

PE Rabbit anti-Human CD13/ANPEP mAb

Catalog No.: A26514

Basic Information

Observed MW

Refer to figures

Calculated MW

110kDa

Category

Primary antibody

Applications

FC

Cross-Reactivity

Human

CloneNo number

ARC53703

Conjugate

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

Recommended Dilutions

FC 5 µl per 10⁶ cells in
100 µl volume

Background

Aminopeptidase N is located in the small-intestinal and renal microvillar membrane, and also in other plasma membranes. In the small intestine aminopeptidase N plays a role in the final digestion of peptides generated from hydrolysis of proteins by gastric and pancreatic proteases. Its function in proximal tubular epithelial cells and other cell types is less clear. The large extracellular carboxyterminal domain contains a pentapeptide consensus sequence characteristic of members of the zinc-binding metalloproteinase superfamily. Sequence comparisons with known enzymes of this class showed that CD13 and aminopeptidase N are identical. The latter enzyme was thought to be involved in the metabolism of regulatory peptides by diverse cell types, including small intestinal and renal tubular epithelial cells, macrophages, granulocytes, and synaptic membranes from the CNS. This membrane-bound zinc metalloprotease is known to serve as a receptor for the HCoV-229E alphacoronavirus as well as other non-human coronaviruses. This gene has also been shown to promote angiogenesis, tumor growth, and metastasis and defects in this gene are associated with various types of leukemia and lymphoma.

Immunogen Information

Gene ID

290

Swiss Prot

P15144

Immunogen

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

Synonyms

APN; AP-M; AP-N; CD13; LAP1; P150; PEPN; hAPN; GP150

Contact

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Product Information

Source

Rabbit

Isotype

IgG

Purification

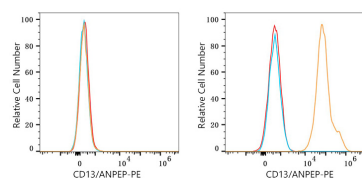
Affinity purification

Storage

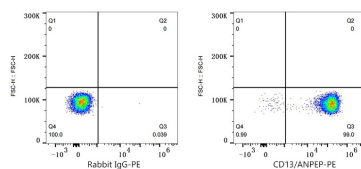
Store at 2-8°C. Avoid freeze.

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

Validation Data



Flow cytometry: 1×10^6 Jurkat cells (negative control, left) and Hep G2 cells (right) were surface-stained with PE Rabbit anti-Human CD13/ANPEP mAb (A26514, 5 μ l/Test, orange line) or PE Rabbit IgG isotype control (A24172, 5 μ l/Test, blue line). Non-fluorescently stained cells were used as blank control (red line).



Flow cytometry: 1×10^6 Human PBMC were surface-stained with PE Rabbit IgG isotype control (A24172, 5 μ l/Test, left) or PE Rabbit anti-Human CD13/ANPEP mAb (A26514, 5 μ l/Test, right). Cells in the Monocyte gate were used for analysis.