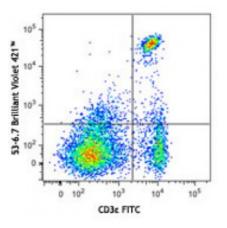
Product Data Sheet

Brilliant Violet 421[™] anti-mouse CD8a

Catalog # / Size:	1103685 / 125 μl 1103690 / 500 μl
	1103765 / 50 μg
Clone:	53-6.7
Isotype:	Rat IgG2a, к
Immunogen:	Mouse thymus or spleen
Reactivity:	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 421 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 421 [™] and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).
Concentration:	microg sizes: 0.2 mg/ml microL sizes: lot-specific



C57BL/6 mouse splenocytes were stained with CD3 ϵ FITC and CD8a (clone 53-6.7) Brilliant Violet 421^m. Quadrant gating was based on the rat IgG2a, κ Brilliant Violet 421^m isotype control.

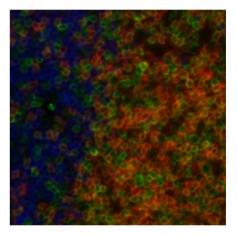
Applications:

Applications:	Flow Cytometry
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Each lot of this antibody is quality Recommended Usage: control tested by immunofluorescent staining with flow cytometric analysis. For immunofluorescent staining using the microg size, the suggested use of this reagent is ≤ 0.5 microg per million cells in 100 microL volume. For immunofluorescent staining using microL sizes, the suggested use of this reagent is ≤ 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

> Brilliant Violet 421[™] excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421[™] is a trademark of Sirigen Group Ltd.

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BL6 mouse lymph nodes, fixed O/N in PLP, blocked with 10% rat serum, stained with CD8a-BV421 (red), B220-Alexa Fluor® 647 (blue), and TCR β -Alexa Fluor® 488 (green) in 1% BSA and 0.1% Tween-20 in PBS. Images were acquired with an automated wid

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Application Clone 53-6.7 antibody competes with clone 5H10-1 antibody for binding to Notes: thymocytes3. The 53-6.7 antibody has been reported to block antigen presentation via MHC class I and inhibit T cell responses to IL-2. This antibody has also been used for depletion of CD8a⁺ cells. Additional reported applications (for the relevant formats) include: immunoprecipitation^{1,3}, in vivo and *in vitro* cell depletion^{2,10,15}. inhibition of CD8 T cell proliferation3, cytotoxicity^{3,4}, blocking of and immunohistochemical staining^{5,6} of acetone-fixed frozen sections and zincfixed paraffin-embedded sections. Clone 53-6.7 is not recommended for immunohistochemistry of formalin-fixed paraffin sections. The LEAF[™] purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 100716). For in vivo studies or highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 100746) with a lower endotoxin limit than standard LEAF[™] purified antibodies (Endotoxin <0.01 EU/microg). Application 1. Ledbetter JA, et al. 1979. Immunol. Rev. 47:63. (IHC, IP) **References:** 2. Hathcock KS. 1991. *Current Protocols in Immunology.* 3.4.1. (Deplete) 3. Takahashi K, et al. 1992. P. Natl. Acad. Sci. USA 89:5557. (Block, IP) 4. Ledbetter JA, et al. 1981. J. Exp. Med. 153:1503. (Block) 5. Hata H, et al. 2004. J. Clin. Invest. 114:582. (IHC) 6. Fan WY, et al. 2001. Exp. Biol. Med. 226:1045. (IHC) 7. Shih FF, et al. 2006. J. Immunol. 176:3438. (FC) 8. Kamimura D, et al. 2006. J. Immunol. 177:306. 9. Bouwer HGA, et al. 2006. P. Natl. Acad. Sci. USA 103:5102. (FC, Deplete) 10. Kao C, et al. 2005. Int. Immunol. 17:1607. PubMed 11. Ko SY, et al. 2005. J. Immunol. 175:3309. (FC) PubMed 12. Rasmussen JW, et al. 2006. Infect. Immun. 74:6590. PubMed 13. Lee CH, et al. 2009. Clin. Cancer Res. PubMed 14. Geiben-Lynn R, et al. 2008. Blood 112:4585. (Deplete) PubMed 15. Kingeter LM, et al. 2008. J. Immunol. 181:6244. PubMed 16. Guo Y, et al. 2008. Blood 112:480. PubMed 17. Andrews DM, et al. 2008. J. Virol. 82:4931. PubMed 18. Britschqui MR, et al. 2008. J. Immunol. 181:7681. PubMed 19. Kenna TJ, et al. 2008. Blood 111:2091. PubMed 20. Jordan JM, et al. 2008. Infect. Immun. 76:3717. PubMed 21. Todd DJ, *et al.* 2009. *J. Exp. Med.* 206:2151. <u>PubMed</u> 22. Bankoti J, *et al.* 2010. *Toxicol. Sci.* 115:422. (FC) <u>PubMed</u> 23. Medyouf H, et al. 2010. Blood 115:1175. PubMed 24. Riedl P, et al. 2009. J. Immunol. 183:370. PubMed 25. Apte SH, et al. 2010. J. Immunol. 185:998. PubMed 26. Bankoti J, et al. 2010. Toxicol. Sci. 115:422. (FC) PubMed 27. del Rio ML, et al. 2011. Transpl. Int. 24:501. (FC) PubMed 28. Chen Z, et al. 2014. Cancer Immunol Res. 2:911. PubMed 29. Ballet R, et al. 2014. PLoS Pathog. 10:1004550. PubMed 30. Hu-Lieskovan S, et al. 2015. Sci Transl Med. 7:279.

Description: CD8, also known as Lyt-2, Ly-2, or T8, consists of disulfide-linked

For research use only. Not for diagnostic use. Not for resale. Sony Biotechnology Inc. will not be held responsible for patent infringement or other violations that may occur with the use of our products. Sony Biotechnology Inc. 1730 North First Street, San Jose, CA 95112 www.sonybiotechnology.com α and β chains that form the α(CD8a)/β(CD8b) heterodimer and α/α homodimer. CD8a is a 34 kD protein that belongs to the immunoglobulin family. The CD8 α/β heterodimer is expressed on the surface of most thymocytes and a subset of mature TCR α/β T cells. CD8 expression on mature T cells is non-overlapping with CD4. The CD8 α/α homodimer is expressed on a subset of γ/δ TCR-bearing T cells, NK cells, intestinal intraepithelial lymphocytes, and lymphoid dendritic cells. CD8 is an antigen-presenting cells or epithelial cells. CD8 promotes T cell activation through its association with the TCR complex and protein tyrosine kinase lck.

Antigen References:

- Academic Press. 2. Zamoyska R. 1994. *Immunity* 1:243.
- 3. Ellmeier W, et al. 1999. Annu. Rev. Immunol. 17:523.