abcam

ab275752

Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free

Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752) is a rabbit recombinant monoclonal antibody provided in a PBS only buffer for easy conjugation. Suitable for **Western Blot, Flow Cytometry (Intra), IHC-P, IHC-Fr, ICC/IF** in **Human, Mouse, Rat**.

- KO validated for confirmed specificity
- BSA, sodium azide, and glycerol-free for easy conjugation
- Biophysical QC for unrivalled batch-batch consistency

Recombinant | RabMAb | KO Validated

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	Constituents: PBS
Form	Liquid
Clonality	Monoclonal
Immunogen	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	EPR23630-49
Purification technique	Affinity purification Protein A
Concentration	0.951 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

ICC/IF

Tested

Species	Mouse, Human
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

IP

Not recommended

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

WB

Tested

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

IHC-P

Tested

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

IHC-Fr

Tested

Species	Mouse
Dilution info	-
Notes	Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

Expected

Species	Human, Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Flow Cyt (Intra)

Tested

Species	Mouse, Human
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Target data

See full target information ATXN2

Function	Involved in EGFR trafficking, acting as negative regulator of endocytic EGFR internalization at the plasma membrane.

Storage

Shipped at conditions	Blue Ice

Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C

Notes

What is this antibody validated in?
Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752) is a rabbit recombinant monoclonal antibody and is validated for use in Western Blot (WB), Flow Cytometry (Intra), Flow Cytometry (Flow Cyt), Immunohistochemistry (IHC-P), Immunohistochemistry (IHC-Fr), Immunocytochemistry/immunofluorescence (ICC/IF) in Human, Mouse, Rat samples.

What is the molecular weight of SCA2?
Anti-SCA2 [EPR23630-49] - BSA and Azide free (ab275752) specifically detects a band for SCA2 (UniProt: Q99700) at a molecular weight of 140kDa.

Specificity confirmed
The specificity of Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752) has been confirmed by Western blot testing in ATXN2 Knockout HAP1 cells.

Other related products
We have a range of other formats of antibody clone [EPR23630-49] also available for your convenience: ab254362, Carrier free - ab275752, Alexa Fluor® 647 - ab286816, Alexa Fluor® 488 - ab300055

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

What are the advantages of a recombinant monoclonal antibody?
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 batch production

For more information, read more on recombinant antibodies.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based 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.

Compatibility
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.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	SCA2 also known as Spinocerebellar Ataxia Type 2 refers to a protein primarily affected by mutations in the ATXN2 gene. It weighs around 150 kDa. SCA2 expresses predominantly in the brain with significant presence in Purkinje cells of the cerebellum as well as regions like the thalamus and hippocampus. It functions as an essential component within neuron-specific structures and may associate with RNA processing events.
Biological function summary	SCA2 shows significant interactions within the neuronal RNA processing machinery. This protein is part of a large complex involved in regulating RNA stability and translation. Its role contributes significantly to maintaining neuron function and viability affecting synaptic signaling and plasticity which are critical for cognitive processes and motor control. SCA2 influences neural communication through modulation and degradation of messenger RNAs.
Pathways	SCA2 plays a role in the mRNA metabolic process related cellular pathways. It interacts in pathways like RNA transport and degradation linking it functionally to proteins such as TDP-43 and FUS which also have prominent roles in RNA processing. These associations support communication across neurons and ensure accurate protein synthesis necessary for neuronal health and function.
Associated diseases and disorders	Malfunction or mutation in SCA2 relates directly to spinocerebellar ataxia type 2 a degenerative disorder affecting movement. The imbalance in related pathways often leads to pathological changes contributing to dysfunctions in Ataxin-2 often seen alongside interactions with proteins like Huntingtin which elucidate its involvement in neurological diseases. Such connections imply a substantial impact on neuronal survival and disease progression.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

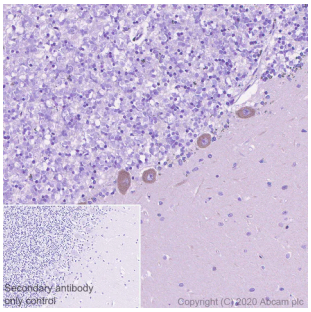
We do not recommend this combination. It is not covered by our product promise.

We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

Full details and terms and conditions can be found here:
Terms & Conditions.

11 product images



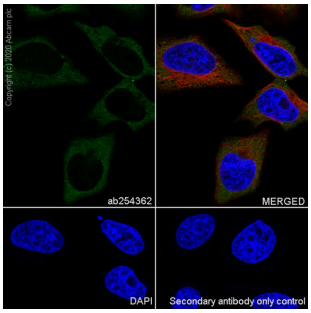
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded human cerebellum tissue labeling SCA2 with ab254362 at 1/1000 dilution, followed by a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining on Purkinje cells in human cerebellum (PMID: 9989626). The section was incubated with ab254362 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

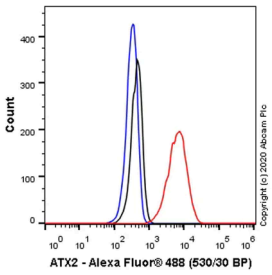


Immunocytochemistry/ Immunofluorescence - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (human epithelial cell line from cervix adenocarcinoma) cells labeling SCA2 with ab254362 at 1/50 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing cytoplasmic staining on HeLa cell line. The nuclear counterstain is DAPI (blue).
Tubulin is detected with ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) at 1/200 dilution (red).

The negative control is secondary antibody only.

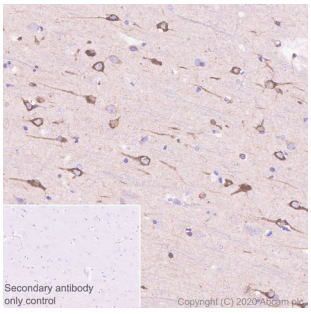


Flow Cytometry (Intracellular) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Flow Cytometry (Intracellular) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol permeabilized HeLa (human epithelial cell line from cervix adenocarcinoma) cells labeling SCA2 with ab254362 at 1/500 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (blue).

Goat Anti-Rabbit IgG H&L (Alexa Fluor®488) at 1/2000 dilution was used as the secondary antibody.

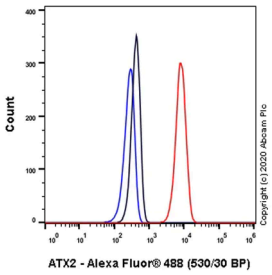


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded human cerebrum tissue labeling SCA2 with ab254362 at 1/1000 dilution, followed by a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining on neurons of human cerebrum (PMID: 9989626). The section was incubated with ab254362 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.

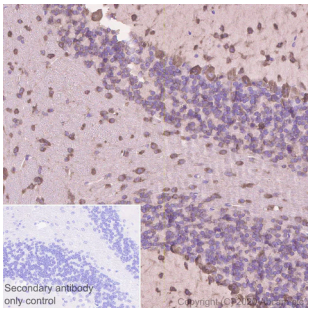


Flow Cytometry (Intracellular) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Flow Cytometry (Intracellular) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Intracellular flow cytometric analysis of 4% paraformaldehyde-fixed, 90% methanol permeabilized NIH/3T3 (mouse embryo fibroblast cell line) cells labeling SCA2 with ab254362 at 1/500 dilution (red) compared with a Rabbit IgG, monoclonal [EPR25A] - Isotype Control (ab172730) (black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (blue).

Goat Anti-Rabbit IgG H&L (Alexa Fluor®488) at 1/2000 dilution was used as the secondary antibody.



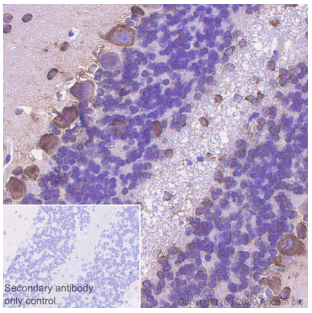
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded mouse cerebellum tissue labeling SCA2 with ab254362 at 1/1000 dilution, followed by a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining in mouse cerebellum (PMID: 26868665). The section was incubated with ab254362 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.



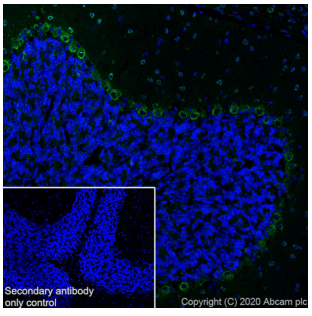
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded rat cerebellum tissue labeling SCA2 with ab254362 at 1/1000 dilution, followed by a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101). Cytoplasmic staining in rat cerebellum (PMID: 26868665). The section was incubated with ab254362 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use secondary from Rabbit specific IHC polymer detection kit HRP/DAB (ab209101).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution2) for 20 mins.



Immunohistochemistry (Frozen sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

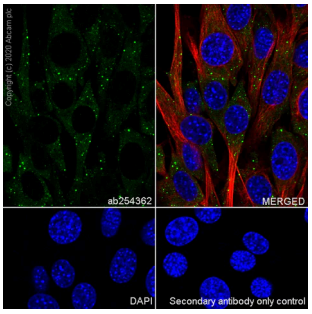
SCA2 Immunohistochemistry (Frozen sections) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse cerebellum tissue labeling SCA2 with ab254362 at 1/50 dilution, followed by ab150077 Goat Anti-

Rabbit IgG H&L (Alexa Fluor® 488) at a 1/1000 dilution. Positive staining on mouse cerebellum is observed. Nuclear counterstain is DAPI.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) at a 1/1000 dilution.

Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).

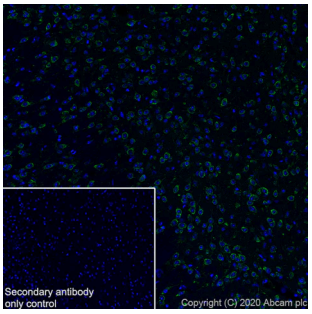


Immunocytochemistry/ Immunofluorescence - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH/3T3 (mouse embryo fibroblast cell line) cells labeling SCA2 with ab254362 at 1/50 dilution, followed by Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) (ab150077) secondary antibody at 1/1000 dilution (green). Confocal image showing cytoplasmic staining on NIH/3T3 cell line. The nuclear counterstain is DAPI (blue).
Tubulin is detected with ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) at 1/200 dilution (red).

The negative control is secondary antibody only.



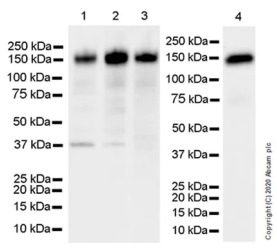
Immunohistochemistry (Frozen sections) - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

SCA2 Immunohistochemistry (Frozen sections) staining using rabbit Anti-SCA2 antibody

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of 4% PFA-fixed, 0.2% Triton X-100 permeabilized frozen mouse cerebrum tissue labeling SCA2 with ab254362 at 1/50 dilution, followed by ab150077 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) at a 1/1000 dilution. Positive staining on mouse cerebrum is observed. Nuclear counterstain is DAPI.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is ab150077 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) at a 1/1000 dilution.

Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).



Western blot - Anti-SCA2 antibody [EPR23630-49] - BSA and Azide free (ab275752)

This data was developed using ab254362, the same antibody clone in a different buffer formulation.
The molecular weight observed is consistent with what has been described in the literature (PMID: 9989626).
Lysates should be made freshly and used in WB immediately to minimize protein degradation.

Blocking/Dilution buffer: 5% NFDM/TBST.

Exposure times: Lanes 1-3: 10 seconds; Lane 4: 15 seconds.
All lanes:
Western blot - Anti-SCA2 antibody [EPR23630-49] (ab254362) at 1/1000 dilution

Lane 1:
HeLa (human cervix adenocarcinoma epithelial cell) whole cell lysate at 20 µg

Lane 2:
HEK-293T (human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate at 20 µg

Lane 3:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate at 20 µg

Lane 4:
PC-12 (rat adrenal gland pheochromocytoma) whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 140 kDa

Observed band size: 145 kDa

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.