abcam

Product datasheet

Human CDC42 knockout HEK-293T cell lysate ab256868

3 图像

概述

产品概述

Knockout cell lysate achieved by CRISPR/Cas9.

Parental Cell Line HEK293T

Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, Homozygous: 2 bp insertion in exon 3.

Passage number <20

Knockout validation Sanger Sequencing, Western Blot (WB)

Reconstitution notesTo use as WB control, resuspend the lyophilizate in 50 μL of LDS* Sample Buffer to have a final

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

DTT.

*Usage of SDS sample buffer is not recommended with these lyophilized lysates.

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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经测试应用 适用于: WB

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存放说明

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

组件	1 kit
ab262823 - Human CDC42 knockout HEK293T cell lysate	1 x 100µg
ab255553 - Human wild-type HEK293T cell lysate	1 x 100µg

Cell type epithelial

STR Analysis Amelogenin X D5S818: 8, 9 D13S317: 12, 14 D7S820: 11 D16S539: 9, 13 vWA: 16, 19 TH01:

7, 9.3 TPOX: 11 CSF1PO: 11, 12

靶标

功能 Plasma membrane-associated small GTPase which cycles between an active GTP-bound and an

inactive GDP-bound state. In active state binds to a variety of effector proteins to regulate cellular responses. Involved in epithelial cell polarization processes. Causes the formation of thin, actin-

rich surface projections called filopodia.

序列相似性 Belongs to the small GTPase superfamily. Rho family. CDC42 subfamily.

翻译后修饰 AMPylation at Tyr-32 and Thr-35 are mediated by bacterial enzymes in case of infection by

H.somnus and V.parahaemolyticus, respectively. AMPylation occurs in the effector region and leads to inactivation of the GTPase activity by preventing the interaction with downstream effectors, thereby inhibiting actin assembly in infected cells. It is unclear whether some human enzyme mediates AMPylation; FICD has such ability in vitro but additional experiments remain to

be done to confirm results in vivo.

细**胞定位** Cell membrane.

形式 There are 2 isoforms produced by alternative splicing. Isoform 1 also known as: Brain; Isoform 2

also known as: Placental.

应用

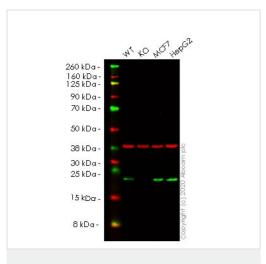
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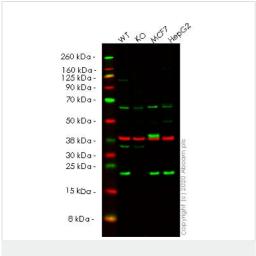
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应 用	Ab评论	说明
WB		Use at an assay dependent concentration. Predicted molecular weight: 21 kDa.

图片



Western blot - Human CDC42 knockout HEK293T cell lysate (ab256868)



Western blot - Human CDC42 knockout HEK293T cell lysate (ab256868)

Lane 1: Wild-type HEK-293T cell lysate (20 µg)

Lane 2: CDC42 knockout HEK-293T cell lysate (20 µg)

Lane 3: MCF7 cell lysate (20 µg)

Lane 4: HepG2 cell lysate (20 µg)

Lanes 1-4: Merged signal (red and green). Green - <u>ab187643</u> observed at 20 kDa. Red - loading control <u>ab8245</u> observed at 37 kDa.

ab187643 Anti-CDC42 antibody [EPR15620] was shown to specifically react with CDC42 in wild-type HEK-293T cells. Loss of signal was observed when knockout cell line ab266522 (knockout cell lysate ab256868) was used. Wild-type and CDC42 knockout samples were subjected to SDS-PAGE. ab187643 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 lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

Lane 1: Wild-type HEK-293T cell lysate (20 µg)

Lane 2: CDC42 knockout HEK-293T cell lysate (20 µg)

Lane 3: MCF7 cell lysate (20 µg)

Lane 4: HepG2 cell lysate (20 µg)

Lanes 1-4: Merged signal (red and green). Green - <u>ab64533</u> observed at 20 kDa. Red - loading control <u>ab8245</u> observed at 37 kDa.

<u>ab64533</u> Anti-CDC42 antibody was shown to specifically react with CDC42 in wild-type HEK-293T cells. Loss of signal was observed when knockout cell line <u>ab266522</u> (knockout cell lysate ab256868) was used. Wild-type and CDC42 knockout samples were subjected to SDS-PAGE. <u>ab64533</u> and Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) were incubated overnight at 4°C at 1 μg/ml and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye[®] 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye[®] 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.

Homozygous: 2 bp insertion in exon 3

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