# abcam

## Product datasheet

# Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free ab238938



#### 8 Images

#### Overview

**Product name** Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free

**Description** Rabbit monoclonal [EPR19386] to Prolactin/PRL - BSA and Azide free

**Host species** Rabbit

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

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

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

Positive control IHC-P: Mouse adenohypophysis tissue.

**General notes** ab238938 is the carrier-free version of ab183967.

> 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**.

#### **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

Clonality Monoclonal
Clone number EPR19386

**Isotype** IgG

#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab238938 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
WB		Use at an assay dependent concentration. Detects a band of approximately 25 kDa (predicted molecular weight: 25 kDa).
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.
IHC-Fr		Use at an assay dependent concentration.  Perform antigen retrieval with heated citrate solution (10mM citrate pH 6.0 + 0.05% Tween-20).
ICC/IF		Use at an assay dependent concentration.

### **Target**

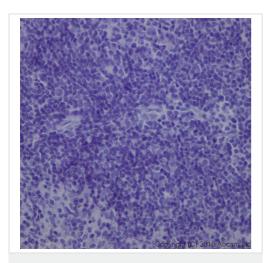
**Relevance** Various hormones are secreted from the anterior pituitary during development and growth.

Prolactin is a growth factor secreted by the anterior pituitary that is necessary for the proliferation

and differentiation of the mammary glands.

Cellular localization Secreted

### **Images**



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Prolactin/PRL antibody
[EPR19386] - BSA and Azide free (ab238938)

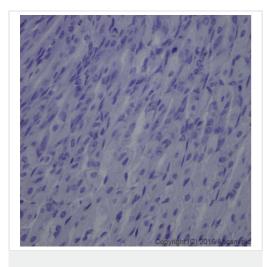
Immunohistochemical analysis of paraffin-embedded Mouse spleen tissue labeling Prolactin/PRL with <u>ab183967</u> at 1/8000 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. No staining was observed on mouse spleen.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is **ab97051** at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab183967</u>).

Heat mediated antigen retrieval was performed with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Prolactin/PRL antibody
[EPR19386] - BSA and Azide free (ab238938)

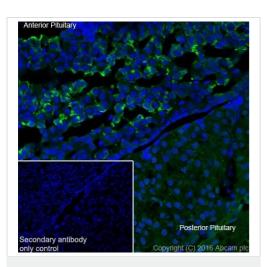
Immunohistochemical analysis of paraffin-embedded Mouse stomach tissue labeling Prolactin/PRL with <u>ab183967</u> at 1/8000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. No staining was observed on mouse stomach.

Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab97051</u> at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab183967).

Heat mediated antigen retrieval was performed with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Frozen sections) - Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free (ab238938)

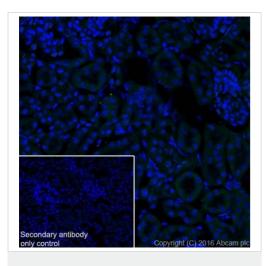
Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen Mouse pituitary tissue labeling Prolactin/PRL with <u>ab183967</u> at 1/200 dilution, followed by Goat Anti-Rabbit lgG (Alexa Fluor<sup>®</sup> 488) (<u>ab150077</u>) secondary antibody at 1/1000 dilution (green).

The result showed cytoplasmic staining on mouse anterior pituitary.

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab150077</u> 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 (ab183967).



Immunohistochemistry (Frozen sections) - Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free (ab238938)

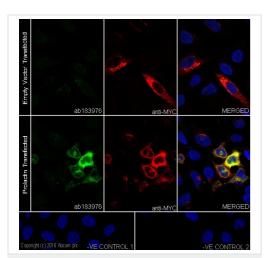
Immunohistochemical analysis of 4% paraformaldehyde-fixed, 0.2% Triton X-100 permeabilized frozen Mouse kidney tissue labeling Prolactin/PRL with <u>ab183967</u> at 1/200 dilution, followed by Goat Anti-Rabbit lgG (Alexa Fluor<sup>®</sup> 488) (<u>ab150077</u>) secondary antibody at 1/1000 dilution (green).

No staining was observed on mouse kidney.

The nuclear counterstain is DAPI (blue).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is <u>ab150077</u> 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 (ab183967).



Immunocytochemistry/ Immunofluorescence - Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free (ab238938)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling Prolactin/PRL with <a href="mailto:ab183967">ab183967</a> at 1/100 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor<sup>®</sup> 488) (<a href="mailto:ab150077">ab150077</a> econdary antibody at 1/1000 dilution (green).

The results show positive staining on HeLa cells transfected with myc-tagged Prolactin.

The nuclear counter stain is DAPI (blue).

Myc is detected with anti-c-Myc (mouse mAb) at 1/1000 dilution and Goat Anti-Mouse IgG H&L (Alexa Fluor® 594) preadsorbed (ab150120) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: <u>ab183967</u> at 1/100 dilution followed by <u>ab150120</u> at 1/1000 dilution.

-ve control 2: Anti-c-Myc at 1/1000 dilution followed by **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 (<u>ab183967</u>).

DAPI MERGED

Ab 183967 DAPI MERGED

Ab 183967 DAPI MERGED

Copyright (c) 2019 Abcam pic .VE CONTROL 1

Immunocytochemistry/ Immunofluorescence - Anti-Prolactin/PRL antibody [EPR19386] - BSA and Azide free (ab238938)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling Prolactin/PRL with <a href="mailto:ab183967">ab183967</a> at 1/100 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor<sup>®</sup> 488) (<a href="mailto:ab150077">ab150077</a>) secondary antibody at 1/1000 dilution (green).

The results show positive staining on HeLa cells transfected with myc-tagged Prolactin.

The nuclear counter stain is DAPI (blue).

Tubulin is detected with Anti-alpha Tubulin antibody [DM1A] - Loading Control (**ab7291**) at 1/1000 dilution and Goat Anti-Mouse lgG H&L (Alexa Fluor® 594) preadsorbed (**ab150120**) at 1/1000 dilution (red).

The negative controls are as follows:

-ve control 1: <u>ab183967</u> at 1/100 dilution followed by <u>ab150120</u> at 1/1000 dilution.

-ve control 2:  $\underline{ab7291}$  at 1/1000 dilution followed by  $\underline{ab150077}$  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 (ab183967).

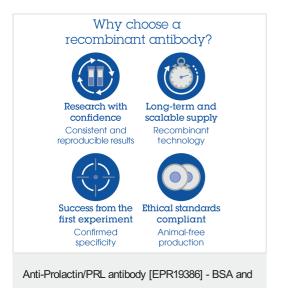
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Prolactin/PRL antibody
[EPR19386] - BSA and Azide free (ab238938)

Immunohistochemical analysis of paraffin-embedded Mouse adenohypophysis tissue labeling Prolactin/PRL with <u>ab183967</u> at 1/8000 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Cytoplasm staining on mouse adenohypophysis was observed. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary

antibody, secondary antibody is <u>ab97051</u> at 1/500 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab183967).

Heat mediated antigen retrieval was performed with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Azide free (ab238938)

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