# PE Rabbit anti-Mouse CD83 mAb

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ABclomal

Catalog No.: A27505

### **Basic Information**

#### **Observed MW**

### **Calculated MW**

21KDa

## Category

Primary antibody

## **Applications**

FC

### **Cross-Reactivity**

Mouse

#### CloneNo number

ARC61465

### Conjugate

PE. Ex:565nm. Em:574nm.

# **Background**

Acts upstream of or within positive regulation of CD4-positive, alpha-beta T cell differentiation; regulation of cytokine production; and response to organic cyclic compound. Located in external side of plasma membrane. Is expressed in alimentary system; genitourinary system; hemolymphoid system gland; and nervous system. Orthologous to human CD83 (CD83 molecule).

## **Recommended Dilutions**

FC

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

# **Immunogen Information**

**Gene ID** 12522

**Swiss Prot** 

088324

### **Immunogen**

Recombinant protein (or fragment). This information is considered to be commercially sensitive.

## **Synonyms**

mCD83

## **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.













Flow cytometry: 1X10^6 C57BL/6 mouse splenocytes (untreated,left) and C57BL/6 mouse splenocytes (treated with 10µg/mL LPS for 4h,right) were surface-stained with PE Rabbit anti-Mouse CD83 mAb (A27505,5 µl/Test,orange line) or PE Rabbit IgG isotype control (A24172,5 µl/Test,blue line). Nonfluorescently stained cells were used as blank control (red line).

Flow cytometry:  $1X10^6$  C57BL/6 mouse splenocytes (untreated,left) and C57BL/6 mouse splenocytes(treated with  $10\mu g/mL$  LPS for 4h,right) were surface-stained with PE Rabbit anti-Mouse CD83 mAb (A25505,5  $\mu$ l/Test).

Flow cytometry: 1X10^6 293T cells (negative control,left) and 293T (Transfection,right) cells were surface-stained with PE Rabbit anti-Mouse CD83 mAb (A27505,5  $\mu$ l/Test,orange line) or PE Rabbit lgG isotype control (A24172,5  $\mu$ l/Test,blue line). Non-fluorescently stained cells were used as blank control (red line).