abcam

Product datasheet

Anti-eIF4Al antibody ab31217

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Overview

Product name Anti-elF4A1 antibody

Description Rabbit polyclonal to elF4A1

Host species Rabbit

Specificity Replenishment batches of our polyclonal antibody, ab31217 are tested in WB. Previous batches

were additionally validated in ICC/IF, IHC-P and IP. These applications are still expected to work and are covered by our Abpromise guarantee. You may also be interested in our alternative

recombinant antibody, ab185946.

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

Species reactivity Reacts with: Mouse, Rat, Cow, Human, African green monkey

Predicted to work with: Monkey

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

Positive control

This antibody gave a positive signal in the following whole cell lysates: HeLa (Human epithelial

carcinoma cell line) Jurkat (Human T cell lymphoblast-like cell line) A431 (Human epithelial carcinoma cell line) NIH 3T3 (Mouse embryonic fibroblast cell line) MEF1 (Mouse embryonic fibroblast cell line) PC12 (Rat adrenal pheochromocytoma cell line) This antibody gave a positive

signal in the following tissue lysates: Liver (Mouse) Pancreas (Mouse) Testis (Mouse)

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). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

1

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee

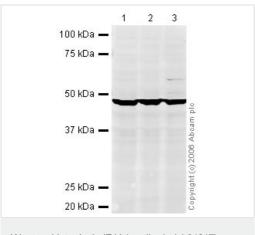
Our <u>Abpromise guarantee</u> covers the use of ab31217 in the following tested applications.

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

Application	Abreviews	Notes
WB	****(4)	Use a concentration of 1 µg/ml. Detects a band of approximately 47 kDa (predicted molecular weight: 47 kDa).
IP	★★★★ <u>(1)</u>	Use at an assay dependent concentration.
ICC/IF	*** <u>*</u>	Use a concentration of 1 µg/ml.
IHC-P		Use at an assay dependent concentration.

ATP-dependent RNA helicase which is a subunit of the elF4F complex involved in cap recognition and is required for mRNA binding to ribosome. In the current model of translation initiation, elF4A unwinds RNA secondary structures in the 5'-UTR of mRNAs which is necessary to allow efficient binding of the small ribosomal subunit, and subsequent scanning for the initiator codon.
Belongs to the DEAD box helicase family. elF4A subfamily. Contains 1 helicase ATP-binding domain. Contains 1 helicase C-terminal domain.

Images



Western blot - Anti-elF4A1 antibody (ab31217)

All lanes: Anti-elF4A1 antibody (ab31217) at 1 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2 : Jurkat whole cell lysate (<u>ab7899</u>)
Lane 3 : A-431 whole cell lysate (<u>ab7909</u>)

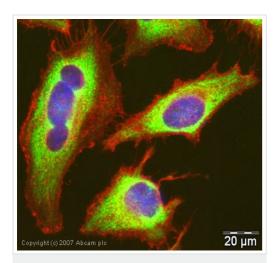
Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat polyclonal to Rabbit lgG (Alexa Fluor® 680) at 1/10000 dilution

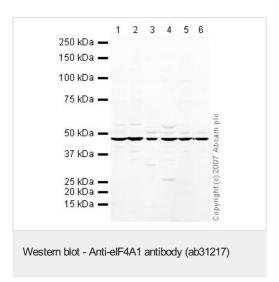
Performed under reducing conditions.

Predicted band size: 47 kDa **Observed band size:** 47 kDa



Immunocytochemistry/ Immunofluorescence - AntieIF4A1 antibody (ab31217)

ICC/IF image of ab31217 stained human HeLa cells. The cells were methanol fixed (5 min) and incubated with the antibody (ab31217, 1µg/ml) for 1h at room temperature. The secondary antibody (green) was Alexa Fluor® 488 goat antirabbit IgG (H+L) used at a 1/1000 dilution for 1h. Image-iTTM FX Signal Enhancer was used to quench autofluorescence. 5% BSA (in TBS-T) was used for all other blocking steps. DAPI was used to stain the cell nuclei (blue). Alexa Fluor® 594 WGA was used to label plasma membranes (red).



All lanes: Anti-elF4A1 antibody (ab31217) at 1 µg/ml

Lane 1: NIH/3T3 whole cell lysate (ab7179)

Lane 2 : MEF1 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lane 3: Liver (Mouse) Tissue Lysate - normal tissue

Lane 4: Mouse pancreas tissue lysate - total protein (ab29363)

Lane 5: Testis (Mouse) Tissue Lysate - normal tissue

Lane 6: PC12 (Rat adrenal pheochromocytoma cell line) Whole

Cell Lysate

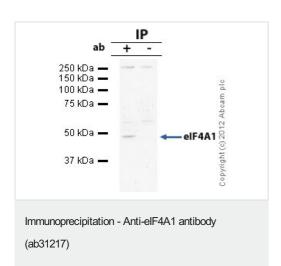
Lysates/proteins at 10 µg per lane.

Secondary

All lanes : IRDye 680 Conjugated Goat Anti-Rabbit lgG (H+L) at 1/10000 dilution

Performed under reducing conditions.

Predicted band size: 47 kDa **Observed band size:** 47 kDa



eIF4A1 was immunoprecipitated using 0.5mg Hela whole cell extract, 5µg of Rabbit polyclonal to eIF4A1 and 50µl of protein G magnetic beads (+). No antibody was added to the control (-). The antibody was incubated under agitation with Protein G beads for 10min, Hela whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation.

Proteins were eluted by addition of $40\mu l$ SDS loading buffer and incubated for 10min at $70^{o}C$; $10\mu l$ of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with ab31217.

Secondary: Mouse monoclonal [SB62a] Secondary Antibody to Rabbit lgG light chain (HRP) (ab99697).

Band: 47kDa: elF4A1

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

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