

## **Product datasheet for SM2201F**

### OriGene Technologies, Inc.

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## **HLA Class II DR Rat Monoclonal Antibody [Clone ID: YD1/63.4.10]**

**Product data:** 

**Product Type:** Primary Antibodies

Clone Name: YD1/63.4.10

Applications: FC

**Recommended Dilution:** Flow cytometry.

Immunohistochemistry on cryostat and Paraffin-embedded sections.

Reactivity: Human

Host: Rat

**Isotype:** IgG2a

Clonality: Monoclonal Immunogen: DAUDI cells

Donor: immunized DA rat spleen cells Fusion Partner: Y3 Ag1.2.3 rat myeloma

**Specificity:** Anti-human HLA-DR monoclonal antibody recognizes the HLA-DR (MHC class II) antigen.

Formulation: PBS containing 0.02% sodium azide (NaN3) as preservative and EIA grade BSA as a stabilizing

protein to bring total protein concentration to 4-5 mg/ml.

Label: FITC

State: Liquid purified Ig fraction

Label: Fluorescein isothiocyanate isomer 1

**Concentration:** lot specific

**Purification:** Affinity chromatography on Protein G

Conjugation: FITC

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

Synonyms: HLA-DR, HLA class II histocompatibility antigen DR, MHC class II antigen DR





Note:

#### Protocol: FLOW CYTOMETRYANALYSIS:

#### Method:

- 1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte®-H cell separation medium.
- 2. Wash 2 times.
- 3. Resuspend the cells to a concentration of 2x10e7 cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain 1x10e6 cells, representing 1 test).
- 4. To each tube, add 50µl of a 0.2-0.1µg dilution of this antibody per 10e6 cells.
- 5. Vortex the tubes to ensure thorough mixing of antibody and cells.
- 6. Incubate the tubes for 30 minutes at 4°C.
- 7. Wash 2 times at 4°C.
- 8. Resuspend the cell pellet in 50 µl ice cold media B.
- 9. Transfer to suitable tubes for flow cytometric analysis containing 15  $\mu$ l of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

#### Media:

A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).

B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100  $\mu$ l of 2M sodium azide in 100 mls).

#### **Results:**

**Tissue Distribution by Flow Cytometry Analysis:** 

**Cell Concentration: 1x10e6 cells per test** 

Antibody Concentration Used: 0.1 μg/10e6 cells

**Isotypic Control: FITC Rat IgG2a** 

# **Product images:**

