abcam

Product datasheet

Human KRT8 (Cytokeratin 8) knockout HeLa cell lysate ab263785

5 图像

概述

产品概述

Knockout cell lysate achieved by CRISPR/Cas9.

Parental Cell Line HeLa

Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, 1 bp insertion in exon 2 and 2 bp deletion in exon 2

and 4 bp deletion in exon 2.

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.

说明

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

性能

存放说明 Store at -80°C. Please refer to protocols.

组件	1 kit
ab255504 - Human KRT8 knockout HeLa cell lysate	1 x 100µg
ab255552 - 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

靶标

功能 Together with KRT19, helps to link the contractile apparatus to dystrophin at the costameres of

striated muscle.

组织特异性 Observed in muscle fibers accumulating in the costameres of myoplasm at the sarcolemma

membrane in structures that contain dystrophin and spectrin. Expressed in gingival mucosa and

hard palate of the oral cavity.

疾病相关 Cirrhosis

序列相似性 Belongs to the intermediate filament family.

翻译后修饰 Phosphorylation on serine residues is enhanced during EGF stimulation and mitosis. Ser-74

phosphorylation plays an important role in keratin filament reorganization.

O-glycosylated. O-GlcNAcylation at multiple sites increases solubility, and decreases stability by

inducing proteasomal degradation.

O-glycosylated (O-GlcNAcylated), in a cell cycle-dependent manner.

细胞定位 Cytoplasm. Nucleus, nucleoplasm. Nucleus matrix.

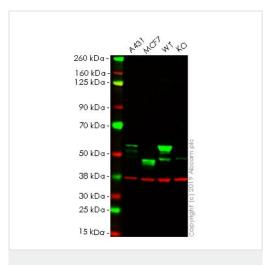
应用

The Abpromise guarantee Abpromise™承诺保证使用ab263785于以下的经测试应用

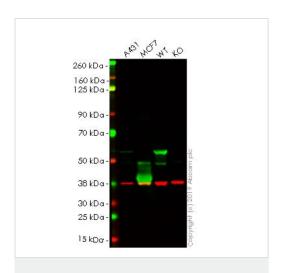
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
WB		Use at an assay dependent concentration.

图片



Western blot - Human KRT8 knockout HeLa cell lysate (ab263785)



Western blot - Human KRT8 knockout HeLa cell lysate (ab263785)

Lane 1: A431 cell lysate (20 µg)

Lane 2: MCF7 cell lysate (20 µg)

Lane 3: Wild-type HeLa cell lysate (20 µg)

Lane 4: KRT8 knockout HeLa cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab53280</u> observed at 55 kDa. Red - loading control, <u>ab8245</u> observed at 37 kDa.

ab53280 was shown to react with Cytokeratin 8 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab255400 (knockout cell lysate ab263785) was used. Wild-type and Cytokeratin 8 knockout samples were subjected to SDS-PAGE. ab53280 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 10000 (For unpurified use at 1/25,000 - 1/50,000) 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.

Lane 1: A431 cell lysate (20 µg)

Lane 2: MCF7 cell lysate (20 µg)

Lane 3: Wild-type HeLa cell lysate (20 µg)

Lane 4: KRT8 knockout HeLa cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab9023</u> observed at 55 kDa. Red - loading control, <u>ab181602</u> observed at 37 kDa.

ab9023 was shown to react with Cytokeratin 8 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab255400 (knockout cell lysate ab263785) was used. Wild-type and Cytokeratin 8 knockout samples were subjected to SDS-PAGE. ab9023 and Anti-GAPDH antibody EPR16891] - Loading Control (ab181602) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Mouse IgG H&L (IRDye® 800CW) preadsorbed (ab216772) and Goat Anti-Rabbit IgG H&L (IRDye® 680RD) preadsorbed (ab216777) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Allele-1: 4 bp deletion in exon 2



Allele-2: 2 bp deletion in exon 2

Sanger Sequencing - Human KRT8 knockout HeLa cell lysate (ab263785)

Mut GGCCCGGGGGCCAGAGGTGGACACCTTGTAAGGACTTCTGGGTCACCCTGATGGACATGG

WT GGCCCGGGGGCCAGAGGTGGACACCTTGTA GGACTTCTGGGTCACCCTGATGGACATGG

Sengger Seguragaing Library MCTTR Argentagat Hold o

Allele-3: 1 bp insertion in exon 2

Sanger Sequencing - Human KRT8 knockout HeLa cell lysate (ab263785)

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