## **Product Data Sheet**

isotype control (open histogram).

## Brilliant Violet 510<sup>™</sup> anti-human HLA-DR

Catalog # / Size:	2138230 / 100 tests 2138225 / 25 tests	1
Clone:	L243	A
Isotype:	Mouse IgG2a, κ	ig j
<b>Reactivity:</b>	Human	2
Preparation:	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 510 <sup>™</sup> under optimal conditions. The solution is free of unconjugated Brilliant Violet 510 <sup>™</sup> and unconjugated antibody.	Human peripheral blood lymphocytes were stained with HLA- DR (clone L243) Brilliant Violet 510 <sup>™</sup> (filled histogram) or mouse lgG2a, κ Brilliant Violet 510 <sup>™</sup>
Formulation:	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).	
Concentration:	Lot-specific	

## **Applications:**

Applications: Flow Cyt	tometry
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**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 510<sup>™</sup> excites at 405 nm and emits at 510 nm. The bandpass filter 510/50 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 510<sup>™</sup> is a trademark of Sirigen Group Ltd.

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## Application<br/>Notes:The L243 monoclonal antibody reacts with the HLA-DR antigen, a member of MHC<br/>class II molecules. It does not cross react with HLA-DP and HLA-DQ. Clone L243<br/>binds a conformational epitope on HLA-DR $\alpha$ which depends on the correct folding<br/>of the $\alpha\beta$ heterodimer.<sup>19</sup>

Additional reported applications (for the relevant formats) include: immunoprecipitation<sup>8</sup>, Western blotting<sup>8</sup>, *in vitro* blocking of mixed lymphocyte reactions<sup>9,10</sup>, depeletion of MHC class II cells<sup>7</sup>, and immunohistochemical staining of acetone-fixed frozen sections<sup>4,5</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. 307612). For highly sensitive assays, we recommend Ultra-LEAF<sup>m</sup> purified antibody (Cat. No. 307648) with a lower endotoxin limit than standard LEAF<sup>m</sup> purified antibodies (Endotoxin <0.01 EU/microg).

Application References:	<ol> <li>Brodsky F. 1984. <i>Immunogenetics</i> 19:179.</li> <li>Robbins P, <i>et al.</i> 1987. <i>Human Immunol.</i> 18:301.</li> <li>Stites D, <i>et al.</i> 1986. <i>Clin. Immunol. Immunopathol.</i> 38:161.</li> <li>Warnke R, <i>et al.</i> 1980. <i>J. Histochem. Cytochem.</i> 28:771. (IHC)</li> <li>Engleman E, <i>et al.</i> 1981. <i>P. Natl. Acad. Sci. USA</i> 78:1791. (IHC)</li> <li>Zipf T, <i>et al.</i> 1981. <i>Cancer Res.</i> 41:4786.</li> <li>Goodier M, <i>et al.</i> 2000. <i>J. Immunol.</i> 165:139. (Depletion)</li> <li>Esser M, <i>et al.</i> 2001. <i>J. Virol.</i> 75:6173. (IP, WB)</li> <li>Kalka-Moll WM, <i>et al.</i> 2002. <i>J. Immunol.</i> 169:6149. (Block)</li> <li>Wang RF, <i>et al.</i> 1999. <i>Science</i> 284:1351. (Block)</li> <li>Vang RF, <i>et al.</i> 2007. <i>J. Exp. Med.</i> 204:3183. PubMed</li> <li>Fujita H, <i>et al.</i> 2000. <i>P. Natl. Acad. Sci. USA</i> 106:21795. PubMed</li> <li>Charles N, <i>et al.</i> 2010. <i>Infect. Immun.</i> 78:4763. PubMed</li> <li>Yoshino N, <i>et al.</i> 2000. <i>Exp. Anim. (Tokyo)</i> 49:97. (FC)</li> <li>Kim WK, <i>et al.</i> 2001. <i>Leuk. Lymphoma</i> 52:273.</li> <li>Galkowska H, <i>et al.</i> 1996. <i>Vet. Immunol.</i> 169:819. PubMed</li> <li>Salkowska H, <i>et al.</i> 2014. <i>Mol Immunol.</i> 59:19. PubMed</li> </ol>
Description:	HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD $lpha$

Description:	HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD $lpha$
	(heavy) chain and a 27 kD $\beta$ (light) chain. It is expressed on B cells, activated T
	cells, monocytes/macrophages, dendritic cells, and other non-professional APCs.
	In conjunction with the CD3/TCR complex and CD4 molecules, HLA-DR is critical
	for efficient peptide presentation to CD4 <sup>+</sup> T cells.

Antigen	1. Levacher M, <i>et al.</i> 1990. <i>Clin. Exp. Immunol.</i> 81:177.	
<b>References:</b>	2. Terstappen L, <i>et al.</i> 1990. <i>J. Leukocyte Biol.</i> 48:138.	
	3. Edwards JA, <i>et al.</i> 1986. <i>J. Immunol.</i> 137:490.	
	4. van Es A, <i>e</i>	

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