

## **Monoclonal Antibody to Ly6G / GR1 Neutrophil Marker - FITC**

<b>Alternate names:</b>	Gr-1 Granulocyte marker
<b>Catalog No.:</b>	CL046F
<b>Quantity:</b>	0.1 mg
<b>Concentration:</b>	0.1 mg/ml
<b>Background:</b>	GR-1 is a 25-30 kDa cell surface antigen and is expressed on myeloid cells but not lymphoid or erythroid cells. The expression of the Gr-1 antigen increases with granulocyte maturation (3) as shown by the distinct populations of bone-marrow cells this monoclonal antibody labels: negative, low positive and high positive. Expression is transient on cells of monocytic lineage (3).
<b>Uniprot ID:</b>	<a href="#">P35461</a>
<b>NCBI:</b>	<a href="#">10090</a>
<b>Host / Isotype:</b>	Rat / IgG2b
<b>Clone:</b>	RB6-8C5
<b>Format:</b>	<b>State:</b> Liquid Ig fraction <b>Purification:</b> Protein G chromatography <b>Buffer System:</b> PBS, 0.02% Na <sub>3</sub> N and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml <b>Label:</b> FITC <i>Absorption / Emission:</i> 495 nm / 528 nm
<b>Applications:</b>	Flow cytometry (1,2,3). Western blot (5). Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.
<b>Specificity:</b>	This antibody reacts with the myeloid differentiation antigen GR-1 (1,2). <b>Species:</b> Mouse. Other species not tested.
<b>Storage:</b>	Store the antibody at 2 - 8 °C for up to one month. For long term storage, aliquot and freeze unused portion at -20 °C in volumes appropriate for single usage. Avoid repeated freezing and thawing This product is photosensitive and should be protected from light. Shelf life: one year from despatch.
<b>General References:</b>	1. Spangrude, G.J., et al.. Purification and characterization of mouse hematopoietic stem cells. Science 241:58-62, 1991. 2. Muller, C. E. et al. Isolation of two early B lymphocyte progenitors from mouse marrow: a committed pre-pre-B cell and a clonogenic thy-1 lo hematopoietic stem cell. Cell

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3. H. Kjetil et al. Characterization and regulation of RB6-8C5 antigen expression on murine bone marrow cells. 1991. J. Immunol. 147: 22-28.

4. Brummer, Elmer et al. Immunological activation of polymorphonuclear neutrophils for fungal killing: Studies with murine cells and blastomyces dermatitidis in vitro. J. Leuko. Bio 36:505-520, 1984.

5. Jutila, M.A. et al. Ly-6C is a monocyte/macrophage and endothelial cell differentiation antigen regulated by interferon-gamma. J. Immunol. 18: 1819-1826, 1988.

6. Haestdal, K., F.W. Ruscetti, J.N. Ihle, S.E.W. Jacobsen, C.M. Dubois, W.C. Kopp, D.L. and J.R. Keller 1991. Characterization and regulation of RB6-8C5 antigen expression on murine bone marrow cell. J.Immunol. 147:22

7. Fleming, T.J., M.L. Fleming, and T.R. Malek. 1993. Selective Expression of Ly-6G on myeloid lineage cells in bone marrow. RB6-85 mAb to granulocyte-differentiation antigen (Gr-1) detects members of the Ly-6 family. J Immunol. 151:139

### Protocols:

#### FLOW CYTOMETRY ANALYSIS:

##### Method:

1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population.
2. Wash 2 times.
3. Resuspend the cells to a concentration of  $2 \times 10^7$  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 0.1 - 0.5  $\mu$ g of antibody per  $10^6$  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 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 by Flow Cytometry Analysis:

Mouse Strain: CBA/J

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

Antibody Concentration Used: 0.1  $\mu$ g/ $10^6$  cells

Isotypic Control: FITC Rat IgG2b

Cell Source Percentage of cells stained above control:

Thymus 1.5%

Whole Blood Monocytes 87.2%

Bone Marrow Macrophages 90.0%

(see picture below)

Strain Distribution by Flow Cytometry Analysis:

Procedure: see above

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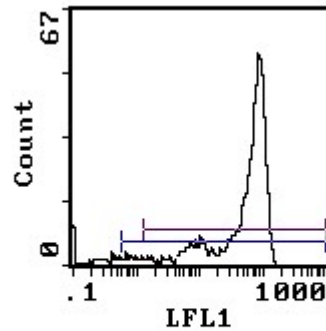
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Cell Concentration: 1x10<sup>6</sup> cells per test  
Antibody Concentration Used: 0.1 µg/10<sup>6</sup> cells  
Strains Tested: BALB/c, C57BL/6, CBA, C3H/he, AKR  
Positive: BALB/c, C57BL/6, CBA, C3H/he, AKR  
Negative: none

Pictures:



Cell Source: Bone Marrow Macrophages  
Percentage of cells stained above control: 90.0%

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