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# TECHNICALLY *Speaking*

Place your order with CEDARLANE® or your local distributor.  
Please contact CEDARLANE® for lot specific information.

## Purified Anti-Mouse CD19 Antigen Monoclonal Antibody

CL8914AP  
LOT: 21000208

### DESCRIPTION:

Cedarlane's anti-mouse CD19 antigen monoclonal antibody is a pan B cell marker. It will not detect plasma cells.

This antibody is suitable for use in flow cytometry.

### PRESENTATION:

200 µg purified Ig buffered in PBS and 0.02% NaN<sub>3</sub>.

### STORAGE/STABILITY:

Store at 4°C. For long term storage, aliquot and freeze unused portion at -20°C in volumes appropriate for single usage. Avoid freeze/thaw cycles. If the reagent is being diluted, it is recommended that the quantity to be used within one week be diluted.

### SPECIFICATIONS:

Clone: 6D5

Specificity: Mouse CD19

Ig Class: Rat IgG<sub>2a</sub>

Antibody Concentration: 0.2 mg/ml

For more information or to place an order please contact...

**CEDARLANE®**  
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website: [www.cedarlanelabs.com](http://www.cedarlanelabs.com) • e-mail: [info@cedarlanelabs.com](mailto:info@cedarlanelabs.com)

## **FLOW CYTOMETRY ANALYSIS:**

### **Method:**

1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte<sup>®</sup>-M cell separation medium (CL5030).
2. Wash 2 times.
3. Resuspend the cells to a concentration of  $2 \times 10^7$  cells/ml in media A. Add 50  $\mu$ l of this suspension to each tube (each tube will then contain  $1 \times 10^6$  cells, representing 1 test).
4. To each tube, add  $\sim 1.0 \mu\text{g}^*$  of **CL8914AP**.
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. Add 100  $\mu$ l of secondary antibody **CLCC40001** (FITC Goat anti-rat IgG (H+L)) at 1:500 dilution.
9. Incubate the tubes at 4°C for 30-60 minutes.  
(It is recommended that the tubes are protected from light since most fluorochromes are light sensitive).
10. Wash 2 times at 4°C in media B.
11. Resuspend the cell pellet in 50  $\mu$ l ice cold media B.
12. 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).

### **REFERENCES:**

1. Krop, I. et al (1996). Antibody to CD19 suppresses self-renewal of B-1 lymphocytes. Eur. J. Immunol. 26:238

**FOR RESEARCH USE ONLY**

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