Product Data Sheet

Brilliant Violet 785[™] anti-human CD69

Catalog # / Size:	2154655 / 25 tests 2154660 / 100 tests	sê. A
Clone:	FN50	Human peripheral blood lymphocytes were stimulated with PMA + ionomycin for 6 hours and then stained with CD69 (clone FN50) Brilliant Violet 785 TM (filled histogram) or mouse IgG1, K Brilliant Violet 785 TM isotype control
Isotype:	Mouse lgG1, κ	
Reactivity:	Human	
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 785 [™] under optimal conditions. The solution is free of unconjugated Brilliant Violet 785 [™] and unconjugated antibody.	
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).	
Workshop Number:	IV A91	
Concentration:	Lot-specific	

Applications:

Applications: Flow Cytometry

```
Each lot of this antibody is quality control tested by immunofluorescent staining
Recommended
                   with flow cytometric analysis. For flow cytometric staining, the suggested use of
        Usage:
                   this reagent is \leq 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 785[™] excites at 405 nm and emits at 785 nm. The bandpass filter 780/60 nm is recommended for detection, although filter optimization may be required depending on other fluorophores used. Be sure to verify that your cytometer configuration and software setup are appropriate for detecting this channel. Refer to your instrument manual or manufacturer for support. Brilliant Violet 785[™] is a trademark of Sirigen Group Ltd.

(open histogram).

This product is subject to proprietary rights of Sirigen Inc. and is made and sold under license from Sirigen Inc. The purchase of this product conveys to the buyer a non-transferable right to use the purchased product for research purposes only. This product may not be resold or incorporated in any manner into another product for resale. Any use for therapeutics or diagnostics is strictly prohibited. This product is covered by U.S. Patent(s), pending patent applications and foreign equivalents.

Application Additional reported applications (for the relevant formats) include: Notes: immunohistochemical staining of acetone-fixed frozen tissue sections2 and immunofluorescence microscopy3.

Application	1. Knapp WB, <i>et al.</i> 1989. Leucocyte Typing IV. Oxford University Press. New York.
References:	2. Sakkas LI, et al. 1998. Clin. and Diag. Lab. Immunol. 5:430. (IHC)
	3. Kim JR, <i>et al.</i> 2005. <i>BMC Immunol.</i> 6:3. (IF)
	4. Verjans GM, <i>et al.</i> 2007. <i>P. Natl. Acad. Sci. USA</i> 104:3496.

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

5. Lu H, *et al.* 2009. *Toxicol Sci.* 112:363. (FC) <u>PubMed</u> 6. Thakral D, *et al.* 2008. *J. Immunol.* 180:7431. (FC) <u>PubMed</u>

7. Yoshino N, *et al.* 2000. *Exp. Anim. (Tokyo)* 49:97. (FC)

Description: CD69 is a 27-33 kD type II transmembrane protein also known as activation inducer molecule (AIM), very early activation antigen (VEA), and MLR3. It is a member of the C-type lectin family, expressed as a disulfide-linked homodimer. Other members of this receptor family include NKG2, NKR-P1 CD94, and Ly49. CD69 is transiently expressed on activated leukocytes including T cells, thymocytes, B cells, NK cells, neutrophils, and eosinophils. CD69 is constitutively expressed by a subset of medullary mature thymocytes, platelets, mantle B cells, and certain CD4⁺ T cells in germinal centers of normal lymph nodes. CD69 is involved in early events of lymphocyte, monocyte, and platelet activation, and has a functional role in redirected lysis mediated by activated NK cells.

Antigen1. Schlossman S, et al. Eds. 1995. Leucocyte Typing V. Oxford University Press.References:New York.

2. Testi R, et al. 1994. Immunol. Today 15:479.