abcam

Product datasheet

Anti-ALDH1A1 antibody [EP1933Y] - C-terminal ab52492





重组 RabMAb

★★★★★ 22 Abreviews 170 References 14 图像

概述

产品名称 Anti-ALDH1A1抗体[EP1933Y] - C-terminal

描述 兔单克隆抗体[EP1933Y] to ALDH1A1 - C-terminal

宿主 Rabbit

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

经测试应用 适用于: Flow Cyt (Intra), ICC/IF, WB, IP, IHC-P

种属反应性 与反应: Mouse, Human

免疫原 Synthetic peptide within Human ALDH1A1 aa 300-400 (C terminal). The exact sequence is

proprietary.

Database link: P00352

阳性对照 WB: HepG2 and NIH/3T3 cell lysate; Human small intestine and mouse liver tissue lysate. IHC:

Human liver, kidney, lung and brain tissue, human breast carcinoma. ICC/IF: HepG2 cells. Flow

Cyt (intra): HepG2 cells. IP: HepG2 cell lysate

常规说明 The mouse recommendation is based on the WB results. We do not guarantee IHC-P for

mouse.

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**.

Rat: We have preliminary internal testing data to indicate this antibody may not react with this

species. Please contact us for more information.

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 pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol, 0.05% BSA

纯**度** Protein A purified

同种型 lgG

应用

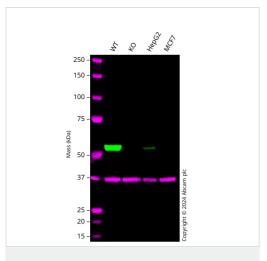
The Abpromise guarantee Abpromise™承诺保证使用ab52492于以下的经测试应用

"应用说明"<mark>部分 下显示的</mark>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 |
|------------------|------------------|---|
| Flow Cyt (Intra) | | 1/20. For unpurified use at 1/1000. <u>ab172730</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody. |
| ICC/IF | ★★★★★ (3) | 1/100 - 1/250. |
| WB | ★★★★ | 1/500 - 1/2000. Detects a band of approximately 55 kDa (predicted molecular weight: 55 kDa). |
| IP | | 1/20 - 1/30. |
| IHC-P | ★★★★☆ (5) | 1/50. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. See IHC antigen retrieval protocols . The mouse recommendation is based on the WB results. We do |
| | | not guarantee IHC-P for mouse. |

| 靶 标 | | |
|------------|--|--|
| 功能 | Binds free retinal and cellular retinol-binding protein-bound retinal. Can convert/oxidize retinaldehyde to retinoic acid. | |
| 通路 | Cofactor metabolism; retinol metabolism. | |
| 序列相似性 | Belongs to the aldehyde dehydrogenase family. | |
| 细胞定位 | Cytoplasm. | |

图片



Western blot - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

All lanes : Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492) at 1/1000 dilution

Lane 1: Wild-type A549 cell lysate

Lane 2: ALDH1A1 knockout A549 cell lysate

Lane 3 : HepG2 cell lysate

Lane 4 : MCF7 cell lysate

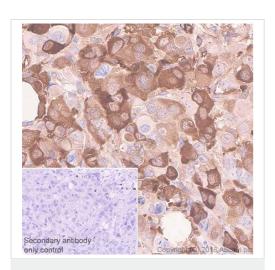
Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat anti-Rabbit lgG H&L 800CW and Goat anti-Mouse lgG H&L 680RD at 1/20000 dilution

Predicted band size: 55 kDa **Observed band size:** 55 kDa

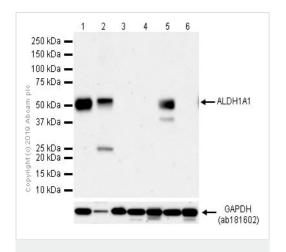
Western blot: Anti-ALDH1A1 antibody [EP1933Y] (ab52492) staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in magenta. In Western blot, ab52492 was shown to bind specifically to ALDH1A1. A band was observed at 55 kDa in wild-type A549 cell lysates with no signal observed at this size in ALDH1A1 knockout cell line. To generate this image, wild-type and ALDH1A1 knockout A549 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L 800CW and Goat anti-Mouse IgG H&L 680RD at 1/20000 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody
[EP1933Y] - C-terminal (ab52492)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast carcinoma tissue sections labeling ALDH1A1 with purified ab52492 at 1/50 dilution (3.54 µg/ml). Perform heat mediated antigen retrieval using **ab93684** (Tris/EDTA buffer, pH 9.0). ImmunoHistoProbe one step HRP Polymer (ready to use) was used as the secondary antibody.

Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.



Western blot - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

All lanes : Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492) at 1/1000 dilution (Purified)

Lane 1 : HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysates at 20 μg

Lane 2: Human small intestine lysates

Lane 3 : PC-3 (Human prostate adenocarcinoma epithelial cell) whole cell lysates at 20 µg

Lane 4 : MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysates at 20 μg

Lane 5: Mouse liver lysates at 20 µg

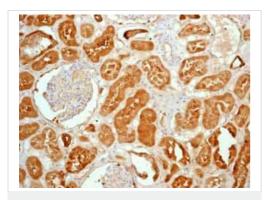
Lane 6 : NIH/3T3 (Mouse embryonic fibroblast) whole cell lysates at $20 \ \mu g$

Secondary

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

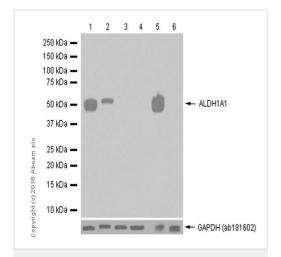
Predicted band size: 55 kDa

Observed band size: 55 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody
[EP1933Y] - C-terminal (ab52492)

Immunohistochemical analysis of Formalin/PFA-fixed paraffinembedded normal human kidney tissue sections labeling ALDH1A1 with ab52492 (unpurified).



Western blot - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

All lanes : Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492) at 1/1000 dilution (unpurified)

Lane 1 : HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysate

Lane 2: Human small intestine tissue lysate

Lane 3 : PC-3 (Human prostate adenocarcinoma epithelial cell) whole cell lysate

Lane 4 : MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate

Lane 5: Mouse liver tissue lysate

Lane 6: NIH/3T3 (Mouse embryonic fibroblast) whole cell lysate

Lysates/proteins at 15 µg per lane.

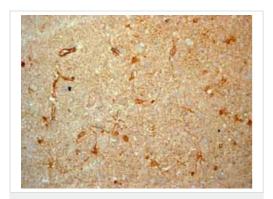
Secondary

All lanes : Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG at 1/2000 dilution

Predicted band size: 55 kDa
Observed band size: 55 kDa

Exposure time: 1 minute

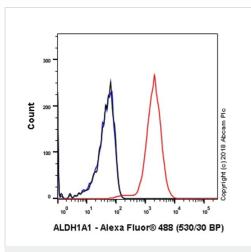
Blocking/Diluting buffer 5% NFDM/TBST



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody

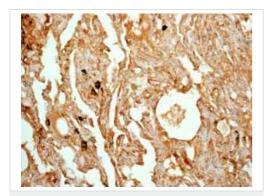
[EP1933Y] - C-terminal (ab52492)

Immunohistochemical analysis of Formalin/PFA-fixed paraffinembedded normal human brain tissue sections labeling ALDH1A1 with ab52492 (unpurified).



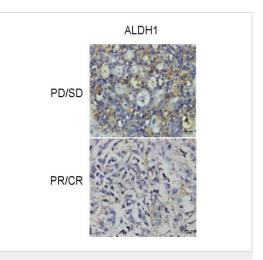
Flow Cytometry (Intracellular) - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

Intracellular Flow Cytometry analysis of HepG2 (Human hepatocellular carcinoma epithelial cell) cells labeling ALDH1A1 with purified ab52492 at 1/20 dilution (10µg/ml) (red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit IgG (Alexa Fluorr® 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody
[EP1933Y] - C-terminal (ab52492)

Immunohistochemical analysis of Formalin/PFA-fixed paraffinembedded normal human lung tissue sections labeling ALDH1A1 with ab52492 (unpurified).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody
[EP1933Y] - C-terminal (ab52492)

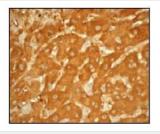
Gong et al PLoS One. 2010 Dec 20;5(12):e15630. doi: 10.1371/journal.pone.0015630. Fig 1. Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/

Tumor tissues of primary invasive ductal carcinomas of the breast were obtained from 192 female patients with stage IIB and III prior to pre-operative neoadjuvant chemotherapy.

(progressive or stable disease, PD/SD)

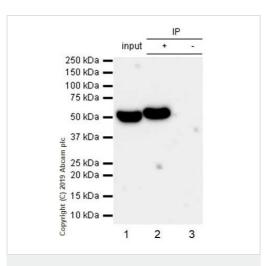
(partial or complete remission, PR/CR)

The level of ALDH1 was tested by immunohistochemistry staining in paraffin-embedded tissue sections. Rabbit monoclonal ALDH1A1 antibody (unpurified ab52492, Abcam) used at a 1:100 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ALDH1A1 antibody
[EP1933Y] - C-terminal (ab52492)

Immunohistochemical analysis of paraffin-embedded human liver tissue sections labeling ALDH1A1 with ab52492 (unpurified) at 1/100 dilution.



Immunoprecipitation - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492) ab52492 (purified) at 1/20 dilution (2µg) immunoprecipitating ALDH1A1 in HepG2 (Human hepatocellular carcinoma epithelial cell) whole cell lysates.

Lane 1: HepG2 whole cell lysates 10µg.

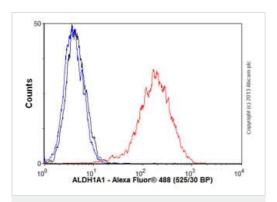
Lane 2: ab52492 IP in HepG2 whole cell lysates.

 $\label{lambda} \textbf{Lane 3:} \ \ \text{Rabbit monoclonal lgG } (\underline{\textbf{ab172730}}) \ \ \text{instead of ab52492 in} \\ \ \ \text{HepG2 whole cell lysates}.$

For western blotting, VeriBlot for IP Detection Reagent (HRP)

(ab131366) was used at 1/1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.

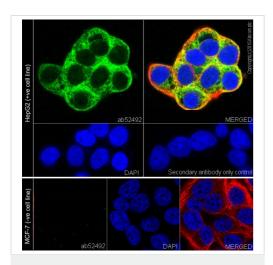


Flow Cytometry (Intracellular) - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

Overlay histogram showing HepG2 (Human liver hepatocellular carcinoma cell line) cells stained with ab52492 (unpurified) (red line).

The cells were fixed with 80% methanol (5 minutes) and then permeabilized with 0.1% PBS-Tween for 20 minutes. 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 (ab52492, 1/1000 dilution) for 30 minutes at 22°C. The secondary antibody used was Alexa Fluorr® 488 goat anti-rabbit lgG (H&L) (ab150077) at 1/2000 dilution for 30 minutes at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (0.1 μ g/1x10⁶ cells) used under the same conditions. Unlabeled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



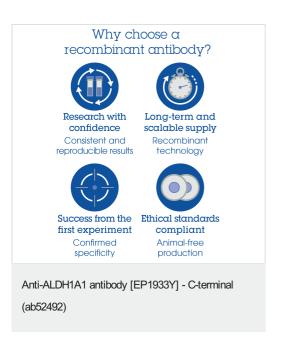
Immunocytochemistry/ Immunofluorescence - Anti-ALDH1A1 antibody [EP1933Y] - C-terminal (ab52492)

Immunocytochemistry/ Immunofluorescence analysis of HepG2 (Human liver hepatocellular carcinoma cell line) cells labeling ALDH1A1 with ab52492 (purified) at 1/500 dilution (4 µg/ml).

Cells were fixed in 100% methanol. ab150077, an AlexaFluor[®]488 Goat anti-Rabbit secondary antibody was used at 1/1000 dilution (2 µg/ml). ab195889, Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor[®] 594) was used to counterstain at 1/200 dilution (2.5 µg/ml). DAPI was used as nuclear counterstain.

Confocal image showing cytoplasmic staining on HepG2 cell line.

Negative control: No staining on MCF-7 cell line.



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