Strain Distribution by Flow Cytometry Analysis:

Cell Concentration : 1×10^6 cells per test.

Antibody Concentration Used: 1.0 μ g / 10^6 cells.

Product images:



Cell Source: Spleen. Percentage of cells stained above control: 75.4%

Haplotype

Strain	K	A	B	J	E	C	S	G	D	+/-
A.TH	s	s	s	s	s	s	s	s	d	-
A.TL	s	k	k	k	k	k	k	k	d	+
B10.A	k	k	k	k	k	d	d	d	d	+
B10.A(4R)	k	k	b	b	b	b	b	b	b	+
B10.BR	k	k	k	k	k	k	k	k	k	+
B10.D2	d	d	d	d	d	d	d	d	d	-
C3H/He	k	k	k	k	k	k	k	k	k	+
C57BL/6		b	b	b	b	b	b	b	b	-