## Technical Data Sheet

# V450 Mouse Anti-Human HLA-DR

#### **Product Information**

561359 **Material Number:** 

MHC class II antigen; HLA class II histocompatibility antigen **Alternate Name:** 

Size: 5 μl Vol. per Test: G46-6 Clone:

Mouse IgG2a, κ Isotype: Reactivity: QC Testing: Human

Workshop:

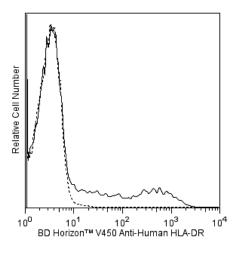
Storage Buffer: Aqueous buffered solution containing protein stabilizer and ≤0.09% sodium

azide

### Description

The G46-6 monoclonal antibody specifically binds to HLA-DR, a major histocompatibility complex (MHC) class II antigen. HLA-DR antigens are encoded by genes within the Human Leukocyte Antigen (HLA) Complex located on chromosome 6. HLA-DR is a transmembrane heterodimeric glycoprotein composed of an α chain (36 kDa) and a β subunit (27 kDa) expressed primarily on antigen presenting cells: B cells, dendritic cells, monocytes, macrophages, and thymic epithelial cells. HLA-DR is also expressed on activated T cells. This molecule plays a major role in mediating cellular interactions during antigen presentation to CD4-positive T cells.

The antibody is conjugated to BD Horizon<sup>TM</sup> V450, which has been developed for use in multicolor flow cytometry experiments and is available exclusively from BD Biosciences. It is excited by the Violet laser Ex max of 406 nm and has an Em Max at 450 nm. Conjugates with BD Horizon<sup>TM</sup> V450 can be used in place of Pacific Blue<sup>TM</sup> conjugates.



Flow cytometric analysis of HLA-DR expression on human peripheral blood lymphocytes. Whole blood was stained with either BD Horizon™ V450 Mouse anti-Human HLA-DR antibody (Cat. No. 561359; solid line histogram) or with a BD Horizon™ V450 Mouse IgG2a, κ Isotype Control (Cat. No. 560550; dashed line histogram). The erythrocytes were lysed with BD PharmLyse™ Lysing Buffer (Cat. No. 555899). The fluorescence histograms were derived from events with the forward and side light-scatter characteristics of viable lymphocytes. Flow cytometry was performed using a BD™ LSR II Flow Cytometer System.

### **Preparation and Storage**

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. The antibody was conjugated with BD Horizon™ V450 under optimum conditions, and unreacted BD Horizon™ V450 was removed.

### **Application Notes**

Application

Flow cytometry Routinely Tested

### **BD Biosciences**

bdbiosciences.com

**United States** Europe 32.53.720.550 877.232.8995 888.268.5430 0120.8555.90 65.6861.0633 0800.771.7157

Store undiluted at 4°C and protected from prolonged exposure to light. Do not freeze.

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### **Suggested Companion Products**

| Catalog Number | Name Name                           | Size   | <u>Clone</u> |  |
|----------------|-------------------------------------|--------|--------------|--|
| 560550         | V450 Mouse IgG2a, κ Isotype Control | 0.1 mg | G155-178     |  |
| 555899         | Lysing Buffer                       | 100 ml | (none)       |  |
| 554656         | Stain Buffer (FBS)                  | 500 ml | (none)       |  |

#### **Product Notices**

- This reagent has been pre-diluted for use at the recommended Volume per Test. We typically use 1 × 10<sup>6</sup> cells in a 100-µl experimental sample (a test).
- 2. An isotype control should be used at the same concentration as the antibody of interest.
- 3. BD Horizon™ V450 has a maximum absorption of 406 nm and maximum emission of 450 nm. Before staining with this reagent, please confirm that your flow cytometer is capable of exciting the fluorochrome and discriminating the resulting fluorescence.
- 4. Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 5. Pacific Blue<sup>TM</sup> is a trademark of Molecular Probes, Inc., Eugene, OR.
- 6. For fluorochrome spectra and suitable instrument settings, please refer to our Fluorochrome Web Page at www.bdbiosciences.com/colors.
- 7. Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.

#### References

Barclay NA, Brown MH, Birkeland ML, et al, ed. The Leukocyte Antigen FactsBook. San Diego, CA: Academic Press; 1997. (Biology)

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