abcam

Product datasheet

Anti-Galectin 3 antibody [A3A12] ab2785

★★★★★ 16 Abreviews 61 References 画像数 9

製品の概要

製品名 Anti-Galectin 3 antibody [A3A12]

製品の詳細 Mouse monoclonal [A3A12] to Galectin 3

由来種 Mouse

特異性 By Western blot, this antibody detects an ~30 kDa protein representing Galectin 3 from Jurkat

cells transfected with human Galectin 3. Immunohistochemical staining of Galectin 3 in rat olfactory bulb yields a pattern consistent with nuclear and plasma membrane staining.

アプリケーション 適用あり: WB, ICC/IF, IHC-P

種交差性 交差種: Mouse, Human

免疫原 Recombinant full length protein corresponding to Human Galectin 3.

エピトープ The epitope for this antibody has been mapped to the first 58 amino acids of Galectin 3.

ポジティブ・コントロール WB: HeLa, MCF7 whole cell lysates. IHC-P: Human ovarian, colon and mouse colon tissue. ICC:

HeLa, MCF7 and NIH-3T3 cells.

特記事項

This antibody has been shown to potentiate the binding of Galectin 3 to lgG.

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

パッファー Preservative: 0.05% Sodium azide

Constituent: 99% PBS

精製度 Affinity purified

1

一次抗体 備考 This antibody has been shown to potentiate the binding of Galectin 3 to IgG.

ポリ/モノ モノクローナル

クローン名 A3A12

アイソタイプ IgG1

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab2785の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	★★★★ (3)	Use a concentration of 1 µg/ml. Detects a band of approximately 30 kDa (predicted molecular weight: 26 kDa).
ICC/IF	★★★★★ (5)	1/100 - 1/200.
IHC-P	★★★★★ (3)	1/20 - 1/200.

ターゲット情報

機能 Galactose-specific lectin which binds IgE. May mediate with the alpha-3, beta-1 integrin the

stimulation by CSPG4 of endothelial cells migration. Together with DMBT1, required for terminal

differentiation of columnar epithelial cells during early embryogenesis.

組織特異性 A major expression is found in the colonic epithelium. It is also abundant in the activated

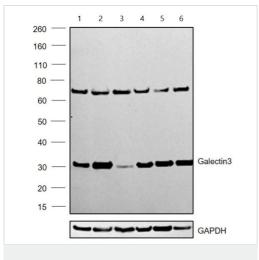
macrophages.

配列類似性 Contains 1 galectin domain.

細胞内局在 Nucleus. Cytoplasmic in adenomas and carcinomas. May be secreted by a non-classical

secretory pathway and associate with the cell surface.

画像



Western blot - Anti-Galectin 3 antibody [A3A12] (ab2785)

All lanes : Anti-Galectin 3 antibody [A3A12] (ab2785) at 1/1000 dilution

Lane 1 : HeLa (human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lane 2: MDA-MB-231 (human breast adenocarcinoma cell line) whole cell lysate

Lane 3 : A549 (human lung carcinoma cell line) whole cell lysate

Lane 4 : PC-3 (human prostate adenocarcinoma cell line) whole
cell lysate

Lane 5 : U-87 MG (human glioblastoma-astrocytoma epithelial cell line) whole cell lysate

Lane 6 : A431 (human epidermoid carcinoma cell line) whole cell lysate

Lysates/proteins at 30 µg per lane.

Secondary

All lanes: Goat anti-Mouse IgG (H+L) (HRP) at 1/4000 dilution

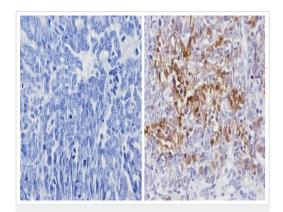
Developed using the ECL technique.

Predicted band size: 26 kDa Observed band size: 30 kDa

Additional bands at: 70 kDa. We are unsure as to the identity of

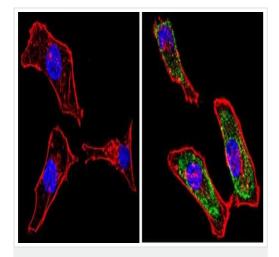
these extra bands.

Whole cell extracts were electrophoresed using a 4-12% Bis-Tris Protein Gel. Resolved proteins were then transferred onto a Nitrocellulose membrane using a dry blotting system.



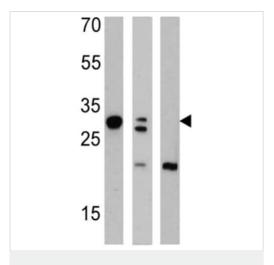
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Galectin 3 antibody
[A3A12] (ab2785)

Immunohistochemical analysis of human ovarian carcinoma tissue (right) labeling Galectin 3 in the nucleus and cytoplasm with ab2785, compared with a negative control (left) by Immunohistochemistry (formalin/PFA-fixed paraffin-embedded sections). To expose target proteins, antigen retrieval method was performed using 10mM sodium citrate (pH 6.0) microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature. Tissue sections were incubated with the primary antibody (1:100 in 3% BSA-PBS) overnight at 4°C. A HRP-conjugated anti-mouse IgG was used as the secondary antibody, followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.



Immunocytochemistry/ Immunofluorescence - Anti-Galectin 3 antibody [A3A12] (ab2785)

Immunofluorescent analysis of HeLa (Human epithelial cell line from cervix adenocarcinoma) cells labeling Galectin 3 (green) in the cytoplasm and nucleus with ab2785. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with a Galectin 3 monoclonal antibody (Product # MA1-940) in 3% BSA-PBS at a dilution of 1:100 and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight-conjugated secondary antibody in PBS at room temperature in the dark. F-actin (red) was stained with a fluorescent red phalloidin and nuclei (blue) were stained with Hoechst or DAPI. Images were taken at a magnification of 60x



Western blot - Anti-Galectin 3 antibody [A3A12] (ab2785)

All lanes : Anti-Galectin 3 antibody [A3A12] (ab2785) at 1/1000 dilution

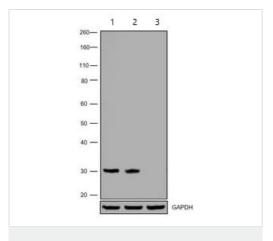
Lane 1 : MCF7 (Human breast adenocarcinoma cell line) whole cell lysate

Lane 2 : HeLa (Human epithelial cell line from cervix adenocarcinoma) whole cell lysate

Lane 3: NIH/3T3 (Mouse embryo fibroblast cell line) whole cell lysate

Lysates/proteins at 25 µg/ml per lane.

Predicted band size: 26 kDa Observed band size: 30 kDa



Western blot - Anti-Galectin 3 antibody [A3A12] (ab2785)

All lanes : Anti-Galectin 3 antibody [A3A12] (ab2785) at 1/1000 dilution

Lane 1 : Wild-Type HeLa (human epithelial cell line from cervix adenocarcinoma) membrane extract

Lane 2: HeLa CAS9 membrane extract

Lane 3: HeLa LGALS3 knockout membrane extract

Lysates/proteins at 30 µg per lane.

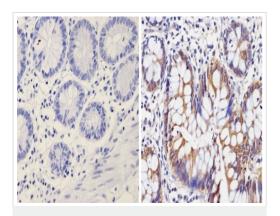
Secondary

All lanes: Goat anti-Mouse IgG (H+L) (HRP) at 1/4000 dilution

Developed using the ECL technique.

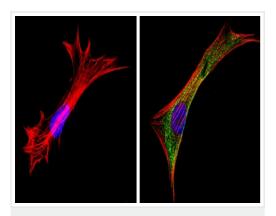
Predicted band size: 26 kDa

Knockout of LGALS3 was achieved by CRISPR-Cas9 genome editing using Lentiviral sgRNA and Cas9 Lentivirus. The samples were electrophoresed using 4-12% Bis-Tris Protein Gel. Resolved proteins were then transferred onto a nitrocellulose membrane using a dry blotting system.



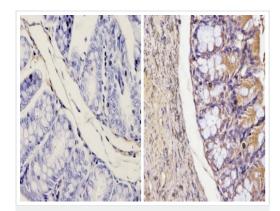
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Galectin 3 antibody
[A3A12] (ab2785)

Immunohistochemical analysis (formalin/PFA-fixed paraffinembedded sections) human colon tissue (right) labelling Galectin 3 in the nucleus and cytoplasm with ab2785, compared with a negative control (left). To expose target proteins, antigen retrieval method was performed using 10mM sodium citrate (pH 6.0) microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature. Tissue sections were incubated with the primary antibody (1:100 in 3% BSA-PBS) overnight at 4°C. A HRP-conjugated anti-mouse IgG was used as the secondary antibody, followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.



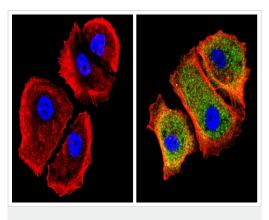
Immunocytochemistry/ Immunofluorescence - Anti-Galectin 3 antibody [A3A12] (ab2785)

Immunofluorescent analysis of NIH/3T3 (Mouse embryo fibroblast cell line) cells labelling Galectin 3 (green) in the cytoplasm and nucleus with ab2785. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with the primary antibody (1:100 in 3% BSA-PBS) overnight at 4 °C. A DyLight-conjugated anti-mouse was used as the secondary antibody. Red (phalloidin) - F-actin, Blue - nuclei. Images were taken at a magnification of 60x.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Galectin 3 antibody
[A3A12] (ab2785)

Immunohistochemical analysis of (formalin/PFA-fixed paraffinembedded sections) mouse colon tissue (right) labelling Galectin 3 in the nucleus and cytoplasm with ab2785, compared with a negative control (left) by . To expose target proteins, antigen retrieval method was performed using 10mM sodium citrate (pH 6.0) microwaved for 8-15 min. Following antigen retrieval, tissues were blocked in 3% H2O2-methanol for 15 min at room temperature. Tissue sections were incubated with the primary antibody (1:20 in 3% BSA-PBS) overnight at 4°C. A HRP-conjugated anti-mouse IgG was used as the secondary antibody, followed by colorimetric detection using a DAB kit. Tissues were counterstained with hematoxylin and dehydrated with ethanol and xylene to prep for mounting.



Immunocytochemistry/ Immunofluorescence - Anti-Galectin 3 antibody [A3A12] (ab2785)

Immunofluorescent analysis of MCF7 (Human breast adenocarcinoma cell line) cells labelling Galectin 3 (green) in the cytoplasm and nucleus with ab2785. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were incubated with the primary antibody (1:100 in 3% BSA-PBS) overnight at 4 °C. A DyLight-conjugated anti-mouse was used as the secondary antibody. Red (phalloidin) - F-actin, Blue - nuclei. Images were taken at a magnification of 60x.

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