# PE Rabbit-anti Human CD3 mAb

Catalog No.: A27110



### **Basic Information**

#### **Observed MW**

Calculated MW 23kDa

Category

Primary antibody

**Applications** 

**Cross-Reactivity** 

Human

CloneNo number

ARC68274-PE

Conjugate

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

# **Background**

The protein encoded by this gene is the CD3-epsilon polypeptide, which together with CD3-gamma, -delta and -zeta, and the T-cell receptor alpha/beta and gamma/delta heterodimers, forms the T-cell receptor-CD3 complex. This complex plays an important role in coupling antigen recognition to several intracellular signal-transduction pathways. The genes encoding the epsilon, gamma and delta polypeptides are located in the same cluster on chromosome 11. The epsilon polypeptide plays an essential role in T-cell development. Defects in this gene cause immunodeficiency. This gene has also been linked to a susceptibility to type I diabetes in women.

## **Recommended Dilutions**

FC

5 μl per 10^6 cells in 100 μl volume

# Immunogen Information

Gene ID 916 **Swiss Prot** 

P07766

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-126 of human CD3 (NP\_000724.1).

## **Synonyms**

T3E; TCRE; IMD18; CD3epsilon

## **Contact**

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### **Product Information**

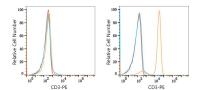
SourceIsotypePurificationRabbitIgGAffinity purification

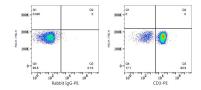
#### **Storage**

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 Raji cells (negative control,left) and Human PBMC (right) were surface-stained with PE Rabbit-anti Human CD3 mAb (A27110,5 µl/Test,orange line) or PE Rabbit IgG isotype control (A24172,5 µl/Test,blue line). Non-fluorescently stained cells were used as blank control (red line). Cells in the lymphocyte gate were used for analysis.

Flow cytometry: 1X10^6 Human PBMC were surface-stained with PE Rabbit IgG isotype control (A24172,5  $\mu$ I/Test,Ieft) or PE Rabbit-anti Human CD3 mAb (A27110,5  $\mu$ I/Test,right). Cells in the lymphocyte gate were used for analysis.