abcam

Product datasheet

Anti-Peroxiredoxin 3/PRDX3 antibody ab73349



★★★★★ 7 Abreviews 35 References 画像数 4

製品の概要

製品名 Anti-Peroxiredoxin 3/PRDX3 antibody

製品の詳細 Rabbit polyclonal to Peroxiredoxin 3/PRDX3

由来種 Rabbit

特異性 Replenishment batches of our polyclonal antibody, ab73349 are tested in WB. Previous batches

were additionally validated in ICC/IF and IHC-P. These applications are still expected to work and

are covered by our Abpromise guarantee. You may also be interested in our alternative

recombinant antibody, ab129206.

アプリケーション 適用あり: IHC-P, ICC/IF, WB

種交差性 交差種: Mouse, Rat, Human

交差が予測される動物種: Chicken, Cow, Xenopus laevis, Chimpanzee, Zebrafish, Orangutan

A

免疫原 Synthetic peptide corresponding to Human Peroxiredoxin 3/PRDX3 aa 200 to the C-terminus

conjugated to keyhole limpet haemocyanin.

(Peptide available as ab85815)

ポジティブ・コントロール Recombinant human Peroxiredoxin 3/PRDX3 protein (ab85559) can be used as a positive

control in WB. This antibody gave a positive signal in the following lysates: Human, Mouse and

Rat Liver, Rat Ovary. ICC/IF: 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. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

1

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

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ab73349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	* * * * * <u>(2)</u>	Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
ICC/IF		Use a concentration of 5 µg/ml.
WB	★★★★ (4)	Use a concentration of 1 µg/ml. Detects a band of approximately 23 kDa (predicted molecular weight: 28 kDa).

ターゲット情報

機能 Involved in redox regulation of the cell. Protects radical-sensitive enzymes from oxidative damage

by a radical-generating system. Acts synergistically with MAP3K13 to regulate the activation of

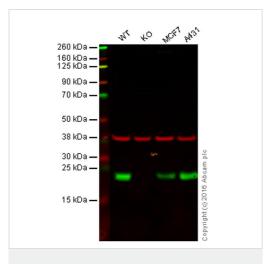
NF-kappa-B in the cytosol.

配列類似性 Belongs to the ahpC/TSA family.

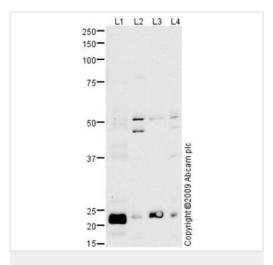
Contains 1 thioredoxin domain.

細胞内局在 Mitochondrion.

画像



Western blot - Anti-Peroxiredoxin 3/PRDX3 antibody (ab73349)



Western blot - Anti-Peroxiredoxin 3/PRDX3 antibody (ab73349)

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

Lane 2: Peroxiredoxin 3/PRDX3 knockout HAP1 cell lysate (20 µg)

Lane 3: MCF7 cell lysate (20 µg)

Lane 4: A431 cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - ab73349 observed at 24 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab73349 was shown to specifically react with Peroxiredoxin 3/PRDX3 in wild-type HAP1 cells. No band was observed when Peroxiredoxin 3/PRDX3 knockout samples were examined. Wild-type and Peroxiredoxin 3/PRDX3 knockout samples were subjected to SDS-PAGE. ab73349 and ab8245 (loading control to GAPDH) were diluted at 1µg/ml and 1/10,000 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 1/10,000 dilution for 1 hour at room temperature before imaging.

All lanes : Anti-Peroxiredoxin 3/PRDX3 antibody (ab73349) at 1 μ g/ml

Lane 1: Human liver tissue lysate - total protein (ab29889)

Lane 2: Liver (Rat) Tissue Lysate

Lane 3: Ovary (Rat) Tissue Lysate

Lane 4: Liver (Mouse) Tissue Lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat polyclonal to Rabbit lgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

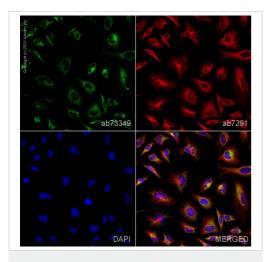
Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 28 kDa **Observed band size:** 23 kDa

Additional bands at: 45 kDa, 52 kDa. We are unsure as to the

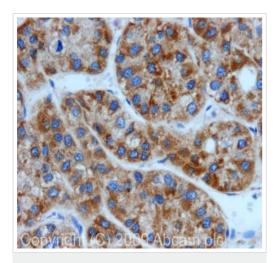
identity of these extra bands.



Immunocytochemistry/ Immunofluorescence - Anti-Peroxiredoxin 3/PRDX3 antibody (ab73349)

ab73349 staining Peroxiredoxin 3/PRDX3 in HeLa cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-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 overnight at 4°C with ab73349 at 5µg/ml and ab7291, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with ab150081, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor[®] 488), pre-adsorbed at 1/1000 dilution (shown in green) and ab150120, Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor[®] 594), pre-adsorbed at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal sections is shown.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Peroxiredoxin 3/PRDX3 antibody (ab73349)

IHC image of Peroxiredoxin 3/PRDX3 staining in human liver carcinoma formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab73349, 5µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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