abcam

Product datasheet

Anti-MELK antibody [EPR3981] - BSA and Azide free ab172889



Recombinant

RabMAb

4 Images

Overview

Product name Anti-MELK antibody [EPR3981] - BSA and Azide free

Description Rabbit monoclonal [EPR3981] to MELK - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: Flow Cyt,ICC/IF,IHC-P or IP

Species reactivity Reacts with: Human

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

Positive control WB: HCT116, HAP1, HeLa, K562, Jurkat and HEK-293T cell lysates.

General notes ab172889 is the carrier-free version of <u>ab108529</u>.

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 **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**.

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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 Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR3981

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise

Our $\underline{\textbf{Abpromise guarantee}}$ covers the use of ab172889 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		Use at an assay dependent concentration. Predicted molecular weight: 75 kDa.

Application notes Is unsuitable for Flow Cyt,ICC/IF,IHC-P or IP.

Target

Function Phosphorylates ZNF622 and may contribute to its redirection to the nucleus. May be involved in

the inhibition of spliceosome assembly during mitosis.

Tissue specificity Expressed in placenta, kidney, thymus, testis, ovary and intestine.

Sequence similarities Belongs to the protein kinase superfamily. CAMK Ser/Thr protein kinase family. SNF1 subfamily.

Contains 1 KA1 (kinase-associated) domain.

Contains 1 protein kinase domain.

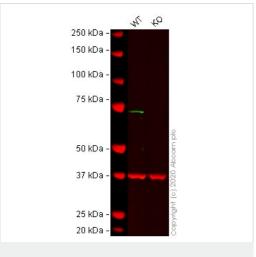
Post-translational

modifications

Autophosphorylated. Thr-478 phosphorylation during mitosis promotes interaction with PPP1R8.

Cellular localization Cytoplasm.

Images



Western blot - Anti-MELK antibody [EPR3981] - BSA and Azide free (ab172889)

All lanes : Anti-MELK antibody [EPR3981] (**ab108529**) at 1/1000 dilution

Lane 1: HCT116 cell lysate

Lane 2: MELK knockout HCT116 cell lysate

Lysates/proteins at 20 µg per lane.

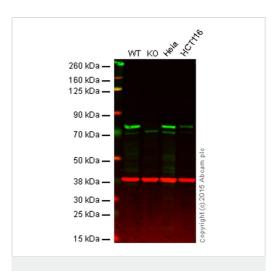
Performed under reducing conditions.

Predicted band size: 75 kDa **Observed band size:** 75 kDa

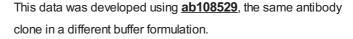
This data was developed using the same antibody clone in a different buffer formulation (<u>ab108529</u>).

Lanes 1 - 2: Merged signal (red and green). Green - <u>ab108529</u> observed at 75 kDa. Red - loading control <u>ab8245</u> (Mouse anti-GAPDH antibody [6C5]) observed at 37kDa.

ab108529 was shown to react with MELK in wild-type HCT116 cells in western blot with loss of signal observed in MELK knockout cell line ab266896 (MELK knockout cell lysate ab257537). Wild-type and MELK knockout HCT116 cell lysates were subjected to SDS-PAGE. Membranes were blocked in 3% milk in TBS-T (0.1% Tween®) before incubation with ab108529 and ab8245 (Mouse anti-GAPDH antibody [6C5]) overnight at 4°C at a 1 in 1000 dilution and a 1 in 20000 dilution respectively. Blots were incubated with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preabsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-MELK antibody [EPR3981] - BSA and Azide free (ab172889)



Lane 1: Wild-type HAP1 cell lysate (20 µg)

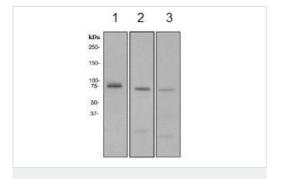
Lane 2: MELK knockout HAP1 cell lysate (20 µg)

Lane 3: HeLa cell lysate (20 µg)

Lane 4: HCT116 cell lysate (20 µg)

Lanes 1 - 4 Merged signal (red and green). Green - <u>ab108529</u> observed at 75 kDa. Red - loading control, <u>ab8226</u>, observed at 42 kDa.

ab108529 was shown to recognize MELK when MELK knockout samples were used, along with additional cross-reactive bands. Wild-type and MELK knockout samples were subjected to SDS-PAGE. **ab108529** and **ab8226** (loading control to beta Actin) were both diluted 1/1000 and incubated overnight at 4°C. 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 1/10 000 dilution for 1 h at room temperature before imaging.



Western blot - Anti-MELK antibody [EPR3981] - BSA and Azide free (ab172889)

All lanes : Anti-MELK antibody [EPR3981] (**ab108529**) at 1/1000 dilution

Lane 1 : K562 cell lysate
Lane 2 : 293T cell lysate
Lane 3 : Jurkat cell lysate

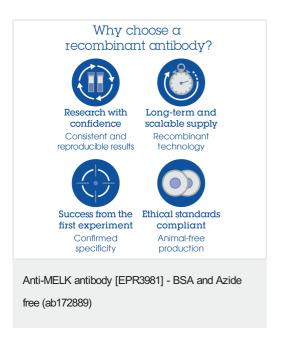
Lysates/proteins at 10 µg per lane.

Secondary

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

Predicted band size: 75 kDa

This data was developed using <u>ab108529</u>, the same antibody clone in a different buffer formulation.



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