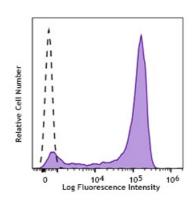
Spark NIR[™] 685 anti-mouse Ly-6G

Catalog # / Size:	1238330 / 100 μg 1238325 / 25 μg
Clone:	1A8
lsotype:	Rat IgG2a, к
Immunogen:	Ly-6G transfected EL-4J cell line.
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with Spark NIR™ 685 under optimal conditions.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide
Workshop Number:	750 under optimal conditions.
Concentration:	0.5 mg/mL

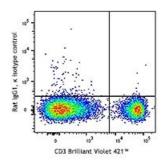


C57BL/6 mouse bone marrow cells were stained with Ly-6G (clone 1A8) Spark NIR[™] 685 (filled histogram) or cells were left unstained (open histogram). Data shown was gated on the myeloid cell population.

Applications:

Applications:	Flow Cytometry
Recommended Usage:	Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is $\leq 0.25 \ \mu$ g per million cells in 100 μ L volume. It is recommended that the reagent be titrated for optimal performance for each application.

* Spark NIR [™] 685 has a maximum excitation of 665 nm and a maximum emission of 685 nm.



C57BL/6 mouse bone marrow cells were stained with CD150 (SLAM) (clone TC15-12F12.2) APC/Fire[™] 750 (filled histogram) or rat IgG2a, κ APC/Fire[™] 750 isotype control (open histogram).

Application Notes:	While 1A8 recognizes only Ly-6G, clone RB6-8C5 recognizes both Ly-6G and Ly-6C. Clone RB6-8C5 binds with high affinity to mouse Ly-6G molecules and to a lower extent to Ly-6C ¹⁵ . Clone RB6-8C5 impairs the binding of anti-mouse Ly-6G clone 1A8 ¹⁵ . However, clone RB6-8C5 is able to stain in the presence of anti- mouse Ly-6C clone HK1.4 ¹⁶ . Additional reported applications (for the relevant formats) include: immunohistochemistry ⁹ of frozen sections ¹⁰ and paraffin-embedded sections ¹¹ , and depletion ⁴ , ¹²⁻¹⁴ . The Ultra-LEAF TM purified antibody (Endotoxin < 0.01 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for <i>in vivo</i> studies or highly sensitive assays (Cat. No. 127632, 127649,
Application References:	 127650, 127661 and 127662). Fleming TJ, et al. 1993. J. Immunol. 151:2399. (FC) Daley JM, et al. 2008. J. Leukocyte Biol. 83:1. (FC) Dietlin TA, et al. 2007. J. Leukocyte Biol. 81:1205. (FC) Daley J, et al. 2007. J. Leukocyte Biol. doi:10.1189. (Deplete) PubMed Tadagavadi RK, et al. 2010. J. Immunol. 185:4904. PubMed Sumagin R, et al. 2010. J. Immunol. 185:7057. PubMed Guiducci C, et al. 2010. J. Immunol. 185:7057. PubMed Guiducci C, et al. 2010. J. Exp Med. 207:2931. PubMed Fujita M, et al. 2011. Cancer Res. 71:2664. PubMed Fujita M, et al. 2010. P. Natl. Acad. Sci. USA 107:21248. [supplementary data] (IHC) Kowanetz M, et al. 2010. J. Gen. Virol. 91:2158. (FC, Deplete) Jaeger BN, et al. 2010. J. Exp. Med. 209:565. (Deplete) Jaeger BN, et al. 2012. J. Exp. Med. 209:565. (Deplete) Bibechini E, et al. 2009. Eur. J. Immunol. 13:65 (FC, Deplete) Ribechini E, et al. 2009. Eur. J. Immunol. 39:3538. Ng LG, et al. 2011. J Invest. Dermatol. 131:2058. PubMed Ma C, et al. 2012. J. Leukoc. Biol. 92:1199. McCartney-Francis, N, et al. 2014. J Leukoc. Biol. 96:917. PubMed Her Z, et al. 2014. EMBO Mol. Med. 7:24. PubMed
Description:	Lymphocyte antigen 6 complex, locus G (Ly-6G), a 21-25 kD GPI-anchored protein, is expressed on the majority of myeloid cells in bone marrow and peripheral granulocytes.
Antigen References:	Fleming TJ, et al. 1993. J. Immunol. 151:2399.