[KO Validated] LAMP2 Rabbit mAb

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Catalog No.: A25646 KO Validated Recombinant

Basic Information

Observed MW

100-130kDa

Calculated MW

45kDa

Category

Primary antibody

Applications

WB,IHC-P,IF/ICC,FC (intra),ELISA

Cross-Reactivity

Human

CloneNo number

ARC54767

Background

The protein encoded by this gene is a member of a family of membrane glycoproteins. This glycoprotein provides selectins with carbohydrate ligands. It may play a role in tumor cell metastasis. It may also function in the protection, maintenance, and adhesion of the lysosome. Alternative splicing of this gene results in multiple transcript variants encoding distinct proteins.

Recommended Dilutions

WB 1:20000 - 1:120000

1:200 - 1:2000 **IHC**

IF/ICC 1:200 - 1:800

1:500 - 1:1000 FC (intra)

ELISA Recommended starting

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

Immunogen Information

Gene ID Swiss Prot 3920 P13473

Immunogen

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

Synonyms

DND; LAMPB; CD107b; LAMP-2; LGP-96; LGP110

Contact

<u>a</u>	400-999-6126
×	cn.market@abclonal.com.cn
<u></u>	www.abclonal.com.cn

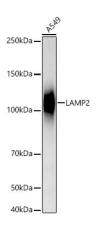
Product Information

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 lysates from A549 cells using [KO Validated] LAMP2 Rabbit mAb (A25646) at 1:20000 dilution.

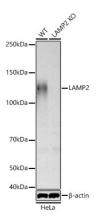
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25 µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time:10s.



Western blot analysis of lysates from wild type (WT) and LAMP2 knockout (KO) HeLa cells using [KO Validated] LAMP2 Rabbit mAb (A25646) at 1:20000 dilution.

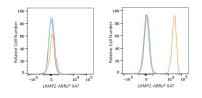
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

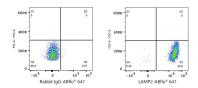
Lysates/proteins: 25 µg per lane.

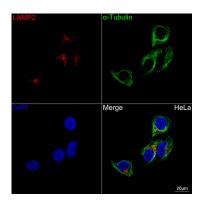
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



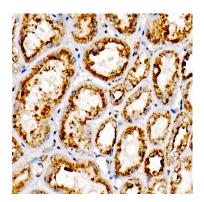




Flow cytometry: 1X10^6 knockout (KO) HeLa cells (negative control,left) and Hela cells (right) were intracellularly-stained with [KO Validated] LAMP2 Rabbit mAb (A25646,2 µg/mL,orange line) or Rabbit IgG isotype control (AC042,2 µg/mL,blue line), followed by Alexa Fluor® 647 conjugated goat antirabbit pAb staining. Non-fluorescently stained cells were used as blank control (red line).

Flow cytometry: 1X10^6 knockout (KO) HeLa cells (negative control,left) and Hela cells (right) were intracellularly-stained with Rabbit IgG isotype control (AC042,2 µg/mL,left) or [KO Validated] LAMP2 Rabbit mAb (A25646,2 µg/mL,right), followed by Alexa Fluor® 647 conjugated goat anti-Rabbit pAb staining.

Confocal imaging of HeLa cells using [KO Validated] LAMP2 Rabbit mAb (A25646, 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 $\alpha\text{-Tubulin}$ 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.



Immunohistochemistry analysis of paraffinembedded Human kidney tissue using [KO Validated] LAMP2 Rabbit mAb (A25646) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.