abcam

Product datasheet

Anti-RHEB antibody [EPR2971] ab92313

重组 RabMAb

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概述	
产品名称	Anti-RHEB抗体[EPR2971]
描述	兔 单 克隆抗体 [EPR2971] to RHEB
宿主	Rabbit
经 测 试应 用	适用于: Flow Cyt (Intra), WB, IHC-P, ICC/IF 不适用于: IP
种属反 应性	与反应: Human
免疫原	Synthetic peptide within Human RHEB aa 50-150. The exact sequence is proprietary. (Peptide available as <u>ab194764</u>)
阳性 对照	WB: Raji and SH-SY5Y lysates. IHC-P: human glioma and brain tissue. ICC/IF: Raji cells. Flow Cyt (intra): Raji cells.
常 规说 明	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 <u>see here</u> . Our RabMAb [®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u> .

性能	
形式	Liquid
存 放 说明	Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.
存储溶液	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 59% PBS, 40% Glycerol, 0.05% BSA
纯 度	Protein A purified
克隆	单 克隆
克 隆 编号	EPR2971

应用

The Abpromise guarantee Abpromise™承诺保证使用ab92313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。

Ab评论	说明
	1/40. <u>ab172730</u> - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody. For unpurified use at 1/100 - 1/1000
	1/2000 - 1/5000. Predicted molecular weight: 20 kDa.Can be blocked with Human RHEB peptide (ab194764) .
* * * * * <u>(2)</u>	 1/500. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. Heat mediated antigen retrieval and use of an HRP/AP polymerized secondary antibody recommended. See <u>IHC antigen retrieval protocols</u> For unpurified use at 1/100 - 1/250
	1/100 - 1/250.

应用说明

Is unsuitable for IP.

靶 标	
功能	Stimulates the phosphorylation of S6K1 and EIF4EBP1 through activation of mTORC1 signaling. Activates the protein kinase activity of mTORC1. Has low intrinsic GTPase activity.
组织 特异性	Ubiquitous. Highest levels observed in skeletal and cardiac muscle.
序列相似性	Belongs to the small GTPase superfamily. Rheb family.
翻 译后 修 饰	Farnesylation is important for efficiently activating mTORC1-mediated signaling.
细 胞定位	Cell membrane.

图片



Anti-RHEB antibody [EPR2971] (ab92313) at 1/2000 dilution + Raji (Human Burkitt's lymphoma B lymphocyte) whole cell lysates at 15 µg

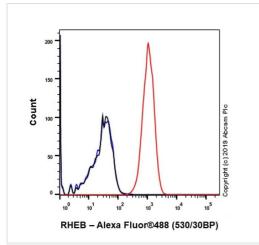
Secondary

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

Predicted band size: 20 kDa

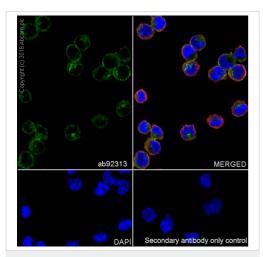
Secondary antibody only control

Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-RHEB antibody [EPR2971] (ab92313) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human glioma tissue sections labeling RHEB with purified ab92313 at 1:500 dilution (0.73 µg/ml). Heat mediated antigen retrieval was performed using Bond [™] Epitope Retrieval Solution 2 (pH 9.0). Rabbit specific IHC polymer detection kit HRP/DAB (**ab209101**) was used as the secondary antibody. Negative control: PBS instead of the primary antibody. Hematoxylin was used as a counterstain.

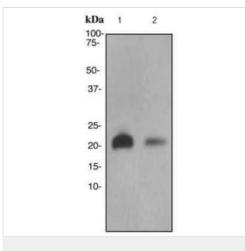


Intracellular Flow Cytometry analysis of Raji (Human Burkitt's lymphoma B lymphocyte) cells labeling RHEB with purified ab92313 at 1/40 dilution (10 µg/ml) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit IgG (Alexa Fluor[®] 488, **ab150077**) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal IgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).

Flow Cytometry (Intracellular) - Anti-RHEB antibody [EPR2971] (ab92313)



Immunocytochemistry/ Immunofluorescence - Anti-RHEB antibody [EPR2971] (ab92313) Immunocytochemistry/ Immunofluorescence analysis of Raji (Human Burkitt's lymphoma B lymphocyte) cells labeling RHEB with purified ab92313 at 1:250 dilution (1.46 µg/ml). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1:200 (2.5 µg/ml). Goat anti rabbit IgG (Alexa Fluor® 488, **ab150077**) was used as the secondary antibody at 1:1000 (2 µg/ml) dilution. DAPI (blue) was used as nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



All lanes : Anti-RHEB antibody [EPR2971] (ab92313) at 1/2000 dilution

Lane 1 : Raji cell Isyate Lane 2 : SH-SY5Y cell Isyate

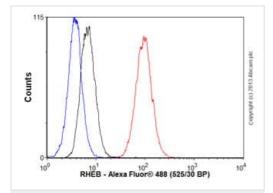
Lysates/proteins at 10 µg per lane.

Secondary

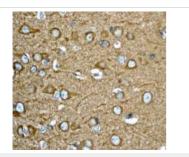
All lanes : HRP labelled goat anti-rabbit at 1/2000 dilution

Western blot - Anti-RHEB antibody [EPR2971] (ab92313)





Flow Cytometry (Intracellular) - Anti-RHEB antibody [EPR2971] (ab92313) Overlay histogram showing Raji cells stained with unpurifiedab92313 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific proteinprotein interactions followed by the antibody (ab92313, 1/1000 dilution) for 30 min at 22°C. The secondary antibody used was Alexa Fluor[®] 488 goat anti-rabbit lgG (H&L) (ab150077) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was rabbit IgG (monoclonal) (0.1µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter. This antibody gave a positive signal in Raji cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-RHEB antibody [EPR2971] (ab92313)



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Unpurified ab92313, at 1/100 dilution, staining RHEB in paraffinembedded Human brain tissue, by Immunohistochemistry.

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

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