

CD74 Rabbit mAb

Catalog No.: A24027

Recombinant

1 Publications

Basic Information

Observed MW

35kDa

Calculated MW

18kDa/24kDa/26kDa/31kDa/34kDa

Category

Primary antibody

Applications

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

Cross-Reactivity

Human

CloneNo number

ARC56788

Background

The protein encoded by this gene associates with class II major histocompatibility complex (MHC) and is an important chaperone that regulates antigen presentation for immune response. It also serves as cell surface receptor for the cytokine macrophage migration inhibitory factor (MIF) which, when bound to the encoded protein, initiates survival pathways and cell proliferation. This protein also interacts with amyloid precursor protein (APP) and suppresses the production of amyloid beta (Abeta). Multiple alternatively spliced transcript variants encoding different isoforms have been identified.

Recommended Dilutions

WB 1:100000 - 1:400000**IHC-P** 1:10000 - 1:50000**IF/ICC** 1:200 - 1:800**FC** 1:500 - 1:1000

ELISA Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID

972

Swiss Prot

P04233-2


Immunogen

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

Synonyms

II; p33; CLIP; DHLAG; HLADG; Ia-GAMMA

Contact

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

Product Information

Source

Rabbit

Isotype

IgG

Purification

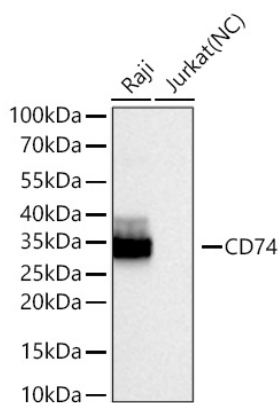
Affinity purification

Storage

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

Buffer: PBS 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



Western blot analysis of various lysates using CD74 Rabbit mAb (A24027) at 1:200000 dilution incubated overnight at 4°C.

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

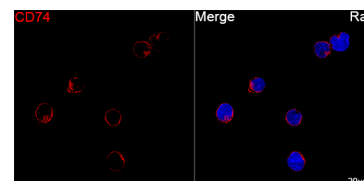
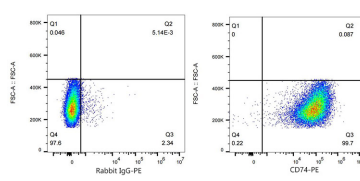
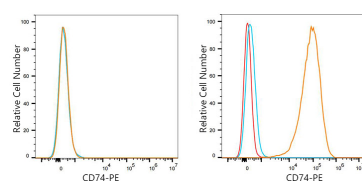
Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Negative control (NC): Jurkat

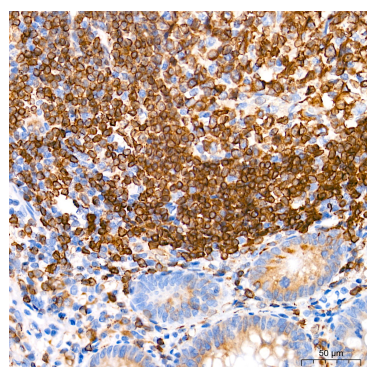
Exposure time: 10s.



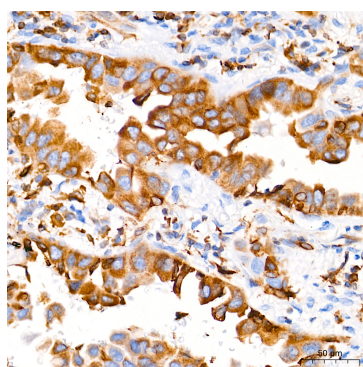
Flow cytometry: 1×10^6 Jurkat cells (negative control, left) and Raji cells (right) were surface-stained with CD74 Rabbit mAb (A24027, 2 µg/mL, orange line) or PE Rabbit IgG isotype control (5 µl/Test, blue line), followed by PE Donkey anti-rabbit Antibody staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry: 1×10^6 Raji cells were surface-stained with PE Rabbit IgG isotype control (5 µl/Test, left) or CD74 Rabbit mAb (A24027, 2 µg/mL, right).

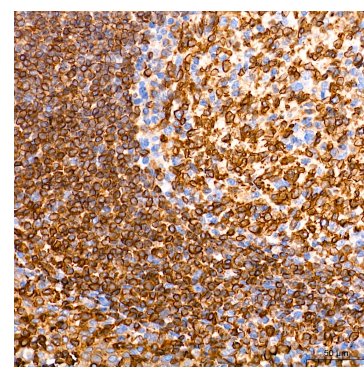
Confocal imaging of Raji cells using CD74 Rabbit mAb (A24027, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.



Immunohistochemistry analysis of paraffin-embedded Human appendix tissue using CD74 Rabbit mAb (A24027) at a dilution of 1:40000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Human lung cancer tissue using CD74 Rabbit mAb (A24027) at a dilution of 1:40000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Human tonsil tissue using CD74 Rabbit mAb (A24027) at a dilution of 1:40000 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.