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## PE/Cyanine7 Rabbit anti-Human CR2/CD21 mAb

Catalog No.: A26631

## **Basic Information**

#### **Observed MW**

## **Calculated MW**

110kDa/113kDa/118kDa/119kDa

## Category

Primary antibody

## **Applications**

FC

## **Cross-Reactivity**

Human

## CloneNo number

ARC56643-PE-Cy7

## Conjugate

PE-Cy7. Ex:565nm. Em:778nm.

## **Background**

This gene encodes a membrane protein, which functions as a receptor for Epstein-Barr virus (EBV) binding on B and T lymphocytes. Genetic variations in this gene are associated with susceptibility to systemic lupus erythematosus type 9 (SLEB9). Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

## **Recommended Dilutions**

FC

5  $\mu$ l per 10^6 cells in 100  $\mu$ l volume

## **Immunogen Information**

Gene ID 1380 **Swiss Prot** 

P20023

### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 21-971 of human CR2/CD21 (NP\_001868.2).

## **Synonyms**

CR; C3DR; CD21; CVID7; SLEB9

## **Contact**

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## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

#### Storage

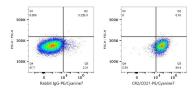
Store at 2-8°C. Avoid freeze.

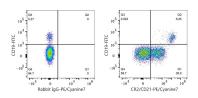
Buffer: PBS with 0.09% Sodium azide, 0.2% BSA, pH7.3.

## **Validation Data**









Flow cytometry: 1X10^6 HAP1 cells (negative control,left) and Daudi cells (right) were surface-stained with PE/Cyanine7 Rabbit anti-Human CR2/CD21 mAb (A26631,5 µl/Test,orange line) or PE/Cyanine7 Rabbit IgG isotype control (5 µl/Test,blue line). Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry: 1X10^6 Daudi cells were surface-stained with PE/Cyanine7 Rabbit IgG isotype control (5 µl/Test,left) or PE/Cyanine7 Rabbit anti-Human CR2/CD21 mAb (A26631,5 µl/Test,right).

Flow cytometry:1X10^6 Human PBMC were surface-stained with FITC Mouse anti-Human CD19 mAb (A22815,5  $\mu$ I/Test) and PE Rabbit IgG isotype control (A24172,5  $\mu$ I/Test,left) or PE/Cyanine7 Rabbit anti-Human CR2/CD21 mAb (A26631,5  $\mu$ I/Test,right). Cells in the lymphocyte gate were used for analysis.