abcam

Product datasheet

Anti-Blooms Syndrome Protein Blm antibody ab5446

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Overview

Product name Anti-Blooms Syndrome Protein Blm antibody

Description Goat polyclonal to Blooms Syndrome Protein Blm

Host species Goat

Tested applications

Suitable for: WB, IP

Species reactivity

Reacts with: Human

Predicted to work with: Chimpanzee

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Immunogen Immunogen was a synthetic peptide, which represented a portion of human Bloom Syndrome

encoded within exon 2.

Positive control WB: HeLa, HEK-293T and Jurkat whole cell lysates. IP: HeLa whole cell lysate.

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. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7

Constituents: 0.021% PBS, 1.764% Sodium citrate, 1.815% Tris

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab5446 in the following tested applications.

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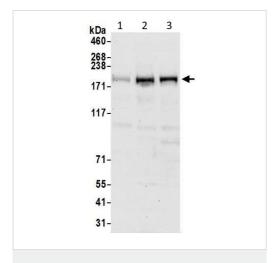
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB	*** <u>*</u>	1/10000 - 1/25000. Predicted molecular weight: 159 kDa.
IP		Use at 2-10 µg/mg of lysate.

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Function	Participates in DNA replication and repair. Exhibits a magnesium-dependent ATP-dependent DNA-helicase activity that unwinds single- and double-stranded DNA in a 3'-5' direction.
Involvement in disease	Defects in BLM are the cause of Bloom syndrome (BLM) [MIM:210900]. BLM is an autosomal recessive disorder characterized by proportionate pre- and postnatal growth deficiency, sunsensitive telangiectatic hypo- and hyperpigmented skin, predisposition to malignancy, and chromosomal instability.
Sequence similarities	Belongs to the helicase family. RecQ subfamily. Contains 1 helicase ATP-binding domain. Contains 1 helicase C-terminal domain. Contains 1 HRDC domain.
Post-translational modifications	Phosphorylated in response to DNA damage. Phosphorylation requires the FANCA-FANCC-FANCE-FANCF-FANCG protein complex, as well as the presence of RMI1.
Cellular localization	Nucleus.

Images



Western blot - Anti-Blooms Syndrome Protein Blm antibody (ab5446)

All lanes : Anti-Blooms Syndrome Protein Blm antibody (ab5446) at $0.04~\mu g/ml$

Lane 1 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

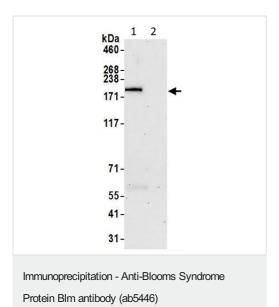
Lane 2: HEK-293T (Human epithelial cell line from embryonic kidney transformed with large T antigen) whole cell lysate

Lane 3: Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lysates/proteins at 50 µg per lane.

Predicted band size: 159 kDa

Exposure time: 30 seconds



Blooms Syndrome Protein Blm was immunoprecipitated from HeLa (human epithelial cell line from cervix adenocarcinoma) whole cell lysate (1 mg for IP, 20% of IP loaded) with ab5446 at 6 μ g/mg lysate. Western blot was performed from the immunoprecipitate using ab5446 at 1 μ g/ml.

Lane 1: ab5446 IP in HeLa whole cell lysate.

Lane 2: Control lgG IP in HeLa whole cell lysate.

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

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