

Anti-P Glycoprotein antibody [EPR10364-57] ab170904

敲除验证
重组 RabMAb

★★★★☆
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概述

产品名称	Anti-P Glycoprotein抗体[EPR10364-57]
描述	兔单克隆抗体[EPR10364-57] to P Glycoprotein
宿主	Rabbit
特异性	<p>P-glycoprotein 1 (also known as Multidrug resistance protein 1) has a predicted molecular weight of 141 kDa, however it has 3 potential glycosylation sites (N-linked) which may affect the migration of the protein. In our hands this antibody detects a predominant protein band migrating in the region of 180-200 kDa and typically will demonstrate a smear on the membrane in the region of the 150-300 kDa due to the glycosylation profile of the protein. It may be necessary to optimise your cell or tissue lysis protocol to efficiently extract P-glycoprotein 1 as it is a multi-pass membrane protein. Abcam recommends not boiling the sample after lysis. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.</p>
经测试应用	<p>适用于: IHC-P, WB</p> <p>不适用于: ICC/IF</p>
种属反应性	与反应: Mouse, Rat, Human
免疫原	<p>Recombinant fragment within Human P Glycoprotein aa 350-750. The exact sequence is proprietary.</p> <p>Database link: P08183</p>
阳性对照	<p>WB: HEK-293T, C6, HeLa, HepG2 and HEK-293T cell lysates; human fetal brain tissue lysate; mouse and rat brain and kidney tissue lysates. Wild-type HAP1 cell lysate; IHC-P: Human hepatocellular carcinoma, brain and kidney tissues.</p>
常规说明	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

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 long term. Avoid freeze / thaw cycle.
存储溶液	Preservative: 0.01% Sodium azide Constituents: 40% Glycerol, 0.05% BSA, PBS
纯度	Protein A purified
克隆	单克隆
克隆编号	EPR10364-57
同种型	IgG

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用 ab170904 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
IHC-P	★★★★★ (1)	1/1200. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See IHC antigen retrieval protocols . For unpurified use at 1/20 - 1/100 The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.
WB	★★★★★ (4)	1/1000 - 1/5000. Predicted molecular weight: 141 kDa. For optimal detection Abcam recommends not boiling the sample after lysis.

应用说明 Is unsuitable for ICC/IF.

靶标

功能	Energy-dependent efflux pump responsible for decreased drug accumulation in multidrug-resistant cells.
组织特异性	Expressed in liver, kidney, small intestine and brain.
疾病相关	Genetic variations in ABCB1 are associated with susceptibility to inflammatory bowel disease type 13 (IBD13) [MIM:612244]. Inflammatory bowel disease is characterized by a chronic relapsing intestinal inflammation. It is subdivided into Crohn disease and ulcerative colitis phenotypes. Crohn disease may involve any part of the gastrointestinal tract, but most frequently the terminal ileum and colon. Bowel inflammation is transmural and discontinuous; it may contain granulomas or be associated with intestinal or perianal fistulas. In contrast, in ulcerative colitis, the inflammation is continuous and limited to rectal and colonic mucosal layers; fistulas and granulomas are not observed. Both diseases include extraintestinal inflammation of the skin, eyes, or joints. Crohn disease and ulcerative colitis are commonly classified as autoimmune diseases.
序列相似性	Belongs to the ABC transporter superfamily. ABCB family. Multidrug resistance exporter (TC 3.A.1.201) subfamily.

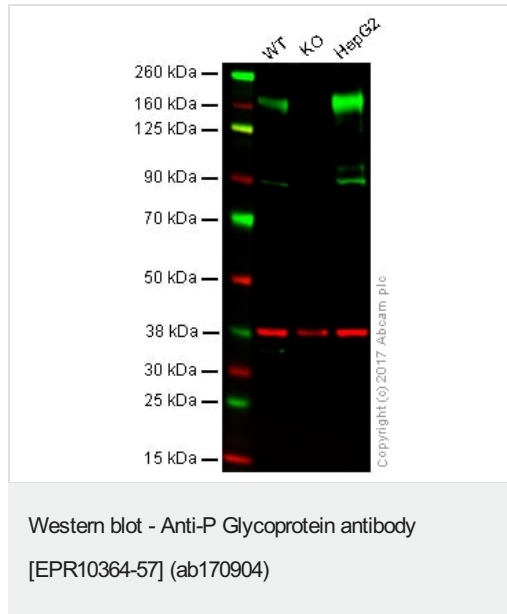
Contains 2 ABC transmembrane type-1 domains.

Contains 2 ABC transporter domains.

细胞定位

Membrane.

图片



Lane 1: Wild type HAP1 whole cell lysate (20 µg)

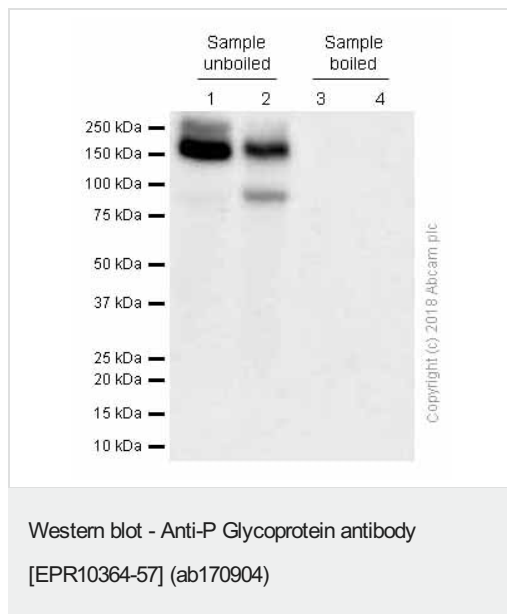
Lane 2: P Glycoprotein knockout HAP1 whole cell lysate (20 µg)

Lane 3: HepG2 whole cell lysate (20 µg)

Lanes 1 - 3: Merged signal (red and green). Green - ab170904 observed at 160 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

ab170904 was shown to specifically react with P Glycoprotein when P Glycoprotein knockout samples were used. Wild-type and P Glycoprotein knockout samples were subjected to SDS-PAGE. ab170904 and **ab8245** (Mouse anti GAPDH loading control) were incubated overnight at 4°C at 1 µg/ml and 1/10000 dilution respectively.

Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed **ab216773** and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed **ab216776** secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 0.1 µg/ml (purified)

Lanes 1 & 3 : C6 (Rat glial tumor glial cell) whole cell lysates prepared using RIPA lysis method

Lanes 2 & 4 : C6 (Rat glial tumor glial cell) whole cell lysates prepared using 1%SDS Hot lysis method

Lysates/proteins at 15 µg per lane.

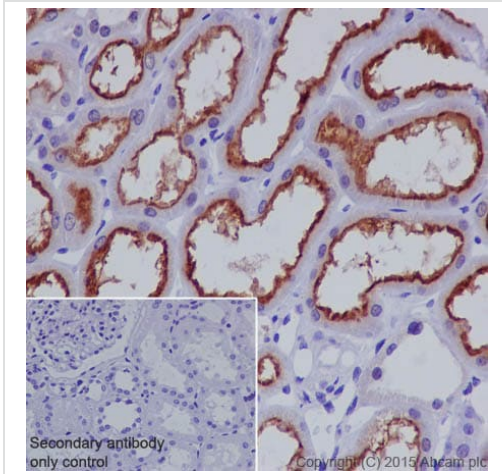
Secondary

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

Predicted band size: 141 kDa

Observed band size: 180 kDa

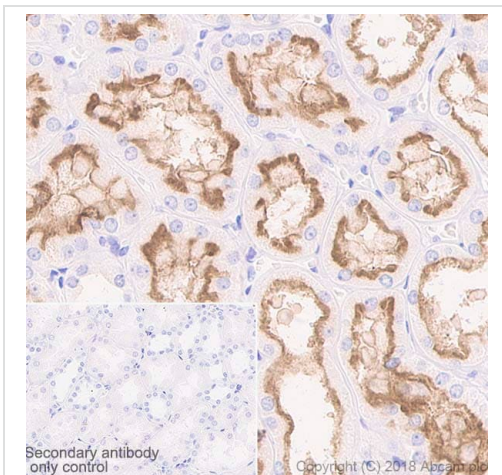
Blocking and diluting buffer: 5% NFDm/TBST.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemical staining of paraffin embedded human kidney with purified ab170904 at a working dilution of 1/100. The secondary antibody used is HRP goat anti-rabbit IgG H&L ([ab97051](#)) at 1/500. Counterstained with hematoxylin. Antigen retrieval was performed using Tris-EDTA buffer, pH 9.0.

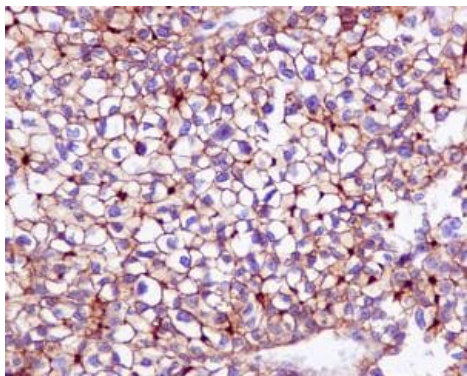
PBS was used instead of the primary antibody as the negative control (inset).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue sections labeling P Glycoprotein with purified ab170904 at 1:1200 dilution (0.24 µg/ml). Heat mediated antigen retrieval was performed using [ab93684](#) (Tris/EDTA buffer, pH 9.0) ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody. Hematoxylin was used as a counterstain.

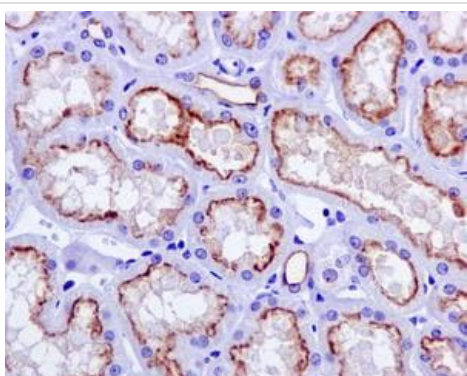
Negative control: PBS instead of the primary antibody (inset).



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human hepatocellular carcinoma tissue labeling P Glycoprotein with unpurified ab170904 at 1/250 dilution.

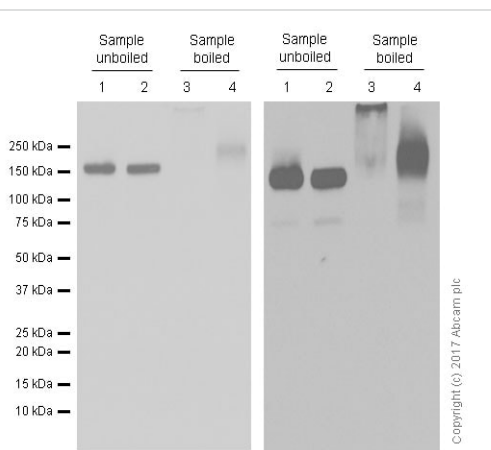
Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis human kidney tissue labeling P Glycoprotein with unpurified ab170904 at 1/250 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/2000 dilution (Purified)

Lanes 1 & 3 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates prepared using RIPA lysis method

Lanes 2 & 4 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate prepared using 1% SDS hot lysis method

Lysates/proteins at 15 µg per lane.

Secondary

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

Predicted band size: 141 kDa

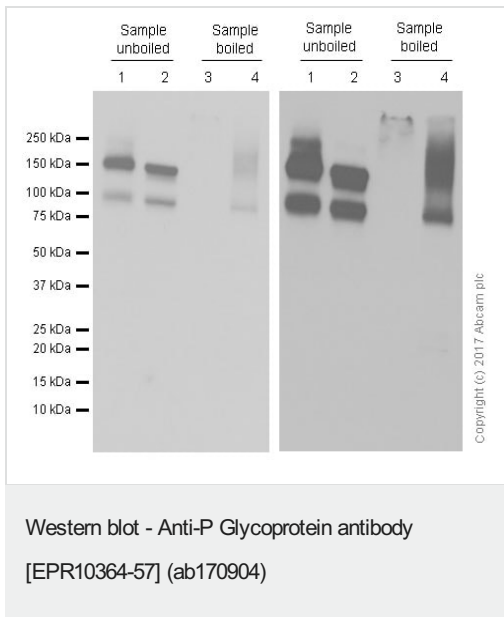
Observed band size: 180 kDa

Blocking and diluting buffer: 5% NFDm/TBST

Exposure time: Left image -10 seconds; Right image - 1 minute.

We suggest **not to boil** the sample after lysis.

For 1% SDS Hot Lysates preparation protocol, please refer to the protocol book in the protocol section and/or [here \(downloadable copy\)](#).



All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/2000 dilution (Purified)

Lanes 1 & 3 : Raw264.7 (Mouse Abelson murine leukemia virus-induced tumor macrophage) whole cell lysates prepared using RIPA lysis method

Lanes 2 & 4 : Raw264.7 (Mouse Abelson murine leukemia virus-induced tumor macrophage) whole cell lysate prepared using 1%SDS lysis method

Lysates/proteins at 15 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/20000 dilution

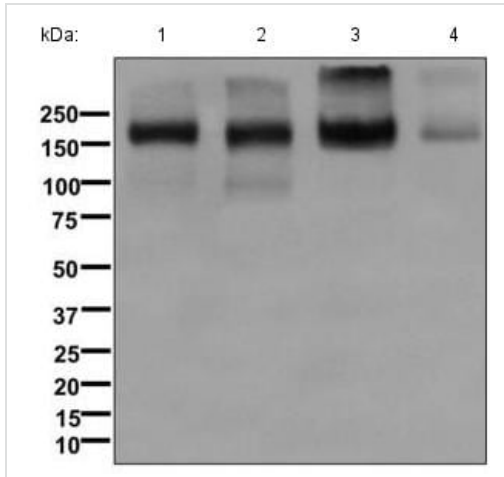
Predicted band size: 141 kDa

Observed band size: 180 kDa

Blocking and diluting buffer: 5% NFDm/TBST

Exposure time: Left image - 3 seconds; Right image - 10 seconds.

We suggest **not to boil** the sample after lysis.



Western blot - Anti-P Glycoprotein antibody
[EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57]
(ab170904) at 1/1000 dilution (unpurified)

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

Lane 2 : HepG2 (Human liver hepatocellular carcinoma cell line)
cell lysate

Lane 3 : HEK-293T (Human epithelial cell line from embryonic
kidney transformed with large T antigen) cell lysate

Lane 4 : Human fetal brain tissue lysate

Lysates/proteins at 10 µg per lane.

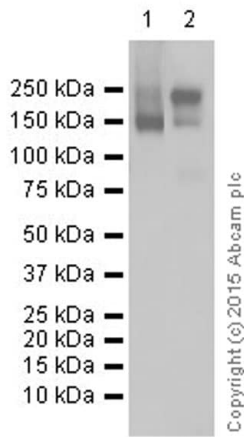
Secondary

Lanes 1-3 : HRP-conjugated goat anti-rabbit IgG at 1/2000 dilution

Lane 4 : Standard HRP labeled goat anti-rabbit at 1/2000 dilution

Developed using the ECL technique.

Predicted band size: 141 kDa



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/1000 dilution (purified)

Lane 1 : Mouse brain lysate

Lane 2 : C6 (Rat glial tumor cell line) cell lysate

Lysates/proteins at 10 µg per lane.

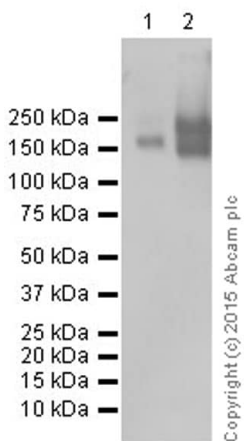
Secondary

All lanes : Goat Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 141 kDa

Observed band size: 180 kDa

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/2000 dilution (purified)

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

Lane 2 : HepG2 (human hepatocellular carcinoma) whole cell lysate

Lysates/proteins at 10 µg per lane.

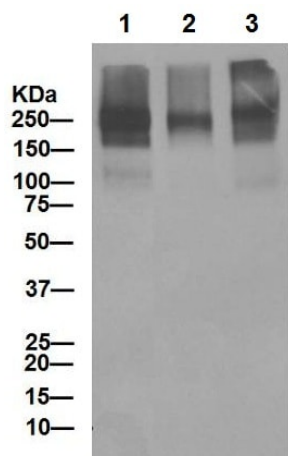
Secondary

All lanes : Goat Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 141 kDa

Observed band size: 180 kDa

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/200 dilution (unpurified)

Lane 1 : HepG2 (Human liver hepatocellular carcinoma cell line) cell lysate

Lane 2 : Mouse brain tissue lysate

Lane 3 : Rat brain tissue lysate

Lysates/proteins at 20 µg per lane.

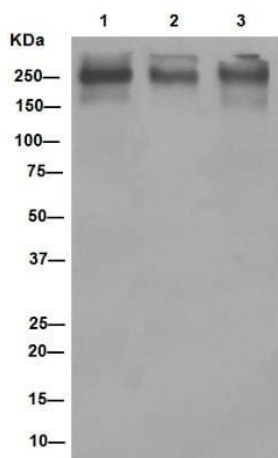
Secondary

All lanes : Peroxidase-conjugated goat anti-rabbit IgG (H+L) at 1/1000 dilution

Predicted band size: 141 kDa

Observed band size: 180 kDa

Blocking/Dilution buffer and concentration: 5% NFDM/TBST.



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/1000 dilution (purified)

Lane 1 : HepG2 (Human liver hepatocellular carcinoma cell line) cell lysate

Lane 2 : Mouse brain tissue lysate

Lane 3 : Rat brain tissue lysate

Lysates/proteins at 20 µg per lane.

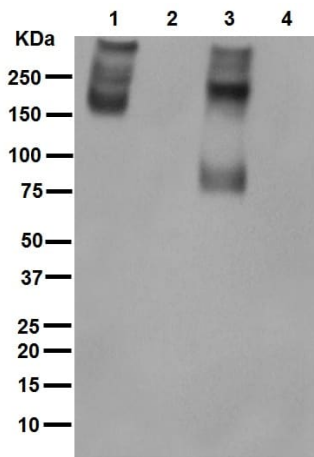
Secondary

All lanes : Peroxidase-conjugated goat anti-rabbit IgG (H+L) at 1/1000 dilution

Predicted band size: 141 kDa

Observed band size: 180 kDa

Blocking/Dilution buffer and concentration: 5% NFDM/TBST.



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/1000 dilution (unpurified)

Lane 1 : Rat brain tissue lysate

Lane 2 : Rat heart tissue lysate

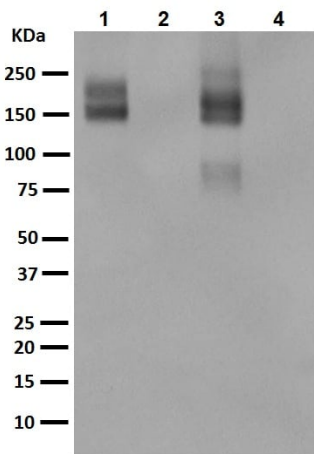
Lane 3 : Rat kidney tissue lysate

Lane 4 : Rat spleen tissue lysate

Predicted band size: 141 kDa

Observed band size: 180 kDa

Exposure time: 2 minutes



Western blot - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

All lanes : Anti-P Glycoprotein antibody [EPR10364-57] (ab170904) at 1/1000 dilution (unpurified)

Lane 1 : Mouse brain tissue lysate

Lane 2 : Mouse heart tissue lysate

Lane 3 : Mouse kidney tissue lysate

Lane 4 : Mouse spleen tissue lysate

Secondary

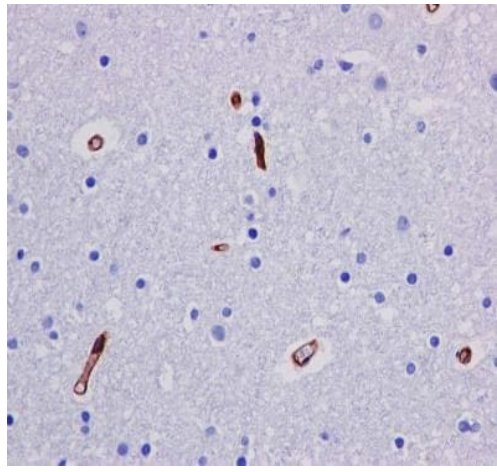
All lanes : HRP-conjugated goat anti-rabbit IgG (H+L) at 1/2000 dilution

Developed using the ECL technique.

Predicted band size: 141 kDa

Observed band size: 180 kDa

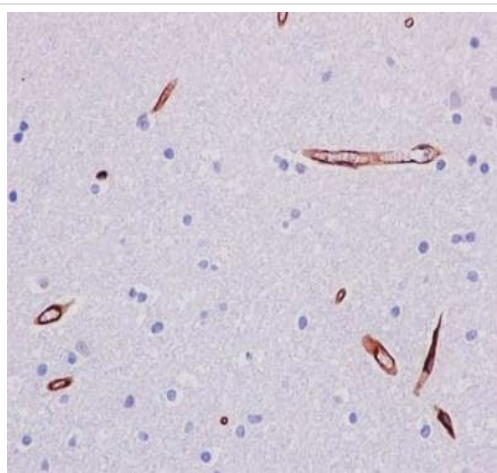
Exposure time: 2 minutes



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human brain tissue labeling P Glycoprotein with unpurified ab170904 at 1/20. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody.

Counterstained with hematoxylin.

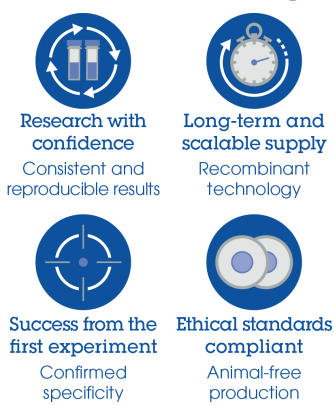


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-P Glycoprotein antibody [EPR10364-57] (ab170904)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human brain tissue labeling P Glycoprotein with purified ab170904 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit IgG was used as the secondary antibody.

Counterstained with hematoxylin.

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Anti-P Glycoprotein antibody [EPR10364-57]
(ab170904)

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