



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

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

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

[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

Technically  
Speaking

**CEDARLANE**<sup>®</sup>  
[www.cedarlanelabs.com](http://www.cedarlanelabs.com)



Conveniently Delivering You Today's Innovations  
for the Science of Tomorrow™

**Anti-Rat CD4  
Monoclonal Antibody**

Catalogue#	Format	Size	Concentration	Isotype Control
<b>CL003A</b>	Ascites	0.5ml	NA	CLCMG100
<b>CL003AP/-2</b>	Purified	250µg/500µg	1.0 mg/ml	CLCMG100
<b>CL003NA</b>	Purified	1.0ml	1.0 mg/ml	CLCMG100
<b>CL003B/-5</b>	Biotin	100µg/500µg	0.1 mg/ml	CLCMG115
<b>CL003F/-5</b>	FITC	100µg/ 500µg	0.1 mg/ml	CLCMG101
<b>CL003APC</b>	APC	100µg	0.1 mg/ml	CLCMG105
<b>CL003PE/-4</b>	PE	50µg/ 200µg	0.1 mg/ml	CLCMG104
<b>CL003AF4</b>	Alexa Fluor <sup>®</sup> 488	100 µg	0.1 mg/ml	N/A
<b>CL003AF7</b>	Alexa Fluor <sup>®</sup> 700	100 µg	0.1 mg/ml	N/A

Alexa Fluor<sup>®</sup> is a registered trademark of Life Technologies Corporation.

Isotype: Mouse IgG<sub>1</sub>

**DESCRIPTION:**

Cedarlane's anti-rat CD4 monoclonal antibody recognizes a determinant on the majority of thymocytes (90-95%), a subset of peripheral T cells and peritoneal macrophages. (1,2,3,10) The antigen recognized by this antibody is a surface glycoprotein of Mr 48,000-52,000 and is the homologue of the human CD4 and the mouse L3/T4 antigen. CL003 labels the rat T helper subset, which mediates the helper activity for B and T cells, graft vs. host (GVH) reactivity and produces IL-2 in the mixed lymphocyte reaction (MLR). (2,4,6.) Addition of CL003 to the MLR, inhibited proliferation and blocks the production of IL-2. (4,5,6,9) T cells which mediate cytotoxicity and suppressor functions are not labelled. (Thus, cells labelled by this antibody are not labelled by MRC OX-8.)

CL003 is invaluable for separating T cell subsets for functional studies and for labelling cells in tissue sections. It has been used in studying the role of T lymphocytes in graft rejections (7) and in studying the subsets of T cells in the rat which mediate graft vs. host disease. (8)

This particular antibody is also one of three antibodies which labels T lymphocyte populations in the rat. These clones include W3/13 (CL002), which labels all T cells as well as MRC OX-8 (CL004) and W3/25 (CL003) which label non-overlapping T cell subpopulations. These monoclonal antibodies used in concert are being employed extensively to investigate cellular aspects of the immune response in rats and prove to be useful as markers for functionally distinct subpopulations of lymphocytes.

This clone is reported to work with frozen and paraffin sections (11) and in functional testing (12).

**PRESENTATION:**

**Ascites:** Lyophilized

**Purified:** Purified IgG buffered in PBS and 0.02% NaN<sub>3</sub>. (Purified from ascitic fluid via Protein G Chromatography). For maximum recovery of contents, spin down tube before use.

**Biotin, FITC, PE, APC, AF488 and AF700:** Biotin/FITC/PE/APC/AF488/AF700 conjugated IgG 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.

**No Azide:** Purified Ig buffered in PBS, no preservative, 0.2µm sterile filtered.

*Continued Overleaf....*

Visit our website for your local distributor.

**CEDARLANE**<sup>®</sup>



[www.cedarlanelabs.com](http://www.cedarlanelabs.com)

An ISO 9001:2015 and ISO 13485:2016  
registered company.

In CANADA: Toll Free: 1-800-268-5058

4410 Paletta Court, Burlington, ON L7L 5R2 ph: (289) 288-0001, fax: (289) 288-0020  
e-mail: [general@cedarlanelabs.com](mailto:general@cedarlanelabs.com)

In the USA: Toll Free: 1-800-721-1644

1210 Turrentine Street, Burlington, NC 27215 ph: (336) 513-5135, fax: (336) 513-5138  
e-mail: [service@cedarlanelabs.com](mailto:service@cedarlanelabs.com)

## **STORAGE/STABILITY:**

Store **Ascites** at -20°C. For all other formats, store at 4°C. **DO NOT FREEZ APC, PE, AF488 and AF700** conjugates. For long term storage (**Purified, No Azide, Biotin and FITC**), aliquot and freeze unused portion at -20°C in volumes appropriate for single usage. Avoid freeze/thaw cycles.

## **SPECIFICATIONS:**

Clone: W3/25

Hybridoma Production:

Immunization: Immunogen: Rat thymocyte membrane

Donor: BALB/c spleen

Fusion Partner: P3-NSI-1-Ag4 (NSI/1)

Specificity: Rat CD4

## **TEST RESULTS:**

Tissue Distribution by Flow Cytometry Analysis:

Rat Strain: Buffalo

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

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

Cell Source

Thymus

Spleen

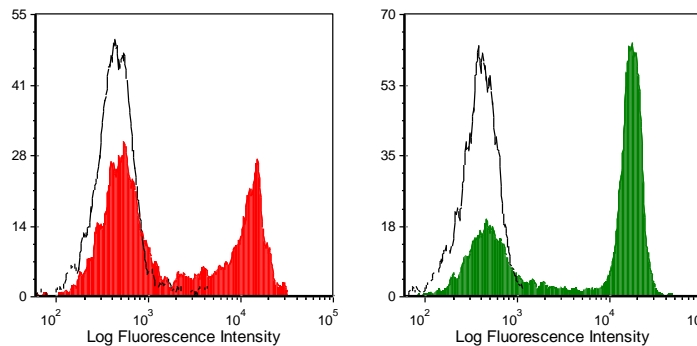
Lymph Node

Percentage of cells stained above control:

97.0%

37.0%

67.0%



Wistar rat splenocytes (left) or lymph nodes (right) were stained with anti-CD4 (clone: W3/25) (filled histogram) or mouse IgG1 isotype control (open histogram).

**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.**

## **REFERENCES:**

1. Williams, A.F., Galfre, G. and C. Milstein. (1977) Cell, 12, 633-673. Analysis of cell surfaces by xenogenic myeloma-hybrid antibodies: Differentiation antigens of rat lymphocytes.
2. Brideau, R.J., Carter, P.B., McMaster, W.R., Mason, D.W. and A.F. Williams. (1980) Eur. J. of Immunol., 10, 609-615. Two subsets of Rat T Lymphocytes defined with monoclonal antibodies.
3. Barclay, A.N. (1981) Immunology, 42, 593-600. The Localization of populations of lymphocytes defined with monoclonal antibodies in rat lymphoid tissues.
4. Cantrell, D.A., Robins, R.A. and R.W. Baldwin. (1982) Cell Immunol. 70, 367-372. Rat lymphocyte subsets: Cellular requirements for the generation of T-cell growth factors.
5. Mason, D.W., Pugh, C.W. and M. Webb. (1981) Immunology, 44, 75-87. The Rat Mixed Lymphocyte Reaction: Roles of a dendritic cell in intestinal lymph and T-Cell subsets defined by monoclonal antibodies.
6. Webb, M., Mason, D.W. and A.F. Williams. (1979) Nature, 282, 841-843. Inhibition of mixed lymphocyte response by monoclonal antibody specific for rat T lymphocyte subset.
7. Dallman, M.J., Mason, D.W. and M. Webb. (1982) Eur. J. Immunol. 12, 511-518. The role of host and donor cells in the rejection of skin allografts by T cell deprived rats injected with syngenic T cells.
8. Mason, D.W. (1981) Transplantation, 32, 222-226. Subsets of T cells in the rat mediating lethal graft vs. host disease.
9. White, R.A.H., Mason, D.W., Williams, A.F., Galfre, G. and C. Milstein. (1978) J. Exp. Med. 148, 644-673. T Lymphocyte heterogeneity in the rat: Separation of functional sub-populations using a monoclonal antibody.
10. Jefferies, W.A., Green, J.R. and A.F. Williams. (1985) J. Exp. Med. 162, 117-127. Authentic T Helper CD4 (W3/25) antigen on rat peritoneal macrophages.
11. Whiteland, J.L et al (1995). Immunohistochemical detection of T cell subsets and other leukocytes in paraffin embedded rat and mouse tissues with monoclonal antibodies. J. Histochem. Cytochem. 43: 313-320.
12. Tamatani, T. et al. (1991) Characterization of the rat leukocyte integrin, CD11/CD18, by the use of LFA-1 subunit-specific monoclonal antibodies. Eur. J. Immunol. 21:627-633.

**FOR RESEARCH USE ONLY**

® is a Registered Trademark of Cedarlane