abcam

Product datasheet

Anti-Ku80 antibody [EPR3467] - BSA and Azide free ab247495



4 Images

Overview

Product name Anti-Ku80 antibody [EPR3467] - BSA and Azide free

Description Rabbit monoclonal [EPR3467] to Ku80 - BSA and Azide free

Host species Rabbit

Specificity The immunogen for this antibody has 78.6% homology with mouse and rat ku80, therefore we do

not guarantee reactivity in these species.

Tested applications Suitable for: Flow Cyt (Intra), IHC-P, WB

Unsuitable for: IP

Species reactivity Reacts with: Human

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

General notes ab247495 is the carrier-free version of ab79391.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our **conjugation kits** for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

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. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EPR3467

Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab247495 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)		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Detects a band of approximately 83 kDa (predicted molecular weight: 83 kDa).

Application notes

Is unsuitable for IP.

Target

Function

Single stranded DNA-dependent ATP-dependent helicase. Has a role in chromosome translocation. The DNA helicase II complex binds preferentially to fork-like ends of double-stranded DNA in a cell cycle-dependent manner. It works in the 3'-5' direction. Binding to DNA may be mediated by XRCC6. Involved in DNA non-homologous end joining (NHEJ) required for double-strand break repair and V(D)J recombination. The XRCC5/6 dimer acts as regulatory subunit of the DNA-dependent protein kinase complex DNA-PK by increasing the affinity of the catalytic subunit PRKDC to DNA by 100-fold. The XRCC5/6 dimer is probably involved in stabilizing broken DNA ends and bringing them together. The assembly of the DNA-PK complex to DNA ends is required for the NHEJ ligation step. In association with NAA15, the XRCC5/6 dimer binds to the osteocalcin promoter and activates osteocalcin expression. The XRCC5/6

dimer probably also acts as a 5'-deoxyribose-5-phosphate lyase (5'-dRP lyase), by catalyzing the beta-elimination of the 5' deoxyribose-5-phosphate at an abasic site near double-strand breaks. XRCC5 probably acts as the catalytic subunit of 5'-dRP activity, and allows to 'clean' the termini of abasic sites, a class of nucleotide damage commonly associated with strand breaks, before such broken ends can be joined. The XRCC5/6 dimer together with APEX1 acts as a negative

regulator of transcription.

Belongs to the ku80 family.

Contains 1 Ku domain.

Developmental stage Expression increases during promyelocyte differentiation.

DomainThe EEXXXDDL motif is required for the interaction with catalytic subunit PRKDC and its

recruitment to sites of DNA damage.

Post-translational modifications

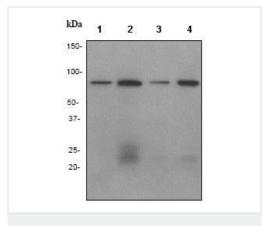
Sequence similarities

Phosphorylated on serine residues. Phosphorylation by PRKDC may enhance helicase activity.

Sumoylated.

Cellular localization Nucleus. Chromosome.

Images



Western blot - Anti-Ku80 antibody [EPR3467] - BSA and Azide free (ab247495)

All lanes: Anti-Ku80 antibody [EPR3467] (ab79391) at 1/2000

dilution

Lane 1: A549 cell lysate, treated with FBS

Lane 2: HeLa cell lysate

Lane 3: U937 (Human histiocytic lymphoma cell line) cell lysate

Lane 4: HepG2 cell lysate

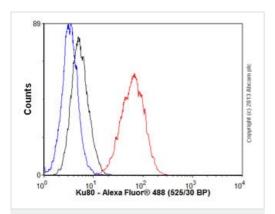
Lysates/proteins at 10 µg per lane.

Secondary

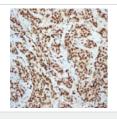
All lanes: goat anti-rabbit HRP at 1/2000 dilution

Predicted band size: 83 kDa **Observed band size:** 83 kDa

This data was developed using <u>ab79391</u>, the same antibody clone in a different buffer formulation.



Flow Cytometry (Intracellular) - Anti-Ku80 antibody [EPR3467] - BSA and Azide free (ab247495)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Ku80 antibody

[EPR3467] - BSA and Azide free (ab247495)

This data was developed using <u>ab79391</u>, the same antibody clone in a different buffer formulation. Overlay histogram showing HeLa cells stained with <u>ab79391</u> (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab79391</u>, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor[®] 488 goat anti-rabbit lgG (H+L) (<u>ab150077</u>) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit lgG (monoclonal) (1?g/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.

This data was developed using <u>ab79391</u>, the same antibody clone in a different buffer formulation.<u>ab79391</u> at 1/250 dilution staining Ku80 in human breast carcinoma by Immunohistochemistry, Paraffin-embedded tissue. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.



Anti-Ku80 antibody [EPR3467] - BSA and Azide free (ab247495)

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

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