# abcam

## Product datasheet

# Anti-ADAR1 antibody [EPR7033] - BSA and Azide free ab240029



Recombinant

RabMAb

## 4 Images

#### Overview

Product name Anti-ADAR1 antibody [EPR7033] - BSA and Azide free

**Description** Rabbit monoclonal [EPR7033] to ADAR1 - BSA and Azide free

Host species Rabbit

**Specificity** The immunogen is designed to detect the p150 isoform and not the p110.

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

Unsuitable for: ICC/IF or IP

Species reactivity Reacts with: Human

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

**General notes** ab240029 is the carrier-free version of **ab126745**.

Our <u>carrier-free</u> 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 cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our <u>conjugation kits</u> 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<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> 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<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**<sup>®</sup> **patents**.

1

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

ClonalityMonoclonalClone numberEPR7033

**Isotype** IgG

#### **Applications**

#### The Abpromise guarantee

Our Abpromise guarantee covers the use of ab240029 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
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Predicted molecular weight: 136 kDa.
Flow Cyt (Intra)		Use at an assay dependent concentration. <u>ab199376</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.

**Application notes** 

Is unsuitable for ICC/IF or IP.

#### **Target**

#### **Function**

Converts multiple adenosines to inosines and creates I/U mismatched base pairs in double-helical RNA substrates without apparent sequence specificity. Has been found to modify more frequently adenosines in AU-rich regions, probably due to the relative ease of melting A/U base pairs as compared to G/C pairs. Functions to modify viral RNA genomes and may be responsible for hypermutation of certain negative-stranded viruses. Edits the messenger RNAs for glutamate receptor (GLUR) subunits by site-selective adenosine deamination. Produces low-level editing at the GLUR-B Q/R site, but edits efficiently at the R/G site and HOTSPOT1. Binds to short interfering RNAs (siRNA) without editing them and suppresses siRNA-mediated RNA interference. Binds to ILF3/NF90 and up-regulates ILF3-mediated gene expression.

**Tissue specificity** Ubiquitously expressed, highest levels were found in brain and lung.

**Involvement in disease** Defects in ADAR are a cause of dyschromatosis symmetrical hereditaria (DSH) [MIM:127400];

also known as reticulate acropigmentation of Dohi. DSH is a pigmentary genodermatosis of

autosomal dominant inheritance characterized by a mixture of hyperpigmented and hypopigmented macules distributed on the dorsal parts of the hands and feet.

**Sequence similarities**Contains 1 A to I editase domain.

Contains 2 DRADA repeats.

Contains 3 DRBM (double-stranded RNA-binding) domains.

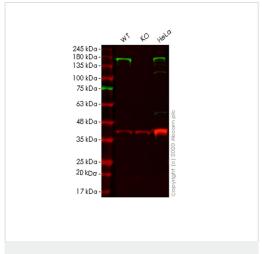
Post-translational modifications

Sumoylation reduces RNA-editing activity.

Cytoplasm. Nucleus > nucleolus. Isoform 1 is found predominantly in cytoplasm but appears to

shuttle between the cytoplasm and nucleus. Isoform 5 is found exclusively in the nucleolus.

#### **Images**



Western blot - Anti-ADAR1 antibody [EPR7033] - BSA and Azide free (ab240029)

**All lanes :** Anti-ADAR1 antibody [EPR7033] (<u>ab126745</u>) at 1/1000 dilution

Lane 1: Wild-type HEK293T cell lysate

Lane 2: ADAR knockout HEK293T cell lysate

Lane 3: HeLa cell lysate

Lysates/proteins at 20 µg per lane.

#### Secondary

**All lanes :** Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) at 1/10000 dilution

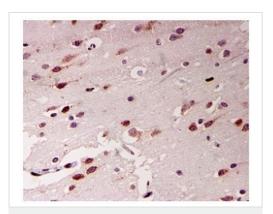
**Predicted band size:** 136 kDa **Observed band size:** 130 kDa

This data was developed using the same antibody clone in a different buffer formulation (<u>ab126745</u>).

**Lanes 1-3:** Merged signal (red and green). Green - <u>ab126745</u> observed at 130 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

<u>ab126745</u> Anti-ADAR1 antibody [EPR7033] was shown to specifically react with ADAR1 in wild-type HEK293T cells. Loss of signal was observed when knockout cell line <u>ab266846</u> (knockout cell lysate <u>ab257131</u>) was used. Wild-type and ADAR1 knockout samples were subjected to SDS-PAGE. <u>ab126745</u> and Anti-

GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



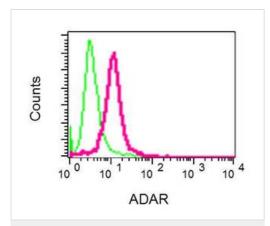
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-ADAR1 antibody

[EPR7033] - BSA and Azide free (ab240029)

<u>ab126745</u>, at 1/50 dilution, staining ADAR1 in paraffin-embedded Human brain tissue by Immunohistochemistry.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab126745</u>).

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



Flow Cytometry (Intracellular) - Anti-ADAR1 antibody [EPR7033] - BSA and Azide free (ab240029)

Intracellular flow cytometric analysis of permeabilized Ramos cells, staining ADAR1 (red) with <u>ab126745</u>. 1x10<sup>6</sup> cells were collected and washed with blocking buffer. Cells were fixed with 2% paraformaldehyde, permeabilized with 1X FACS permeabilizing solution and blocked with blocking buffer for 30 minutes at room temperature. Cells were incubated with primary antibody (1/10) for 30 minutes at room temperature before a Fluorescently-conjugated secondary antibody or 30 min at room temperature. A rabbit lgG was used as a negative control (green). This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab126745</u>).



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