

US office: Acris Antibodies, Inc. San Diego, CA UNITED STATES Phone: +1-858-888-7900 Fax: +1-858-888-7904 US-info@acris-antibodies.com CL039R Acris Antibodies GmbH

Schillerstr. 5 32052 Herford GERMANY Phone: +49-5221-34606-0 Fax: +49-5221-34606-11 info@acris-antibodies.com



# Monoclonal Antibody to CD90 - PE

| Alternate names: | CDw90, THY1, Thy-1, Thy-1 membrane glycoprotein   |
|------------------|---|
| Catalog No.:     | CL039R  |
| Quantity:        | 50 µg   |
| Concentration:   | 0.1 mg/ml   |
| Background:      | CD90 (Thy1) antigen is a GPI linked glycoprotein member of the Immunoglobulin<br>superfamily. It is expressed on murine T cells, thymocytes, neural cells, cells of<br>granulocytic lineage, early hematopoietic progenitors, fibroblasts, neurons and Kupffer's<br>cells. Thy1 may play a role in cell to cell or cell to ligand interactions during synaptogenesis<br>and other events in the brain. It is found in most mouse strains except AKR/J, A, Thy1.1 and<br>B6.PL (74NS) expressing Thy1.1. |
| Uniprot ID:      | <u>P01831</u>   |
| NCBI:            | <u>NP_033408.1</u>  |
| GenelD:          | <u>21838</u>  |
| Host / Isotype:  | Mouse / IgG2b   |
| Clone:           | 5a-8  |
| Immunogen:       | CBA/J.<br>Donor: AKR/J Spleen.<br>Fusion Partner: Spleen from immunized recipient fused with myeloma P3-NSI-1-Ag4-1.  |
| Format:          | State: Liquid purified IgG fraction.<br>Purification: Protein G Chromatography.<br>Buffer System: PBS containing 0.02% Sodium Azide as preservative and EIA grade BSA as<br>a stabilizing protein to bring total protein concentration to 4-5 mg/ml.<br>Label: PE   |
| Applications:    | <b>Flow Cytometry.</b><br>Other applications not tested. Optimal dilutions are dependent on conditions and should<br>be determined by the user.   |
| Specificity:     | This monoclonal antibody reacts with all T lymphocytes from mouse strains expressing the<br>Thy 1.2 phenotype (e.g. C57BL/6, C3H/He, DBA/2, CBA/J, BALB/c), but does not react with<br>lymphocytes expressing the Thy 1.1 phenotype [e.g. AKR/J, B6.PL(74NS)].<br><b>Species:</b> Mouse.<br>Other species not tested.   |
| Storage:         | Store the antibody undiluted at 2-8°C.<br><b>DO NOT FREEZE!</b><br>This product is photosensitive and should be protected from light.<br>Shelf life: one year from despatch.  |
|                  | For research and in vitro use only. Not for diagnostic or therapeutic work.   |

**For research and in vitro use only. Not for diagnostic or therapeutic work.** Material Safety Datasheets are available at www.acris-antibodies.com or on request.

Antibody Hotline - Technical Questions - Antibody Location Service Free Call: 0800-2274746 (Germany only) - www.acris-antibodies.com



General References: 1. Krieg, A., Gourley, M. and Steinberg, A. 1991. Association of Murine Lupus and Thymic Full-Length Endeneous Retroviral Expression Maps To A Bone Marrow Stem Cell. J. Immunol. 146:3002-3005.

2. Haba, S. and Nisonoff, A., 1991. Induction of Tolerance To Syngeneic IgE In Neonatal Mice. J. Immunol. 146:807-811. 3. Miyajima, H., Takao, H., et al. 1991. Suppression By IL-2 Of IgE Production By B Cells Stimulated By IL-4. J. Immunol. 146:457- 462.

4. Kruger, M. and Riley, R. 1990. The Age-Dependent Loss Of Bone Marrow B Cell Precursors In Autoimmune NZ Mice Results From Decreased Mitotic Activity, But Not From Inherent Stromal Cell Defects. J. Immunol. 144:103-110.

5. Fine, J., Siverstone, A. and Gasiewicz, T. 1990. Impairment Of Prothymocyte Activity By 2,3,7,8-Tetrachlorocibenzo-p-Dioxin. J. Immunol. 144:1169-1176.

#### Protocols: 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 1x10e6 cells, representing 1 test). 4. To each tube, add 0.5 µg\* of this Ab.

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  $\mu$ 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:**

<u>Mouse Strain</u>: BALB/c <u>Cell Concentration</u>: 1x10e6 cells per test <u>Antibody Concentration Used</u>: 0.5 µg/10e6 cells <u>Isotypic Control</u>: PE-Mouse IgG2b,k

## Cell Source Percentage of cells stained above control:

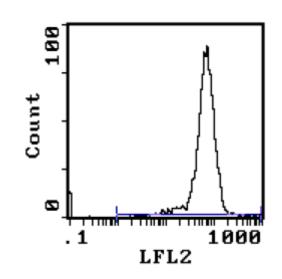
Thymus: 98.4% Spleen: 28.8%

## **Results - Strain Distribution:**

<u>Cell Concentration</u>: 1x10e6 cells per test <u>Antibody Concentration Used</u>: 0.5 µg/10e6 cells <u>Strains Tested</u>: C57BL/6, C3H/He, CBA/J, BALB/c, ATL, AKR/J <u>Positive</u>: C57BL/6, C3H/He, CBA/J, BALB/c, ATL <u>Negative</u>: AKR/J



**Pictures:** 



Cell Source: Thymus - Percentage of cells stained above control: 98.4%