abcam

Product datasheet

Human EMC10 (C19orf63) knockout HeLa cell line ab265783

4 Images

Overview

Product name Human EMC10 (C19orf63) knockout HeLa cell line

Parental Cell Line HeLa
Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, 55 bp deletion in exon 1 and 73 bp deletion in exon 1

Passage number <20

Knockout validation Sanger Sequencing, Western Blot (WB)

Tested applications Suitable for: WB

Biosafety level 2

General notesRecommended control: Human wild-type HeLa cell line (<u>ab255928</u>). Please note a wild-type cell line is not automatically included with a knockout cell line order, if required please add

recommended wild-type cell line at no additional cost using the code WILDTYPE-TMTK1.

 $\textbf{Cryopreservation cell medium:} \ \ \textbf{Cell Freezing Medium-DMSO Serum free media, contains}$

8.7% DMSO in MEM supplemented with methyl cellulose.

Culture medium: DMEM (High Glucose) + 10% FBS

Initial handling guidelines: Upon arrival, the vial should be stored in liquid nitrogen vapor phase and not at -80°C. Storage at -80°C may result in loss of viability.

- 1. Thaw the vial in 37°C water bath for approximately 1-2 minutes.
- 2. Transfer the cell suspension (0.8 mL) to a 15 mL/50 mL conical sterile polypropylene centrifuge tube containing 8.4 mL pre-warmed culture medium, wash vial with an additional 0.8 mL culture medium (total volume 10 mL) to collect remaining cells, and centrifuge at 201 x g (rcf) for 5 minutes at room temperature. 10 mL represents minimum recommended dilution. 20 mL represents maximum recommended dilution.
- 3. Resuspend the cell pellet in 5 mL pre-warmed culture medium and count using a haemocytometer or alternative cell counting method. Based on cell count, seed cells in an appropriate cell culture flask at a density of 2x10⁴ cells/cm². Seeding density is given as a guide only and should be scaled to align with individual lab schedules.
- 4. Incubate the culture at 37°C incubator with 5% CO₂. Cultures should be monitored daily.

Subculture guidelines:

All seeding densities should be based on cell counts gained by established methods. A guide seeding density of $2x10^4$ cells/cm² is recommended.

1

A partial media change 24 hours prior to subculture may be helpful to encourage growth, if

Cells should be passaged when they have achieved 80-90% confluence.

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We will provide viable cells that proliferate on revival.

Properties

Number of cells 1 x 10⁶ cells/vial, 1 mL

Adherent /Suspension Adherent
Tissue Cervix
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

Mycoplasma free Yes

Storage instructions Shipped on Dry Ice. Store in liquid nitrogen.

Storage buffer Constituents: 8.7% Dimethylsulfoxide, 2% Cellulose, methyl ether

Target

Tissue specificity Present in serum (at protein level). Expressed in the pituitary gland; very low levels in other brain

regions.

Sequence similarities Belongs to the EMC10 family.

Post-translational

modifications

Glycosylated.

Cellular localization Secreted and Membrane.

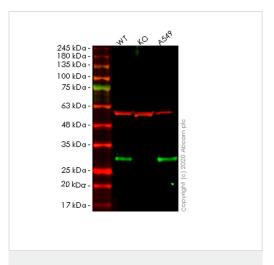
Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab265783 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		Use at an assay dependent concentration. Predicted molecular weight: 27 kDa.

Images



Western blot - Human EMC10 knockout HeLa cell line (ab265783)

All lanes : Anti-C19orf63 antibody [EPR13223] (<u>ab181209</u>) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: EMC10 knockout HeLa cell lysate

Lane 3: A549 cell lysate

Lysates/proteins at 20 µg per lane.

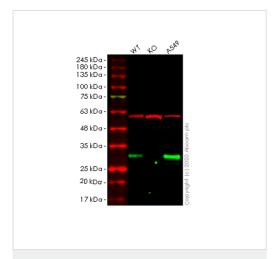
Secondary

All lanes: Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/10000 dilution

Predicted band size: 27 kDa Observed band size: 27 kDa

Lanes 1-4: Merged signal (red and green). Green - <u>ab181209</u> observed at 27 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab181209 Anti-C19orf63 antibody [EPR13223] was shown to specifically react with C19orf63 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265783 (knockout cell lysate ab257939) was used. Wild-type and C19orf63 knockout samples were subjected to SDS-PAGE. ab181209 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 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.



Western blot - Human EMC10 knockout HeLa cell line (ab265783)

All lanes : Anti-C19orf63 antibody [EPR13223-65] (**ab180148**) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: EMC10 knockout HeLa cell lysate

Lane 3: A549 cell lysate

Lysates/proteins at 20 µg per lane.

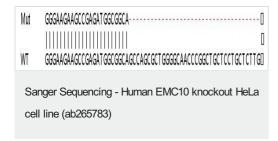
Secondary

All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (<u>ab216773</u>) at 1/10000 dilution

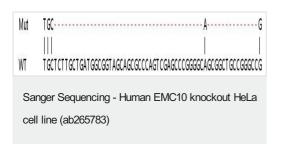
Predicted band size: 27 kDa Observed band size: 27 kDa

Lanes 1-3: Merged signal (red and green). Green - <u>ab180148</u> observed at 27 kDa. Red - loading control <u>ab7291</u> observed at 50 kDa.

ab180148 Anti-C19orf63 antibody [EPR13223-65] was shown to specifically react with C19orf63 in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265783 (knockout cell lysate ab257939) was used. Wild-type and C19orf63 knockout samples were subjected to SDS-PAGE. ab180148 and Anti-alpha Tubulin antibody [DM1A] - Loading Control (ab7291) were incubated overnight at 4°C at 1 in 1000 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.



Allele-1: 73 bp deletion in exon 1.



Allele-2: 55 bp deletion in exon 1.

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