# **CD3E Rabbit pAb**

Catalog No.: A1753 8 Publications



## **Basic Information**

#### **Observed MW**

23kDa

#### **Calculated MW**

23kDa

#### Category

Primary antibody

### **Applications**

WB,IHC-P,IF/ICC,IP,FC,ELISA

#### **Cross-Reactivity**

Human, Mouse, Rat

## **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**

**WB** 1:500 - 1:1000

**IHC-P** 1:50 - 1:200

**IF/ICC** 1:50 - 1:200

IP 0.5μg-4μg antibody for 200μg-400μg extracts of

whole cells

FC 1:100 - 1:500

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

## **Immunogen Information**

**Gene ID**916

Swiss Prot
P07766

#### **Immunogen**

Recombinant fusion protein containing a sequence corresponding to amino acids 1-126 of human CD3E (NP\_000724.1).

## Synonyms

T3E; TCRE; IMD18; CD3epsilon; CD3E

## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

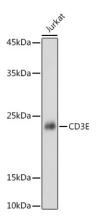
#### **Storage**

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.

## Contact

2	400-999-6126
$\bowtie$	cn.market@abclonal.com.cr
•	www.abclonal.com.cr

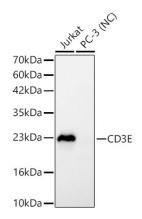


Western blot analysis of lysates from Jurkat cells, using CD3E Rabbit pAb (A1753) at 1:1000 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins:  $25\mu g$  per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 1s.



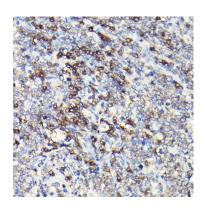
Western blot analysis of various lysates using Human CD3E Rabbit pAb (A1753) at 1:900 dilution. Secondary antibody: HRP-conjugated Goat anti-Rabbit lgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

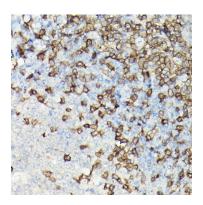
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020). Negative control (NC): PC-3.

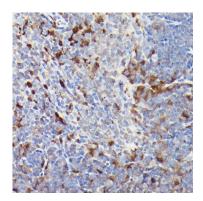
Exposure time: 1s.



Immunohistochemistry analysis of paraffinembedded Human extranodal NK-T cell lymphoma using CD3E Rabbit pAb (A1753) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.

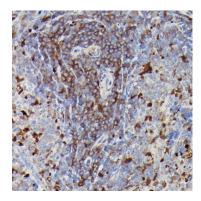


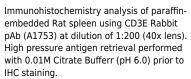
Immunohistochemistry analysis of paraffinembedded Human tonsil using CD3E Rabbit pAb (A1753) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.

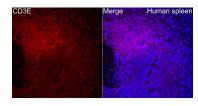


Immunohistochemistry analysis of paraffinembedded Mouse spleen using CD3E Rabbit pAb (A1753) at dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Bufferr (pH 6.0) prior to IHC staining.

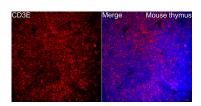
## **Validation Data**



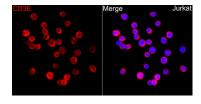


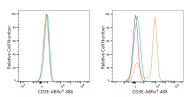


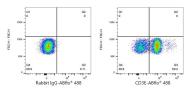
Immunofluorescence analysis of paraffinembedded Human spleen tissue using CD3E Rabbit pAb (A1753) at a dilution of 1:100 (40x lens). Secondary antibody:Cy3 Goat Anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining. Perform high pressure antigen retrieval with 0.01 M citrate buffer (pH 6.0) prior to IF staining.



Immunofluorescence analysis of paraffinembedded Mouse thyums tissue using CD3E Rabbit pAb(A1753) at a dilution of 1:100 (40x lens). Secondary antibody:Cy3 Goat Anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining. Perform high pressure antigen retrieval with 0.01 M citrate buffer (pH 6.0) prior to IF staining.







Immunofluorescence analysis of Jurkat cells using Human CD3E Rabbit pAb (A1753) at dilution of 1:100 (40x lens). Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.

Flow cytometry: 1X10^6 Raji cells (negative control,left) and Human PBMC (right) were surface-stained with CD3E Rabbit pAb (A1753,2 µg/mL,orange line) or Rabbit IgG isotype control (AC042,2 µg/mL,blue line), followed by FITC conjugated goat anti-Rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry: 1X10^6 Human PBMC were surface-stained with Rabbit IgG isotype control (AC042,2  $\mu$ g/mL,Ieft) or CD3E Rabbit pAb (A1753,2  $\mu$ g/mL,right), followed by FITC conjugated goat anti-Rabbit pAb staining.