abcam

Product datasheet

Human TYROBP knockout THP-1 cell lysate ab273774

2 Images

Overview

Product name Human TYROBP knockout THP-1 cell lysate

Product overview Western blot data indicates that the CRISPR gene edit may have resulted in a truncation of the

protein of interest. Please see data images.

Parental Cell Line THP-1

Organism Human

Mutation description Knockout achieved by CRISPR/Cas9; X = 1 bp insertion; Frameshift = 99%

Passage number <20

Knockout validation Next Generation Sequencing (NGS)

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 **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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Tested applications Suitable for: WB

1

Properties

Storage instructions

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

Components	1 kit
ab280624 - Human TYROBP knockout THP-1 cell lysate	1 x 100µg
ab269602 - Human wild-type THP-1 cell lysate	1 x 100µg

Cell type acute monocytic leukemia

Disease Acute Monocytic Leukemia

Gender Male

Target

Function

 $Non-covalently\ associates\ with\ activating\ receptors\ of\ the\ CD300\ family.\ Cross-linking\ of\ CD300-linking\ of\ CD300$

TYROBP complexes results in cellular activation.

Tissue specificity

Expressed at low levels in the early development of the hematopoietic system and in the promonocytic stage and at high levels in mature monocytes. Expressed in hematological cells and tissues such as peripheral blood leukocytes and spleen. Also found in bone marrow, lymph nodes, placenta, lung and liver. Expressed at lower levels in different parts of the brain especially in the basal ganglia and corpus callosum.

Involvement in disease

Defects in TYROBP are a cause of polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy (PLOSL) [MIM:221770]; also called presenile dementia with bone cysts or Nasu-Hakola disease (NHD). PLOSL is a recessively inherited disease characterized by a combination of psychotic symptoms rapidly progressing to presenile dementia and bone cysts restricted to wrists and ankles. PLOSL has a global distribution, although most of the patients have been diagnosed in Finland and Japan, with an estimated population prevalence of 2x10(-6)

in the Finns.

Sequence similarities

Belongs to the TYROBP family.

Post-translational

modifications

Tyrosine phosphorylated.

Cellular localization

Membrane.

Applications

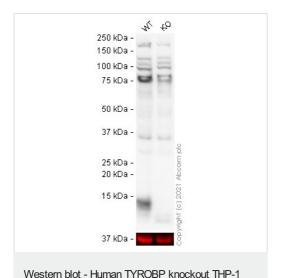
The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab273774 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: 12 kDa. Western blot data indicates that the CRISPR gene edit may have resulted in a truncation of the protein of interest. Please see data images.

Images



cell lysate (ab273774)

Lane 1: Wild-type THP-1 cell lysate 40 µg

Lane 2: TYROBP knockout THP-1 cell lysate 40 µg False colour image of Western blot: Anti-DAP12 antibody staining at 1 µg/ml, shown in black; Mouse anti-GAPDH antibody [6C5] (ab8245) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab93846 was shown to bind specifically to DAP12. A band was observed at 12 kDa in wild-type THP-1 cell lysates with no signal observed at this size in TYROBP knockout cell line ab273726 (knockout cell lysate ab273774). The band observed in the knockout lysate lane below 12 kDa is likely to represent a truncated form of DAP12. This has not been investigated further and the functional properties of the gene product have not been determined. To generate this image, wild-type and TYROBP knockout THP-1 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 5 % BSA 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 before development with Optiblot (ECL reagent ab133456) and imaged with 2 minutes exposure time. Secondary antibodies used were HRP conjugated Goat anti-Rabbit (H+L) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.

CTEACCCTSTGTGCAGNTGCAGTTGCT-CTACGSTGGGCCGGGCCTGCTGGCAGGG Reference

CTGACCCTGTGTGCAGNTGCAGTTGCT-CTACGSTGGGCCGGGCCTGTGGCAGGG Insertion, 35429 reads, 50.774

CTGACCCTGTGTGCAGNTGCAGTTGCT-CTACGSTGGGCCGGGCCTGTGGCAGGG Reference

CTGACCCTGTGTGCAGNTGCAGTTGCT-CTACGSTGGGCCCGGGCCTGTGGCAGGG Reference

Next Generation Sequencing - Human TYROBP

knockout THP-1 cell lysate (ab273774)

Knockout achieved by CRISPR/Cas9; X = 1 bp insertion; Frameshift = 99%

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