# abcam

## Product datasheet

# Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [Y34] ab237432

Recombinant RabMAb

### 3 Images

#### Overview

**Product name** Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [Y34]

**Description** Alexa Fluor® 488 Rabbit monoclonal [Y34] to Cleaved PARP1

**Host species** Rabbit

Conjugation Alexa Fluor® 488, Ex: 495nm, Em: 519nm

Specificity This antibody is specific for p85 cleaved form of PARP1.

**Tested applications** Suitable for: ICC/IF, Flow Cyt (Intra)

Species reactivity Reacts with: Human

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

Positive control ICC/IF: MCF7 cells treated with Staurosporine (1µM, 5 hr). Flow Cyt (intra): Staurosporine

treated MCF7 cells.

**General notes** This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our  $\mathsf{RabMAb}^{\mathsf{®}}$  technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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Life Technologies Corporation, 5781 Van Allen Way, Carlsbad, CA 92008 USA or **outlicensing@thermofisher.com**.

#### **Properties**

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

Purity Protein A purified

**Clonality** Monoclonal

Clone number Y34 lsotype lgG

#### **Applications**

#### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab237432 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
ICC/IF		1/1000. This product gave a positive signal in MCF7 cells treated with Staurosporine (1µM, 5 hr) fixed with 4% formaldehyde (10 min)
Flow Cyt (Intra)		1/5000.

#### **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. With EEF1A1 and TXK, forms a complex that acts as a T-helper 1 (Th1) cell-specific transcription factor and binds the promoter of IFN-gamma to directly regulate its transcription, and is thus involved importantly in Th1 cytokine production. Required for PARP9 and DTX3L recruitment to DNA damage sites. PARP1-dependent PARP9-DTX3L-mediated ubiquitination promotes the rapid and specific recruitment of 53BP1/TP53BP1, UIMC1/RAP80, and BRCA1 to DNA damage sites.

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 and TXK.

modifications

 ${\hbox{Poly-ADP-ribosylated by PARP2. Poly-ADP-ribosylation mediates the recruitment of CHD1L to} \\$ 

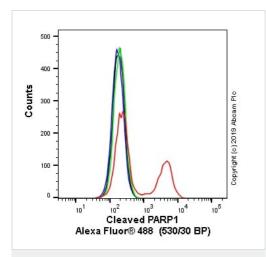
DNA damage sites.

S-nitrosylated, leading to inhibit transcription regulation activity.

**Cellular localization** 

Nucleus. Nucleus, nucleolus. Localizes at sites of DNA damage.

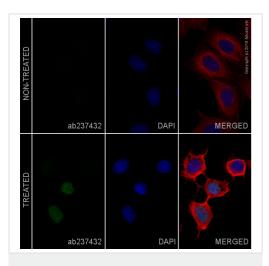
#### **Images**



Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [Y34] (ab237432)

Overlay histogram showing MCF7 cells treated (red line) or untreated (green line) with 1  $\mu$ M Staurosporine for 5 hrs and subsequent staining with ab237432. The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS/10% normal Goat serum to block non-specific protein-protein interactions followed by the antibody (ab237432) (1x  $10^6$  in 100  $\mu$ l at 0.1  $\mu$ g/ml (1/5000 dilution)) for 30 min at 22°C.

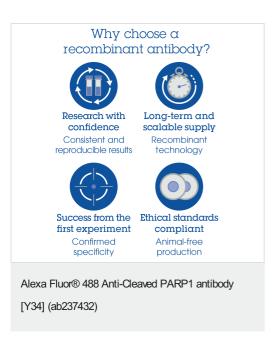
Isotype control antibody (black line) was Rabbit IgG (monoclonal) Alexa Fluor<sup>®</sup> 488 (<u>ab199091</u>) used at the same concentration and conditions as the primary antibody. Unlabeled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 50 mW Blue laser (488nm) and 530/30 bandpass filter.



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [Y34] (ab237432)

ab237432 staining Cleaved PARP1 in MCF7 cells Staurosporine-treated (1 $\mu$ M, 5 hr). The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab237432 at 1/1000 dilution (shown in green) and <u>ab195884</u>, Rat monoclonal to Tubulin (Alexa Fluor<sup>®</sup> 647), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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