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# TECHNICALLY *Speaking*

Place your order with CEDARLANE® or your local distributor.

*Please contact CEDARLANE® for lot specific information.*

## **FITC Anti-Rat $\gamma\delta$ TCR Monoclonal Antibody**

**CL056F**

**CL056F-5**

**LOT: 5631**

### **DESCRIPTION:**

Cedarlane's anti-rat  $\gamma\delta$  T cell receptor monoclonal antibody detects a T cell-specific heterodimeric 48 and 50 kD cell surface protein that is expressed on greater than 90% of CD3<sup>+</sup>  $\alpha\beta$ TCR<sup>-</sup> rat peripheral T lymphocytes, and identifies a dense network of dendritic cells in the epidermis as  $\gamma\delta$  T cells. Immobilized this antibody induces a strong proliferative response in  $\gamma\delta$  T cell cultures supplemented with either IL-2 or IL-4.

Applications: Flow cytometry, immunohistochemical staining on frozen sections.

### **PRESENTATION:**

100  $\mu$ g (CL056F) or 500  $\mu$ g (CL056F-5) FITC conjugated Ig buffered in PBS, 0.02% NaN<sub>3</sub> 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...

**CEDARLANE®**  
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**SPECIFICATIONS:**

Clone: V65

Hybridoma Production:

Immunization: Immunogen: Rat T Blasts  
Donor: BALB/c spleen

Fusion Partner: X63-Ag 8.653

Specificity: Rat  $\gamma\delta$  TCR

Ig Class: Mouse IgG<sub>1</sub>

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<sup>®</sup>-Rat cell separation medium (CL5040).
2. Wash 2 times.
3. Resuspend the cells to a concentration of  $2 \times 10^7$  cells/ml in media A. Add 50  $\mu$ l of this suspension to each tube (each tube will then contain  $1 \times 10^6$  cells, representing 1 test).
4. To each tube, add 0.2 - 0.5  $\mu$ g\* of **CL056F or CL056F-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  $\mu$ l 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 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:

Rat Strain: Fischer

Cell Concentration :  $1 \times 10^6$  cells per test

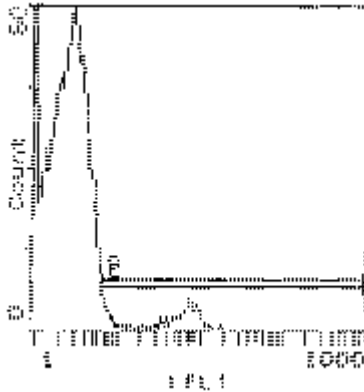
Antibody Concentration Used: 0.2 µg/ $10^6$  cells

Isotypic Control: FITC Mouse IgG<sub>1</sub>

Cell SourcePercentage of cells stained above control:

Thymus	4.1%
Splenic T Cells*	6.1%

\*(T cells isolated with CL102 - Cedarlane's Rat T Cell Recovery Column Kit)



Cell Source: Splenic T cells

Percentage of cells stained above control: 6.1%

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

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

**REFERENCES:**

1. Kühnlein, P., Park, H-J., Herrmann, T., Elbe, A., and Hünig, T. (1994). Identification and Characterization of Rat  $\gamma\delta$  T-Lymphocytes in Peripheral Lymphoid Organs, Small Intestine, and Skin With a Monoclonal Antibody to a Constant Determinant of the  $\gamma\delta$  T cell Receptor. *J. Immunol.* 153:979-986.

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