abcam

Product datasheet

Anti-GRP78 BiP antibody [EPR4041(2)] ab108615

יובעבלי RabMAb

★★★★★ 3 Abreviews 60 References 画像数 11

製品の概要

製品名 Anti-GRP78 BiP antibody [EPR4041(2)]

Rabbit monoclonal [EPR4041(2)] to GRP78 BiP 製品の詳細

由来種 Rabbit

特異性 Stimulation may be required to allow detection of the target protein due to low levels of

endogenous expression in some samples. Please see images below for recommended

treatment conditions and positive controls.

The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for

mouse and rat.

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

適用なし: ICC/IF

種交差性 交差種: Mouse, Rat, Human

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

ポジティブ・コントロール WB: HeLa, NIH/3T3, LnCaP, MCF7, 293T, U-87 MG, THP-1, MDA-MB-231 and HepG2 whole

> cell lysate (ab7900); Human liver lysates, Mouse heart, lung, liver and testis lysates; Rat liver lysates. Huvec and RAW 264.7 cells treated with tunicamycin. IHC-P: Human breast carcinoma

Tissue Flow Cyt (intra): HeLa cells

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 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.

Stable for 12 months at -20°C.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol, 0.05% BSA, PBS

精製度 Protein A purified

ポリ/モノ モノクローナル **ウローン名** EPR4041(2)

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ab108615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
Flow Cyt (Intra)		1/200.
WB	**** (1)	1/1000 - 1/10000. Predicted molecular weight: 72 kDa. Stimulation may be required to allow detection of the target protein due to low levels of endogenous expression in some samples. Please see images for recommended treatment conditions and positive controls.
IHC-P	****(1)	1/100 - 1/500. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. See IHC antigen retrieval protocols. For unpurified, use 1/50 - 1/100. The mouse and rat recommendation is based on the WB results. We do not guarantee IHC-P for mouse and rat.

追加情報 Is unsuitable for ICC/IF.

ターゲット情報

機能 Probably plays a role in facilitating the assembly of multimeric protein complexes inside the endoplasmic reticulum. Involved in the correct folding of proteins and degradation of misfolded proteins via its interaction with DNAJC10, probably to facilitate the release of DNAJC10 from its substrate.

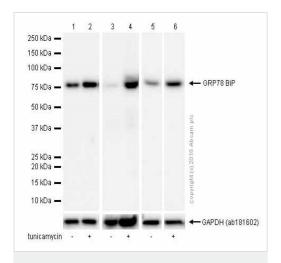
関連疾患 Autoantigen in rheumatoid arthritis.

配列類似性 Belongs to the heat shock protein 70 family.

細胞内局在 Endoplasmic reticulum lumen. Melanosome. Cytoplasm. Identified by mass spectrometry in

melanosome fractions from stage I to stage IV.

画像



Western blot - Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615)

All lanes : Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution (purified)

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates

Lane 2 : HeLa (Human cervix adenocarcinoma epithelial cell) treated with 10ug/ml

tunicamycin for 12 hours whole cell lysates

Lane 3 : MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysates

Lane 4 : MCF-7 (Human breast adenocarcinoma epithelial cell) treated with 10ug/ml

tunicamycin for 48 hours whole cell lysates

Lane 5 : MEF (Mouse embryonic fibroblast) whole cell lysatesLane 6 : MEF (Mouse embryonic fibroblast) treated with 10ug/ml tunicamycin for 48 hours 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

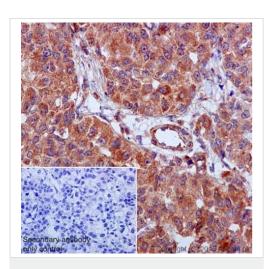
Predicted band size: 72 kDa

Blocking and diluting buffer: 5% NFDM/TBST.

Tunicamycin induces the expression of GRP78 BiP reported by PMID 22859938

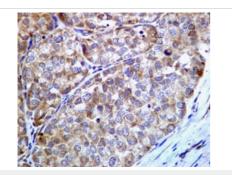
Exposure time:

Lanes 1 and 2: 20 seconds Lanes 3 and 4: 30 seconds Lanes 5 and 6: 5 seconds



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GRP78 BiP antibody
[EPR4041(2)] (ab108615)

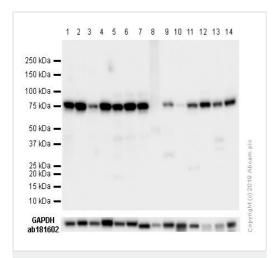
Immunohistochemical staining of paraffin embedded human hepatocellular carcinoma with purified ab108615 at a working dilution of 1/500. The secondary antibody used is HRP goat antirabbit lgG H&L (ab97051) at 1/500. The sample is counter-stained with hematoxylin. Antigen retrieval was perfomed using Tris-EDTA buffer, pH 9.0. PBS was used instead of the primary antibody as the negative control, and is shown in the inset.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-GRP78 BiP antibody
[EPR4041(2)] (ab108615)

Unpurified ab108615 at 1/50 dilution staining GRP78 BiP in Human breast carcinoma tissue by Immunohistochemistry Formalin-fixed, Paraffin-embedded tissue.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Western blot - Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615)

All lanes : Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution

Lane 1 : MCF7 (Human breast adenocarcinoma epithelial cell) whole cell lysates

Lane 2 : HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysates

Lane 3: 293T (Human embryonic kidney epithelial cell) whole cell lysates

Lane 4 : U-87 MG (Human glioblastoma-astrocytoma epithelial cell) whole cell lysates

Lane 5 : THP-1 (Human monocytic leukemia monocyte) whole cell lysates

Lane 6 : MDA-MB-231 (Human breast adenocarcinoma epithelial cell) whole cell lysates

Lane 7: NIH/3T3 (Mouse embryonic fibroblast) whole cell lysates

Lane 8 : Human uterus lysates
Lane 9 : Human liver lysates
Lane 10 : Mouse heart lysates

Lane 11 : Mouse lung lysatesLane 12 : Mouse testis lysatesLane 13 : Mouse liver lysates

Lane 14: Rat liver lysates

Lysates/proteins at 20 µg per lane.

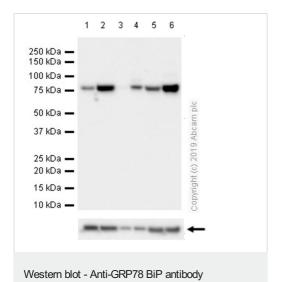
Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution (Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugated)

Predicted band size: 72 kDa **Observed band size:** 78 kDa

Exposure time: 26 seconds

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



[EPR4041(2)] (ab108615)

All lanes : Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysates

Lane 2 : HeLa treated with 2.5 μ g/ml tunicamycin for 24h whole cell lysates

Lane 3: HUVEC (Human umbilical vein endothelial cell) whole cell lysates

Lane 4 : HUVEC treated with 10 μ g/ml tunicamycin for 48h whole cell lysates

Lane 5 : Raw 264.7 (Mouse Abelson murine leukemia virusinduced tumor macrophage) whole cell lysates

Lane 6 : Raw 264.7 treated with 5 μ g/ml tunicamycin for 18h whole cell lysates

Lysates/proteins at 20 µg per lane.

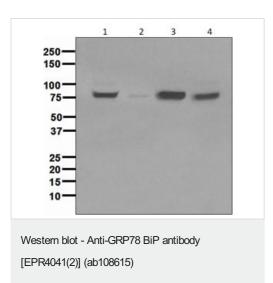
Secondary

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

Predicted band size: 72 kDa **Observed band size:** 78 kDa

Exposure time: 3 seconds

Blocking and diluting buffer and concentration: 5% NFDM/TBST



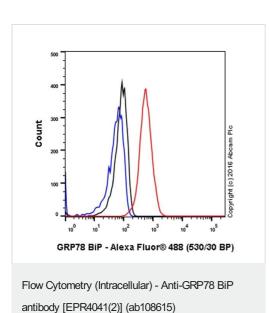
All lanes : Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution (unpurified)

Lane 1: HeLa cell lysates
Lane 2: NIH/3T3 cell lysates
Lane 3: LnCaP cell lysates
Lane 4: HepG2 cell lysates

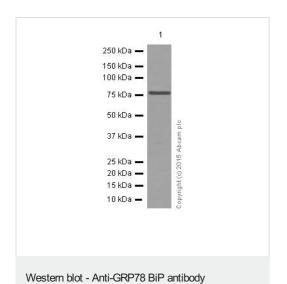
Secondary

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

Predicted band size: 72 kDa



Intracellular Flow Cytometry analysis of HeLa (human cervix adenocarcinoma) cells labeling GRP78 BiP with purified ab108615 at 1/200 dilution (10ug/mL) (red). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. A Goat anti rabbit IgG (Alexa Fluorr[®] 488) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal IgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) were used as the unlabeled control.



[EPR4041(2)] (ab108615)

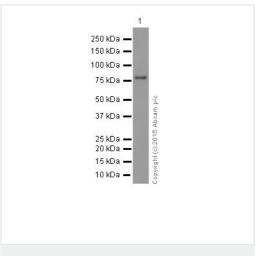
Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution (purified) + Raw264.7 cell lysate at 20 µg

Secondary

HRP goat anti-rabbit lgG (H+L) at 1/50000 dilution

Predicted band size: 72 kDa **Observed band size:** 78 kDa

Blocking buffer: 5% NFDM/TBST Dilution buffer: 5% NFDM/TBST



Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/5000 dilution (purified) + HepG2 at 20 μg

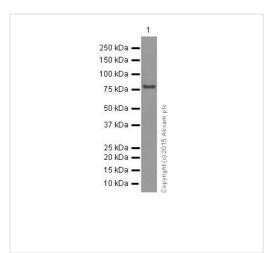
Secondary

HRP goat anti-rabbit lgG (H+L) at 1/50000 dilution

Predicted band size: 72 kDa **Observed band size:** 78 kDa

Western blot - Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615)

Blocking buffer: 5% NFDM/TBST Dilution buffer: 5% NFDM/TBST



Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615) at 1/1000 dilution (purified) + LnCaP at 20 μg

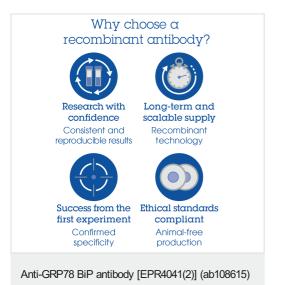
Secondary

HRP goat anti-rabbit lgG (H+L) at 1/50000 dilution

Predicted band size: 72 kDa **Observed band size:** 78 kDa

Western blot - Anti-GRP78 BiP antibody [EPR4041(2)] (ab108615)

Blocking buffer: 5% NFDM/TBST Dilution buffer: 5% NFDM/TBST



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