abcam

Product datasheet

Human PPP2R5E knockout HeLa cell lysate ab258135

4 Images

Overview

Product name	Human PPP2R5E knockout HeLa cell lysate
Product overview	Knockout cell lysate achieved by CRISPR/Cas9.
Parental Cell Line	HeLa
Organism	Human
Mutation description	Knockout achieved by using CRISPR/Cas9, 1 bp deletion in exon2 and Insertion of the selection cassette in exon2.
Passage number	<20
Knockout validation	Sanger Sequencing, Western Blot (WB)
Reconstitution notes	To 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 **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.** 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. This product is subject to limited use licenses from The Broad Institute, ERS Genomics Limited and Sigma-Aldrich Co. LLC, and is developed with patented technology. For full details of the licenses and patents please refer to our **limited use license** and **patent pages**.
Tested applications	**Suitable for:** WB

Properties

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

Components	1 kit
ab262311 - Human PPP2R5E 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

Target

Function The B regulatory subunit might modulate substrate selectivity and catalytic activity, and also might direct the localization of the catalytic enzyme to a particular subcellular compartment.

Sequence similarities Belongs to the phosphatase 2A regulatory subunit B56 family.

Post-translational modifications Phosphorylated on serine residues.

Cellular localization Cytoplasm.

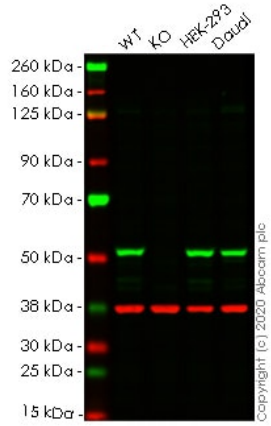
Applications

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab258135 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: 55 kDa.

Images



Western blot - Human PPP2R5E knockout HeLa cell lysate (ab258135)

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

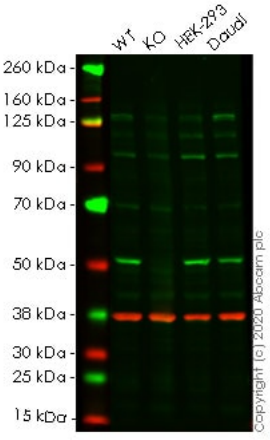
Lane 2: PPP2R5E knockout HeLa cell lysate (20 µg)

Lane 3: HEK-293 cell lysate (20 µg)

Lane 4: Daudi cell lysate (20 µg)

Lanes 1-4: Merged signal (red and green). Green - **ab198500** observed at 55 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab198500 Recombinant Anti-PPP2R5E antibody [EPR17147] - C-terminal was shown to specifically react with PPP2R5E in wild-type HeLa cells. Loss of signal was observed when knockout cell line **ab265637** (knockout cell lysate ab258135) was used. Wild-type and PPP2R5E knockout samples were subjected to SDS-PAGE. **ab198500** 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.



Western blot - Human PPP2R5E knockout HeLa cell lysate (ab258135)

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

Lane 2: PPP2R5E knockout HeLa cell lysate (20 µg)

Lane 3: HEK-293 cell lysate (20 µg)

Lane 4: Daudi cell lysate (20 µg)

Lanes 1-4: Merged signal (red and green). Green - **ab198290** observed at 55 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab198290 Anti-PPP2R5E antibody [EPR17146] was shown to specifically react with PPP2R5E in wild-type HeLa cells. Loss of signal was observed when knockout cell line **ab265637** (knockout cell lysate ab258135) was used. Wild-type and PPP2R5E knockout samples were subjected to SDS-PAGE. **ab198290** and Anti-GAPDH antibody [6C5] - Loading Control (**ab8245**) were incubated overnight at 4°C at 1 in 1000 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.

Sanger Sequencing - Human PPP2R5E knockout HeLa cell lysate (ab258135)

Allele-1: 1 bp deletion in exon2

Sanger Sequencing - Human PPP2R5E knockout HeLa cell lysate (ab258135)

Allele-2: Insertion of the selection cassette in exon2

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