

## Product datasheet

# Human CASP1 (Caspase-1) knockout THP-1 cell lysate ab277988

[2 Images](#)

### Overview

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<b>Product name</b>	Human CASP1 (Caspase-1) knockout THP-1 cell lysate
<b>Product overview</b>	Knockout cell lysate achieved by CRISPR/Cas9.
<b>Parental Cell Line</b>	THP-1
<b>Organism</b>	Human
<b>Passage number</b>	<20
<b>Knockout validation</b>	Western Blot (WB)
<b>Reconstitution notes</b>	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. <i>*Usage of SDS sample buffer is not recommended with these lyophilized lysates.</i>

**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

## Properties

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

Components	1 kit
ab277989 - Human CASP1 (Caspase-1) knockout THP-1 cell lysate	1 x 100µg
ab282895 - Human wild-type THP-1 cell lysate	1 x 100µg

**Cell type** acute monocytic leukemia  
**Disease** Acute Monocytic Leukemia  
**Gender** Male

## Target

**Function** Thiol protease that cleaves IL-1 beta between an Asp and an Ala, releasing the mature cytokine which is involved in a variety of inflammatory processes. Important for defense against pathogens. Cleaves and activates sterol regulatory element binding proteins (SREBPs). Can also promote apoptosis.

**Tissue specificity** Expressed in larger amounts in spleen and lung. Detected in liver, heart, small intestine, colon, thymus, prostate, skeletal muscle, peripheral blood leukocytes, kidney and testis. No expression in the brain.

**Sequence similarities** Belongs to the peptidase C14A family.  
Contains 1 CARD domain.

**Post-translational modifications** The two subunits are derived from the precursor sequence by an autocatalytic mechanism.

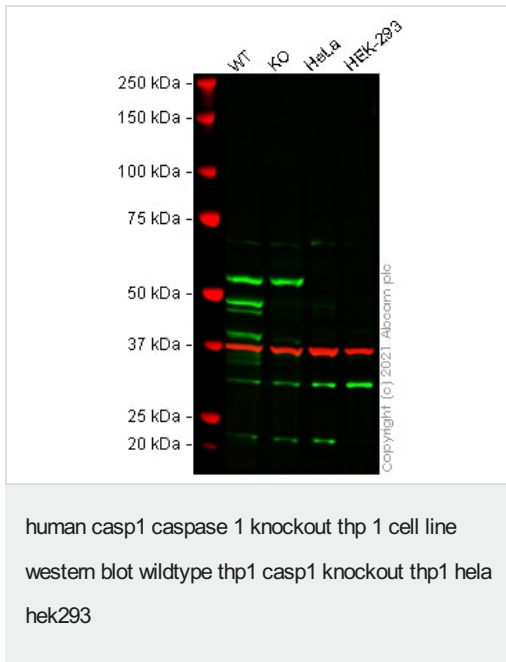
**Cellular localization** Cytoplasm.

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab277988 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



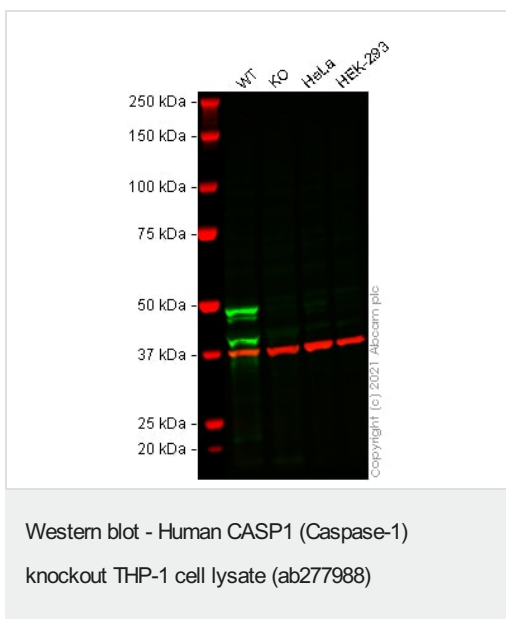
**Lane 1:** Wild-type THP-1 cell lysate 40 µg

**Lane 2:** THP-1 cell lysate 40 µg

**Lane 3:** HeLa cell lysate 20 µg

**Lane 4:** HEK-293 cell lysate 20 µg

False colour image of Western blot: Anti-CASP1 antibody 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, the antibody was shown to bind specifically to CASP1. A band was observed at 35-40, 48 kDa in wild-type THP-1 cell lysates with no signal observed at this size in CASP1 knockout cell line [ab276116](#) (knockout cell lysate ab277988). To generate this image, wild-type and CASP1 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 fluorescent western blot (TBS-based) blocking solution 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.



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

**Lane 2:** CASP1 knockout THP-1 cell lysate 40 µg

**Lane 3:** HeLa cell lysate 20 µg

**Lane 4:** HEK-293 cell lysate 20 µg

False colour image of Western blot: Anti-Caspase-1 antibody [EPR19672] 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, [ab207802](#) was shown to bind specifically to Caspase-1. A band was observed at 22, 35-40, 48 kDa in wild-type THP-1 cell lysates with no signal observed at this size in CASP1 knockout cell line [ab276116](#) (knockout cell lysate ab277988). To generate this image, wild-type and CASP1 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 fluorescent

western blot (TBS-based) blocking solution 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.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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