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Semaphorin-4D/CD100 Rabbit mAb

Catalog No.: A24404 Recombinant

Basic Information

Observed MW

Calculated MW
82kDa/96kDa

Category
Primary antibody

Applications
IF/ICC,FC,ELISA

Cross-Reactivity
Human

CloneNo number
ARC63157

Background

Plexin-B1 is related to axon guidance in the central nervous system. The Sema4D function is also widely detected in the immune system and was found to be the first semaphorin expressed on the surface of many types of immune cells. In the immune system, CD72 is a low-affinity receptor for Sema4D, and studies have shown that Sema4D can not only regulate T cell activation, but also participate in the regulation of B cell survival and differentiation. In the immune system, soluble fragments containing extracellular domains produced by proteolytic cleavage can regulate many physiological functions of Sema4D. Sema4D is also associated with tumorigenesis because studies have confirmed that it is overexpressed in various types of solid tumor cells. To some extent, the role of Sema4D in tumorigenesis is related to its ability to cause tumor angiogenesis, cell invasion, and immunosuppression by enhancing bone marrow-derived suppressor cell function.

Recommended Dilutions

IF/ICC	1:50-1:200
FC	1:50 - 1:200
ELISA	Recommended starting concentration is 1 μg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID
10507

Swiss Prot
Q92854

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

Synonyms
SEMA4D; C9orf164; CD100; M-sema-G; SEMAJ; coll-4; semaphorin-4D; Semaphorin-4D/CD100

Contact

400-999-6126

cn.market@abclonal.com.cn

www.abclonal.com.cn

Product Information

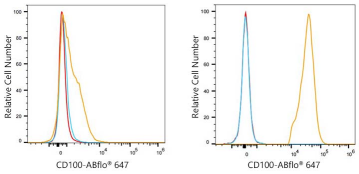
Source
Rabbit

Isotype
IgG

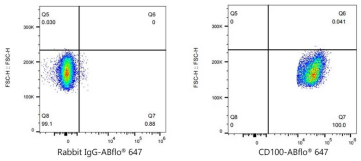
Purification
Affinity purification

Storage
Store at -20℃. Avoid freeze / thaw cycles.
Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

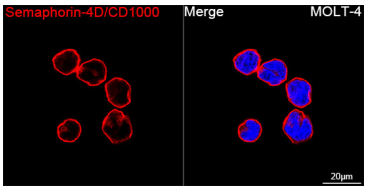
Validation Data



Flow cytometry: 1X10^6 PC-3 cells (Low Expression,left) and MOLT-4 cells (right) were surface-stained with Semaphorin-4D/CD100 Rabbit mAb(A24404,2µg/mL,orange line) or ABflo® 647 Rabbit IgG isotype control (A22070,2µg/mL,blue line),followed by Alexa Fluor®647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).



Flow cytometry:1X10^6 MOLT-4 cells were surface-stained with ABflo® 647 Rabbit IgG isotype control (A22070,2 µg/mL,left) or Semaphorin-4D/CD100 Rabbit mAb(A24404,2 µg/mL,right).



Confocal imaging of MOLT-4 cells using Semaphorin-4D/CD100 Rabbit mAb (A24404, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.