

#### 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 AM31873RP-N

## CD45 / LCA (CD45RC) Rat Monoclonal Antibody [Clone ID: IBL-8]

#### **Product data:**

Product Type:	Primary Antibodies
Clone Name:	IBL-8
Applications:	FC
Recommended Dilution:	Immunohistochemistry on acetone-fixed frozen sections. Flow Cytometry. Immunoprecipitation.
Reactivity:	Mouse
Host:	Rat
lsotype:	lgG1
Clonality:	Monoclonal
Immunogen:	Mouse Spleen Cells <u>Donor:</u> Wistar spleen <u>Fusion Partner:</u> Sp-2/0 Ag14
Specificity:	Anti-mouse CD45RC is against the exon C-dependent RC isoform and reacts strongly with B cells, and less intensely with most CD8+ T cells. It does not recognize CD4+ T cells. Also, myeloid cells do not express the RC isoform.
Formulation:	PBS containing 0.02% sodium azide (NaN3) as preservative and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml. Label: PE State: Liquid purified Ig fraction Label: R - Phycoerythrin
Concentration:	lot specific
Purification:	Affinity purified
Conjugation:	PE
Storage:	Store the antibody undiluted at 2-8°C. <b>DO NOT FREEZE!</b>
Stability:	Shelf life: one year from despatch.
Gene Name:	protein tyrosine phosphatase, receptor type, C



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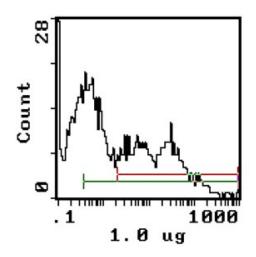
Database Link:	Entrez Gene 19264 Mouse
	<u>P06800</u>
Background:	CD45 (L-CA) is a transmembrane phosphotyrosine phosphatase expressed on leukocytes.
Synonyms:	PTPRC, Leukocyte common antigen, L-CA, T200
Note:	Protocol: FLOW CYTOMETRY ANALYSIS:
	Method:
	1. Prepare a cell suspension in media A. For cell 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 1.0 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 $\mu$ l 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 (10
	μl of 2M sodium azide in 100 mls).
	B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 $\mu$ l c
	2M sodium azide in 100 mls).
	Results: Tissue Distribution by Flow Outomatry Analysis
	Tissue Distribution by Flow Cytometry Analysis: <u>Mouse Strain:</u> BALB/c
	<u>Cell Concentration:</u> 1x10e6 cells per test
	Antibody Concentration Used: 1.0 µg/10e6 cells
	Isotypic Control: PE Bat IgG1

Isotypic Control: PE Rat IgG1

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## **Product images:**



Percentage of cells stained above control: 34.8%

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