SONY

Product Data Sheet

Brilliant Violet 650™ anti-human HLA-DR

Catalog # / Size: 2138245 / 25 tests
2138250 / 100 tests

Clone: L243

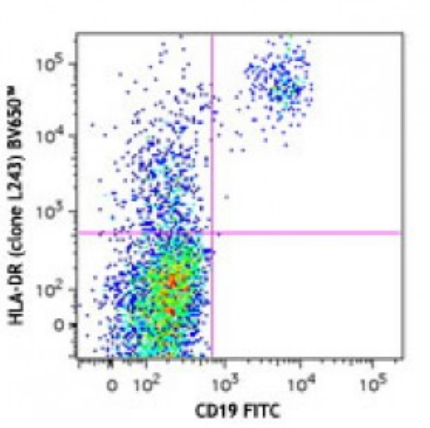
Isotype: Mouse IgG2a, κ

Reactivity: Human

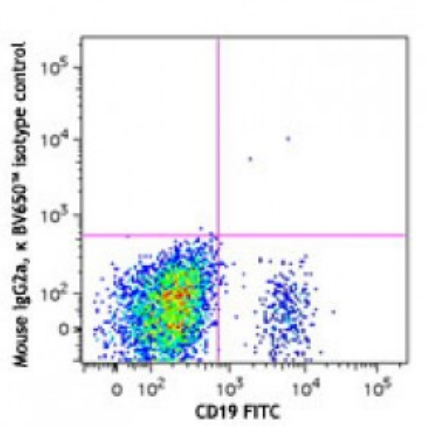
Preparation: The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 650™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 650™ and unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).

Concentration: Lot-specific



Human peripheral blood lymphocytes were stained with CD19 FITC and HLA-DR (clone L243) Brilliant Violet 650™ (top) or mouse IgG2a, κ Brilliant Violet 650™ isotype control (bottom).



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 650™ excites at 405 nm and emits at 645 nm. The bandpass filter 660/20 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 650™ is a trademark of Sirigen Group Ltd.

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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 Notes:

The L243 monoclonal antibody reacts with the HLA-DR antigen, a member of MHC class II molecules. It does not cross react with HLA-DP and HLA-DQ. Clone L243 binds a conformational epitope on HLA-DRα which depends on the correct folding of the αβ heterodimer.[19]

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

Application References:

1. Brodsky F. 1984. *Immunogenetics* 19:179.
2. Robbins P, *et al.* 1987. *Human Immunol.* 18:301.
3. Stites D, *et al.* 1986. *Clin. Immunol. Immunopathol.* 38:161.
4. Warnke R, *et al.* 1980. *J. Histochem. Cytochem.* 28:771. (IHC)
5. Engleman E, *et al.* 1981. *P. Natl. Acad. Sci. USA* 78:1791. (IHC)
6. Zipf T, *et al.* 1981. *Cancer Res.* 41:4786.
7. Goodier M, *et al.* 2000. *J. Immunol.* 165:139. (Depletion)
8. Esser M, *et al.* 2001. *J. Virol.* 75:6173. (IP, WB)
9. Kalka-Moll WM, *et al.* 2002. *J. Immunol.* 169:6149. (Block)
10. Wang RF, *et al.* 1999. *Science* 284:1351. (Block)
11. Zaba LC, *et al.* 2007. *J. Exp. Med.* 204:3183. PubMed
12. Fujita H, *et al.* 2009. *P. Natl. Acad. Sci. USA* 106:21795. PubMed
13. Charles N, *et al.* 2010. *Nat. Med.* 16:701. (FC) PubMed
14. Goncalves RM, *et al.* 2010. *Infect. Immun.* 78:4763. PubMed
15. Yoshino N, *et al.* 2000. *Exp. Anim. (Tokyo)* 49:97. (FC)
16. Kim WK, *et al.* 2006. *Am. J. Pathol.* 168:822. (FC)
17. Stein R, *et al.* 2011. *Leuk. Lymphoma* 52:273.
18. Galkowska H, *et al.* 1996. *Vet. Immunol. Immunopathol.* 53:329.
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20. Lauterbach N, *et al.* 2014. *Mol Immunol.* 59:19. PubMed

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Description:

HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD α (heavy) chain and a 27 kD β (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 $CD4^+$ T cells.

Antigen References:

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