

#### OriGene Technologies, Inc.

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# Product datasheet for AM08081RP-S

## MHC Class I H2 Kd/Dd Mouse Monoclonal Antibody [Clone ID: 34-1-2S]

#### **Product data:**

Product Type:	Primary Antibodies
Clone Name:	34-1-25
Applications:	FC
Recommended Dilution:	Flow Cytometry.
Reactivity:	Mouse
Host:	Mouse
lsotype:	lgG2a
Clonality:	Monoclonal
Immunogen:	BDF splenocytes <u>Donor:</u> C3H spleen <u>Fusion Partner:</u> Sp2/0-Ag14
Specificity:	Anti H-2KdDd mAb reacts with both H-2Kd and H-2Dd products. The antibody also cross reacts with Kbsrpq. This K-D cross reaction indicates the presence of shared specificities between the two separate H-2 regions.
Formulation:	PBS containing 0.02% sodium azide (NaN3) as preservative and EIA grade BSA as a stabilizing protein Label: PE State: Liquid purified lg fraction Label: R - Phycoerythrin
Concentration:	lot specific
Purification:	Affinity chromatography on Protein G
Conjugation:	PE
Storage:	Store the antibody undiluted at 2-8°C. <b>DO NOT FREEZE!</b>
Stability:	Shelf life: one year from despatch.



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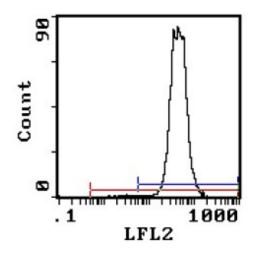
	MHC Class I H2 Kd/Dd Mouse Monoclonal Antibody [Clone ID: 34-1-2S] – AM08081RP-S
Note:	Protocol: FLOW CYTOMETRY ANALYSIS:
	Method:
	1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte®-M cell separation medium. 2. Wash 2 times.
	3. Resuspend the cells to a concentration of 2x10e7 cells/ml in media A. Add 50 μl of this suspension to each tube (each tube will then contain 1 x 10e6 cells, representing 1 test).
	4. To each tube, add 0.2-0.5 μg of this antibody per 10e6 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 µl 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 μl of 2M sodium azide in 100 mls).
	B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100
	μl of 2M sodium azide in 100 mls).
	Results:
	Tissue Distribution by Flow Cytometry Analysis:
	Mouse Strain: BALB/c
	Cell Concentration: 1x10e6 cells per test
	Antibody Concentration Used: 0.5 µg/10e6 cells
	Isotypic Control: PE Mouse IgG2a
	Percentage of cells stained above control:

Thymus 86.3%

Spleen 97.2%

Lymph Node 100%

## **Product images:**



Cell Source: Spleen Percentage of cells stained above control: 97.2%

Strain Distribution	by Flow Cytometry Analysis:	
Procedure: As abov	The second s	
	ation: 0.2 µg/10 <sup>6</sup> cells	
Strains Tested:		
Strain	H-2 Loci Alleles	+/-
	$\underline{K} \underline{A}_{B} \underline{A}_{n} \underline{E}_{B} \underline{E}_{n} \underline{C4} \underline{C4S} \underline{D}$	
C3H/He	k k k k k k k	-
C57BL/6	b b b b b b b	(+/-)
BALB/c	d d d d d d d	+
DBA/1	99999999	+
SJL	<b>S S S S S S S S</b>	+
B10.M	ff f f f f f	(+/-)
A.TH	ssssss sd	+
A.TL	skkkkkd	+
B10.A(3R)	b b b b/k k d d d	+
P/J	ррррррр р	(+/-)

For a more detailed strain distribution - see reference 1.

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