# abcam

# Product datasheet

# Anti-Neurogranin antibody [EPR21152] - BSA and Azide free ab230154



# 8 Images

#### Overview

**Product name** Anti-Neurogranin antibody [EPR21152] - BSA and Azide free

**Description** Rabbit monoclonal [EPR21152] to Neurogranin - BSA and Azide free

**Host species** Rabbit

**Tested applications** Suitable for: IHC-P, WB, IHC-Fr, IP, ICC/IF

**Species reactivity** Reacts with: Mouse, Rat, Human

Predicted to work with: Common marmoset

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

Positive control IHC-P: Rat cerebrum tissue.

General notes ab230154 is the carrier-free version of ab217672.

> 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<sup>®</sup> 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 numberEPR21152

**Isotype** IgG

## **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab230154 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		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 15 kDa (predicted molecular weight: 8 kDa).
IHC-Fr		Use at an assay dependent concentration.  Perform heat mediated antigen retrieval by using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).
IP		Use at an assay dependent concentration.
ICC/IF		Use at an assay dependent concentration.

# **Target**

**Function** Acts as a "third messenger" substrate of protein kinase C-mediated molecular cascades during

synaptic development and remodeling. Binds to calmodulin in the absence of calcium.

**Tissue specificity** In the cerebral cortex, found in the cell bodies of neurons in layers II-VI, and in apical and basal

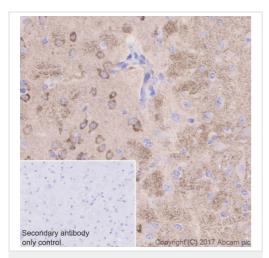
dendrites of pyramidal neurons. Is not found in the dendrites in patients with Alzheimer disease.

**Sequence similarities** Belongs to the neurogranin family.

Contains 1 collagen-like domain.

Contains 1 IQ domain.

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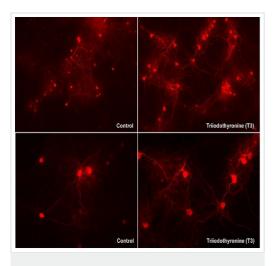
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Neurogranin antibody
[EPR21152] - BSA and Azide free (ab230154)

Immunohistochemical analysis of paraffin-embedded mouse cerebrum tissue labeling Neurogranin with <u>ab217672</u> at 1/5000 dilution, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Mainly cytoplasmic staining on mouse cerebrum (PMID: 26076492) 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 (ab217672).



Immunocytochemistry/ Immunofluorescence - Anti-Neurogranin antibody [EPR21152] - BSA and Azide free (ab230154)

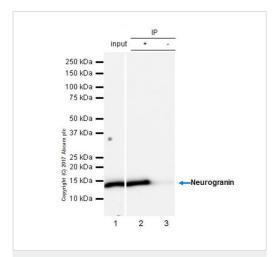
Image courtesy of Ms. Babben Tinner (QBM Cell Science).

Immunocytochemistry/ Immunofluorescence analysis of primary rat cortical neurons labeling Neurogranin with <u>ab217672</u> at 1/1500. The cells were fixed with 4% paraformaldehyde containing 0.2% picric acid in 0.1M phosphate buffer, pH 6.9 for 20 minutes. Permeabilization was with 0.3% Triton-X 100 in PBS (PBSTx). <u>ab150068</u> at 1/200 was used as the secondary antibody

The rat cortical neurons were cultured for 29 days *in vitro*. They were either left untreated (Control) or treated beginning on the 10<sup>th</sup> day with 60ng/mL triiodothyronine (T3), to enhance neurogranin expression.

The cells were visualized with an inverted microscope at 10X magnification (upper panels) or 20X magnification (lower panels).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab217672).



Immunoprecipitation - Anti-Neurogranin antibody [EPR21152] - BSA and Azide free (ab230154)

Secondary antibody only control Copyright (C) 2017 Abcam plc

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

[EPR21152] - BSA and Azide free (ab230154)

Neurogranin was immunoprecipitated from 0.35 mg mouse cerebral cortex lysate with <u>ab217672</u> at 1/30 dilution. Western blot was performed from the immunoprecipitate using <u>ab217672</u> at 1/1000 dilution. VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/5,000 dilution.

Lane 1: Mouse cerebral cortex lysate 10 µg (Input).

Lane 2: ab217672 IP in mouse cerebral cortex lysate (+).

**Lane 3:** Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab217672</u> in mouse cerebral cortex lysate (-).

Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 15 seconds.

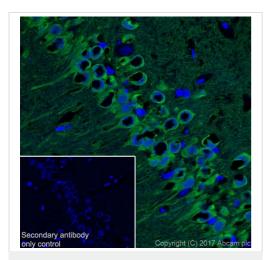
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab217672).

Immunohistochemical analysis of paraffin-embedded human cerebrum tissue labeling Neurogranin with <u>ab217672</u> at 1/5000 dilution, followed by a ready to use Goat Anti-Rabbit lgG H&L (HRP). Mainly cytoplasmic but also weak nuclear staining on human cerebrum (PMID: 26076492; PMID: 21516261) 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).

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Immunohistochemistry (Frozen sections) - Anti-Neurogranin antibody [EPR21152] - BSA and Azide free (ab230154)

Immunohistochemistry (Frozen sections) - Anti-Neurogranin antibody [EPR21152] - BSA and Azide free (ab230154)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen rat hippocampus CA1 tissue labeling Neurogranin with <a href="mailto:ab217672">ab217672</a> at 1/3000 dilution (green), followed by <a href="mailto:ab2150077">ab150077</a> AlexaFluor<sup>®</sup> 488 Goat anti-Rabbit secondary at a 1/1000 dilution. Positive staining in the stratum pyramidal neurons of hippocampus CA1 on rat brain (PMID: 15389631; 21516261) is observed. Counter stained with DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <a href="mailto:ab150077"><u>ab150077</u></a> AlexaFluor<sup>®</sup>488 Goat anti-Rabbit used at a 1/1000 dilution.

Perform heat-mediated antigen retrieval by using sodium citrate buffer (10 mM citrate pH 6.0 + 0.05% Tween-20)

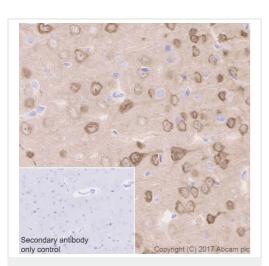
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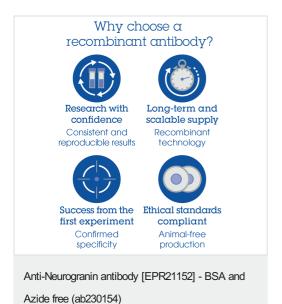
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