abcam

Product datasheet

Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [4B5BD2] ab170171

2 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-Cleaved PARP1抗体[4B5BD2]

描述 Alexa Fluor® 488荧光小鼠单克隆抗体[4B5BD2] to Cleaved PARP1

宿主 Mouse

偶联物 Alexa Fluor® 488. Ex: 495nm, Em: 519nm

经测试应用 适用于: ICC, Flow Cyt (Intra)

种属反应性 与反应: Human

免疫原 Synthetic peptide within Human Cleaved PARP1 aa 200-300 (N terminal). The exact sequence is

proprietary.

Database link: P09874

阳性对照 HeLa cells treated with 1 µM staurosporine for 4 hours.

常规说明

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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.

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

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found below, along with publications, customer reviews and Q&As

Product was previously marketed under the MitoSciences sub-brand.

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C. Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In

the Dark.

存储溶液 Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, 68% PBS

纯**度** Ammonium Sulphate Precipitation

 克隆
 单克隆

 克隆编号
 4B5BD2

 同种型
 IqG1

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab170171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 |
|------------------|------|-------------------------------------------------------------------------------------------------------------------------------------------|
| ICC | | Use a concentration of 1 μ g/ml. Use Antigen Retrieval Buffer (100 mM Tris, 5% urea, pH 9.5) at 95°C for 10 min to boost signal. |
| Flow Cyt (Intra) | | Use a concentration of 1 µg/ml. <u>ab171463</u> - Mouse monoclonal lgG1, is suitable for use as an isotype control with this antibody. |

靶标

功能

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.

序列相似性

Contains 1 BRCT domain.

Contains 1 PARP alpha-helical domain.

Contains 1 PARP catalytic domain. Contains 2 PARP-type zinc fingers.

翻译后修饰

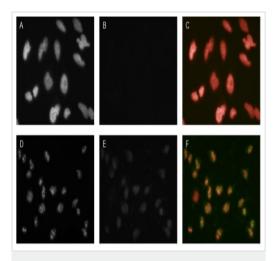
Phosphorylated by PRKDC and TXK.

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.

细胞定位 Nucleus. Nucleus, nucleolus. Localizes at sites of DNA damage.

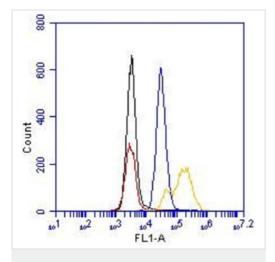
图片



Immunocytochemistry - Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [4B5BD2] (ab170171)

Immunocytochemistry with anti-cleaved PARP1 antibody conjugated to Alexa Fluor[®] 488.

HeLa cells were vehicle-treated (panels A-C) or treated with 1 μ M staurosporine for 4 hours (panels D-F), then fixed. Cells were treated with antigen retrieval buffer (100 mM Tris, 5% urea, pH 9.5) for 10 minutes at 95°C, then permeabilized and blocked. Cells were incubated with 1 μ g/mL of the cleaved PARP1 antibody conjugated to Alexa Fluor® 488, then co-stained with the DNA stain DAPI. Images of DAPI signals (A and D), anti-cleaved PARP1 signal (B and E), and overlays of DAPI (artificially colored red for better contrast) and anti-cleaved PARP1 (colored green) images (C and F) are shown.



Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-Cleaved PARP1 antibody [4B5BD2] (ab170171)

Flow cytometry with anti-cleaved PARP1 antibody conjugated to Alexa Fluor® 488.

Flow cytometric analysis was performed on HeLa vehicle-treated cells and on HeLa cells treated with 1 μ M staurosporine for 4 hours. Cells were fixed with paraformaldehyde and permeablized with methanol. HeLa vehicle-treated cells were stained with 1 μ g/mL of the cleaved PARP1 antibody conjugated to Alexa488 (blue) or a negative, nonreactive Alexa Fluor® 488-conjugated control antibody (black). HeLa staurosporine-treated cells were stained with 1 μ g/mL of the cleaved PARP1 antibody conjugated to Alexa Fluor® 488 (yellow) or a negative, nonreactive Alexa Fluor® 488-conjugated control antibody (red). 1% BSA in PBS was used as the blocking reagent for all blocking and antibody incubation steps.

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