abcam

Product datasheet

Anti-Clusterin alpha chain antibody [EPR17539-95] -BSA and Azide free ab230150

Recombinant

RabMAb

11 Images

Overview

Product name Anti-Clusterin alpha chain antibody [EPR17539-95] - BSA and Azide free

Description Rabbit monoclonal [EPR17539-95] to Clusterin alpha chain - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: IHC-Fr, IP, IHC-P, WB

Unsuitable for: ICC/IF

Species reactivity Reacts with: Mouse. Rat

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Rat adrenal gland tissue.

General notes ab230150 is the carrier-free version of ab184100.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR17539-95

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab230150 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-Fr		Use at an assay dependent concentration. Perform heat-mediated antigen retrieval by using Tris-EDTA buffer (pH9.0) (ab94681).
IP		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Detects a band of approximately 38 kDa (predicted molecular weight: 52 kDa).

Application notes

Is unsuitable for ICC/IF.

Target

Function

Isoform 1 functions as extracellular chaperone that prevents aggregation of nonnative proteins. Prevents stress-induced aggregation of blood plasma proteins. Inhibits formation of amyloid fibrils by APP, APOC2, B2M, CALCA, CSN3, SNCA and aggregation-prone LYZ variants (in vitro). Does not require ATP. Maintains partially unfolded proteins in a state appropriate for subsequent refolding by other chaperones, such as HSPA8/HSC70. Does not refold proteins by itself. Binding to cell surface receptors triggers internalization of the chaperone-client complex and subsequent lysosomal or proteasomal degradation. Secreted isoform 1 protects cells against apoptosis and against cytolysis by complement. Intracellular isoforms interact with ubiquitin and SCF (SKP1-CUL1-F-box protein) E3 ubiquitin-protein ligase complexes and promote the ubiquitination and subsequent proteasomal degradation of target proteins. Promotes proteasomal degradation of COMMD1 and IKBKB. Modulates NF-kappa-B transcriptional activity. Nuclear isoforms promote apoptosis. Mitochondrial isoforms suppress BAX-dependent release of cytochrome c into the

cytoplasm and inhibit apoptosis. Plays a role in the regulation of cell proliferation.

Tissue specificity Detected in blood plasma, cerebrospinal fluid, milk, seminal plasma and colon mucosa. Detected

> in the germinal center of colon lymphoid nodules and in colon parasympathetic ganglia of the Auerbach plexus (at protein level). Ubiquitous. Detected in brain, testis, ovary, liver and pancreas,

and at lower levels in kidney, heart, spleen and lung.

Sequence similarities Belongs to the clusterin family.

Post-translational Isoform 1 is proteolytically cleaved on its way through the secretory system, probably within the modifications

Golgi lumen.

Polyubiquitinated, leading to proteasomal degradation.

Heavily N-glycosylated. About 30% of the protein mass is comprised of complex N-linked

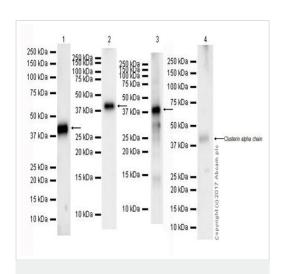
carbohydrate.

Cellular localization Nucleus. Cytoplasm. Mitochondrion membrane. Cytoplasm > cytosol. Microsome. Endoplasmic

> reticulum. Cytoplasmic vesicle > secretory vesicle > chromaffin granule. Isoforms lacking the Nterminal signal sequence have been shown to be cytoplasmic and/or nuclear. Secreted isoforms can retrotranslocate from the secretory compartments to the cytosol upon cellular stress. Detected

in perinuclear foci that may be aggresomes containing misfolded, ubiquitinated proteins. Detected at the mitochondrion membrane upon induction of apoptosis and Secreted. Can retrotranslocate from the secretory compartments to the cytosol upon cellular stress.

Images



Western blot - Anti-Clusterin alpha chain antibody [EPR17539-95] - BSA and Azide free (ab230150)

All lanes: Anti-Clusterin alpha chain antibody [EPR17539-95] (ab184100) at 1/1000 dilution

Lane 1: Mouse serum lysate at 20 µg

Lane 2: Rat serum lysate at 10 µg

Lane 3: Mouse lung lysate at 10 µg

Lane 4: Mouse adrenal gland lysate at 20 µg

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at

1/100000 dilution

Developed using the ECL technique.

Predicted band size: 52 kDa

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).

Blocking/Dilution buffer and concentration: 5% NFDM/TBST.

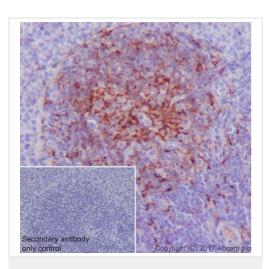
Exposure times:

Lanes 1,3,4: 24 seconds;

Lane 2: 3 seconds.

The molecular mass observed is consistent with what has been described in the literature (PMID: 8373947).

The blot was developed on a BIO-RAD[®] ChemiDoc™ MP instrument



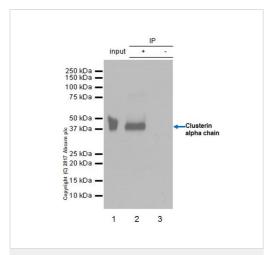
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] - BSA and Azide free (ab230150)

Immunohistochemical analysis of paraffin-embedded rat spleen tissue labeling Clusterin alpha chain with **ab184100** at 1/4000 dilution, followed by a ready to use Goat Anti-Rabbit lgG H&L (HRP). Positive staining on rat spleen (PMID: 24865838) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



Immunoprecipitation - Anti-Clusterin alpha chain antibody [EPR17539-95] - BSA and Azide free (ab230150)

Clusterin alpha chain was immunoprecipitated from 0.35 mg mouse serum lysate with **ab184100** at 1/30 dilution. Western blot was performed from the immunoprecipitate using **ab184100** at 1/1,000 dilution. VeriBlot for IP Detection Reagent (HRP) (**ab131366**), was used for detection at 1/10,000 dilution.

Lane 1: Mouse serum lysate 10 µl (Input).

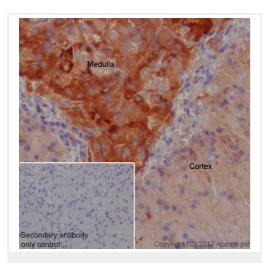
Lane 2: ab184100 IP in mouse serum lysate (+).

Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab184100</u> in mouse serum lysate (-).

Blocking/Dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 1 second.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab184100</u>).



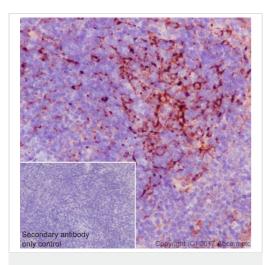
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] - BSA and Azide free (ab230150)

Immunohistochemical analysis of paraffin-embedded mouse adrenal gland tissue labeling Clusterin alpha chain with <u>ab184100</u> at 1/4000 dilution, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Stronger cytoplasmic staining in the medulla of mouse adrenal gland than the cortex (PMID: 16436671) is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit IgG H&L (HRP).

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



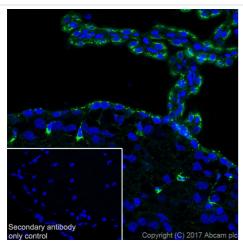
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Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab184100</u>).



Immunohistochemistry (Frozen sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] -BSA and Azide free (ab230150)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen rat testis tissue labeling Clusterin alpha chain with ab184100 at 1/250 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000 dilution. Positive staining in Sertoli cells, immature spermatozoa

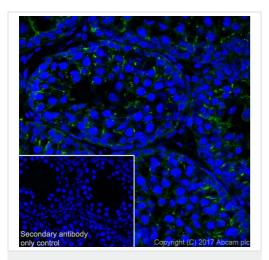
The nuclear counterstain is DAPI (blue).

and Leydig cells (PMID: 22552734) is observed.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 AlexaFluor®488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat mediated antigen retrieval using ab94681 (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



Immunohistochemistry (Frozen sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] -BSA and Azide free (ab230150)

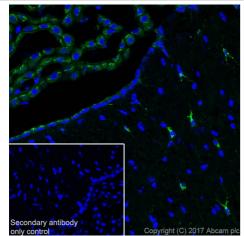
Immunohistochemical analysis of frozen rat brain (lateral ventricle) tissue labeling Clusterin alpha chain with ab184100 at 1/250 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000 dilution. Positive cytoplasmic staining in endothelial cells of choroid plexus and astrocytes in rat brain (PMID:18620027; 21385939) is observed.

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is **ab150077** AlexaFluor[®]488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat mediated antigen retrieval using ab94681 (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



Immunohistochemistry (Frozen sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] -BSA and Azide free (ab230150)

Triton X-100 permeabilized frozen mouse brain (lateral ventricle) tissue labeling Clusterin alpha chain with ab184100 at 1/250 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000 dilution. Positive cytoplasmic staining in endothelial cells of choroid plexus and astrocytes in mouse brain (PMID:18620027; 21385939) is observed.

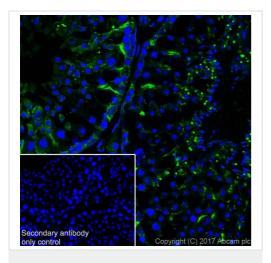
The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is **ab150077** AlexaFluor[®]488 Goat anti-Rabbit used at a 1/1000 dilution.

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2%

Perform heat mediated antigen retrieval using ab94681 (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



Immunohistochemistry (Frozen sections) - Anti-Clusterin alpha chain antibody [EPR17539-95] -BSA and Azide free (ab230150)

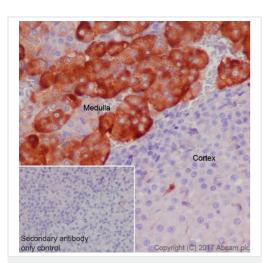
Immunohistochemical analysis of 4% paraformaldehude-fixed, 0.2% Triton X-100 permeabilized frozen mouse testis tissue labeling Clusterin alpha chain with ab184100 at 1/250 dilution (green), followed by ab150077 AlexaFluor®488 Goat anti-Rabbit secondary at a 1/1000 dilution. Positive staining in Sertoli cells, immature spermatozoa and Leydig cells is observed; also observe particles deposited on sperm membrane (PMID: 22552734).

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 AlexaFluor®488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat mediated antigen retrieval using ab94681 (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



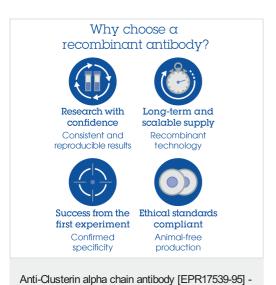
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Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab184100).



BSA and Azide free (ab230150)

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