

# Produktinformation



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# Lieferung & Zahlungsart

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Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

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Please contact CEDARLANE® for lot specific information.

### FITC Anti-Mouse γδ TCR **Monoclonal Antibody**

**CL7201F** CL7201F-3 LOT: 7131

#### **DESCRIPTION:**

Cedarlane's anti-mouse  $\gamma\delta$  T cell receptor monoclonal antibody reacts with the surface on all γδ TCR bearing cells and does not react with receptors on αβ TCR positive cells. It is thought that this clone may be specific for a determinant present on Cδ<sup>7</sup>. The γδ T cell receptors are present on murine CD4-CD8- thymocytes, peripheral T cells, intestinal CD8<sup>+</sup> intraepithelial lymphocytes and Thy 1<sup>+</sup> dendritic epidermal cells in the skin <sup>1</sup>.

Use of this antibody in conjunction with an anti-CD3 monoclonal antibody (Cedarlane's anti-CD3E Monoclonal Antibody CL7202F) allows for accurate measurements of the mutually exclusive sub-populations of  $\gamma\delta$  TCR and  $\alpha\beta$  TCR bearing T cells. Cedarlane's anti mouse γδ TCR monoclonal antibody has also been used successfully for the characterization of murine intraepithelial lymphocytes.

This clone is reported to work with frozen sections<sup>6</sup>.

#### PRESENTATION:

100 μg (CL7201F) or 300 μg (CL7201F-3) 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.

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### **SPECIFICATIONS:**

Clone: GL-3

**Hybridoma Production**:

Immunization: Immunogen:C57BL/6 intraepithelial lymphocytes

Donor: Armenian Hamster.

Fusion Partner: Murine myeloma cell line SP2/0

Specificity: Mouse γδ T cell receptor

Ig Class: Hamster IgG

<u>Format</u>: 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. (Purified from ascitic fluid via Protein G Chromatography)

Antibody Concentration: 0.1 mg/ml

### FLOW CYTOMETRY ANALYSIS:

#### Method:

- Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte<sup>®</sup>-M cell separation medium (CL5030).
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of  $2x10^7$  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 1.0 μg\* of **CL7201F or CL7201F-3** per 10<sup>6</sup> 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 flurochromes 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 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 (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:

### <u>Tissue Distribution by Flow Cytometry Analysis:</u>

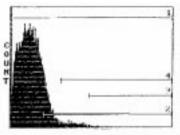
Mouse Strain: CBA/J

Cell Concentration :  $1x10^6$  cells per tests Antibody Concentration Used:  $1.0 \mu g/10^6$  cells

Isotypic Control: FITC Hamster IgG

Cell Source	Percentage of cells stained above control:
Thymus	3.7%
Splenic T Cells*	3.9 %

\*(T cells isolated with CL101 - Cedarlane's Mouse T Cell Recovery Column Kit)



LFL1

Cell Source: Splenic T Cells Percentage of cells stained above control: 3.9%

# 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 :  $1x10^6$  cells per tests Antibody Concentration Used:  $1.0 \mu g/10^6$  cells

Strains Tested: C57BL/6, CBA/J, BALB/c, AKR, C3H/He Positive: C57BL/6, CBA/J, BALB/c, AKR, C3H/He

Negative: none

#### **REFERENCES:**

- 1. Brenner et al. 1986. Indentification of a putative Second T Cell receptor. Nature (Lond.) 322:145.
- Cron. R & et al. 1988. A functional subpopulation of peripheral murine T lymphocytes which express a novel T Cell Structure. J. Immunol. 141:1074.
- 3. Nakawishii, N.K. et al. 1987. Tγ protein is expressed on fetal thymocytes as a disulphide linked heterodimer. Nature (Lond.) 325:720.
- 4. Sowder et al. 1988. A large subpopulation of avian T cells express a homologue of the mammalian  $T\gamma/\delta$  receptor. J. Exp. Med. 167:315.
- Goodman, T & L. Lefrancois. 1988. Expression of the γδ TCR on intestinal CD8<sup>+</sup> intraepithelial lymphocytes. Nature (Lond.) 333:855.
- 6. Skarstein, K. et al. 1994. Oligoclonality of T cells in salivary glands of autoimmune MRL/*lpr* mice. Immunology. **81**:497-501.
- 7. Goodman, T & L. Lefrancois. 1989. Intraepithelial Lymphocytes. J. Exp. Med. Vol. 170:1569-1581.

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