CD24 Rabbit mAb

Catalog No.: A25473 Recombinant



Basic Information

Observed MW

37kDa

Calculated MW

8kDa

Category

Primary antibody

Applications

ELISA,WB,IF/ICC,FC

Cross-Reactivity

Mouse

CloneNo number

ARC66569

Background

CD24, also know as heat stable antigen HSA, is a P-selectin ligand involved in adhesion. It is a GPI-anchored glycoprotein expressed on many types of cells, including hematopoietic cells, neural cells, and epithelial cells. CD24 is widely used to delineate stages of lymphocyte development . It also binds to Siglec-10 in humans or Siglec-G in mice . CD24 is frequently used as a marker to identify and isolate cancer stem cells in various cancer types .

Recommended Dilutions

WB 1:1000 - 1:2000

IF/ICC 1:50 - 1:200

FC 1:500 - 1:1000

Immunogen Information

Gene IDSwiss Prot
12484
P24807

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 27-53 of mouse CD24 (NP_033976.1).

Synonyms

HSA; Cd24; Ly-52; nectadrin

Contact

6	400-999-6126
\bowtie	cn.market@abclonal.com.cn
$\overline{\bullet}$	www.abclonal.com.cn

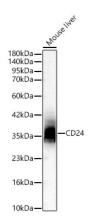
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

Storage

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

Buffer: PBS with 0.05% proclin300,0.05% BSA,50% glycerol,pH7.3.

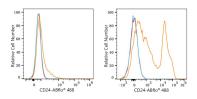


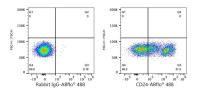
Western blot analysis of lysates from Mouse liver using CD24 Rabbit mAb(A25473) at 1:1000 dilution. Secondary antibody:HRP Goat Anti-Rabbit lgG (H+L)(AS014) at 1:10000 dilution. Lysates/proteins: 25 μ g per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection:ECL Basic Kit (RM00020).

Exposuretime:30s.



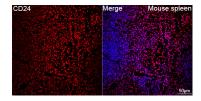


DAPI Merge Neuro-2a

Flow cytometry: 1X10^6 RAW 264.7 cells (Low Expression,left) and C57BL/6 mouse Splenocytes (right) were surface-stained with CD24 Rabbit mAb (A25473,2 µg/mL,orange line) or ABflo® 488 Rabbit IgG isotype control (A22069,5 µl/Test,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 C57BL/6 mouse Splenocytes cells were surface-stained with ABflo® 488 Rabbit IgG isotype control (A22069,5 µl/Test,left) or CD24 Rabbit mAb (A25473,2 µg/mL,right).

Confocal imaging of Neuro-2a cells using CD24 Rabbit mAb (A25473, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.



Confocal imaging of paraffin-embedded Mouse speeln tissue using CD24 Rabbit mAb (A25473, 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: 40x. Perform high pressure antigen retrieval with 0.01 M citrate buffer (pH 6.0) prior to IF staining.