ICAM3/CD50 Rabbit mAb

Catalog No.: A25722 Recombinant



Basic Information

Observed MW

140kDa

Calculated MW

60kDa

Category

SMab Recombinant Monoclonal Antibody

Applications

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

Cross-Reactivity

Human

Background

The protein encoded by this gene is a member of the intercellular adhesion molecule (ICAM) family. All ICAM proteins are type I transmembrane glycoproteins, contain 2-9 immunoglobulin-like C2-type domains, and bind to the leukocyte adhesion LFA-1 protein. This protein is constitutively and abundantly expressed by all leucocytes and may be the most important ligand for LFA-1 in the initiation of the immune response. It functions not only as an adhesion molecule, but also as a potent signalling molecule. Alternative splicing results in multiple transcript variants encoding different isoforms.

Recommended Dilutions

WB	1:1000 - 1:5000
IF/ICC	1:200 - 1:800
IF-P	1:200 - 1:800
IHC-P	1:200 - 1:2000
FC	1:500 - 1:1000
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific

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assay requirements.

Immunogen Information

Gene ID	Swiss Prot
3385	P32942

Immunogen

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

Synonyms

CD50; CDW50; ICAM-R

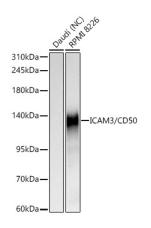
Product Information

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

Storage

Store at -20°C. 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.



Western blot analysis of various lysates using ICAM3/CD50 Rabbit mAb (A25722)at 1:1600 dilution incubated overnight at 4°C.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000

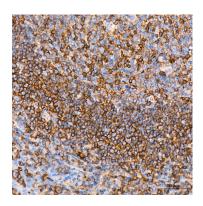
dilution.

Lysates/proteins: 25µg per lane.

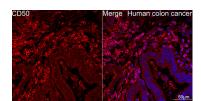
Blocking buffer: 3% nonfat dry milk in TBST.

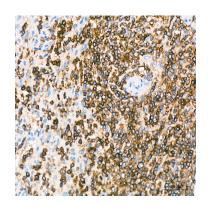
Detection: ECL Basic Kit (RM00020) Negative control (NC): Daudi

Exposure time: 1s.

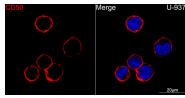


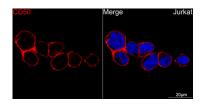
Immunohistochemistry analysis of paraffin-embedded Human tonsil tissue using ICAM3/CD50 Rabbit mAb (A25722) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



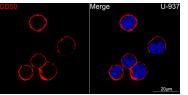


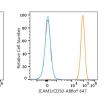
Immunohistochemistry analysis of paraffin-embedded Human spleen tissue using ICAM3/CD50 Rabbit mAb (A25722) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.





Confocal imaging of Jurkat cells using ICAM3/CD50 Rabbit mAb (A25722, 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.



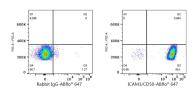


Confocal imaging of paraffinembedded Human colon cancer tissue using ICAM3/CD50 Rabbit mAb (A25722, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer

Confocal imaging of U-937 cells using ICAM3/CD50 Rabbit mAb (A25722, 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.

Flow cytometry: 1X10^6 Daudi cells (negative control,left) and U-937 cells (right) were surface-stained with ICAM3/CD50 Rabbit mAb (A25722,2 μg/mL, or ange line), or Rabbit IgG isotype control (AC042,2 µg/mL,blue line), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining. Non-fluorescently stained cells were used as blank control (red

(pH 6.0) prior to IF staining. Objective: 40x. line).



Flow cytometry: 1X10^6 U-937 cells were surface-stained with Rabbit IgG isotype control (AC042,2 µg/mL,left) or ICAM3/CD50 Rabbit mAb (A25722,2 µg/mL,right), followed by Alexa Fluor® 647 conjugated goat anti-rabbit pAb staining.