Leader in Biomolecular Solutions for Life Science

PerCP Mouse anti-Human CD3 mAb

Catalog No.: A26828



Basic Information

Observed MW

Calculated MW 23kDa

Category Primary antibody

Applications FC

Cross-Reactivity Human

CloneNo number AMC50002

Conjugate PerCP. Ex:482nm. Em:678nm.

Recommended Dilutions

FC

5 μl per 10^6 cells in 100 μl volume

Background

The protein encoded by this gene is the CD3-epsilon polypeptide, which together with CD3gamma, -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.

Immunogen Information

Gene ID 916 Swiss Prot P07766

Immunogen

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

Synonyms

T3E; TCRE; IMD18; CD3epsilon

Contact

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|-----------|---------------------------|
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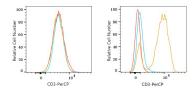
Product Information

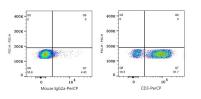
Source Mouse **Isotype** IgG **Purification** Affinity purification

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 PerCP Mouse anti-Human CD3 mAb (A26828, μ I/Test,orange line) or PerCP Mouse IgG 2a isotype control (A26378,5 μ /Test,blue line). Non-fluorescently stained cells were used as blank control (red line).Cells in the Lymphocytes gate were used for analysis.

Flow cytometry: 1X10^6 Human PBMC were surface-stained with PerCP Mouse IgG 2a isotype control (A26378,5 μ I/Test,left) or PerCP Mouse anti-Human CD3 mAb (A26828,5 μ I/Test,right).Cells in the Lymphocytes gate were used for analysis.