abcam

Product datasheet

Anti-CREB antibody [LB9] ab178322

敲除 <mark>验证</mark>

★★★★★ <u>1 Abreviews</u> <u>15 References</u> 8 图像

概述

产 品名称	Anti-CREB抗体[LB9]
描述	小鼠单克隆抗体[LB9] to CREB
宿主	Mouse
经测试应 用	适用于: ICC/IF, WB, IP, IHC-P
种属反应性	与反应: Mouse, Rat, Human, African green monkey
	预测可用于: Non human primates 🛛 📤
免疫原	Recombinant full length protein corresponding to Human CREB aa 1-341. Sequence:
	MTMESGAENQ QSGDAAVTEA ENQQMTVQAQ
	PQIATLAQVS MPAAHATSSA PTVTLVQLPN
	GQTVQVHGVI QAAQPSVIQS PQVQTVQSSC
	KDLKRLFSGT QISTIAESED SQESVDSVTD
	SQKRREILSR RPSYRKILND LSSDAPGVPR
	IEEEKSEEET SAPAITTVTV PTPIYQTSSG
	QYIAITQGGA IQLANNGTDG VQGLQTLTMT
	NAAATQPGTT ILQYAQTTDG QQILVPSNQV
	VVQAASGDVQ TYQIRTAPTS TIAPGVVMAS
	SPALPTQPAE EAARKREVRL MKNREAAREC
	RRKKKEYVKC LENRVAVLEN QNKTLIEELK
	ALKDLYCHKS D
	Database link: <u>P16220</u>
	Run BLAST with Run BLAST with
阳性 对照	HeLa cells; Human bladder and rectal cancer tissues; Human tonsil tissue; Human Jurkat, NIH/3T3, C2C12, MDA-MB-231, T98G, HeLa, 293T, HepG2, A431, A549, K562, MCF7 and U2OS whole cell lysates; Mouse MEF, 3T3L1 and C2C12 whole cell lyastes; Rat NRK whole cell lysate; Monkey COS7 whole cell lysate.
常 规说 明	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. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term. Avoid freeze / thaw cycle.
存储溶液	Preservative: 0.05% Sodium azide Constituents: 0.1% BSA, 99% PBS
纯 度	Protein A purified
克隆	单 克隆
克隆 编号	LB9
同种型	lgG1

应用

The Abpromise guarantee Abpromise™承诺保证使用ab178322于以下的经测试应用

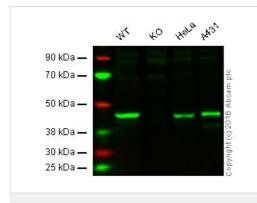
"应用说明"部分下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
ICC/IF		1/500.
WB	★★★★ <u></u> (<u>1)</u>	1/500. Detects a band of approximately 43 kDa (predicted molecular weight: 36 kDa).
IP		Use at an assay dependent concentration. Suggested use 2 $\mu g.$
IHC-P		1/200. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

靶 标	
功能	This protein binds the cAMP response element (CRE), a sequence present in many viral and cellular promoters. CREB stimulates transcription on binding to the CRE. Transcription activation is enhanced by the TORC coactivators which act independently of Ser-133 phosphorylation. Implicated in synchronization of circadian rhythmicity.
疾病相关	Defects in CREB1 may be a cause of angiomatoid fibrous histiocytoma (AFH) [MIM:612160]. A distinct variant of malignant fibrous histiocytoma that typically occurs in children and adolescents and is manifest by nodular subcutaneous growth. Characteristic microscopic features include lobulated sheets of histiocyte-like cells intimately associated with areas of hemorrhage and cystic pseudovascular spaces, as well as a striking cuffing of inflammatory cells, mimicking a lymph node metastasis. Note=A chromosomal aberration involving CREB1 is found in a patient with angiomatoid fibrous histiocytoma. Translocation t(2;22)(q33;q12) with CREB1 generates a EWSR1/CREB1 fusion gene that is most common genetic abnormality in this tumor type.

序列相似性	Belongs to the bZIP family. Contains 1 bZIP domain. Contains 1 KID (kinase-inducible) domain.
翻 译后 修 饰	Stimulated by phosphorylation. Phosphorylation of both Ser-133 and Ser-142 in the SCN regulates the activity of CREB and participates in circadian rhythm generation. Phosphorylation of Ser-133 allows CREBBP binding (By similarity). Phosphorylated upon DNA damage, probably by ATM or ATR. Sumoylated by SUMO1. Sumoylation on Lys-304, but not on Lys-285, is required for nuclear localization of this protein. Sumoylation is enhanced under hypoxia, promoting nuclear localization and stabilization.
细 胞定位	Nucleus.

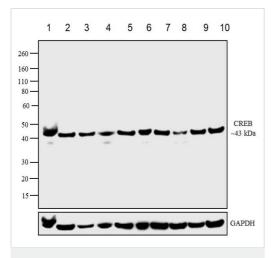
图片



Western blot - Anti-CREB antibody [LB9] (ab178322)

Lane 1: Wild-type HAP1 cell lysate (20 µg) Lane 2: CREB knockout HAP1 cell lysate (20 µg) Lane 3: HeLa cell lysate (20 µg) Lane 4: A431 cell lysate (20 µg) Lanes 1 - 4: Merged signal (red and green). Green - ab178322 observed at 44 kDa. Red - loading control, <u>ab18058</u>, observed at 124 kDa.

ab178322 was shown to recognize CREB when CREB knockout samples were used, along with additional cross-reactive bands. Wild-type and CREB knockout samples were subjected to SDS-PAGE. Ab178322 and **ab18058** (loading control to Vinculin) were diluted at 1/250 and 1/10,000 dilution respectively and incubated overnight at 4C. Blots were developed with IRDye® 800CW Goat anti-Rabbit IgG (H + L) and IRDye® 680 Goat anti-Mouse IgG (H + L) secondary antibodies at 1/10,000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-CREB antibody [LB9] (ab178322)

All lanes : Anti-CREB antibody [LB9] (ab178322) at 1/500 dilution

Lane 1 : Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate

Lane 2 : HeLa (Human cervix adenocarcinoma epithelial cell) whole cell lysate

Lane 3 : K-562 (Human chronic myelogenous leukemia lymphoblast cell line) whole cell lysate

Lane 4 : Hep G2 (Human liver hepatocellular carcinoma cell line) whole cell lysate

Lane 5 : NIH/3T3 (Mouse embryonic fibroblast cell line) whole cell lysate

Lane 6 : C2C12 whole cell lysate

Lane 7 : U-2 OS (Human bone osteosarcoma epithelial cell line) whole cell lysate

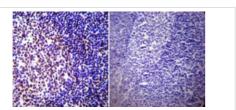
Lane 8: A549 (Human lung carcinoma cell line) whole cell lysate

Lane 9 : MDA-MB-231 (Human breast adenocarcinoma cell line) whole cell lysate

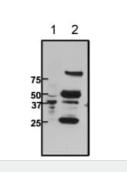
Lane 10 : T98G whole cell lysate

Lysates/proteins at 30 µg per lane.

Predicted band size: 36 kDa Observed band size: 43 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CREB antibody [LB9] (ab178322) Immunohistochemical analysis of deparaffinized Human tonsil tissue labeling CREB with ab178322 at 1/100 dilution followed by DAB staining.



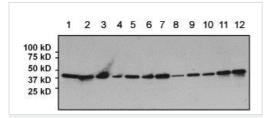
Immunoprecipitation - Anti-CREB antibody [LB9] (ab178322)

Immunoprecipitation analysis of CREB was performed on untreated 293T cells. The antigen:antbody complex was formed by binding 500µg whole cell lysate with 2µg ab178322 and captured on 50µl Protein A/G Plus Agarose.

Lane 1: 293T lysate

Lane 2: IP of 293T lysate

WB detection used ab178322 at 1/1000 followed with a goat antimouse-HRP secondary antibody at 1/20,000.



Western blot - Anti-CREB antibody [LB9] (ab178322)

All lanes : Anti-CREB antibody [LB9] (ab178322) at 1/500 dilution

Lane 1 : HeLa whole cell lysate Lane 2 : 293T whole cell lysate Lane 3 : A431 whole cell lysate Lane 4 : A549 whole cell lysate Lane 5 : MEF whole cell lysate Lane 6 : MCF7 whole cell lysate Lane 7 : U2OS whole cell lysate Lane 8 : K562 whole cell lysate Lane 9 : COS7 whole cell lysate Lane 10 : 3T3L1 whole cell lysate Lane 11 : C2C12 whole cell lysate Lane 12 : NRK whole cell lysate

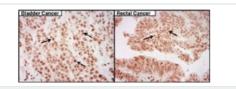
Lysates/proteins at 25 µg per lane.

Secondary

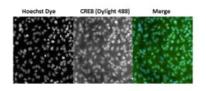
All lanes : goat anti-mouse-HRP at 1/20000 dilution

Developed using the ECL technique.

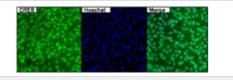
Predicted band size: 36 kDa



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CREB antibody [LB9] (ab178322) Immunohistochemical analysis of deparaffinized Human bladder cancer (left) and rectal cancer (right) tissues labeling CREB with ab178322 at 1/200 dilution followed by DAB staining.



Immunocytochemistry/ Immunofluorescence - Anti-CREB antibody [LB9] (ab178322) Immunofluorescent analysis of untreated HeLa cells (formalin-fixed, 0.1% Triton X-100 permeabilized) labeling CREB with ab178322 at 1/400 dilution (green) followed with DyLight 488 goat anti-mouse IgG secondary antibody at 1/400 dilution. Nuclei (blue) were stained with Hoechst 33342 dye.



Immunocytochemistry/ Immunofluorescence - Anti-CREB antibody [LB9] (ab178322) Immunofluorescent analysis of untreated HeLa cells (formalin-fixed, 0.1% Triton X-100 permeabilized) labeling CREB with ab178322 at 1/500 dilution (green) followed with DyLight 488 goat anti-mouse IgG secondary antibody at 1/400 dilution. Nuclei (blue) were stained with Hoechst 33342 dye.

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