abcam

Product datasheet

Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)] ab282009



5 Images

Overview

Product name Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)]

Description Alexa Fluor® 555 Rabbit monoclonal [EPR4207(2)] to LAMP2A

Host species Rabbit

Conjugation Alexa Fluor® 555. Ex: 555nm. Em: 565nm

Tested applications Suitable for: IHC-P Species reactivity Reacts with: Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Human liver and prostate carcinoma tissue.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.2

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, 68.98% PBS

Purity Protein A purified

ClonalityMonoclonalClone numberEPR4207(2)

Isotype IgG

Applications

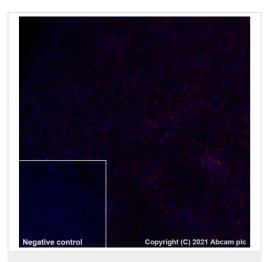
Images

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab282009 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Target		
Function	Implicated in tumor cell metastasis. May function in protection of the lysosomal membrane from autodigestion, maintenance of the acidic environment of the lysosome, adhesion when expressed on the cell surface (plasma membrane), and inter- and intracellular signal transduction. Protects cells from the toxic effects of methylating mutagens.	
Tissue specificity	Isoform LAMP-2A is highly expressed in placenta, lung and liver, less in kidney and pancreas, low in brain and skeletal muscle. Isoform LAMP-2B is highly expressed in skeletal muscle, less in brain, placenta, lung, kidney and pancreas, very low in liver.	
Involvement in disease	Danon disease	
Sequence similarities	Belongs to the LAMP family.	
Post-translational modifications	O- and N-glycosylated; some of the 16 N-linked glycans are polylactosaminoglycans.	
Cellular localization	Cell membrane. Endosome membrane. Lysosome membrane. This protein shuttles between lysosomes, endosomes, and the plasma membrane.	
Form	Alternative splicing produces 3 isoforms.	

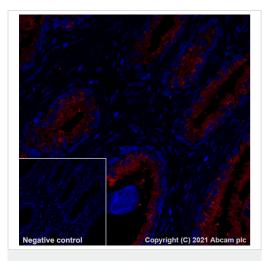


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)] (ab282009)

Immunohistochemical analysis of paraffin-embedded Human liver tissue labeling LAMP2A with ab282009 at 1/100 dilution.

Positive staining on human liver is observed. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab282009 at 1/100 dilution (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount®. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)] (ab282009)

Immunohistochemical analysis of paraffin-embedded Human prostate carcinoma tissue labeling LAMP2A with ab282009 at 1/100 dilution.

Positive staining on human prostate carcinoma is observed. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab282009 at 1/100 dilution (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). Heat mediated antigen retrieval using **ab93684** (Tris/EDTA buffer,

Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)] (ab282009)

Immunohistochemical analysis of paraffin-embedded Human cerebrum tissue labeling LAMP2A with ab282009 at 1/100 dilution.

Negative control: no staining on human cerebrum. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab282009 at 1/100 dilution (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. Nonspecific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

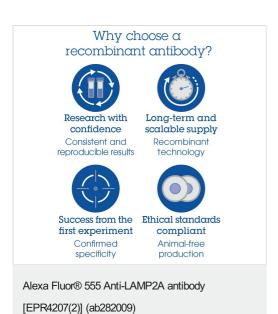
Heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 555 Anti-LAMP2A antibody [EPR4207(2)] (ab282009)

Immunohistochemical analysis of paraffin-embedded Human skeletal muscle tissue labeling LAMP2A with ab282009 at 1/100 dilution.

Negative control: no staining on human skeletal muscle. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab282009 at 1/100 dilution (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount®. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). Heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).



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