abcam

Product datasheet

Anti-53BP1 antibody [EPR2173(2)] ab175188





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Overview

Product name Anti-53BP1 antibody [EPR2173(2)]

Description Rabbit monoclonal [EPR2173(2)] to 53BP1

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, ICC/IF

Unsuitable for: Flow Cyt or IP

Species reactivity Reacts with: Human

Immunogen Synthetic peptide within Human 53BP1 aa 900-1000 (Cysteine residue). The exact sequence is

proprietary.

Database link: Q12888

Positive control WB: HAP1, HeLa, and HepG2 lysates. Human fetal brain and heart lysates. IHC-P: Human

cervical carcinoma and prostate hyperplasia tissue. ICC: HepG2 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 RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb patents**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

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 long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

Purity Tissue culture supernatant

Clonality Monoclonal
Clone number EPR2173(2)

Isotype IgG

Applications

The Abpromise quarantee Our Abpromise quarantee covers the use of ab175188 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		1/50000 - 1/200000. Predicted molecular weight: 214 kDa.
IHC-P		1/50 - 1/100. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC/IF	★★★★★ (1)	1/100 - 1/250.

Application notes Is unsuitable for Flow Cyt or IP.

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Function May have a role in checkpoint signaling during mitosis. Enhances TP53-mediated transcriptional

activation. Plays a role in the response to DNA damage.

Involvement in diseaseNote=A chromosomal aberration involving TP53BP1 is found in a form of myeloproliferative

disorder chronic with eosinophilia. Translocation t(5;15)(q33;q22) with PDGFRB creating a

TP53BP1-PDGFRB fusion protein.

Sequence similaritiesContains 2 BRCT domains.

Post-translational

modifications

Asymmetrically dimethylated on Arg residues by PRMT1. Methylation is required for DNA binding.

Phosphorylated at basal level in the absence of DNA damage. Hyper-phosphorylated in an ATM-

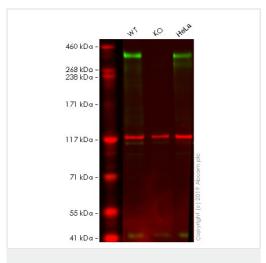
dependent manner in response to DNA damage induced by ionizing radiation. Hyperphosphorylated in an ATR-dependent manner in response to DNA damage induced by UV

irradiation.

Cellular localization Nucleus. Chromosome > centromere > kinetochore. Associated with kinetochores. Both nuclear

and cytoplasmic in some cells. Recruited to sites of DNA damage, such as double stand breaks. Methylation of histone H4 at 'Lys-20' is required for efficient localization to double strand breaks.

Images



Western blot - Anti-53BP1 antibody [EPR2173(2)] (ab175188)

All lanes : Anti-53BP1 antibody [EPR2173(2)] (ab175188) at 1/50000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: TP53BP1 knockout HAP1 whole cell lysate

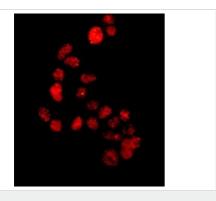
Lane 3: HeLa whole cell lysate

Lysates/proteins at 20 µg per lane.

Predicted band size: 214 kDa

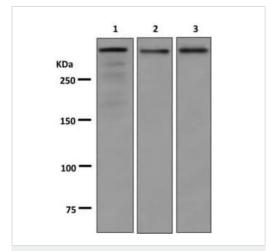
Lanes 1 - 3: Merged signal (red and green). Green - ab175188 observed at 213 kDa. Red - loading control, **ab130007**, observed at 130 kDa.

ab175188 was shown to recognize 53BP1 in wild-type HAP1 cells as signal was lost at the expected MW in TP53BP1 knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and TP53BP1 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3% Milk. Ab175188 and ab130007 (Mouse anti Vinculin loading control) were incubated overnight at 4°C at 1/50000 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.



Immunocytochemistry/ Immunofluorescence - Anti-53BP1 antibody [EPR2173(2)] (ab175188)

Immunofluorescence analysis of HepG2 cells labeling 53BP1 with ab175188 at a 1/100 dilution.



Western blot - Anti-53BP1 antibody [EPR2173(2)] (ab175188)

All lanes : Anti-53BP1 antibody [EPR2173(2)] (ab175188) at 1/50000 dilution

Lane 1: Human fetal heart tissue lysate

Lane 2: HepG2 cell lysate

Lane 3: Human fetal brain tissue lysate

Lysates/proteins at 10 µg per lane.

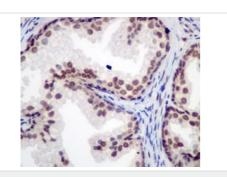
Predicted band size: 214 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-53BP1 antibody
[EPR2173(2)] (ab175188)

Immunohistochemical analysis of paraffin-embedded
Human cervical carcinoma tissue labeling 53BP1 with ab175188 at a 1/50 dilution.

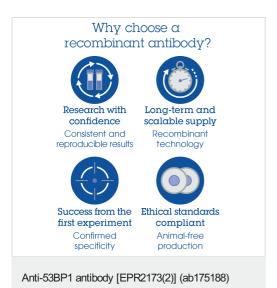
Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-53BP1 antibody
[EPR2173(2)] (ab175188)

Immunohistochemical analysis of paraffin-embedded Human prostate hyperplasia tissue labeling 53BP1 with ab175188 at a 1/50 dilution.

Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.



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