abcam

Product datasheet

Anti-PARP1 antibody [A6.4.12] ab110915

★★★★★ 1 Abreviews 13 References 3 Images

Overview

Product name Anti-PARP1 antibody [A6.4.12]

Description Mouse monoclonal [A6.4.12] to PARP1

Host species Mouse

Tested applications
Suitable for: IHC-P
Species reactivity
Reacts with: Human

Predicted to work with: Mouse, Rat, Hamster, Xenopus laevis, Drosophila melanogaster

edicted to work with Mode, Nat, Hamster, Acropus lacvis, Diosophila melanogasti

Immunogen Full length native protein (purified) corresponding to Human PARP1.

Positive control Human placenta, skin and spleen tissues.

General notes

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

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

Properties

Form Liquid

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

Storage buffer Preservative: 0.09% Sodium azide

Purity Protein G purified

Clonality Monoclonal
Clone number A6.4.12
Isotype IqG1

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab110915 in the following tested applications.

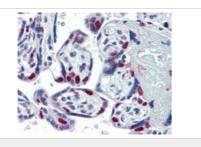
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		Use a concentration of 10 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Target

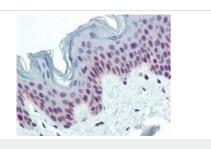
Function	Involved in the base excision repair (BER) pathway, by catalyzing the poly(ADP-ribosyl)ation of a limited number of acceptor proteins involved in chromatin architecture and in DNA metabolism. This modification follows DNA damages and appears as an obligatory step in a detection/signaling pathway 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 modifications	Phosphorylated by PRKDC. Phosphorylated upon DNA damage, probably by ATM or ATR. 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.

Images



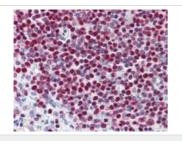
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PARP1 antibody
[A6.4.12] (ab110915)

ab110915, at 10 μ g/ml, staining PARP1 in formalin fixed, paraffin embedded Human placenta by Immunohistochemistry. Biotinylated anti mouse IgG secondary antibody, alkaline phosphatase streptavidin and chromogen was used to develop staining.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PARP1 antibody
[A6.4.12] (ab110915)

ab110915, at 10 μ g/ml, staining PARP1 in formalin fixed, paraffin embedded Human skin by Immunohistochemistry. Biotinylated anti mouse IgG secondary antibody, alkaline phosphatase streptavidin and chromogen was used to develop staining.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PARP1 antibody
[A6.4.12] (ab110915)

ab110915, at 10 μ g/ml, staining PARP1 in formalin fixed, paraffin embedded Human spleen by Immunohistochemistry. Biotinylated anti mouse IgG secondary antibody, alkaline phosphatase streptavidin and chromogen was used to develop staining.

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

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