abcam

Product datasheet

Anti-Calreticulin antibody [EPR3924] - ER Marker ab92516



ייבער RabMAb

★★★★ 6 Abreviews 76 References 画像数 18

製品の概要

製品名 Anti-Calreticulin antibody [EPR3924] - ER Marker

製品の詳細 Rabbit monoclonal [EPR3924] to Calreticulin - ER Marker

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P, Flow Cyt (Intra), ICC/IF

種交差性 交差種: Mouse, Rat, Human, African green monkey

交差が予測される動物種: Monkey 🔷

免疫原 Synthetic peptide within Human Calreticulin aa 50-150. The exact sequence is proprietary.

Database link: P27797

(Peptide available as ab180826)

ポジティブ・コントロール WB: SH-SY5Y, HL-60, HepG2, HeLa, Fetal kidney and Fetal brain lysates; Human kidney tissue;

> Mouse and Rat brain lysates. ICC/IF: HAP1 cells (HAP1-CALR as negative cell line) IHC-P: Human colon, kidney, liver, placenta, stomach, breast carcinoma and Papillary carcinoma of

thyroid gland tissues; Mouse liver and Rat lung tissues.

特記事項 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**.

製品の特性

製品の状態

保存方法 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.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

1

supernatant

精製度 Protein A purified

ポリ/モノ モノクローナル

クローン名 EPR3924

アイソタイプ IgG

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ab92516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	**** <u>(1)</u>	1/1000 - 1/10000. Predicted molecular weight: 48 kDa.
IHC-P		1/250 - 1/500. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. The use of a HRP/AP polymerized secondary antibody is recommended for enhanced staining.
Flow Cyt (Intra)		1/10 - 1/100. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
ICC/IF	★★★☆☆(3)	1/500.

ターゲット情報

機能 Molecular calcium-binding chaperone promoting folding, oligomeric assembly and quality control

in the ER via the calreticulin/calnexin cycle. This lectin interacts transiently with almost all of the monoglucosylated glycoproteins that are synthesized in the ER. Interacts with the DNA-binding

domain of NR3C1 and mediates its nuclear export.

配列類似性 Belongs to the calreticulin family.

ドメイン Can be divided into a N-terminal globular domain, a proline-rich P-domain forming an elongated

arm-like structure and a C-terminal acidic domain. The P-domain binds one molecule of calcium with high affinity, whereas the acidic C-domain binds multiple calcium ions with low affinity. The interaction with glycans occurs through a binding site in the globular lectin domain.

The zinc binding sites are localized to the N-domain.

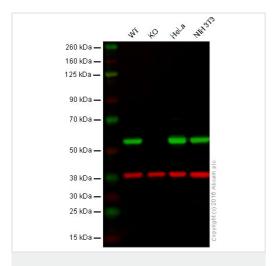
Associates with PDIA3 through the tip of the extended arm formed by the P-domain.

細胞内局在 Endoplasmic reticulum lumen. Cytoplasm > cytosol. Secreted > extracellular space > extracellular

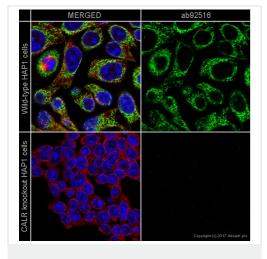
matrix. Cell surface. Also found in cell surface (T cells), cytosol and extracellular matrix.

Associated with the lytic granules in the cytolytic T-lymphocytes.

画像



Western blot - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)



Immunocytochemistry/ Immunofluorescence - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

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

Lane 2: Calreticulin knockout HAP1 cell lysate (20 µg)

Lane 3: HeLa cell lysate (20 µg)

Lane 4: NIH3T3 cell lysate (20 µg)

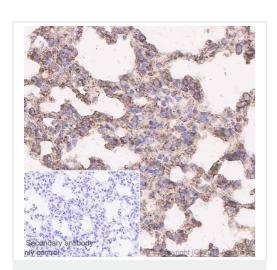
Lanes 1 - 4: Merged signal (red and green). Green - ab92516 observed at 55 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab92516 was shown to specifically react with Calreticulin when Calreticulin knockout samples were used. Wild-type and Calreticulin knockout samples were subjected to SDS-PAGE. ab92516 and ab8245 (loading control to GAPDH) were diluted 1/1000 and 1/10000 respectively and incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.

ab92516 staining Calreticulin in wild-type HAP1 cells (top panel) and CALR knockout HAP1 cells (bottom panel). The cells were fixed with 100% methanol (5min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated with ab92516 at 1/500 and ab195889 at 1/250 dilution (shown in pseudocolour red) overnight at +4°C, followed by a further incubation at room temperature for 1h with a goat secondary antibody to Rabbit lgG (Alexa Fluor[®] 488) (ab150081) at 2 μg/ml (shown in green). Nuclear DNA was labelled in blue with DAPI.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Alexa Fluor[®] 488 (**ab196158**) and Alexa Fluor[®] 647 (**ab196159**) conjugated versions are available for this clone.



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

[EPR3924] - ER Marker (ab92516)

Immunohistochemical analysis of paraffin-embedded Rat lung tissue labeling Calreticulin with ab92516, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Cytoplasmic staining on rat lung. The section was incubated with ab229902 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit IgG H&L (HRP).



Western blot - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

All lanes : Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516) at 1/10000 dilution

Lane 1 : HepG2 (Human hepatocellular carcinoma epithelial cell) Whole cell lysates

Lane 2: HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates

Lane 3: Mouse brain lysates

Lane 4: Rat brain lysates

Lane 5: COS-1 (African green monkey kidney fibroblast-like) whole cell lysates

Lysates/proteins at 15 µg per lane.

Secondary

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

Predicted band size: 48 kDa

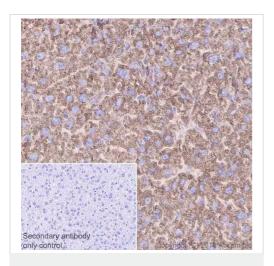
Observed band size: 55 kDa

Blocking/Diluting buffer and concentration: 5% NFDM/TBST Exposure time: Lane 1 to 3: 10 seconds; Lane 4 and 5: 130 seconds

Flow Cytometry (Intracellular) - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

Overlay histogram showing HAP1 wildtype (green line) and HAP1-CALR knockout cells (red line) stained with ab92516. The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab92516, 1µg/ml) for 30 min at 22°C. The secondary antibody used was Alexa Fluorr[®] 488 goat anti-rabbit lgG (H&L) preadsorbed (ab150081) at 1/2000 dilution for 30 min at 22°C. A rabbit lgG isotype control antibody (ab172730) was used at the same concentration and conditions as the primary antibody (HAP1 wildtype - black line, HAP1-CALR knockout - grey line). Unlabelled sample was also used as a control (this line is not shown for the purpose of simplicity). Acquisition of >5,000 events were collected using a 50 mW Blue laser (488nm) and 530/30 bandpass filter.

Alexa Fluorr[®]488 (<u>ab196158</u>) and Alexa Fluorr[®]647 (<u>ab196159</u>) conjugated versions are available for this clone.

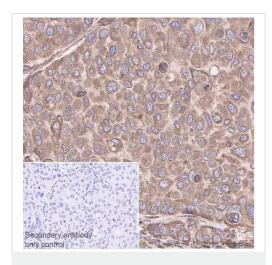


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Immunohistochemical analysis of paraffin-embedded Mouse liver tissue labeling Calreticulin with ab92516, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Cytoplasmic staining on mouse liver. The section was incubated with ab229902 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

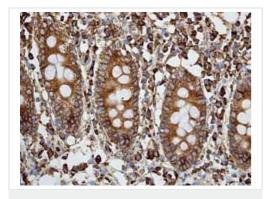


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Immunohistochemical analysis of paraffin-embedded Human breast carcinoma tissue labeling Calreticulin with ab92516, followed by a ready to use Goat Anti-Rabbit IgG H&L (HRP). Cytoplasmic staining human breast carcinoma. The section was incubated with ab229902 for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin. Heat mediated antigen retrieval using ab93684 (Tris/EDTA buffer, pH 9.0).

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is a ready to use Goat Anti-Rabbit lgG H&L (HRP).

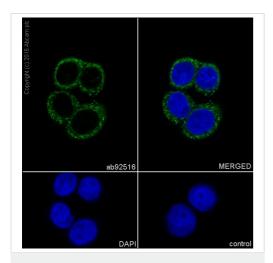


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Formalin-fixed, paraffin-embedded normal human colon tissue stained for Calreticulin using ab92516 at 1/250 dilution in immunohistochemical analysis.

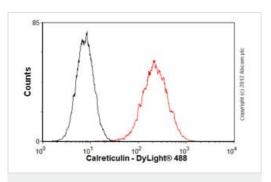
Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

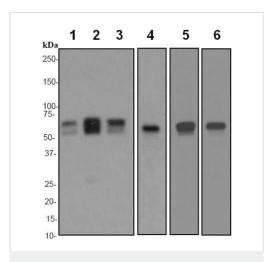
Immunocytochemistry/Immunofluorescence analysis of HT-29 (human colorectal adenocarcinoma) labelling Calreticulin with purified ab92516 at 1/500. Cells were fixed with 100% methanol. An Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/1000) was used as the secondary antibody (Ab150077). Nuclei counterstained with DAPI (blue).

Control: PBS only



Flow Cytometry (Intracellular) - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

Overlay histogram showing HeLa cells stained with ab92516 (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 (ab92516, 1/100 dilution) 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) (1 μ g/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed.



Western blot - Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516)

All lanes : Anti-Calreticulin antibody [EPR3924] - ER Marker (ab92516) at 1/1000 dilution

Lane 1: SH-SY5Y cell lysate

Lane 2: HL-60 cell lysate

Lane 3: HepG2 cell lysate

Lane 4: HeLa cell lysate

Lane 5 : Human fetal kidney lysate

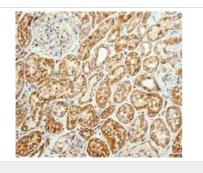
Lane 6: Human fetal brain lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 48 kDa

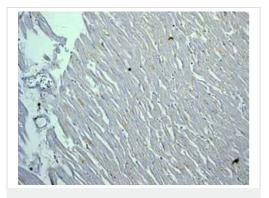


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

ab92516, at 1/250 dilution, staining Calreticulin in paraffin embedded Human kidney tissue by Immunohistochemistry.

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

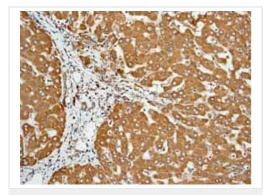


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

ab92516 showing negative staining in Normal human heart tissue.

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

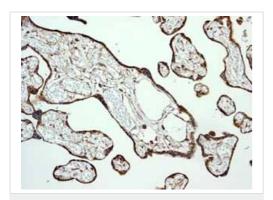


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Formalin-fixed, paraffin-embedded normal human liver tissue stained for Calreticulin using ab92516 at 1/250 dilution in immunohistochemical analysis.

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

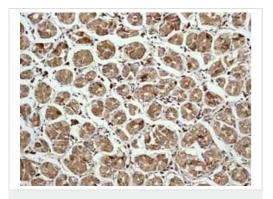


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Formalin-fixed, paraffin-embedded normal human placenta tissue stained for Calreticulin using ab92516 at 1/250 dilution in immunohistochemical analysis.

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

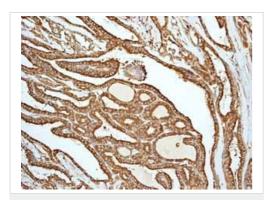


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Formalin-fixed, paraffin-embedded normal human stomach tissue stained for Calreticulin using ab92516 at 1/250 dilution in immunohistochemical analysis.

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

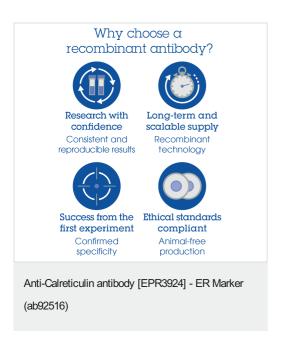


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Calreticulin antibody

[EPR3924] - ER Marker (ab92516)

Formalin-fixed, paraffin-embedded Papillary carcinoma of human thyroid gland tissue.stained for Calreticulin using ab92516 at 1/250 dilution in immunohistochemical analysis.

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



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