abcam

Product datasheet

Anti-S6K1 (phospho T229) antibody ab5231

14 References 1 图像

概述

产品名称 Anti-S6K1 (phospho T229)抗体

描述 兔多克隆抗体to S6K1 (phospho T229)

宿主 Rabbit

经测试应用 适用于: WB

种属反应性 与反应: Human

免疫原 Synthetic peptide corresponding to Human S6K1 (phospho T229). The sequence is conserved in

rat.

阳性对照 Hek293 cells +/- EGF; truncated p70S6K protein, an N-terminus His-tagged fusion protein

corresponding to amino acids 1-421 of human p70S6K containing a T412E mutation to become

constitutively active.

常规说明 70 kDa ribosomal protein S6 kinase (p70S6K or p70S6K beta 1) is a member of a

serine/threonine kinase family that phosphorylates the 40S ribosomal protein S6, thereby modulating the translation of ribosomal proteins and translation elongation factors. p70S6K is activated in response to the phosphatidylinositol 3-kinase (PI3K) and mitogen-activated protein kinase (MAPK) pathways, and is required for progression through the cell cycle and for cell growth. Activation of p70S6K is regulated by phosphorylation of seven different residues distributed throughout the protein, with the critical sites being threonines 229, 389, 421 and serine

424. Threonine 229 is located within the activation loop.

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

性能

形式 Liquid

存放说明 Shipped at 4°C. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

存储溶液 pH: 7.30

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Preservative: 0.05% Sodium azide Constituents: PBS, 0.1% BSA

BSA is IgG and protease free

纯**度** Immunogen affinity purified

纯**化**说明 Purified from rabbit serum by sequential epitope-specific chromatography. The antibody has

been negatively preadsorbed using a non-phosphopeptide corresponding to the site of phosphorylation to remove antibody that is reactive with non-phosphorylated p70S6K. The final

product is generated by affinity chromatography using a p70S6K derived peptide that is

phosphorylated at threonine 229.

Primary antibody说明 70 kDa ribosomal protein S6 kinase (p70S6K or p70S6K beta 1) is a member of a

serine/threonine kinase family that phosphorylates the 40S ribosomal protein S6, thereby modulating the translation of ribosomal proteins and translation elongation factors. p70S6K is activated in response to the phosphatidylinositol 3-kinase (PI3K) and mitogen-activated protein kinase (MAPK) pathways, and is required for progression through the cell cycle and for cell growth. Activation of p70S6K is regulated by phosphorylation of seven different residues distributed throughout the protein, with the critical sites being threonines 229, 389, 421 and serine

424. Threonine 229 is located within the activation loop.

克隆 多克隆

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab5231于以下的经测试应用

"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说 明
WB		Use a concentration of 0.1 - 1 μg/ml. Predicted molecular weight: 70 kDa.

靶标

功能

Acts to integrate nutrient and growth factor signals in regulation of protein synthesis, cell proliferation, cell growth, cell cycle progression and cell survival. Downstream effector of the mTOR signaling pathway. Phosphorylates specifically ribosomal protein S6 in response to insulin or several classes of mitogens. During translation initiation, the inactive form associatess with the eIF-3 complex under conditions of nutrient depletion. Mitogenic stimulation leads to phosphorylation and dissociation from the eIF-3 complex and the free activated form can phosphorylate other translational targets including EIF4B. Promotes protein synthesis by phosphorylating PDCD4 at 'Ser-67' and targeting it for degradation. Phosphorylates RICTOR leading to regulation of mammalian target of rapamycin complex 2 (mTORC2) signaling; probably phosphorylates RICTOR at 'Thr-1135'. Phosphorylates IRS1 at multiple serine residues coupled with insulin resistance; probably phosphorylates IRS1 at 'Ser-270'. Required for TNF-alpha induced IRS-1 degradation. Phosphorylates EEF2K in response to IGF1 and inhibits EEF2K activity. Phosphorylates BAD at 'Ser-99' in response to IGF1 leading to BAD inactivation and inhibition of BAD-induced apoptosis. Phosphorylates mitochondrial RMP leading to dissociation of a RMP:PPP1CC complex; probably phosphorylates RMP at 'Ser-99'. The free mitochondrial

PPP1CC can dephosphorylate RPS6KB1 at Thr-412 which is proposed to be a negative feed back mechanism for the RPS6KB1 antiapoptotic function. Phosphorylates GSK3B at 'Ser-9' under conditions leading to loss of the TSC1-TSC2 complex. Phosphorylates POLDIP3.

组织特异性 Widely expressed.

序列相似性 Belongs to the protein kinase superfamily. AGC Ser/Thr protein kinase family. S6 kinase

subfamily.

Contains 1 AGC-kinase C-terminal domain.

Contains 1 protein kinase domain.

结**构域** The autoinhibitory domain is believed to block phosphorylation within the AGC-kinase C-terminal

domain and the activation loop.

The TOS (TOR signaling) motif is essential for activation by mTORC1.

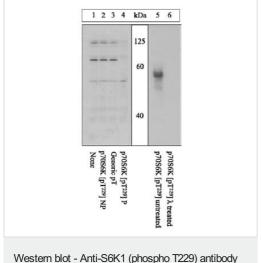
翻译后修饰 Phosphorylation at Thr-412 is regulated by mTORC1. The phosphorylation at this site is

maintained by an agonist-dependent autophosphorylation mechanism.

细胞定位 Cytoplasm: Nucleus. Cytoplasm and Cell junction > synapse > synaptosome. Mitochondrion outer

membrane.

图片



Western blot - Anti-S6K1 (phospho T229) antibody (ab5231)

Extracts prepared from Hek293 cells (1-4) or activated p70 S6 kinase protein (5, 6) were resolved on a 10% polyacrylamide gel and transferred to PVDF. Membranes were either untreated (1-5), or treated with lambda (ë) phosphatase (6), blocked with a 5% BSA-TBST buffer overnight at 4°C, then incubated with 0.50 µg/mL ab5231 for two hours at room temperature in a 3% BSA-TBST buffer, following its prior incubation with: nopeptide (1, 5, 6), the non-phosphorylated peptide corresponding to the phosphopeptide (2), a generic phosphothreonine-containing peptide (3), or, the phosphopeptide immunogen (4). After washing, membranes were incubated with goat F(ab')2 anti-rabbit lgG alkaline phosphatase and the signal was detected using the Tropix WesternStar method. The data show that only the phosphopeptide corresponding to p70S6K [pT229] blocks the antibody signal, demonstrating the specificity of the antibody. The data also show that phosphatase stripping eliminates the si

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