



# SZABO SCANDIC

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## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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

**Anti-Mouse Complement Component  
C3 Monoclonal Antibody**

Catalogue#	Format	Size	Concentration	Isotype Control
<b>CL7503AP</b>	Purified	250µg	1.0 mg/ml	CLCR2A00
<b>CL7503AP-2</b>	Purified	500µg	1.0 mg/ml	CLCR2A00
<b>CL7503NA</b>	Purified	1.0mg	1.0 mg/ml	CLCR2A00
<b>CL7503B</b>	Biotin	100µg	0.1 mg/ml	CLCR2A15
<b>CL7503F</b>	FITC	100µg	0.1 mg/ml	CLCR2A01
<b>CL7503APC</b>	APC	100µg	0.1 mg/ml	CLCR2A05
<b>CL7503PE</b>	PE	50µg	0.1 mg/ml	CLCR2A04
<b>CL7503TRC</b>	TRITC	50µg	0.1 mg/ml	CLCR2A04
<b>CL7503AF4</b>	Alexa Fluor <sup>®</sup> 488	100 µg	0.1 mg/ml	N/A

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

Isotype: Rat IgG<sub>2a</sub>

**DESCRIPTION:**

Cedarlane's Anti-Mouse Complement Component C3 monoclonal antibody reacts with mouse C3 as well as the breakdown products C3b, iC3b and C3dg. C3 is the most abundant complement protein in serum. C3 and its cleavage products, C3a and C3b, play a central role in the complement activation cascade. C3b forms an integral part of the C3 and C5 convertases as it promotes complement activation and the subsequent formation of the membrane attack complex. C3a possesses anaphylatoxic as well as various immunoregulatory properties.

Also, C3 has been implicated in developmental and non-inflammatory process such as hematopoiesis, skeletal and vascular development and reproduction.

Reported applications of this antibody include immunofluorescence<sup>2, 5</sup>, ELISA<sup>4</sup>, Western blots<sup>1</sup>, flow cytometry and immunohistochemistry<sup>1, 3</sup> on acetone fixed frozen sections.

**PRESENTATION:**

**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.

**APC, Biotin, FITC, PE, TRITC and AF488** 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.

**STORAGE/STABILITY:**

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

*Continued.....*

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## **SPECIFICATIONS:**

Clone: RMC11H9

### Hybridoma Production:

#### Immunization:

Immunogen: C57BL/6 thymocytes incubated with a rat IgG2b anti-murine Thy-1 (RmT1) and C57BL/6 serum (as a source for C3)

Donor: Lou/c rat spleen cells

Fusion Partner: Mouse myeloma cell line P3X63 Ag. 8.653

Specificity: Mouse Complement Component C3

## **TEST RESULTS:**

### Tissue Distribution by Flow Cytometry Analysis:

Mouse Strain: C<sub>3</sub>H/He

Cell Concentration: 1x10<sup>6</sup> cells per test

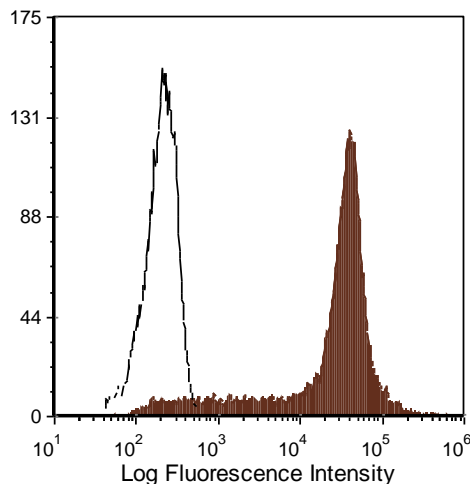
Antibody Concentration Used: 0.2 µg/10<sup>6</sup> cells

#### Cell Source:

Thymocytes incubated with anti-mouse CD90 (CL8600AP), then incubated with fresh mouse serum (same strain).

Percentage of cells stained above control:

88%



C3H/He mouse thymocytes incubated with anti-mouse CD90 (Thy 1.2) (clone: 5a-8) and fresh mouse serum were stained with anti-mouse C3 (clone: RmC11H9) (filled histogram) or rat IgG2a 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. Kremmer, E et al. (1990). Hybridoma. 9 (4):309-17.
2. Robson, M et al. (2001). J. Immunol. 166(11): 6820-6828.
3. Mastellos et al. (2004). Molecular Immunology 40: 1213-1221.
4. Zhili Xu et al. (2008). J. Virology. 82(23): 11705-13.
5. Binstadt, B et al. (2009). PNAS. 106(39): 16758-63.

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