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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Technically
Speaking

CEDARLANE[®] 
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Conveniently Delivering You Today's Innovations
for the Science of Tomorrow™

**Anti-Rat CD25 (IL-2R)
Monoclonal Antibody**

| Catalogue# | Format | Size | Concentration | Isotype Control |
|------------|----------|-------------|---------------|-----------------|
| CL039A | Ascites | 0.5ml | NA | CLCMG100 |
| CL039AP/-2 | Purified | 250µg/500µg | 1.0 mg/ml | CLCMG100 |
| CL039B/-5 | Biotin | 100µg/500µg | 0.1 mg/ml | CLCMG115 |
| CL039F/-5 | FITC | 100µg/500µg | 0.1 mg/ml | CLCMG101 |
| CL039PE/-4 | PE | 50µg/200µg | 0.1 mg/ml | CLCMG104 |
| CL039NA | No Azide | 1.0mg | 1.0 mg/ml | CLCMG100 |

Isotype: Mouse IgG₁

DESCRIPTION:

Cedarlane's anti-rat Interleukin-2 receptor monoclonal antibody recognizes the smaller (alpha subunit) 55kD chain of the IL-2 receptor found on activated rat T cells, thymic dendritic cells but not resting lymphocytes (1). CL039 binds to the rat interleukin-2 receptor and has proven to be an important marker for activated T cells (2).

This clone is reported to work with frozen and paraffin sections (3).

PRESENTATION:

Ascites: Lyophilized.

Purified: Purified IgG buffered in PBS and 0.02% NaN₃. (Purified from ascitic fluid via Protein G Chromatography). For maximum recovery of contents, spin down tube before use.

Biotin, FITC and PE: Biotin/FITC/PE conjugated IgG buffered in PBS, 0.02% NaN₃ 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 with no preservative; 0.2µm sterile filtered.

STORAGE/STABILITY:

Store **Ascites** at -20°C. For all other formats, store at 4°C. DO NOT FREEZE **PE** conjugates. Handle NA format under aseptic conditions. For long term storage (**Purified, Biotin** and **FITC**), aliquot and freeze unused portions at -20°C in volumes appropriate for single usage. Avoid freeze/thaw cycles.

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CEDARLANE[®] 

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An ISO 9001:2000 and ISO 13485:2003
registered company.

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SPECIFICATIONS:

Clone: MRC OX-39

Hybridoma Production:

Immunization: Immunogen: T blasts from a mixed lymphocyte reaction between purified CD4 positive T cells and irradiated spleens.
Donor: BALB/c spleen
Fusion Partner: NSO/1

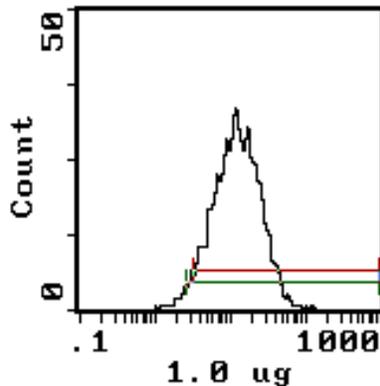
Specificity: Rat CD25 (IL-2 Receptor)

TEST RESULTS:

Tissue Distribution by Flow Cytometry Analysis:

Rat Strain: Wistar
Cell Concentration: 1x10⁶ cells per test
Antibody Concentration Used: 1.0µg/10⁶ cells

| <u>Cell Source</u> | <u>Percentage of cells stained above control:</u> |
|--------------------|---|
| T-cell blasts | 95.3% |
| Thymus | 0.59% |



Cell Source: Con-A activated thymocytes (T-cell blasts)

**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. Paterson, D.J., Jeffries, W.A., Green, J.R., Brandon, M.R., Corthesy, P., Puklavec, M. and A.F. Williams. (1987) Mol. Immunol. 24, 1281-1290. Antigen of Activated Rat T Lymphocytes Including a Molecule of 50,000 Mr Detected Only on CD4 Positive T Blasts.
2. Barclay, A.N. (1981) Immunology. 42 593-600 The Localization of populations of lymphocytes defined with monoclonal antibodies in rat lymphoid tissues.
3. 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.

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