Product Data Sheet

Brilliant Violet 785[™] anti-human CD4

Catalog # / Size:	2102765 / 25 tests 2102770 / 100 tests	E
Clone:	RPA-T4	
Isotype:	Mouse IgG1, κ	Interpret text text text text text text text t
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 T114	
Concentration:	0.2	

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 ≤ 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.	
Application Notes:	The RPA-T4 antibody binds to the D1 domain of CD4 (CDR1 and CDR3 epitopes) and can block HIV gp120 binding and inhibit syncytia formation. Additional reported applications (for the relevant formats) include: immunohistochemistry of acetone-fixed frozen sections ^{3,4,5} , and blocking of T cell activation ^{1,2} . This clone was tested in-house and does not work on formalin fixed paraffin-embedded (FFPE) tissue. The LEAF TM purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 300516).	
Application References:	 Knapp W, <i>et al.</i> 1989. Leucocyte Typing IV. Oxford University Press. New York. (Activ) Moir S, <i>et al.</i> 1999. <i>J. Virol.</i> 73:7972. (Activ) Deng MC, <i>et al.</i> 1995. <i>Circulation</i> 91:1647. (IHC) Friedman T, <i>et al.</i> 1999. <i>J. Immunol.</i> 162:5256. (IHC) Mack CL, <i>et al.</i> 2004. <i>Pediatr. Res.</i> 56:79. (IHC) Lan RY, <i>et al.</i> 2006. <i>Hepatology</i> 43:729. Zenaro E, <i>et al.</i> 2009. <i>J. Leukoc. Biol.</i> 86:1393. (FC) <u>PubMed</u> Yoshino N, <i>et al.</i> 2000. <i>Exp. Anim. (Tokyo)</i> 49:97. (FC) 	

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 **Description:** CD4, also known as T4, is a 55 kD single-chain type I transmembrane glycoprotein expressed on most thymocytes, a subset of T cells, and monocytes/macrophages. CD4, a member of the Ig superfamily, recognizes antigens associated with MHC class II molecules, and participates in cell-cell interactions, thymic differentiation, and signal transduction. CD4 acts as a primary receptor for HIV, binding to HIV gp120. CD4 has also been shown to interact with IL-16.

Antigen 1. Center D, *et al.* 1996. *Immunol. Today* 17:476.
References: 2. Gaubin M, *et al.* 1996. *Eur. J. Clin. Chem. Clin. Biochem.* 34:723.

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