

A23549

Leader in Biomolecular Solutions for Life Science



CD45 Rabbit mAb

Catalog No.: A23549

Recombinant

3 Publications

Basic Information

Observed MW

180-250kDa

Calculated MW

145kDa

Category

Monoclonal Antibody

Applications

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

Cross-Reactivity

Mouse

CloneNo number

ARC60962

Background

Enables several functions, including heparan sulfate proteoglycan binding activity; heparin binding activity; and protein tyrosine phosphatase activity. Involved in several processes, including lymphocyte differentiation; positive regulation of macromolecule metabolic process; and regulation of signal transduction. Acts upstream of or within several processes, including lymphocyte differentiation; positive regulation of lymphocyte activation; and regulation of protein phosphorylation. Located in external side of plasma membrane; focal adhesion; and membrane raft. Is expressed in several structures, including 3rd branchial arch; alimentary system; cardiovascular system; hemolymphoid system; and placenta. Used to study systemic lupus erythematosus. Human ortholog(s) of this gene implicated in hepatitis C; multiple sclerosis; severe combined immunodeficiency; and severe combined immunodeficiency, autosomal recessive, T cell-negative, B cell-positive, Nk cell-positive. Orthologous to human PTPRC (protein tyrosine phosphatase receptor type C).

Recommended Dilutions

WB	1:500 - 1:1000
IHC-P	1:200 - 1:800
IF	1:100 - 1:800
FC	1:100 - 1:500
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Contact

www.abclonal.com

Immunogen Information

Gene ID

19264

Swiss Prot

P06800

Immunogen

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

Synonyms

loc; B220; Cd45; L-CA; Ly-5; T200; CD45R; Lyt-4; CD45

Product Information

Source

Rabbit

Isotype

IgG

Purification

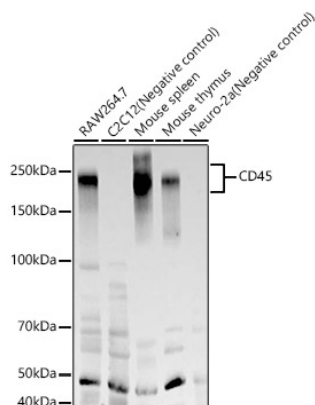
Affinity purification

Storage

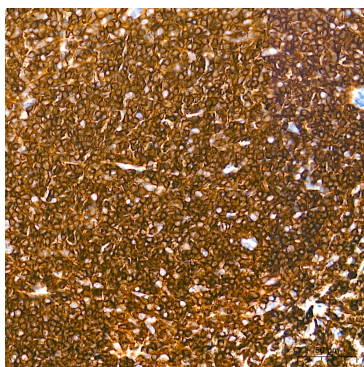
Store at -20°C. Avoid freeze / thaw cycles.

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

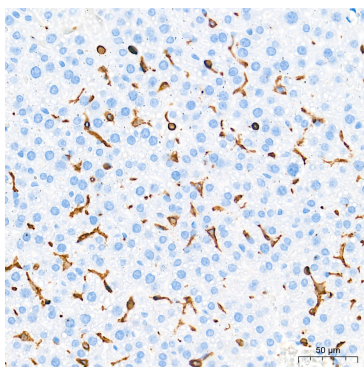
Validation Data



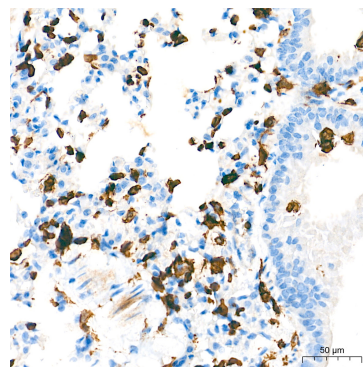
Western blot analysis of various lysates, using CD45 Rabbit mAb (A23549) at 1:1000 dilution.
 Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
 Lysates/proteins: 25µg per lane.
 Blocking buffer: 3% nonfat dry milk in TBST.
 Detection: ECL Enhanced Kit (RM00021).
 Exposure time: 120s.



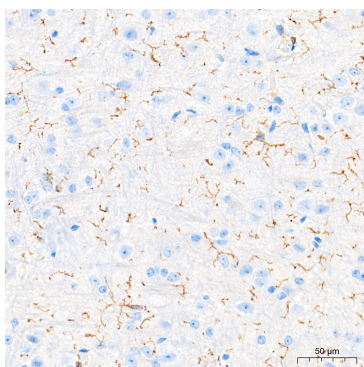
Immunohistochemistry analysis of paraffin-embedded Mouse spleen tissue using CD45 Rabbit mAb (A23549) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



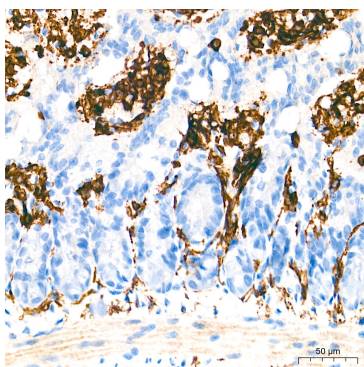
Immunohistochemistry analysis of paraffin-embedded Mouse liver tissue using CD45 Rabbit mAb (A23549) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



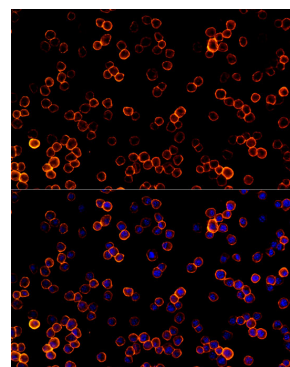
Immunohistochemistry analysis of paraffin-embedded Mouse lung tissue using CD45 Rabbit mAb (A23549) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Mouse brain tissue using CD45 Rabbit mAb (A23549) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

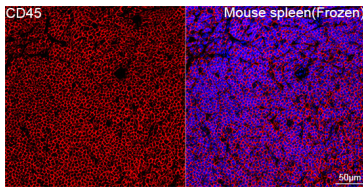


Immunohistochemistry analysis of paraffin-embedded Mouse colon tissue using CD45 Rabbit mAb (A23549) at a dilution of 1:200 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

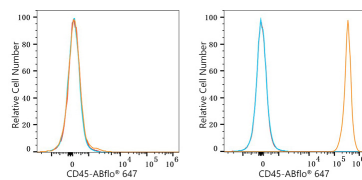


Immunofluorescence analysis of RAW264.7 cells using CD45 Rabbit mAb (A23549) at dilution of 1:100 (40x lens). Secondary antibody: Cy3-conjugated Goat anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.

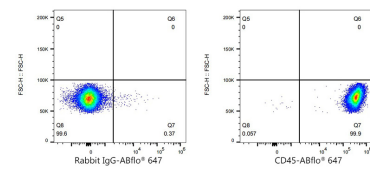
Validation Data



Confocal imaging of frozen sections
Mouse spleen tissue using CD45
Rabbit mAb (A23549, 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).
Microwave antigen retrieval
performed with 0.01M Citrate Buffer
(pH 6.0) prior to IF staining.
Objective: 40x.



Flow cytometry: 1×10^6 C2C12 cells
(negative control, left) and C57BL/6
mouse Splenocytes (right) were
surface-stained with CD45 Rabbit
mAb (A23549, 2 µg/mL, orange line)
or ABflo® 647 Rabbit IgG isotype
control (A22070, 5 µl/Test, 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: 1×10^6 C57BL/6
mouse Splenocytes were surface-
stained with ABflo® 647 Rabbit IgG
isotype control (A22070, 5 µl/Test, left)
or CD45 Rabbit mAb (A23549, 2
µg/mL, right).