abcam

Product datasheet

Human ANP32A (PHAP1) knockout HEK-293T cell lysate ab258303

2 图像

概述

产品名称 人ANP32A (PHAP1) knockout HEK-293T cell裂解物

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: 8 bp deletion in exon 1.

Passage number <20

Knockout validation Sanger Sequencing

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.

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

1

性能

存放说明

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

组 件	1 kit
ab260523 - Human ANP32A 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

Implicated in a number of cellular processes, including proliferation, differentiation, caspase-dependent and caspase-independent apoptosis, suppression of transformation (tumor suppressor), inhibition of protein phosphatase 2A, regulation of mRNA trafficking and stability in association with ELAVL1, and inhibition of acetyltransferases as part of the INHAT (inhibitor of histone acetyltransferases) complex. Plays a role in E4F1-mediated transcriptional repression.

组织特异性

Expressed in all tissues tested. Highly expressed in kidney and skeletal muscle, moderate levels of expression in brain, placenta and pancreas, and weakly expressed in lung. Found in all regions of the brain examined (amygdala, caudate nucleus, corpus callosum, hippocampus and

thalamus), with highest levels in amygdala.

序列相似性

Belongs to the ANP32 family.

Contains 4 LRR (leucine-rich) repeats.

Contains 1 LRRCT domain.

翻译后修饰

Phosphorylated on serine residues.

The N-terminus is blocked.

Some glutamate residues are glycylated by TTLL8. This modification occurs exclusively on

glutamate residues and results in a glycine chain on the gamma-carboxyl group.

细胞定位

Nucleus. Cytoplasm. Endoplasmic reticulum. Translocates to the cytoplasm during the process of

neuritogenesis (By similarity). Shuttles between nucleus and cytoplasm.

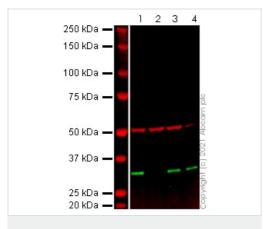
应用

The Abpromise guarantee

Abpromise™承诺保证使用ab258303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.



Western blot - Human ANP32A (PHAP1) knockout HEK-293T cell lysate (ab258303)

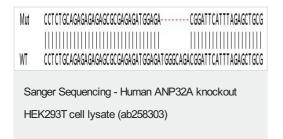
Lane 1: Wild-type HEK-293T cell lysate 20 µg

Lane 2: ANP32A knockout HEK-293T cell lysate 20 µg

Lane 3: HeLa cell lysate 20 µg

Lane 4: THP-1 cell lysate 20 µg

False colour image of Western blot: Anti-PHAP1 antibody - Cterminal staining at 0.5 µg/ml, shown in green; Mouse anti-Alpha Tubulin [DM1A] (ab7291) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab189110 was shown to bind specifically to PHAP1. A band was observed at 33 kDa in wild-type HEK-293T cell lysates with no signal observed at this size in ANP32A knockout cell line ab266148 (knockout cell lysate ab258303). To generate this image, wild-type and ANP32A knockout HEK-293T cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween[®] 20 (TBS-T) 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 IgG H&L (IRDye® 800CW) preabsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Homozygous: 8 bp deletion in exon 1

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