### **Product Data Sheet**

#### Brilliant Violet 750™ anti-human HLA-DR

**Catalog** # / 2138360 / 100 tests

**Size:** 2138355 / 25 tests

Clone: L243

**Isotype:** Mouse IgG2a, κ

Immunogen: Human rheumatoid synovial fluid

cells and fibronectin-purified

monocytes.

Reactivity: Human, Non-human primate, Other

**Preparation:** The antibody was purified by affinity

chromatography and conjugated with Brilliant Violet 750™ under optimal conditions. The solution is free of unconjugated Brilliant Violet 750™

and unconjugated antibody.

**Formulation:** Phosphate-buffered solution, pH 7.2,

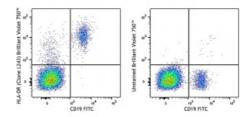
containing 0.09% sodium azide and

BSA (origin USA).

Workshop Number:

VI MA36

Concentration: Lot-specific



Human peripheral blood lymphocytes were stained with CD19 FITC and HLA-DR (Clone L243) Brilliant Violet 750™ (left) or CD19 FITC only (right).

#### **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  $\mu$ l per million cells in 100  $\mu$ l staining volume or 5  $\mu$ l per 100  $\mu$ l of whole blood.

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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-DR $\alpha$  which depends on the correct folding 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>.

# 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.
- 19. Moro M, et al. 2005. BMC Immunol. 6:24.
- 20. Lauterbach N, et al. 2014. Mol Immunol. 59:19. PubMed

#### **Description:**

HLA-DR is a heterodimeric cell surface glycoprotein comprised of a 36 kD  $\alpha$  (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 $^+$ 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, et al. 1984. Transplantation 37:65.
- 5. O'Doherty U, et al. 1994. Immunology 82:487.
- 6. Thomas R, et al. 1994. J. Immunol. 153:4016. 7. Grouard G, et al. 1996. Nature 384:364.