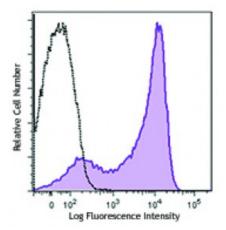
## **Product Data Sheet**

## PE/Dazzle<sup>™</sup> 594 anti-mouse CD62L

Catalog # / Size:	1122240 / 100 μg 1122235 / 25 μg
Clone:	MEL-14
Isotype:	Rat IgG2a, к
Immunogen:	C3H/eb mouse B lymphoma 38C-13
<b>Reactivity:</b>	Mouse
Preparation:	The antibody was purified by affinity chromatography and conjugated with PE/Dazzle <sup>™</sup> 594 under optimal conditions. The solution is free of unconjugated PE/Dazzle <sup>™</sup> 594 and unconjugated antibody.
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide.
<b>Concentration:</b>	0.2



C57BL/6 mouse splenocytes were stained with CD62L (clone MEL-14) PE/Dazzle<sup>m</sup> 594 (filled histogram) or rat IgG2a,  $\kappa$  PE/Dazzle<sup>m</sup> 594 isotype control (open histogram).

## **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 $\leq 0.06$ microg per million cells in 100 microL volume. It is recommended that the reagent be titrated for optimal performance for each application.	
	* PE/Dazzle™ 594 has a maximum excitation of 566 nm and a maximum emission of 610 nm.	
Application Notes:	Additional reported applications (for the relevant formats) include: immunoprecipitation <sup>1-3</sup> , complement-dependent cytotoxicity4, <i>in vivo</i> and <i>in vitro</i> blocking of adhesion <sup>1-3,5</sup> , and immunohistochemical staining of acetone-fixed frozen sections and zinc-fixed paraffin-embedded sections <sup>6</sup> . The LEAF <sup>TM</sup> purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for functional assays (Cat. No. 104416).	
Application References:	<ol> <li>Gallatin WM, <i>et al.</i> 1983. <i>Nature</i> 304:30. (IP, Block)</li> <li>Siegelman MH, <i>et al.</i> 1990. <i>Cell</i> 61:611. (IP, Block)</li> <li>Lewinsohn DM, <i>et al.</i> 1987. <i>J. Immunol.</i> 138:4313. (IP, Block)</li> <li>Iwabuchi K, <i>et al.</i> 1991. <i>Immunobiology</i> 182:161. (CMCD)</li> <li>Pizcueta P, <i>et al.</i> 1994. <i>Am. J. Pathol.</i> 145:461.</li> <li>Reichert RA, <i>et al.</i> 1986. <i>J. Immunol.</i> 136:3535. (IHC, FC)</li> <li>Olver S, <i>et al.</i> 2006. <i>Cancer Res.</i> 66:571.</li> <li>Fukushima A, <i>et al.</i> 2006. <i>Invest. Ophthalmol. Vis. Sci.</i> 47:657. PubMed</li> <li>Benson MJ, <i>et al.</i> 2007. <i>J. Exp. Med.</i> doi:10.1084/jem.20070719. (FC) PubMed</li> <li>Chappaz S, <i>et al.</i> 2007. <i>Blood</i> doi:10.1182/blood-2007-02-074245. (FC)</li> <li>PubMed</li> <li>Lee JW, <i>et al.</i> 2006. <i>Nature Immunol.</i> 8:181.</li> <li>Shigeta A, et al. 2008. <i>Blood</i> 112:4915 (FC) PubMed</li> <li>de Vries VC, <i>et al.</i> 2009. <i>Am. J. Transplant.</i> 9:2270 PubMed</li> </ol>	

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:** CD62L is a 74-95 kD glycoprotein also known as L-selectin, LECAM-1, Ly-22, LAM-1, and MEL-14. It is a member of the selectin family and is expressed on the majority of B and naïve T cells, a subset of memory T cells, monocytes, granulocytes, most thymocytes, and a subset of NK cells. CD62L is important in lymphocyte homing to high endothelial venules (HEV) in peripheral lymph nodes and leukocyte "rolling" on activated endothelium. CD62L also contributes to neutrophil emigration at inflammatory sites. CD62L is rapidly shed from lymphocytes and neutrophils upon cellular activation and the expression levels of CD62L (in conjunction with other markers) have been used to distinguish naïve, effector, and memory T cells. CD62L has been reported to interact with CD34, GlyCAM-1, and MAdCAM-1.

Antigen 1. Barclay AN, et al. 1997. The Leukocyte Antigen FactsBook Academic Press.

References: 2. Kishimoto TK, et al. 1990. P. Natl. Acad. Sci. USA 87:2244.

3. Tedder TF, et al. 1995. J. Exp. Med. 181:2259.