abcam

Product datasheet

Human PRKCD (PKC delta) knockout HeLa cell lysate ab257043

画像数 2

製品の概要

製品名 Human PRKCD (PKC delta) knockout HeLa cell lysate

製品の概要

Knockout cell lysate achieved by CRISPR/Cas9.

Parental Cell Line HeLa
Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, 2 bp deletion in exon5.

Passage number <20

Knockout validation Sanger Sequencing, Western Blot (WB)

 $\label{eq:Reconstitution notes} \textbf{To use as WB control, resuspend the lyophilizate in 50 μL of LDS* Sample Buffer to have a final μL of LDS* Sample$

concentration of 2 mg/ml. For reducing conditions, we recommend a final concentration of 0.1 M

DTT.

 $\hbox{*Usage of SDS sample buffer is not recommended with these lyophilized lysates.}$

特記事項

Lysate preparation: Our lysates are made using RIPA buffer to which we add a protease inhibitor cocktail and phosphatase inhibitor cocktail (ratio: 300:100:10). *This means that the protein of interest is denatured.* If you require a native form of the protein please use the live cell version - found **here**. Please refer to our lysis protocol for further details on how our lysates are prepared.

User storage instructions: Lyophilizate may be stored at 4°C. After reconstitution, store at -20°C for short-term storage or -80°C for long-term storage.

Access thousands of knockout cell lysates, generated from commonly used cancer cell lines. See here for more information on knockout cell lysates.

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アプリケーション **適用あり**: WB

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製品の特性

保存方法

Store at -80°C. Please refer to protocols.

内容	1 kit
ab261984 - Human PRKCD knockout HeLa cell lysate	1 x 100μg
ab255929 - Human wild-type HeLa cell lysate	1 x 100μg

Cell type epithelial

Disease Adenocarcinoma

Gender Female

STR Analysis Amelogenin X D5S818: 11, 12 D13S317: 12, 13.3 D7S820: 8, 12 D16S539: 9, 10 vWA: 16, 18

TH01: 7 TPOX: 8,12 CSF1PO: 9, 10

ターゲット情報

機能

This is calcium-independent, phospholipid-dependent, serine- and threonine-specific enzyme. PKC is activated by diacylglycerol which in turn phosphorylates a range of cellular proteins. PKC also serves as the receptor for phorbol esters, a class of tumor promoters. May play a role in antigen-dependent control of B-cell function. Phosphorylates MUC1 in the C-terminal and regulates the interaction between MUC1 and beta-catenin.

配列類似性

Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. PKC subfamily.

Contains 1 AGC-kinase C-terminal domain.

Contains 1 C2 domain.

Contains 2 phorbol-ester/DAG-type zinc fingers.

Contains 1 protein kinase domain.

ドメイン

The C1 domain, containing the phorbol ester/DAG-type region 1 (C1A) and 2 (C1B), is the

diacylglycerol sensor.

The C2 domain is a non-calcium binding domain. It binds proteins containing phosphotyrosine in

a sequence-specific manner.

翻訳後修飾

Phosphorylated on Thr-507, within the activation loop. Autophosphorylated and/or phosphorylated.

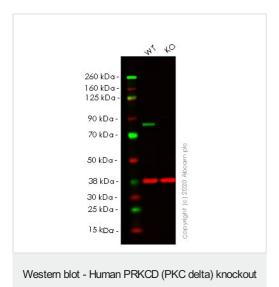
Although the Thr-507 phosphorylation occurs it is not a prerequisite for enzymatic activity.

細胞内局在 Cytoplasm. Membrane.

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ab257043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		Use at an assay dependent concentration. Predicted molecular weight: 78 kDa.



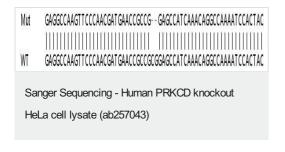
HeLa cell lysate (ab257043)

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

Lane 2: PRKCD knockout HeLa cell lysate (20µg)

Lanes 1-2: Merged signal (red and green). Green - <u>ab182126</u> observed at 80 kDa. Red - loading control <u>ab8245</u> observed at 37 kDa.

ab182126 Anti-PKC delta antibody [EPR17075] was shown to specifically react with PKC delta in wild-type HeLa cells in western blot. Loss of signal was observed when knockout cell line ab265721 (knockout cell lysate ab257043) was used. Wild-type and PKC delta knockout samples were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab182126 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 5000 dilution and 1 in 20000 dilution respectively. 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 in 20000 dilution for 1 hour at room temperature before imaging.



Homozygous: 2 bp deletion in exon5

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