

## GMP APC/Fire™ 750 anti-human HLA-DR Antibody

<b>Catalog# / Size</b>	260170 / 100 tests
<b>Clone</b>	L243
<b>Other Names</b>	Major Histocompatibility Class II, MHC class II
<b>Isotype</b>	Mouse IgG2a, κ
<b>Description</b>	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 <sup>+</sup> T cells.

### Product Details

<b>Reactivity</b>	Human
<b>Antibody Type</b>	Monoclonal
<b>Host Species</b>	Mouse
<b>Formulation</b>	Phosphate-buffered solution, pH 7.2, containing True-Stain Monocyte Blocker™, 0.09% sodium azide and 0.2% (w/v) BSA (origin USA) and a stabilizer.
<b>Preparation</b>	The antibody was purified by affinity chromatography and conjugated with APC/Fire™ 750 under optimal conditions.
<b>Concentration</b>	200 µg/mL
<b>Storage &amp; Handling</b>	The antibody solution should be stored undiluted between 2°C and 8°C, and protected from prolonged exposure to light. <b>Do not freeze.</b>
<b>Application</b>	<a href="#">FC - Quality tested</a>
<b>Recommended Usage</b>	Each lot of this antibody is quality control tested by <a href="#">immunofluorescent staining with flow cytometric analysis</a> . For flow cytometric staining, the suggested use of this reagent is 5 µL per million cells in 100 µL staining volume or 5 µL per 100 µL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.  * APC/Fire™ 750 has a maximum excitation of 650 nm and a maximum emission of 787 nm.
<b>Excitation Laser</b>	Red Laser (633 nm)
<b>Application Notes</b>	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. <sup>19</sup>  Additional reported applications (for the relevant formats) include: immunoprecipitation <sup>8</sup> , Western blotting <sup>8</sup> , <i>in vitro</i> blocking of mixed lymphocyte reactions <sup>9,10</sup> , depletion of MHC class II cells <sup>7</sup> , immunohistochemical staining of acetone-fixed frozen sections <sup>4,5</sup> , and spatial biology (IBEX) <sup>21,22</sup> . For sensitive functional assays, we recommend using the Ultra-LEAF™ purified antibody (Endotoxin < 0.01 EU/µg, Azide-Free, 0.2 µm filtered) (Cat. No. 307648, 307665 - 307669).
<b>Application References</b>	<ol style="list-style-type: none"> <li>1. Brodsky F. 1984. <i>Immunogenetics</i> 19:179.</li> <li>2. Robbins P, et al. 1987. <i>Human Immunol.</i> 18:301.</li> <li>3. Stites D, et al. 1986. <i>Clin. Immunol. Immunopathol.</i> 38:161.</li> <li>4. Warnke R, et al. 1980. <i>J. Histochem. Cytochem.</i> 28:771. (IHC)</li> <li>5. Engleman E, et al. 1981. <i>P. Natl. Acad. Sci. USA</i> 78:1791. (IHC)</li> <li>6. Zipf T, et al. 1981. <i>Cancer Res.</i> 41:4786.</li> <li>7. Goodier M, et al. 2000. <i>J. Immunol.</i> 165:139. (Depletion)</li> <li>8. Esser M, et al. 2001. <i>J. Virol.</i> 75:6173. (IP, WB)</li> <li>9. Kalka-Moll WM, et al. 2002. <i>J. Immunol.</i> 169:6149. (Block)</li> <li>10. Wang RF, et al. 1999. <i>Science</i> 284:1351. (Block)</li> <li>11. Zaba LC, et al. 2007. <i>J. Exp. Med.</i> 204:3183. <a href="#">PubMed</a></li> </ol>
<b>(PubMed link indicates BioLegend citation)</b>	

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13. Charles N, *et al.* 2010. *Nat. Med.* 16:701. (FC) [PubMed](#)
14. Goncalves RM, *et al.* 2010. *Infect. Immun.* 78:4763. [PubMed](#)
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20. Lauterbach N, *et al.* 2014. *Mol Immunol.* 59:19. [PubMed](#)
21. Radtke AJ, *et al.* 2020. *Proc Natl Acad Sci USA.* 117:33455-33465. (SB) [PubMed](#)
22. Radtke AJ, *et al.* 2022. *Nat Protoc.* 17:378-401. (SB) [PubMed](#)

#### Disclaimer

**GMP RUO Flow Cytometry Antibodies.** BioLegend GMP RUO fluorophore conjugated antibodies are manufactured in a dedicated GMP facility and compliant with ISO 13485:2016. For research use only. Not for use in diagnostic or therapeutic procedures. Our processes include:

- Batch-to-batch consistency
- Material traceability
- Documented procedures
- Documented employee training
- Equipment maintenance and monitoring records
- Lot-specific certificates of analysis
- Quality audits per ISO 13485:2016
- QA review of released products

## Antigen Details

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<b>Structure</b>	Ig superfamily, MHC class II, heterodimeric transmembrane protein, 36 kD heavy and 27 kD light chain
<b>Distribution</b>	B cells, activated T cells, monocytes/macrophages, dendritic cells, other APCs
<b>Function</b>	Peptide presentation
<b>Ligand/Receptor</b>	CD3/TCR, CD4
<b>Cell Type</b>	Antigen-presenting cells, B cells, Dendritic cells, Macrophages, Monocytes, T cells, Tregs
<b>Biology Area</b>	Immunology, Innate Immunity
<b>Molecular Family</b>	MHC Antigens
<b>Antigen References</b>	<ol style="list-style-type: none"> <li>1. Levacher M, <i>et al.</i> 1990. <i>Clin. Exp. Immunol.</i> 81:177.</li> <li>2. Terstappen L, <i>et al.</i> 1990. <i>J. Leukocyte Biol.</i> 48:138.</li> <li>3. Edwards JA, <i>et al.</i> 1986. <i>J. Immunol.</i> 137:490.</li> <li>4. van Es A, <i>et al.</i> 1984. <i>Transplantation</i> 37:65.</li> <li>5. O'Doherty U, <i>et al.</i> 1994. <i>Immunology</i> 82:487.</li> <li>6. Thomas R, <i>et al.</i> 1994. <i>J. Immunol.</i> 153:4016.</li> <li>7. Grouard G, <i>et al.</i> 1996. <i>Nature</i> 384:364.</li> </ol>
<b>Gene ID</b>	<a href="#">3122</a> <a href="#">3123</a>

## Related Protocols

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[Cell Surface Flow Cytometry Staining Protocol](#)

## Other Formats

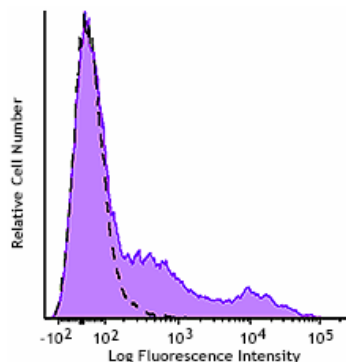
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APC anti-human HLA-DR, FITC anti-human HLA-DR, PE anti-human HLA-DR, PE/Cyanine5 anti-human HLA-DR, Purified anti-human HLA-DR, Biotin anti-human HLA-DR, PE/Cyanine7 anti-human HLA-DR, APC/Cyanine7 anti-human HLA-DR, Alexa Fluor® 488 anti-human HLA-DR, Alexa Fluor® 647 anti-human HLA-DR, Pacific Blue™ anti-human HLA-DR, Alexa Fluor® 700 anti-human HLA-DR, PerCP anti-human HLA-DR, PerCP/Cyanine5.5 anti-human HLA-DR, Brilliant Violet 605™ anti-human HLA-DR, Brilliant Violet 421™ anti-human HLA-DR, Brilliant Violet 570™ anti-human HLA-DR, Brilliant Violet 711™ anti-human HLA-DR, Brilliant Violet 785™ anti-human HLA-DR, Brilliant Violet 510™ anti-human HLA-DR, Ultra-LEAF™ Purified anti-human HLA-DR, Brilliant Violet 650™ anti-human HLA-DR, Purified anti-human HLA-DR (Maxpar® Ready), PE/Dazzle™ 594 anti-human HLA-DR, APC/Fire™ 750 anti-human HLA-DR, TotalSeq™-A0159 anti-human HLA-DR, TotalSeq™-B0159 anti-human HLA-DR, TotalSeq™-C0159 anti-

human HLA-DR, Brilliant Violet 750™ anti-human HLA-DR, APC/Fire™ 810 anti-human HLA-DR, PE/Fire™ 640 anti-human HLA-DR, Spark Violet™ 538 anti-human HLA-DR Antibody, KIRAVIA Blue 520™ anti-human HLA-DR, TotalSeq™-D0159 anti-human HLA-DR, PE/Fire™ 810 anti-human HLA-DR, GMP PE/Dazzle™ 594 anti-human HLA-DR, Spark Violet™ 423 anti-human HLA-DR, GMP FITC anti-human HLA-DR, GMP APC anti-human HLA-DR

## Product Data

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Typical results from human peripheral blood lymphocytes and monocytes stained either with L243 APC/Fire™ 750 used at 5  $\mu$ L/test (filled histogram) or with an isotype control (open histogram).

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