abcam

Product datasheet

Anti-PARP1 antibody [Y17] - BSA and Azide free ab284670



2 Images

Overview

Product name Anti-PARP1 antibody [Y17] - BSA and Azide free

Description Rabbit monoclonal [Y17] to PARP1 - BSA and Azide free

Host species Rabbit

Specificity This antibody is specific to PARP1. It should recognize the intact form (116kDa) and the p25

cleaved form of PARP1.

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF or IHC

Species reactivity Reacts with: Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Jurkat and Camptothecin cell lysate.

General notes ab284670 is the carrier-free version of **ab32378**

Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for

increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our **conjugation kits** for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.

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If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C.

Storage buffer

Constituent: 100% PBS

Carrier free Yes

Clonality Monoclonal

Clone number Y17 Isotype lgG

Applications

Our Abpromise guarantee covers the use of ab284670 in the following tested applications. The Abpromise guarantee

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: 113 kDa.

Application notes Is unsuitable for Flow Cyt,ICC/IF or IHC.

Target

Function Involved in the base excision repair (BER) pathway, by catalyzing the poly(ADP-ribosyl)ation of a

limited number of acceptor proteins involved in chromatin architecture and in DNA metabolism.

This modification follows DNA damages and appears as an obligatory step in a

detection/signaling pathway leading to the reparation of DNA strand breaks. Mediates the poly(ADP-ribosyl)ation of APLF and CHFR. Positively regulates the transcription of MTUS1 and

negatively regulates the transcription of MTUS2/TIP150.

Sequence similarities Contains 1 BRCT domain.

> Contains 1 PARP alpha-helical domain. Contains 1 PARP catalytic domain. Contains 2 PARP-type zinc fingers.

Post-translational

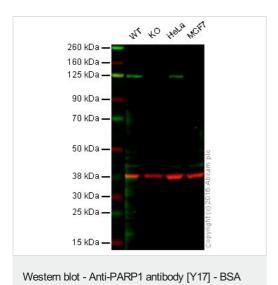
Phosphorylated by PRKDC. Phosphorylated upon DNA damage, probably by ATM or ATR. modifications

Poly-ADP-ribosylated by PARP2. Poly-ADP-ribosylation mediates the recruitment of CHD1L to

DNA damage sites.

S-nitrosylated, leading to inhibit transcription regulation activity.

Cellular localization Nucleus.



and Azide free (ab284670)

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This data was developed using <u>ab32378</u>, the same antibody clone in a different buffer formulation.

Lane 1: Wild type HAP1 whole cell lysate (20 µg)

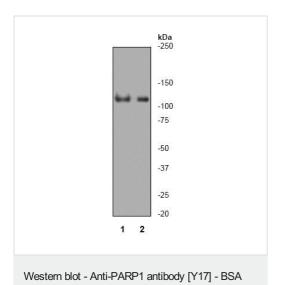
Lane 2: PARP1 knockout HAP1 whole cell lysate (20 µg)

Lane 3: HeLa whole cell lysate (20 µg)

Lane 4: MCF7 whole cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab32378</u> observed at 125 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

<u>ab32378</u> was shown to specifically react with PARP1 when PARP1 knockout samples were used. Wild-type and PARP1 knockout samples were subjected to SDS-PAGE. Ab32378 and <u>ab8245</u> (Mouse anti GAPDH loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/10000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) <u>ab216773</u> and 680CW Goat anti Mouse secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



All lanes: Anti-PARP1 antibody [Y17] (ab32378) at 1/1000 dilution

Lane 1: Jurkat cell lysate.

Lane 2: Jurkat and Camptothecin cell lysate.

Predicted band size: 113 kDa **Observed band size:** 115 kDa

This data was developed using <u>ab32378</u>, the same antibody clone in a different buffer formulation.

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

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