

# PE Rabbit anti-Human CD96/TACTILE mAb

Catalog No.: A26836

## Basic Information

### Observed MW

### Calculated MW

66kDa

### Category

Primary antibody

### Applications

FC

### Cross-Reactivity

Human

### CloneNo number

ARC62525-PE

### Conjugate

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

## Background

The protein encoded by this gene belongs to the immunoglobulin superfamily. It is a type I membrane protein. The protein may play a role in the adhesive interactions of activated T and NK cells during the late phase of the immune response. It may also function in antigen presentation. Alternative splicing generates multiple transcript variants encoding distinct isoforms.

## Recommended Dilutions

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

## Immunogen Information

### Gene ID

10225

### Swiss Prot

P40200

### Immunogen

Recombinant Protein corresponding to a sequence within amino acids 25-519 of human CD96/TACTILE (NP\_937839.1).

### Synonyms

TACTILE

## Contact

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## Product Information

### Source

Rabbit

### Isotype

IgG

### Purification

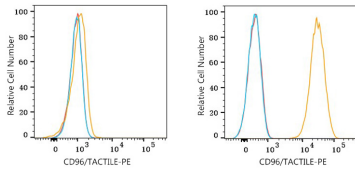
Affinity purification

### Storage

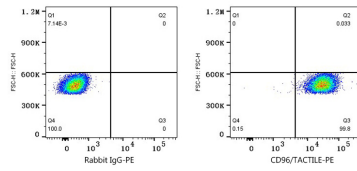
Store at 2-8°C. Avoid freeze.

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

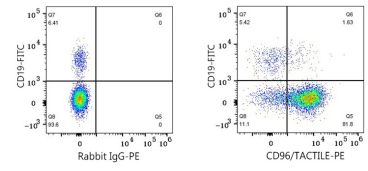
## Validation Data



Flow cytometry: 1X10<sup>6</sup> HeLa cells (negative control, left) and MOLT-4 cells (right) were surface-stained with PE Rabbit anti-Human CD96/TACTILE mAb (A26836, 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).



Flow cytometry: 1X10<sup>6</sup> MOLT-4 cells were surface-stained with PE Rabbit IgG isotype control (A24172, 5 μl/Test, left) or PE Rabbit anti-Human CD96/TACTILE mAb (A26836, 5 μl/Test, right).



Flow cytometry: 1X10<sup>6</sup> Human PBMC were surface-stained with FITC Mouse anti-Human CD19 mAb (A22815, 5 μl/Test) and PE Rabbit IgG isotype control (A24172, 5 μl/Test, left) or PE Rabbit anti-Human CD96/TACTILE mAb (A26836, 5 μl/Test, right). Cells in the lymphocyte gate were used for analysis.