

# Product datasheet for AM31873RP-L

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OriGene Technologies, Inc.

## 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 Isotype: IgG1

Clonality: Monoclonal

Immunogen: Mouse Spleen Cells

**Donor:** Wistar spleen

Fusion Partner: 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.

DO NOT FREEZE!

**Stability:** Shelf life: one year from despatch.

**Gene Name:** protein tyrosine phosphatase, receptor type, C





### CD45 / LCA (CD45RC) Rat Monoclonal Antibody [Clone ID: IBL-8] - AM31873RP-L

Database Link: Entrez Gene 19264 Mouse

P06800

**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  $\mu$ l 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 µl ice cold media B.
- 9. Transfer to suitable tubes for flow cytometric analysis containing 15  $\mu$ 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  $\mu$ l of 2M sodium azide in 100 mls).

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

#### **Results:**

Tissue Distribution by Flow Cytometry Analysis:

Mouse Strain: BALB/c

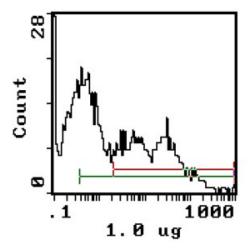
Cell Concentration: 1x10e6 cells per test

Antibody Concentration Used: 1.0 µg/10e6 cells

<u>Isotypic Control:</u> PE Rat IgG1



# **Product images:**



Percentage of cells stained above control: 34.8%