

# Anti-Natriuretic peptides A antibody [EPR20247] - BSA and Azide free ab225873

Recombinant RabMAb

7 Images

### Overview

<b>Product name</b>	Anti-Natriuretic peptides A antibody [EPR20247] - BSA and Azide free
<b>Description</b>	Rabbit monoclonal [EPR20247] to Natriuretic peptides A - BSA and Azide free
<b>Host species</b>	Rabbit
<b>Tested applications</b>	<b>Suitable for:</b> WB, IHC-P, IHC-Fr, IP
<b>Species reactivity</b>	<b>Reacts with:</b> Rat, Human
<b>Immunogen</b>	Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.
<b>Positive control</b>	WB: Human fetal heart and rat heart tissue lysates. IHC-P: Human heart, rat auricle and rat ventricle tissues. IHC-Fr: Rat heart tissue. IP: Rat heart lysate. mlHC: Human cardiac muscle tissue.
<b>General notes</b>	<p>ab225873 is the carrier-free version of <a href="#">ab209232</a>.</p> <p>Our <b>carrier-free</b> 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.</p> <p>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.</p> <p>Use our <b>conjugation kits</b> for antibody conjugates that are ready-to-use in as little as 20 minutes with &lt;1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.</p> <p>Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMAb<sup>®</sup> patents</a>.</p>

### Properties

**Form** Liquid

<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C. Do Not Freeze.
<b>Storage buffer</b>	pH: 7.2 Constituent: PBS
<b>Carrier free</b>	Yes
<b>Purity</b>	Protein A purified
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EPR20247
<b>Isotype</b>	IgG

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab225873 in the following tested applications.

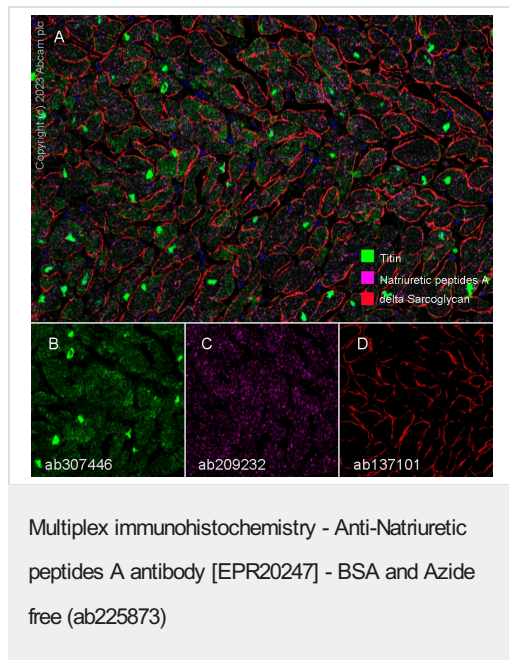
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
<b>WB</b>		Use at an assay dependent concentration. Predicted molecular weight: 16 kDa.
<b>IHC-P</b>		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
<b>IHC-Fr</b>		Use at an assay dependent concentration. Antigen retrieval: Heated citrate solution (10mM citrate pH 6.0 + 0.05% Tween-20).
<b>IP</b>		Use at an assay dependent concentration.

## Target

<b>Function</b>	Hormone playing a key role in cardiovascular homeostasis through regulation of natriuresis, diuresis, and vasodilation. Also plays a role in female pregnancy by promoting trophoblast invasion and spiral artery remodeling in uterus. Specifically binds and stimulates the cGMP production of the NPR1 receptor. Binds the clearance receptor NPR3.
<b>Involvement in disease</b>	Atrial standstill 2 Atrial fibrillation, familial, 6
<b>Sequence similarities</b>	Belongs to the natriuretic peptide family.
<b>Post-translational modifications</b>	Cleaved by CORIN upon secretion to produce the functional hormone. Atrial natriuretic factor: Cleaved by MME. The cleavage initiates degradation of the factor and thereby regulate its activity.
<b>Cellular localization</b>	Secreted.

## Images



Fluorescence multiplex immunohistochemical analysis of the human cardiac muscle (Formalin/PFA-fixed paraffin-embedded sections).

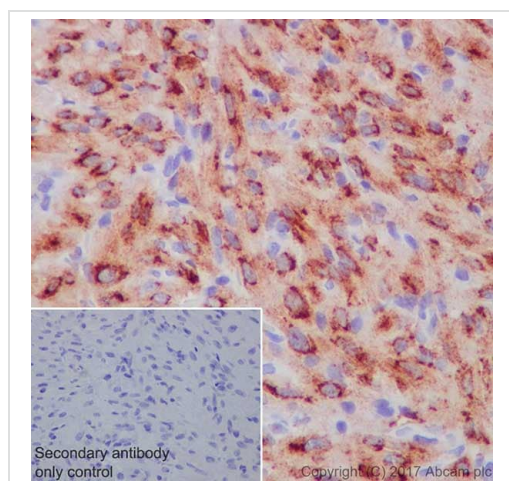
Panel A: merged staining of anti-delta Sarcoglycan ([ab137101](#), red; Opal™690), anti-Titin ([ab307446](#), green; Opal™520) and anti-Natriuretic peptides A ([ab209232](#), magenta; Opal™570) on human cardiac muscle. Panel B: anti-Titin displayed nucleus and cytoplasm expression. Panel C: anti-Natriuretic peptides A displayed granular cytoplasmic expression. Panel D: anti-delta Sarcoglycan displayed membrane expression. Opal Polymer HRP Ms + Rb was used as a secondary antibody.

The section was incubated in three rounds of staining: in the order of [ab137101](#) at 1/1000 (1.043 µg/ml) dilution, [ab307446](#) at 1/500 (0.95 µg/ml) dilution, and [ab209232](#) at 1/3000 (0.241 µg/ml) dilution for 30 mins at room temperature. Each round was followed by a separate fluorescent tyramide signal amplification system. DAPI (blue) was used as a nuclear counter stain.

The immunostaining was performed on a Leica Biosystems BOND® RX instrument with an Opal™ 4-color kit. Image acquisition was performed with Leica SP8 confocal microscope.

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab209232](#)).



Immunohistochemical analysis of paraffin-embedded rat auricle tissue labeling Natriuretic peptides A with [ab209232](#) at 1/8000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use.

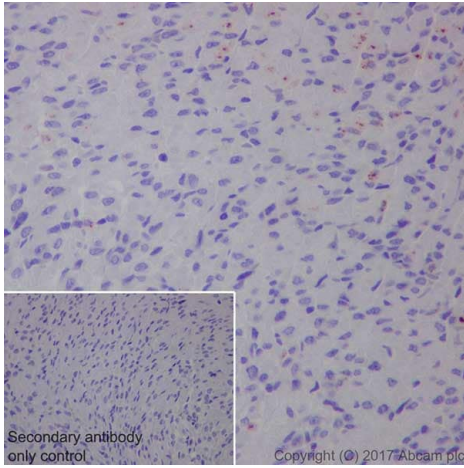
Granularly cytoplasmic and perinuclear staining on rat auricle (PMID: 2942710, PMID: 1824903).

Counter stained with Hematoxylin.

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide ([ab209232](#)).

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



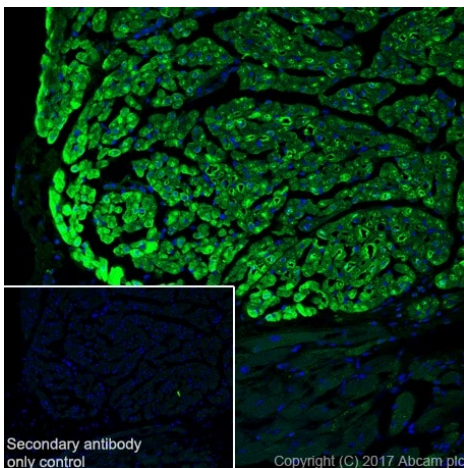
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Natriuretic peptides A antibody [EPR20247] - BSA and Azide free (ab225873)

Immunohistochemical analysis of paraffin-embedded rat ventricle tissue labeling Natriuretic peptides A with **ab209232** at 1/8000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Weak staining on rat ventricle (PMID: 2942710, PMID: 1824903). Counter stained with Hematoxylin.

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab209232**).

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



Immunohistochemistry (Frozen sections) - Anti-Natriuretic peptides A antibody [EPR20247] - BSA and Azide free (ab225873)

Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen rat heart tissue labeling Natriuretic peptides A with **ab209232** at 1/100 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (**ab150077**) secondary antibody at 1/1000 dilution (green).

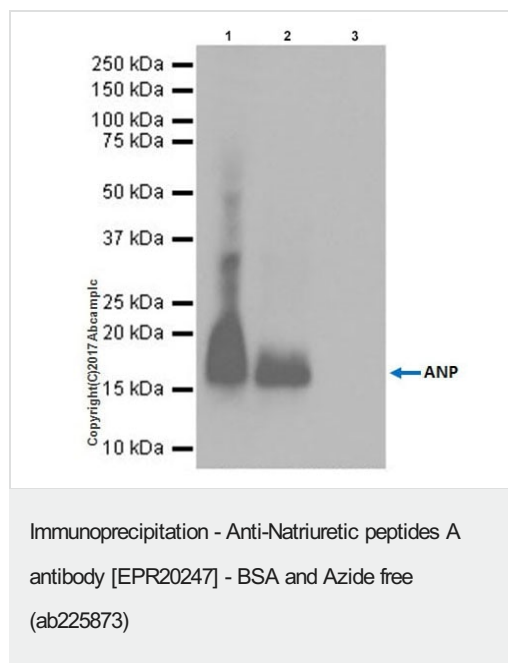
Cytoplasmic staining on the auricula of rat heart (PMID: 2942710, PMID: 1824903).

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat anti-rabbit IgG (Alexa Fluor® 488) (**ab150077**) at 1/1000 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab209232**).

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



Natriuretic peptides A was immunoprecipitated from 0.35 mg of rat heart lysate with **ab209232** at 1/30 dilution.

Western blot was performed from the immunoprecipitate using **ab209232** at 1/500 dilution.

VeriBlot for IP Detection Reagent (HRP) (**ab131366**), was used for detection at 1/1000 dilution.

Lane 1: Rat heart lysate 10 µg (Input).

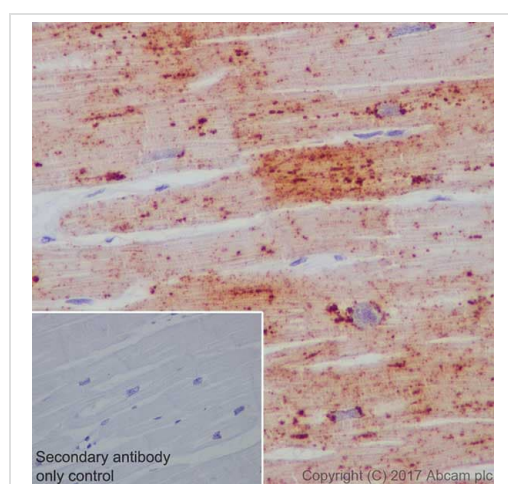
Lane 2: **ab209232** IP in rat heart lysate.

Lane 3: Rabbit monoclonal IgG (**ab172730**) instead of **ab209232** in rat heart lysate.

Blocking and 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 (**ab209232**).



Immunohistochemical analysis of paraffin-embedded human heart tissue labeling Natriuretic peptides A with **ab209232** at 1/4000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use.

Granularly cytoplasmic and perinuclear staining on human heart (PMID: 2942710, PMID: 1824903).

Counter stained with Hematoxylin.

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

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab209232**).

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



### Why choose a recombinant antibody?



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Consistent and reproducible results



**Long-term and scalable supply**  
Recombinant technology



**Success from the first experiment**  
Confirmed specificity



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Animal-free production

Anti-Natriuretic peptides A antibody [EPR20247] -  
BSA and Azide free (ab225873)

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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