abcam

Product datasheet

Alexa Fluor® 488 Anti-53BP1 antibody [EPR2172(2)] ab237174



3 Images

Overview

Product name Alexa Fluor® 488 Anti-53BP1 antibody [EPR2172(2)]

Description Alexa Fluor® 488 Rabbit monoclonal [EPR2172(2)] to 53BP1

Host species Rabbit

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

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

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

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

Positive control ICC/IF: UV treated HeLa cells Flow Cyt (intra): HeLa 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**.

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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. 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 EPR2172(2)

Isotype lgG

Applications

Target

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab237174 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
Flow Cyt (Intra)		1/50.
ICC/IF		1/100. This product gave a positive signal in UV treated HeLa cells fixed with 4% formaldehyde (10 min)

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 disease Note=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 similarities Contains 2 BRCT domains.

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

modifications 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

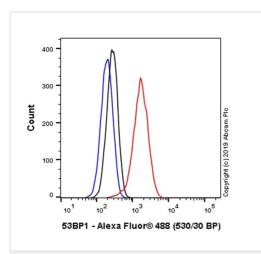
irradiation.

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

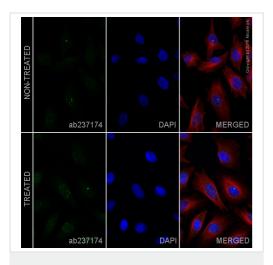


Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-53BP1 antibody [EPR2172(2)] (ab237174)

Overlay histogram showing HeLa cells stained with ab237174 (red line). 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 (ab237174, 1/50 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was Rabbit IgG (monoclonal)
Alexa Fluor[®] 488 (<u>ab199091</u>) used at the same concentration and conditions as the primary antibody. Unlabelled 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-53BP1 antibody [EPR2172(2)] (ab237174)

ab237174 staining 53BP1 in UV treated HeLa cells. 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 ab237174 at 1/100 dilution (shown in green) and ab195884, Rat monoclonal to Tubulin (Alexa Fluor® 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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