Product datasheet

Anti-mTOR (phospho S2448) antibody [EPR426(2)]
ab109268

Overview

Product name: Anti-mTOR (phospho S2448) antibody [EPR426(2)]
Description: Rabbit monoclonal [EPR426(2)] to mTOR (phospho S2448)
Host species: Rabbit
Specificity: This antibody only detects mTOR/FRAP phosphorylated at serine 2448.
Tested applications:
Suitable for: WB, IHC-P, Dot blot
Unsuitable for: Flow Cyt, ICC/IF or IP
Species reactivity:
Reacts with: Mouse, Human, Pig
Immunogen: Synthetic peptide (the amino acid sequence is considered to be commercially sensitive) within Human mTOR (phospho S2448). The exact sequence is proprietary.
Positive control: WB: 293T, HeLa and NIH/3T3 cell lysates. IHC-P: Human breast carcinoma tissue.
General notes: Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents

We are constantly working hard to ensure we provide our customers with best in class antibodies. As a result of this work we are pleased to now offer this antibody in purified format. We are in the process of updating our datasheets. The purified format is designated 'PUR' on our product labels. If you have any questions regarding this update, please contact our Scientific Support team.

This product is a recombinant rabbit monoclonal antibody.

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. Stable for 12 months at -20°C.
Storage buffer: pH: 7.20
Preservative: 0.01% Sodium azide
Constituents: 40% Glycerol, 59% PBS, 0.05% BSA
Purity: Protein A purified
Clonality: Monoclonal
Clone number: EPR426(2)
Isotype: IgG

Applications

Our Abpromise guarantee covers the use of ab109268 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<table>
<thead>
<tr>
<th>Application</th>
<th>Abreviews</th>
<th>Notes</th>
</tr>
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<tbody>
<tr>
<td>IHC-P</td>
<td>![star_rating] 🕒</td>
<td>1/50 - 1/100. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.</td>
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<tr>
<td>Dot blot</td>
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<td>1/1000.</td>
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Application notes: Is unsuitable for Flow Cyt, ICC/IF or IP.

Target

Function: Kinase subunit of both mTORC1 and mTORC2, which regulates cell growth and survival in response to nutrient and hormonal signals. mTORC1 is activated in response to growth factors or amino-acids. Growth factor-stimulated mTORC1 activation involves AKT1-mediated phosphorylation of TSC1-TSC2, which leads to the activation of the RHEB GTPase that potently activates the protein kinase activity of mTORC1. Amino-acid-signaling to mTORC1 requires its relocalization to the lysosomes mediated by the Ragulator complex and the Rag GTPases. Activated mTORC1 up-regulates protein synthesis by phosphorylating key regulators of mRNA translation and ribosome synthesis. mTORC1 phosphorylates EIF4EBP1 and releases it from inhibiting the elongation initiation factor 4E (eIF4E). mTORC1 phosphorylates and activates S6K1 at 'Thr-421', which then promotes protein synthesis by phosphorylating PDCD4 and targeting it for degradation. Phosphorylates MAF1 leading to attenuation of its RNA polymerase III-repressive function. mTORC2 is also activated by growth factors, but seems to be nutrient-insensitive. mTORC2 plays a critical role in AKT1 'Ser-473' phosphorylation, which may facilitate the phosphorylation of the activation loop of AKT1 on 'Thr-308' by PDK1 which is a prerequisite for full activation. mTORC2 regulates the phosphorylation of SGK1 at 'Ser-422'. mTORC2 also modulates the phosphorylation of PRKCA on 'Ser-657'.

Tissue specificity: Expressed in numerous tissues, with highest levels in testis.

Sequence similarities: Belongs to the PI3/P4-kinase family.
Contains 1 FAT domain.
Contains 1 FATC domain.
Contains 7 HEAT repeats.
Contains 1 PI3K/P4K domain.

Post-translational modifications: Autophosphorylated; when part of mTORC1 or mTORC2.
Cellular localization


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268)

Immunohistochemical analysis of paraffin-embedded human breast carcinoma tissue using unpurified ab109268 at a dilution of 1/50.

Western blot - Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268)

All lanes: Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268) at 1/2000 dilution (Purified)

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

Lane 2: HeLa grown in serum-free media overnight, then treated with 200nM PMA for 4 hours whole cell lysate

Lane 3: HeLa grown in serum-free media overnight, then treated with 200nM PMA for 4 hours whole cell lysate. Then the membrane was incubated with alkaline phosphatase

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 289 kDa

Observed band size: 289 kDa

Blocking buffer: 5% NFDM/TBST

Diluting buffer: 5% NFDM/TBST
Dot blot analysis of mTOR (phospho S2448) phospho peptide (Lane 1) and mTOR non-phospho peptide (Lane 2) labeling mTOR (phospho S2448) phospho peptide with purified ab109268 at a dilution of 1/1000 (0.073µg/ml). A Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated (ab97051) was used as the secondary antibody at a dilution of 1/100,000.

Blocking buffer: 5% NFDM/TBST
Diluting buffer: 5% NFDM/TBST

All lanes: Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268) at 1/1000 dilution (purified)

Lane 1: untreated NIH/3T3 cell lysate
Lane 2: NIH/3T3 cell lysate treated with insulin
Lane 3: NIH/3T3 cell lysate treated with insulin, then the membrane treated with alkaline phosphatase

Lysates/proteins at 10 µg per lane.

Secondary
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution (HRP goat anti-rabbit IgG (H+L))

Predicted band size: 289 kDa
Observed band size: 289 kDa

Exposure time: 2 minutes

Blocking buffer: 5% NFDM/TBST
Dilution buffer: 5% NFDM/TBST
Immunohistochemical cytoplasmic and nuclear staining of paraffin embedded human endometrium carcinoma with purified ab109268 at a working dilution of 1 in 100. The secondary antibody used is ab97051, a HRP goat anti-rabbit IgG (H+L), at 1/500. The sample is counter-stained with hematoxylin. Antigen retrieval was performed using Tris-EDTA buffer, pH 9.0. PBS was used instead of the primary antibody as the negative control, and is shown in the inset.

**Western blot**

All lanes: Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268) at 1/1000 dilution (purified)

Lane 1: untreated HeLa cell lysate

Lane 2: HeLa cell lysate treated with insulin

Lane 3: HeLa cell lysate treated with insulin, and the membrane treated with alkaline phosphatase

Lysates/proteins at 10 µg per lane.

**Secondary**

All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/2000 dilution (HRP goat anti-rabbit (H+L))

Predicted band size: 289 kDa

Observed band size: 289 kDa

Exposure time: 1 minute
Blocking buffer: 2% BSA/TBST

Dilution buffer: 2% BSA/TBST

All lanes: Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268) at 1/2000 dilution (purified)

Lane 1: untreated HEK293 cell lysate
Lane 2: HEK293 cell lysate treated with alkaline phosphatase

Lysates/proteins at 10 µg per lane.

Secondary
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution (HRP goat anti-rabbit IgG (H+L))

Predicted band size: 289 kDa
Observed band size: 289 kDa

Exposure time: 3 minutes

Blocking buffer: 5% NFDM/TBST

Dilution buffer: 5% NFDM/TBST

All lanes: Anti-mTOR (phospho S2448) antibody [EPR426(2)] (ab109268) at 1/5000 dilution (unpurified)

Lane 1: untreated HEK293 cell lysate
Lane 2: HEK293 cell lysate treated with alkaline phosphatase

Lysates/proteins at 10 µg per lane.

Secondary
All lanes: Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/2000 dilution (HRP goat anti-rabbit IgG (H+L))

Predicted band size: 289 kDa
Observed band size: 289 kDa

Exposure time: 3 minutes
Blocking buffer: 5% NFDM/TBST
Dilution buffer: 5% NFDM/TBST

Please note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"