abcam

Product datasheet

Human CORO1C (Coronin-1C) knockout HEK-293T cell lysate ab258377

3 Images

Overview

Product name Human CORO1C (Coronin-1C) knockout HEK-293T cell lysate

Product overview

Knockout cell lysate achieved by CRISPR/Cas9.

Parental Cell Line HEK293T

Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, 2 bp insertion in exon 2 and Insertion of the selection

cassette in exon 2.

Passage number <20

Knockout validation Sanger Sequencing

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.

Notes

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 <u>here</u>. 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.

Abcam has not and does not intend to apply for the REACH Authorisation of customers' uses of

products that contain European Authorisation list (Annex XIV) substances.

It is the responsibility of our customers to check the necessity of application of REACH Authorisation, and any other relevant authorisations, for their intended uses.

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Suitable for: WB

Properties

Storage instructions

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

Components	1 kit
ab261222 - Human CORO1C 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

Target

Function May be involved in cytokinesis, motility, and signal transduction.

Tissue specificity Ubiquitous.

Sequence similaritiesBelongs to the WD repeat coronin family.

Contains 4 WD repeats.

Applications

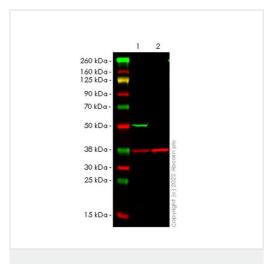
The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab258377 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.

Images



Western blot - Human CORO1C knockout HEK293T cell lysate (ab258377)

Lane 1: Wild-type 293T (human embryonic kidney epithelial cell) whole cell lysate 20 μg **Lane 2:**Coronin-1C knockout HEK-293T whole cell lysate 20 μg

Blocking buffer and

concentration: ½ volume of Odyssey Blocking Buffer (TBS)+ ½

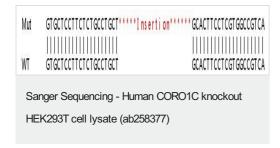
volume of 0.1% TBST

False colour image of Western blot: Anti-Coronin-1C antibody [EPR25365-24] (ab283693) staining at 1/1000 dilution, shown in green; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red.

In Western blot, <u>ab283693</u> was shown to bind specifically to Coronin-1C. A band was observed at 53 kDa in wild-type HEK-293T cell lysates with no signal observed at this size in Coronin-1C knockout cell line <u>ab266381</u> (knockout cell lysate (ab258377).

To generate this image, wild-type and Coronin-1C knockout HEK-293T cell lysates were analyzed. First, samples were run on an SDS-PAGE gel then transferred onto an immobilon-FL PVDF membrane. Membranes were blocked in Intercept[®] (TBS) Blocking Buffer diluted with an equal volume of 0.1% TBS 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 lgG H&L (IRDye $^{\$}$ 800CW) preabsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye $^{\$}$ 680RD) preabsorbed (<u>ab216776</u>) at 1/20000 dilution.



Allele-1: Insertion of the selection cassette in exon 2

Allele-2: 2 bp insertion in exon 2

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