

CD24 Rabbit mAb

Catalog No.: A25383 **Recombinant**

Basic Information

Observed MW

30-45kDa

Calculated MW

8kDa; 13kDa

Category

Primary antibody

Applications

ELISA, WB, FC, IF/ICC

Cross-Reactivity

Human

CloneNo number

ARC66763

Background

This gene encodes a sialoglycoprotein that is expressed on mature granulocytes and B cells and modulates growth and differentiation signals to these cells. The precursor protein is cleaved to a short 32 amino acid mature peptide which is anchored via a glycosyl phosphatidylinositol (GPI) link to the cell surface. This gene was missing from previous genome assemblies, but is properly located on chromosome 6. Non-transcribed pseudogenes have been designated on chromosomes 1, 15, 20, and Y. Alternative splicing results in multiple transcript variants.

Recommended Dilutions

WB	1:500-1:1000
FC	1:500 - 1:1000
IF/ICC	1:50-1:200

Immunogen Information

Gene ID

100133941

Swiss Prot

P25063

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 1-80 of human CD24 (NP_037362.1).

Synonyms

CD24A; CD24

Contact

	400-999-6126
	cn.market@abclonal.com.cn
	www.abclonal.com.cn

Product Information

Source

Rabbit

Isotype

IgG

Purification

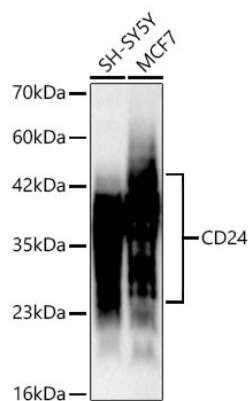
Affinity purification

Storage

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

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

Validation Data



Western blot analysis of various lysates using CD24 Rabbit mAb (A25383) at 1:1000 dilution incubated overnight at 4°C.

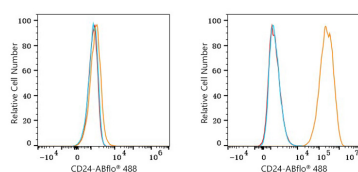
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L)(AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

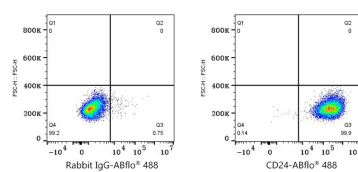
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020)

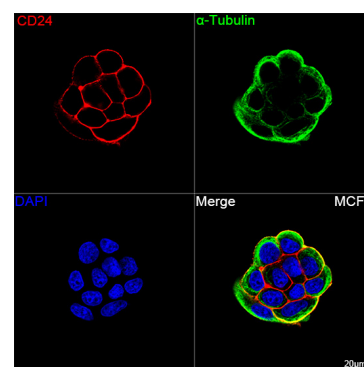
Exposure time: 10 s.



Flow cytometry: 1×10^6 293F cells (Low Expression, left) and MCF7 cells (right) were surface-stained with CD24 Rabbit mAb (A25383, 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: 1×10^6 MCF7 cells were surface-stained with ABflo® 488 Rabbit IgG isotype control (A22069, 5 µL/Test, left) or CD24 Rabbit mAb (A25383, 2 µg/mL, right).



Confocal imaging of MCF7 cells using CD24 Rabbit mAb (A25383, 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.