abcam

Product datasheet

Anti-ERp57 antibody [MaP.Erp57] ab13506



★★★★★ 7 Abreviews 36 References 画像数 6

製品の概要

製品名 Anti-ERp57 antibody [MaP.Erp57]

製品の詳細 Mouse monoclonal [MaP.Erp57] to ERp57

由来種 Mouse

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

種交差性 交差種: Mouse, Human

免疫原 Recombinant full length protein (Human).

ポジティブ・コントロール WB: HAP1, HepG2 and HEK-293 whole cell lysate; mouse pancreatic cancer cell lysate. ICC/IF:

HeLa and HaCaT cells. IHC-P: Muse backskin tissue. IHC-Fr: Mouse duodenum, pancreas, and

pancreatic neoplastic tissue. Flow Cyt: HeLa cells.

特記事項

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

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

ארע"א Preservative: 0.09% Sodium azide

Constituents: PBS, 50% Glycerol

精製度 Protein G purified

特記事項(精製) This antibody is affinity purified.

ポリ/モノ モノクローナル **クローン名** MaP.Erp57

アイソタイプ IgG1

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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ab13506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
IHC-P		1/100.
WB	★★★★★ (3)	1/2000. Predicted molecular weight: 61 kDa. This concentration is sufficient for the detection of ERp57 in 20μg of HeLa heat shocked cell lysate by colorimetric analysis.
ICC/IF		1/100.
Flow Cyt		Use 1-2µg for 10 ⁶ cells. ab170190 - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody.

ターゲット情報

組織特異性 Detected in the flagellum and head region of spermatozoa (at protein level).

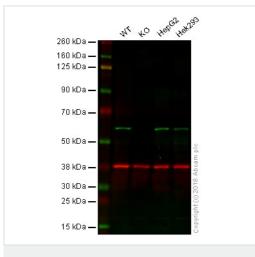
配列類似性 Belongs to the protein disulfide isomerase family.

Contains 2 thioredoxin domains.

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

fractions from stage I to stage IV.

画像



Western blot - Anti-ERp57 antibody [MaP.Erp57] (ab13506)

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

Lane 2: ERp57 knockout HAP1 whole cell lysate (20 µg)

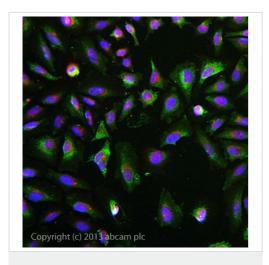
Lane 3: HepG2 whole cell lysate (20 µg)

Lane 4: HEK293 whole cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - ab13506 observed at 57 kDa. Red - loading control, <u>ab181602</u>, observed at 37 kDa.

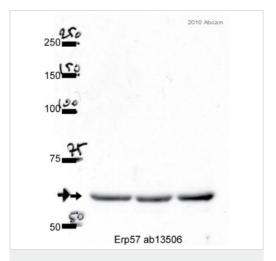
ab13506 was shown to specifically react with ERp57 in wild-type HAP1 cells as signal was lost in ERp57 knockout cells. Wild-type and ERp57 knockout samples were subjected to SDS-PAGE. ab13506 and ab181602 (Rabbit anti-GAPDH loading control) were incubated overnight at 4°C at 0.5 µg/ml and 1/20000 dilution respectively. Blots were developed with Goat anti-Mouse lgG H&L (IRDye® 800CW) preabsorbed ab216772 and Goat anti-Rabbit lgG H&L (IRDye® 680RD) preabsorbed ab216777 secondary antibodies at 1/20000 dilution for 1 hour at room temperature

before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-ERp57 antibody [MaP.Erp57] (ab13506)

ICC/IF image of ab13506 stained HeLa cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody ab13506 at 5μg/ml overnight at +4°C. The secondary antibody (green) was DyLight® 488 goat anti- mouse (ab96879) lgG (H+L) used at a 1/250 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43μM.



Western blot - Anti-ERp57 antibody [MaP.Erp57] (ab13506)

This image is courtesy of an Abreview submitted by Pawel Mazur

ERp57 antibody (ab13506) at 1/1000 dilution (in PBST) for 8 hours at 4° C + lysate of mouse of pancreatic cancer cell line at 25 μ g.

Secondary

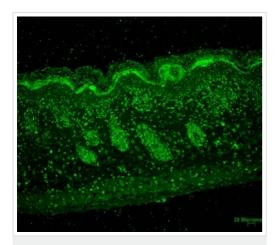
An HRP conjugated Sheep anti-mouse IgG polyclonal at 1/5000 dilution

developed using the ECL technique

Performed under non-reducing conditions.

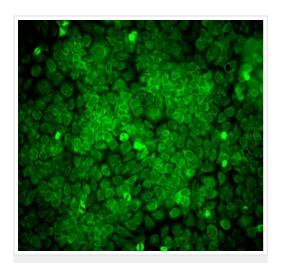
Exposure time: 1 minute

Blocking Step: 10% Milk for 1 hour at room temperature.



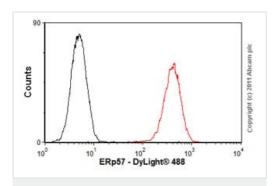
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ERp57 antibody
[MaP.Erp57] (ab13506)

Paraffin-embedded mouse backskin tissue ficed with Bouin's fixative, stained for ERp57 using ab13506 at 1/100 dilution in ICC/IF. Primary antibody was incubated for 1 hour at room temperature. Secondary antibody is a FITC-conjugated goat antimouse (green) incubated at 1/50 dilution for 1 hour at room temperature.



Immunocytochemistry/ Immunofluorescence - Anti-ERp57 antibody [MaP.Erp57] (ab13506)

HaCaT (Human keratinocyte cell line) cells labeling ERp57 using ab13506 at 1/100 dilution in ICC.IF. Cells were fixed using 100% methanol for 10 minutes at -20°C and incubated with primary antibody for 1 hour at room temperature. Secondary antibody is a FITC-conjugated goat anti-mouse (green) incubated at 1/50 dilution for 1 hour at room temperature.



Flow Cytometry - Anti-ERp57 antibody [MaP.Erp57] (ab13506)

Overlay histogram showing HeLa cells stained with ab13506 (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. The cells were then incubated with the antibody (ab13506, 2µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse lgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (ab91353, 2µg/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive signal in HeLa cells fixed with 4% paraformaldehyde (10 min)/permeabilized in 0.1% PBS-Tween used under the same conditions.

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