abcam

Product datasheet

Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker ab51034

重组 RabMAb

★★★★★ 6 Abreviews 16 References 13 图像

概述

产品名称 Anti-pan Cadherin抗体[EPR1792Y] - Intercellular Junction Marker

描述 兔单克隆抗体[EPR1792Y] to pan Cadherin - Intercellular Junction Marker

宿主 Rabbit

特异性 The immunogen used for this product shares 100% homology with N-cadherin, 100% homology

with R-cadherin, 93% homology to K-cadherin, 93% homology to P-cadherin, 93% homology to E-cadherin, and 73% homology to VE-cadherin. Cross-reactivities with these proteins have not

been confirmed experimentally.

经测试应用 适用于: Flow Cyt (Intra), WB, IP, IHC-P, ICC/IF

种属反应性 与反应: Mouse, Rat, Human

免疫原 Synthetic peptide within Human pan Cadherin (C terminal). The exact sequence is proprietary.

Database link: P19022

阳性对照 WB: C6 whole cell lysate. Mouse heart tissue lysate. Human, mouse, and rat brain tissue lysates;

ICC: MCF7 cells; IHC-P: Rat, mouse and human cardiac muscle tissue; human and rat colon and

mouse testis tissue; IP: Rat heart lysate; Flow Cyt (intra): HeLa cells.

常规说明 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**.

性能

形式 Liquid

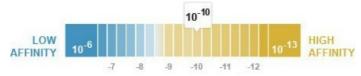
存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle.

1

解离常数(K_D)

 $K_D = 6.10 \times 10^{-10} M$



Learn more about K_D

存储溶液 pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 0.05% BSA, 40% Glycerol, 59% PBS

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR1792Y

同种型 IgG

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab51034于以下的经测试应用

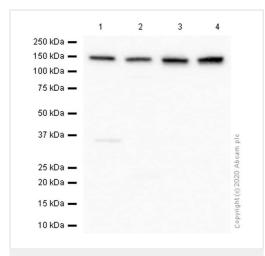
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Flow Cyt (Intra)		1/20. For unpurified use at 1/30. <u>ab172730</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
WB	**** (1)	1/50000. Detects a band of approximately 140 kDa (predicted molecular weight: 100 kDa).
IP		1/20. For unpurified use at 1/40.
IHC-P	★★★★★ (2)	1/500. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols. For unpurified use at 1/100.
ICC/IF	*** <u>*</u> (2)	1/100 - 1/250.

靶标

相关性

Cadherins are members of a multigene family of single chain glycoprotein receptors mediating calcium dependent cell-cell adhesion. They play an important role in the growth and development of cells via the mechanisms of control of tissue architecture and the maintenance of tissue integrity. Cadherins are expressed in a tissue specific manner and and are required for assembly of cells into solid tissue. Individual cadherin molecules are known to co-operate with each other to form a linear cell adhesion zipper. In adhesion junctions cadherins are bound to beta and gamma catenins which in turn bind to alpha catenin, an actin binding protein. Cadherins play an important part in tumor invasion and metastasis.



Western blot - Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034)

All lanes : Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034) at 1/50000 dilution (purified)

Lane 1: C6 (Rat glial tumor glial cell) whole cell lysate

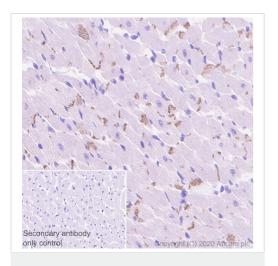
Lane 2 : Human brain lysate
Lane 3 : Mouse brain lysate
Lane 4 : Rat brain lysate

Lysates/proteins at 20 µg per lane.

Secondary

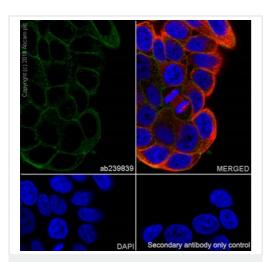
 $\begin{tabular}{ll} \textbf{All lanes:} Goat Anti-Rabbit \ lgG (HRP) with minimal cross-reactivity with human \ lgG \ at 1/2000 \ dilution \end{tabular}$

Predicted band size: 100 kDa Observed band size: 140 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody
[EPR1792Y] - Intercellular Junction Marker
(ab51034)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat cardiac muscle tissue sections labeling pan Cadherin with purified ab51034 at 1/500 dilution (0.26 µg/mL). Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

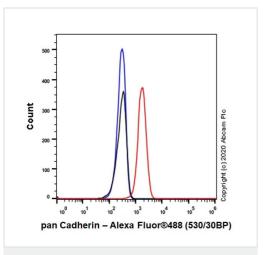


Immunocytochemistry/ Immunofluorescence - Antipan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034)

Immunocytochemistry/Immunofluorescence analysis of MCF7 (human breast adenocarcinoma epithelial cells labeling pan Cadherin with purified ab51034 at 1/250. Cells were fixed with 4% Paraformaldehyde and permeabilised with 0.1% TritonX-100. ab150077, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/1000) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

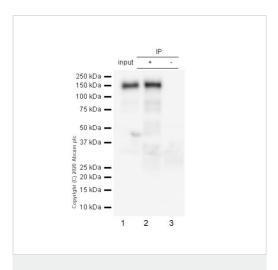
ab195889, Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) 1/200 was used as counterstain antibody.

Confocal image showing membranous staining in MCF7 cells.



Flow Cytometry (Intracellular) - Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034)

Intracellular Flow Cytometry analysis of HeLa (Human cervix adenocarcinoma epithelial cell) cells labeling pan Cadherin with purified ab51034 at 1/20 dilution (10µg/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit IgG (Alexa Fluorr® 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Immunoprecipitation - Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034)

Purified ab51034 at 1/20 dilution (0.6 μg) immunoprecipitating pan Cadherin in Rat heart lysate.

Lane 1 (input): Rat heart lysate 10 µg

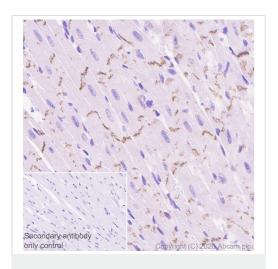
Lane 2 (+): ab51034 + Rat heart lysate.

Lane 3 (-): Rabbit monoclonal lgG (<u>ab172730</u>) instead of ab51034 in Rat heart lysate.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) (1/1000 dilution) was used for Western blotting.

Blocking Buffer and concentration: 5% NFDM/TBST. Diluting buffer and concentration: 5% NFDM/TBST.

Observed band size: 140 kDa

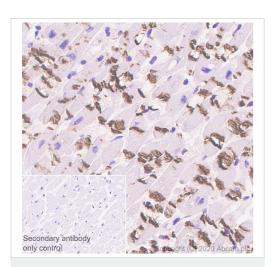


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody

[EPR1792Y] - Intercellular Junction Marker

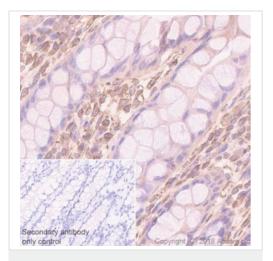
(ab51034)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse cardiac muscle tissue sections labeling pan Cadherin with purified ab51034 at 1/500 dilution (0.26 µg/mL). Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody
[EPR1792Y] - Intercellular Junction Marker
(ab51034)

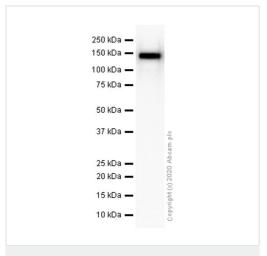
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human cardiac muscle tissue sections labeling pan Cadherin with purified ab51034 at 1/500 dilution (0.26 µg/mL). Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (ab209101) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody
[EPR1792Y] - Intercellular Junction Marker
(ab51034)

Immunohistochemical analysis of Paraffin-embedded human colon tissue sections labeling pan Cadherin with unpurified ab51034 at 1/100. Goat Anti-Rabbit IgG H&L (HRP) was used as the secondary antibody. Sections were counterstained with Hematoxylin. Antigen retrieval was heat mediated using **ab93684** (Tris/EDTA buffer, pH 9.0).

Positive staining on human colon.

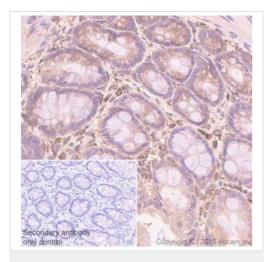


Western blot - Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034) Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034) at 1/50000 dilution (purified) + Mouse heart lysate at 15 μg

Secondary

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

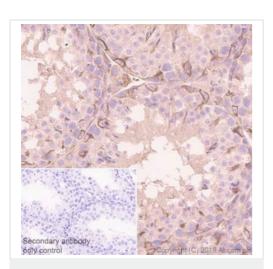
Predicted band size: 100 kDa **Observed band size:** 140 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody
[EPR1792Y] - Intercellular Junction Marker
(ab51034)

Immunohistochemical analysis of Paraffin-embedded rat colon tissue sections labeling pan Cadherin with unpurified ab51034 at 1/100. Goat Anti-Rabbit IgG H&L (HRP) was used as the secondary antibody. Sections were counterstained with Hematoxylin. Antigen retrieval was heat mediated using ab93684 (Tris/EDTA buffer, pH 9.0).

Positive staining on rat colon.



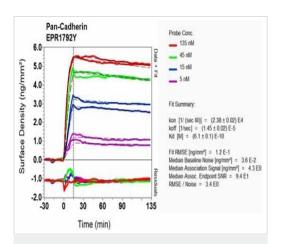
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-pan Cadherin antibody

[EPR1792Y] - Intercellular Junction Marker

(ab51034)

Immunohistochemical analysis of Paraffin-embedded mouse testis tissue sections labeling pan Cadherin with unpurified ab51034 at 1/100. Goat Anti-Rabbit IgG H&L (HRP) was used as the secondary antibody. Sections were counterstained with Hematoxylin. Antigen retrieval was heat mediated using ab93684 (Tris/EDTA buffer, pH 9.0).

Positive staining on mouse testis.



Ol-RD Scanning - Anti-pan Cadherin antibody [EPR1792Y] - Intercellular Junction Marker (ab51034)

Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about KD



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