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Place your order with CEDARLANE® or your local distributor.

Please contact CEDARLANE® for lot specific information.

**FITC Anti-Rat RT1.A^c
Monoclonal Antibody**

**CL027F
CL027F-5
LOT: 2731**

DESCRIPTION:

Cedarlane's anti-rat RT1.A^c recognizes a polymorphic determinant of the MHC class I antigen in the rat ("c" and "n" haplotypes). CL027F is an excellent antibody for labelling cells of donor or host origin in bone marrow chimeras. (1,2)

PRESENTATION:

100 µg (CL027F) or 500 µg (CL027F-5) FITC conjugated Ig buffered in PBS, 0.02% NaN₃ and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml.

STORAGE/STABILITY:

Store at 4°C. For long term storage, aliquot and freeze unused portion at -20°C in volumes appropriate for single usage. Avoid freeze/thaw cycles. Avoid prolonged exposure to light.

For more information or to place an order please contact...

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website: www.cedarlanelabs.com • e-mail: info@cedarlanelabs.com

SPECIFICATIONS:

Clone: MRC OX-27

Hybridoma Production:

Immunization: Immunogen: Phytohaemagglutinin Blasts
Donor: BALB/c spleen

Fusion Partner: P3-NS1-1-AG4 (NS1/1)

Specificity: Rat RT1.A^c (rat MHC Class I polymorphic)

Ig Class: Mouse IgG_{2a}

Format: FITC conjugated Ig buffered in PBS, 0.02% NaN₃ and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml. (Purified from ascitic fluid via Protein G Chromatography)

Antibody Concentration: 0.1 mg/ml

FLOW CYTOMETRY ANALYSIS:

Method:

1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte[®]-Rat cell separation medium (CL5040).
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 0.5-0.2 μ g* of **CL027F** or **CL027F-5** 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:

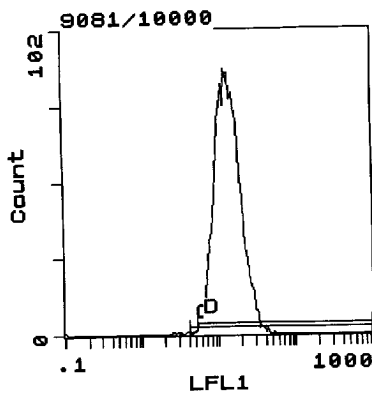
Rat Strain: Brown Norway

Cell Concentration : 1×10^6 cells per test

Antibody Concentration Used: $0.2 \mu\text{g}/10^6$ cells

Isotypic Control: FITC Mouse IgG_{2a}

<u>Cell Source</u>	<u>Percentage of cells stained above control:</u>
Thymus	73.9%
Spleen	99.0%
Lymph Node	100%



Cell Source: Spleen

Percentage of cells stained above control: 99.0%

N.B. Appropriate control samples should always be included in any labeling studies.

*** For optimal results in various applications, it is recommended that each investigator determine dilutions appropriate for individual use.**

Strain Distribution by Flow Cytometry Analysis:

Procedure: see page 2

Cell Concentration : 1×10^6 cells per test

Antibody Concentration Used: $0.5 \mu\text{g}/10^6$ cells

Strains Tested: Wistar, Buffalo, Brown Norway, Fischer, ACI

Positive: Brown Norway

Negative: Wistar, Buffalo, Fischer, ACI

REFERENCES:

- 1) Butcher, G.W. (1987) 19: 3-21. Monoclonal antibodies specific for alloantigens of the rat. Rat Membrane Alloantigens News.
- 2) Jeffries, W.A. et al. (1985) J. Exp. Med. 162: 117-127. Authentic T Helper CD4 (W3/25) antigen on rat peritoneal macrophages.

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