

TECHNICAL DATA SHEET

redFluor™ 710 Anti-Mouse Ly-6G (Gr-1) (RB6-8C5)

Catalog Number: 80-5931

PRODUCT INFORMATION

Contents: redFluor™ 710 Anti-Mouse Ly-6G (Gr-1) (RB6-8C5)

Isotype: Rat IgG2b, kappa

Concentration: 0.2 mg/mL

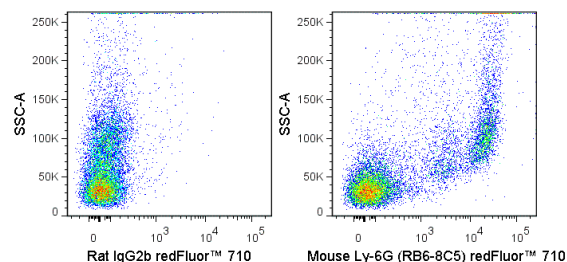
Clone: RB6-8C5

Reactivity: Mouse

Use By: 12 months from date of receipt

Storage Conditions: 2-8°C protected from light

Formulation: 10 mM NaH₂PO₄, 150 mM NaCl, 0.09% NaN₃, pH7.2



C57Bl/6 bone marrow cells were stained with 0.015 ug redFluor™ 710 Anti-Mouse Ly-6G (80-5931) (right panel) or 0.015 ug redFluor™ 710 Rat IgG2b isotype control (left panel).

DESCRIPTION

The RB6-8C5 antibody binds to mouse Ly-6G, commonly known as Gr-1, a member of the Ly-6 superfamily of GPI-anchored cell surface proteins with roles in cell signaling and cell adhesion. Gr-1 is differentially expressed during development and maturation of cells in the myeloid lineage and is expression at varying stages and levels on monocytes, macrophages, granulocytes, and peripheral neutrophils. In the mouse, the RB6-8C5 antibody is typically used in combination with the macrophage labeling antibody M1/70 (Anti-CD11b) for phenotypic analysis of monocytes, macrophages and granulocytes. Note: The RB6-8C5 antibody has been reported to cross-react with Ly-6C on cells expressing this antigen (Fleming et al. 1993. *J. Immunol.* 151:2399-2408 and Sasmono et al. 2007. *J. Leukoc. Biol.* 82: 111-123) and has been cited in the literature for identification of Ly-6G/Ly-6C. Other reports suggest that this antibody is specific for Ly-6G, without cross-reactivity for Ly-6C (Nagendra S. and Schlueter AJ. 2003. *Cytometry A*, 58(2): 195-200.

PREPARATION & STORAGE

This monoclonal antibody was purified from tissue culture supernatant via affinity chromatography. The purified antibody was conjugated under optimal conditions, with unreacted dye removed from the preparation. It is recommended to store the product undiluted at 4°C, and protected from prolonged exposure to light. Do not freeze.

APPLICATION NOTES

This antibody preparation has been quality-tested for flow cytometry using mouse spleen cells, or an appropriate cell type (where indicated). The amount of antibody required for optimal staining of a cell sample should be determined empirically in your system.

redFluor™ 710 dye is excited by the red (633-647 nm) laser and has a peak emission of 710 nm. The recommended band pass filter for this dye is 710/50. redFluor™ 710 can be used as an alternative for Alexa Fluor® 700. Confirm that your cytometer is configured to detect this fluorochrome.

REFERENCES

- Berent-Maoz B, Montecino-Rodriguez E, Signer RAJ, and Dorshkind K. 2012. *Blood*. 119:5715-5721. (Flow cytometry) von Bruhl M-L, Stark K, Steinhart A, et al. 2012. *J. Exp. Med.* 209: 819-835. (Intravital fluorescent microscopy - video) Le HT, Tran VG, Kim W, Kim H, Cho HR, and Kwon B. 2012. *J. Immunol.* 189:287-295. (in vivo neutrophil depletion) Doring Y, Soehnlein O, Drechsler M, Shagdarsuren E, Chaudhari SM, Meiler S, Hartwig H, Hristov M, Koenen RR, Hieronymus T, Zenke M, Weber C, and Zernecke A. 2012. *Arterioscler. Thromb. Vasc. Biol.* 32: 1613-1623. (in vivo depletion) Hickman HD, Li L, Reynoso GV, Rubin EJ, Skon CN, Mays JW, Gibbs J, Schwartz O, Bennink JR, and Yewdell JW. 2011. *J. Exp. Med.* 208: 2511-2524. (Immunohistochemistry – OCT embedded frozen tissue) Wang T, Tian L, Haino M, Gao J-L, Lake R, Ward Y, Wang H, Siebenlist U, Murphy PM, and Kelly K. 2007. *Infect. Immun.* 75(3):1144-1153. (Immunohistochemistry – zinc fixed tissue) Nutt SL, Metcalf D, D'Amico A, Polli M, and Wu L. 2005. *J. Exp. Med.* 201:221-231. (Immunomagnetic bead depletion) Whiteland JL, Nicholls SM, Shimeld C, Easty DL, Williams NA, and Hill TJ. 1995. *J. Histochem. Cytochem.* 43:313-320. (Immunohistochemistry – frozen tissue, paraffin embedded tissue) Fleming TJ, Fleming ML, and Malek TR. 1993. *J. Immunol.* 151:2399-2408. (in vitro blocking, immunoprecipitation)

Tonbo Biosciences tests all antibodies by flow cytometry. Citations are provided as a resource for additional applications that have not been validated by Tonbo Biosciences. Please choose the appropriate format for each application and consult Materials and Methods sections for additional details about the use of any product in these publications.

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