

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Product datasheet for CL060PE

MHC Class I H-2Ld Mouse Monoclonal Antibody [Clone ID: 30-5-7S]

Product data:

Product Type:	Primary Antibodies	
Clone Name:	30-5-7S	
Applications:	FC	
Recommended Dilution:	Flow Cytometry (see protocol).	
Reactivity:	Mouse	
Host:	Mouse	
lsotype:	lgG2a	
Clonality:	Monoclonal	
Immunogen:	Recipient: BALB/c-H-2dm2 Donor: BALB/c Spleen cells Fusion Partner: SP2/0.Ag14	
Specificity:	This monoclonal antibody detects the public specificity H-2.65 of the H-2Ld antigen. This antibody also recognizes H-2Dq and H-2Lq molecules.	
Formulation:	PBS containing 0.02% Sodium Azide as preservative and EIA grade BSA as stabilizing protein to bring the protein concentration to 4-5 mg/ml Label: PE State: Liquid purified IgG fraction	
Conjugation:	PE	
Storage:	Store the antibody undiluted at 2-8°C. DO NOT FREEZE! Avoid prolonged exposure to Light	
Stability:	Shelf life: one year from despatch.	
Database Link:	<u>P01897</u>	
Synonyms:	H2-L	



This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2022 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US

	~ \$%
\cup	ORIGENE

Note:

Protocol: Flow Cytometry Analysis:

Method:

1. Prepare cell suspension in Media A. For cell preparations, deplete the red blood cell population with Lympholyte®-M cell separation media.

2. Wash 2 times.

3. Resuspend the cells to a concentration of 2x10e7 cells/ml in media A. Add 50 μ l of this

suspension to each tube (each tube will then contain $1 \times 10e6$ cells, representing 1 test). 4. To each tube add 2.0 µg of this Antibody CL060PE 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 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 phosphate buffered saline. (This stains dead cells by intercalating DNA.)

MEDIA:

A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 μ l of 2 M sodium azide in 100 mls).

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

Results in Tissue Distribution by Flow Cytometry Analysis:

<u>Mouse Strain</u>: BALB/c <u>Cell Concentration</u>: 1x10e6 cells/test <u>Antibody Concentration Used</u>: 2.0 µg/10e6 cells <u>Isotype Control</u>: PE Mouse IgG2a

Cell Source - % of cells stained above Control:

Spleen: 97.3% Thymus: 87.1% Lymph Node: 98.5%

Strain Distribution:

Procedure: As above Antibody Concentration: 1/20 Strains Tested: See **Figure 2**

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2022 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US 

Product images:

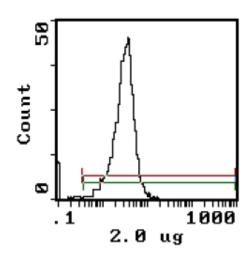


Figure 1.

Strain	<u>Haplotype</u>	+/-
BALB/c	$H-2^{d}$	+
A.TH	H-2K⁵D [₫]	+
A.TL	H-2K⁵D [₫]	+
B.10A(3R)	$H-2K^{b}D^{d}$	+
C3H/He	$H-2^{k}$	-
C57BL/6	H-2 ^b	-

Figure 2.

This product is to be used for laboratory only. Not for diagnostic or therapeutic use. ©2022 OriGene Technologies, Inc., 9620 Medical Center Drive, Ste 200, Rockville, MD 20850, US