abcam

Product datasheet

Anti-N Cadherin antibody - Intercellular Junction Marker ab18203

★★★★★ 47 Abreviews 758 References 画像数 11

製品の概要

製品名 Anti-N Cadherin antibody - Intercellular Junction Marker

製品の詳細 Rabbit polyclonal to N Cadherin - Intercellular Junction Marker

由来種 Rabbit

特異性 Replenishment batches of our polyclonal antibody, ab18203 are tested in WB. Previous batches

were additionally validated in ICC/IF, IHC-P and IP. These applications are still expected to work and are covered by our Abpromise guarantee. You may also be interested in our alternative

recombinant antibody, ab76011.

アプリケーション 適用あり: WB, IHC-P, IP, ICC/IF

種交差性 交差種: Mouse, Rat, Human

交差が予測される動物種: Chicken, Cow, Pig, Xenopus laevis 4

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

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

特記事項

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.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

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精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ lgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab18203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	★★★★☆ (8)	Use a concentration of 1 µg/ml. Detects a band of approximately 125-135 kDa (predicted molecular weight: 100 kDa).
IHC-P	**** (13)	Use a concentration of 1 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
IP		Use at an assay dependent concentration.
ICC/IF	★★★★☆ (7)	Use a concentration of 5 µg/ml.

ターゲット情報

機能 Cadherins are calcium dependent cell adhesion proteins. They preferentially interact with

themselves in a homophilic manner in connecting cells; cadherins may thus contribute to the sorting of heterogeneous cell types. CDH2 may be involved in neuronal recognition mechanism. In

hippocampal neurons, may regulate dendritic spine density.

配列類似性 Contains 5 cadherin domains.

細胞内局在 Cell membrane.

画像



Western blot - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

All lanes : Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203) at 1 μ g/ml

Lane 1 : Brain (Rat) Tissue Lysate at 10 μg **Lane 2 :** Brain (Mouse) Tissue Lysate at 10 μg **Lane 3 :** Brain (Human) Tissue Lysate at 20 μg

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) preadsorbed at 1/50000 dilution

Developed using the ECL technique.

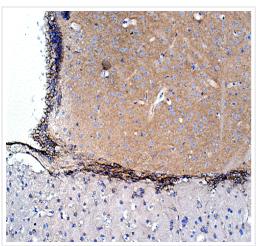
Performed under reducing conditions.

Predicted band size: 100 kDa **Observed band size:** 125 kDa

Exposure time: 1 minute

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 2% Bovine Serum Albumin before being incubated with ab18203 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution ab133406.

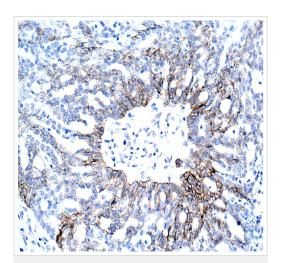
The N Cadherin protein has a predicted molecular weight of 100 kDa, however it is extensively glycosylated and has been shown to run in the 125-135 kDa region (SwissProt data).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

Image courtesy of Mr Carl Hobbs, Kings College London

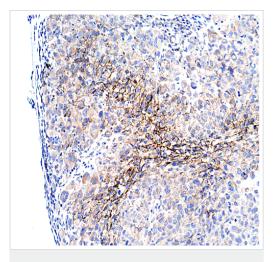
Anti N-cadherin (ab18203) staining of mouse brain using immunohistochemistry (formaldehyde-fixed, paraffin-embedded sections). Heat-mediated antigen retrieval was carried out using citric acid. Samples were incubated with primary antibody (1/1000) for two hours at room temperature. A biotin-conjugated goat antirabbit IgG polyclonal was used as the secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

Image courtesy of Mr Carl Hobbs, Kings College London.

Anti N-cadherin (ab18203) staining of human ovarian cancer tissue using immunohistochemistry (formaldehyde-fixed, paraffinembedded sections). Heat-mediated antigen retrieval was carried out using citric acid. Samples were incubated with primary antibody (1/1000) for two hours at room temperature. A biotin-conjugated goat anti-rabbit IgG polyclonal was used as the secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

Image courtesy of Mr Carl Hobbs, Kings College London.

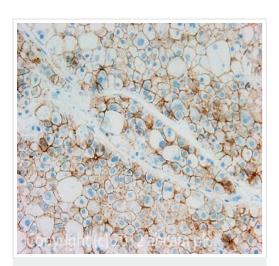
Anti N-cadherin (ab18203) staining in a human melanoma xenograft mouse model using immunohistochemistry (formaldehyde-fixed, paraffin-embedded sections). Heat-mediated antigen retrieval was carried out using citric acid. Samples were incubated with primary antibody (1/1000) for two hours at room temperature. A biotinconjugated goat anti-rabbit IgG polyclonal was used as the secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

Image courtesy of Mr Carl Hobbs, Kings College London.

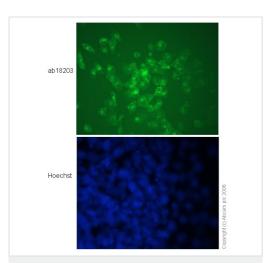
Anti N-cadherin (ab18203) staining of E17 developing rat retina using immunohistochemistry (formaldehyde-fixed, paraffinembedded sections). Heat-mediated antigen retrieval was carried out using citric acid. Samples were incubated with primary antibody (1/1000) for two hours at room temperature. A biotin-conjugated goat anti-rabbit IgG polyclonal was used as the secondary antibody.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

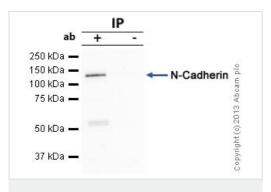
IHC image of N Cadherin staining in Human liver cancer formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab18203, 1µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.



Immunocytochemistry/ Immunofluorescence - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203) stained human embryonic stem cells differentiated into mesoderm.



Immunoprecipitation - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

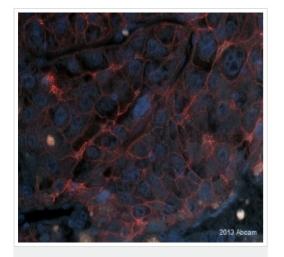
N Cadherin was immunoprecipitated using 0.5mg Mouse Brain whole tissue lysate, 5µg of Rabbit polyclonal to N Cadherin and 50µl of protein G magnetic beads (+). No antibody was added to the control (-).

The antibody was incubated under agitation with Protein G beads for 10min, Mouse Brain whole tissue lysate lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation.

Proteins were eluted by addition of 40μ I SDS loading buffer and incubated for 10min at 70° C; 10μ I of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with ab18203.

Secondary: Mouse monoclonal [SB62a] Secondary Antibody to Rabbit IgG light chain (HRP) (ab99697).

Band: 135kDa: N Cadherin



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

This image is courtesy of an anonymous Abreview

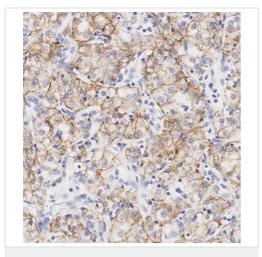
ab18203 staining N Cadherin in Human liver tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffinembedded sections). Tissue was fixed with formaldehyde and blocked with 10% serum for 30 minutes at 25°C; antigen retrieval was by heat mediation in a 10 mM citrate buffer pH6.0. Samples were incubated with primary antibody (1/100 in PBS plus casein) for 90 minutes at 37°C. A Biotin-conjugated Goat anti-rabbit IgG polyclonal (1/200) was used as the secondary antibody.



Paraformaldehyde-fixed, 0.2% Triton X100 permeabilized HaCaT (human keratinocyte cell line) cells stained for N Cadherin (green) using ab18203 at 1/200 dilution in ICC/IF, followed by Donkey anti Rabbit Alexa Fluor 568 at 1/500 dilution.

Immunocytochemistry/ Immunofluorescence - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

This image is courtesy of an Abreview submitted by ${\rm Dr.}$ Ann Wheeler.



with ab18203 at 1µg/ml.

Immunohistochemistry of kidney carcinoma staining N Cadherin

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-N Cadherin antibody - Intercellular Junction Marker (ab18203)

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