abcam

Product datasheet

Anti-ATE1 antibody [EPR13667(2)] - N-terminal ab199423



1 References 9 Images

Overview

Product name Anti-ATE1 antibody [EPR13667(2)] - N-terminal

Description Rabbit monoclonal [EPR13667(2)] to ATE1 - N-terminal

Host species Rabbit

Tested applications Suitable for: Flow Cyt (Intra), IHC-P, WB

Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control WB: HepG2, HeLa and 293 whole cell lysate. Human fetal liver tissue lysate. Mouse brain and

spleen tissue lysate. C6, Raw264.7, PC-12 and NIH/3T3 whole cell lysates. Mouse and Rat kidney tissue lysate. Rat spleen tissue lysate. IHC: Mouse liver tissue, Human hepatocellular carcinoma tissue, Rat kidney tissue. ICC/IF: HepG2 cells. Flow Cyt (intra): HeLa cells

General notesThis 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**.

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 long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

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Clonality Monoclonal

Clone number EPR13667(2)

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab199423 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
Flow Cyt (Intra)		1/220. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
IHC-P		1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		1/1000. Detects a band of approximately 59 kDa (predicted molecular weight: 59 kDa).

Target

Function Involved in the post-translational conjugation of arginine to the N-terminal aspartate or glutamate

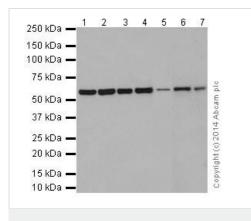
of a protein. This arginylation is required for degradation of the protein via the ubiquitin pathway.

Does not arginylate cysteine residues.

Sequence similarities Belongs to the R-transferase family.

Cellular localization Cytoplasm and Nucleus. Cytoplasm.

Images



Western blot - Anti-ATE1 antibody [EPR13667(2)] -

N-terminal (ab199423)

All lanes : Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423) at 1/1000 dilution

Lane 1: C6 (Rat glial tumor cells) whole cell lysate

Lane 2: Raw264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) whole cell lysate

Lane 3: PC-12 (Rat adrenal gland pheochromocytoma) whole cell

lysate

Lane 4: NIH/3T3 (Mouse embryo fibroblast cells) whole cell lysate

Lane 5 : Mouse kidney tissue lysate

Lane 6: Rat kidney tissue lysate

Lane 7: Rat spleen tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

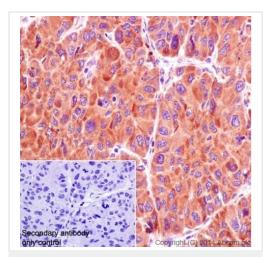
All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Developed using the ECL technique.

Predicted band size: 59 kDa **Observed band size:** 59 kDa

Exposure time: 3 minutes

Blocking and diluting buffer 5% NFDM/TBST



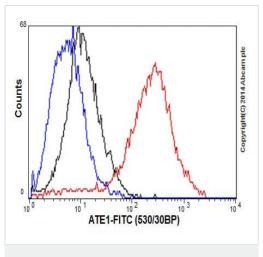
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ATE1 antibody

[EPR13667(2)] - N-terminal (ab199423)

Immunohistochemical analysis of paraffin-embedded human hepatocellular carcinoma tissue labeling ATE1 with ab199423 at 1/250 followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500. Cytoplasmic staining on human hepatocellular carcinoma tissue is observed (Subcellular location: Nucleus and cytoplasm [UniProt]). Counter-stained with hematoxylin.

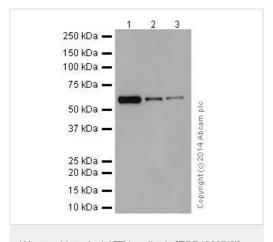
Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423)

Intracellular flow cytometric analysis of 2% paraformaldehyde-fixed HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling ATE1 with ab199423 at 1/220 dilution (red) compared with a rabbit monoclonal lgG isotype control (black) and a unlabelled control (cells without incubation with primary antibody and secondary antibody; blue). Goat anti rabbit lgG (FITC) at 1/150 dilution was used as the secondary antibody.



Western blot - Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423)

All lanes : Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423) at 1/10000 dilution

Lane 1 : HepG2 (Human liver hepatocellular carcinoma) whole cell lysate

Lane 2 : HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

Lane 3: 293 (Human epithelial cells from embryonic kidney) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

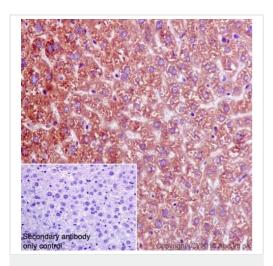
All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Developed using the ECL technique.

Predicted band size: 59 kDa **Observed band size:** 59 kDa

Exposure time: 3 minutes

Blocking and diluting buffer 5% NFDM/TBST

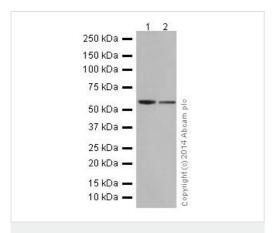


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ATE1 antibody

[EPR13667(2)] - N-terminal (ab199423)

Immunohistochemical analysis of paraffin-embedded Mouse liver tissue labeling ATE1 with ab199423 at 1/250 followed by Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500. Cytoplasm staining on Mouse liver tissue is observed (Subcellular location: Nucleus and Cytoplasm [UniProt]). Counter stained with Hematoxylin. Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423)

All lanes : Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423) at 1/1000 dilution

Lane 1 : Mouse brain tissue lysate

Lane 2 : Mouse spleen tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

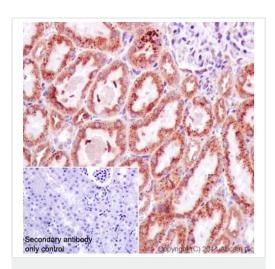
All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Developed using the ECL technique.

Predicted band size: 59 kDa Observed band size: 59 kDa

Exposure time: 1 minute

Blocking and diluting buffer 5% NFDM/TBST



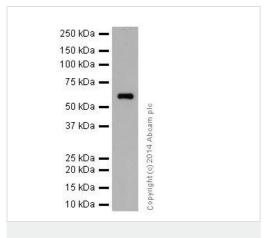
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ATE1 antibody

[EPR13667(2)] - N-terminal (ab199423)

Immunohistochemical analysis of paraffin-embedded Rat kidney tissue labeling ATE1 with ab199423 at 1/250 followed by Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500. Cytoplasm staining on Rat kidney tissue is observed (Subcellular location: Nucleus and Cytoplasm [UniProt]). Counter stained with Hematoxylin.

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423)

Anti-ATE1 antibody [EPR13667(2)] - N-terminal (ab199423) at 1/10000 dilution + Human fetal liver tissue lysate at 10 µg

Secondary

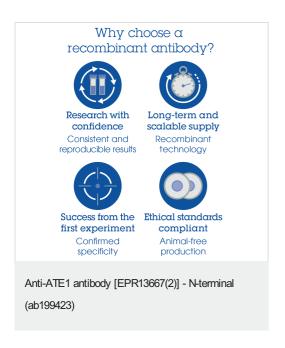
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Developed using the ECL technique.

Predicted band size: 59 kDa **Observed band size:** 59 kDa

Exposure time: 10 seconds

Blocking and diluting buffer 5% NFDM/TBST



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