# abcam

# Product datasheet

# Anti-Galectin 8/Gal-8 antibody [EPR4857] ab109519

Recombinant RabMAb

★★★★ 4 Abreviews 12 References

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#### Overview

**Product name** Anti-Galectin 8/Gal-8 antibody [EPR4857]

**Description** Rabbit monoclonal [EPR4857] to Galectin 8/Gal-8

**Host species** Rahhit

Specificity This antibody has weak cross-reactivity with Galectin 9.

**Tested applications** Suitable for: Flow Cyt (Intra), WB, IP, IHC-P, ICC/IF

Species reactivity Reacts with: Human

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

Positive control HepG2 or LNCaP cell lysate Human prostatic adenocarcinoma tissue IP: HeLa whole cell lysate.

**General notes** 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**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

## **Properties**

**Form** Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Stable for 12 months at -20°C.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 59% PBS, 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal

Clone number EPR4857

**Isotype** IgG

### **Applications**

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab109519 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	
Flow Cyt (Intra)		1/10 - 1/100. <b>ab172730</b> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.	
WB	<b>★★★★ (4)</b>	1/1000 - 1/10000. Predicted molecular weight: 40 kDa.	
IP		1/10 - 1/100.	
IHC-P		1/100 - 1/250. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.  See IHC antigen retrieval protocols.	
ICC/IF		1/50 - 1/100.	

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Tissue specificity Ubiquitous. Selective expression by prostate carcinomas versus normal prostate and benign

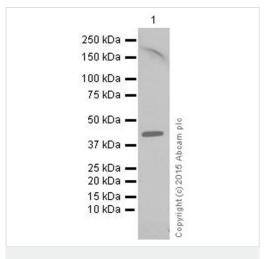
prostatic hypertrophy.

Sequence similarities Contains 2 galectin domains.

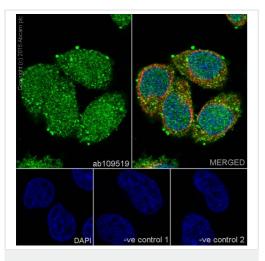
**Domain** Contains two homologous but distinct carbohydrate-binding domains.

Cellular localization Cytoplasm.

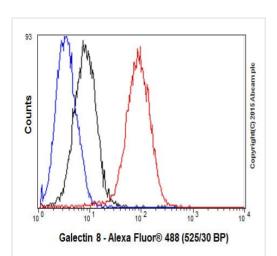
# **Images**



Western blot - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)



Immunocytochemistry/ Immunofluorescence - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)



Flow Cytometry (Intracellular) - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519) at 1/10000 dilution (purified) + LNCAP (human prostat carcinoma) whole cell lysate at 10  $\mu$ g

#### **Secondary**

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

**Predicted band size:** 40 kDa **Observed band size:** 40 kDa

Blocking and diluting buffer 5% NFDM/TBST

ab109519 staining Galectin 8/Gal-8 in HeLa (human cervix adenocarcinoma) cells by ICC/IF

(Immunocytochemistry/immunofluorescence). Cells were fixed with 4% Paraformaldehyde and permeabilized with 0.1% Triton X-100. Samples were incubated with primary antibody at a dilution of 1/100. A goat anti rabbit IgG (Alexa Fluor® 488) (ab150077) was used as the secondary antibody at a dilution of 1/1000. ab7291 and ab150120 were used as counterstains for primary antibody ab109519 and secondary antibody ab150077 respectively and DAPI was used as a nuclear counterstain.

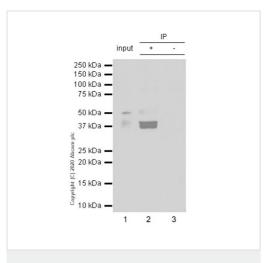
**Negative control 1**: Rabbit primary antibody and anti-mouse secondary antibody (<u>ab150120</u>)

**Negative control 2:** Mouse primary antibody (<u>ab7291</u>) and antirabbit secondary antibody (<u>ab150077</u>)

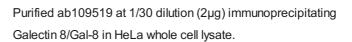
ab109519 staining Galectin 8/Gal-8 in the human cell line HEK293 (human embryonic kidney) by intracellular flow cytometry. Cells were fixed with 4% paraformaldehyde and the sample was incubated with the primary antibody at a dilution of 1/100. A goat anti rabbit IgG (Alexa Fluor<sup>®</sup> 488) at a dilution of 1/500 was used as the secondary antibody.

Isoytype control: Rabbit monoclonal IgG (Black)

Unlabelled control: Cell without incubation with primary antibody and secondary antibody (Blue)



Immunoprecipitation - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)



Lane 1 (input): HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate  $10\mu g$ 

Lane 2 (+): ab109519 + HeLa whole cell lysate.

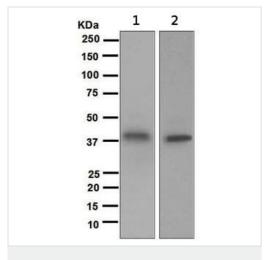
Lane 3 (-): Rabbit monoclonal IgG (ab172730) instead of ab109519 in HeLa whole cell lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (1/10,000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST.

Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 36/40 kDa



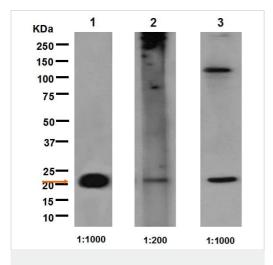
Western blot - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

**All lanes :** Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519) at 1/1000 dilution (unpurified)

Lane 1 : HepG2 cell lysate
Lane 2 : LNCaP cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 40 kDa



Western blot - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

Lanes 1 & 3: unpurified at 1/1000 dilution

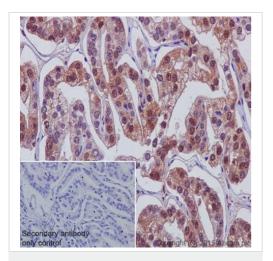
**Lane 2**: Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519) at 1/200 dilution (unpurified)

All lanes: Galectin 9 recombinant protein

Lysates/proteins at 0.02 µg per lane.

Predicted band size: 40 kDa

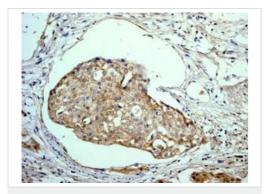
ab109519 has weak cross-reactivity with Galectin 9.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

ab109519 staining Galectin 8/Gal-8 in human prostatic carcinoma tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde and antigen retrieval was by heat mediation in a EDTA buffer. Samples were incubated with primary antibody at a dilution of 1/250. A goat anti-rabbit IgG H&L (HRP) ab97051 was used as the secondary antibody at a dilution of 1/500.

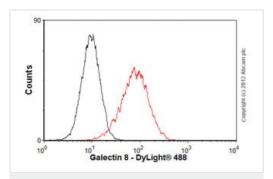
Negative control 1: PBS in place of primary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

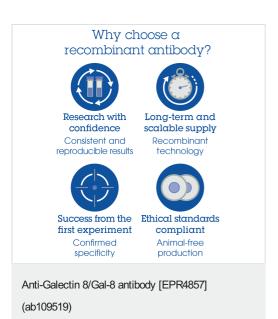
Unpurified ab109519, at a 1/100 dilution, staining Human prostatic adenocarcinoma, using Immunohistochemstry, Formalin/PFA-fixed paraffin-embedded tissue.

Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Flow Cytometry (Intracellular) - Anti-Galectin 8/Gal-8 antibody [EPR4857] (ab109519)

Overlay histogram showing HeLa cells stained with unpurified ab109519 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab109519, 1/100) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-rabbit lgG (H+L) (ab96899) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (1µg/1x106 cells) used under the same conditions. Acquisition of >5,000 events was performed.



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