abcam

Product datasheet

Anti-EBP50/NHERF-1 antibody [EBP-10] ab9526

13 References 2 图像

概述

产品名称	Anti-EBP50/NHERF-1抗体[EBP-10]
描述	小鼠单克隆抗体[EBP-10] to EBP50/NHERF-1
宿主	Mouse
特异性	Human ERM (ezrin/radixin/moesin)-binding phosphoprotein of 50 kDa (EBP50) / Na+/H+ exchange regulatory factor (NHERF-1)
经测试应用	**适用于:** Flow Cyt, WB
种属反应性	**与反应:** Human
免疫原	Recombinant full length protein corresponding to Human EBP50/NHERF-1.
阳性对照	RAJI human lymphoma 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 or -80°C. Avoid freeze / thaw cycle.
存储溶液	pH: 7.40 Preservative: 0.097% Sodium azide Constituent: PBS
纯度	Protein A purified
纯化说明	Purified from hybridoma culture supernatant. Purity >95% by SDS-PAGE.
克隆	单克隆
克隆编号	EBP-10
同种型	IgG2b

应用

The Abpromise guarantee **Abpromise™承诺保证使用ab9526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
Flow Cyt		Use 1µg for 10^6 cells. **ab170192** - Mouse monoclonal IgG2b, is suitable for use as an isotype control with this antibody.
WB		Use at an assay dependent concentration. Recommended dilution: 2µg/ml. Sample preparation: Lysis buffer with maltoside (1%).Application note: Non-reducing conditions with SDS-PAGE (10% separating gel).

靶标

功能 Scaffold protein that connects plasma membrane proteins with members of the ezrin/moesin/radixin family and thereby helps to link them to the actin cytoskeleton and to regulate their surface expression. Necessary for recycling of internalized ADRB2. Was first known to play a role in the regulation of the activity and subcellular location of SLC9A3. Necessary for cAMP-mediated phosphorylation and inhibition of SLC9A3. May enhance Wnt signaling. May participate in HTR4 targeting to microvilli (By similarity). Interacts with MCC.

组织特异性 Detected in liver, kidney, pancreas, prostate, spleen, small intestine and placenta, in particular in the syncytiotrophoblast.

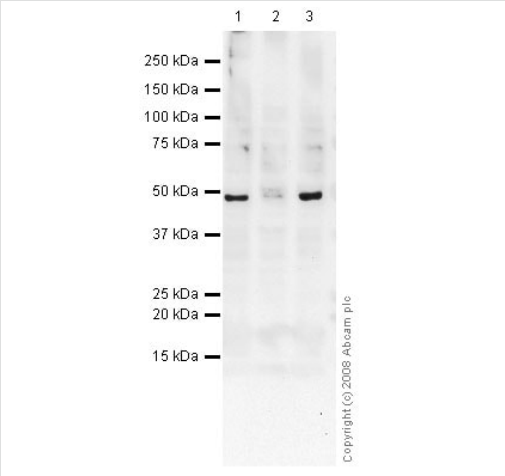
疾病相关 Defects in SLC9A3R1 are the cause of hypophosphatemic nephrolithiasis/osteoporosis type 2 (NPHLOP2) [MIM:612287]. Hypophosphatemia results from idiopathic renal phosphate loss. It contributes to the pathogenesis of hypophosphatemic urolithiasis (formation of urinary calculi) as well to that of hypophosphatemic osteoporosis (bone demineralization).

序列相似性 Contains 2 PDZ (DHR) domains.

翻译后修饰 Phosphorylated on serine residues.

细胞定位 Cytoplasm. Apical cell membrane. Endomembrane system. Cell projection > filopodium. Cell projection > ruffle. Cell projection > microvillus. Translocates from the cytoplasm to the apical cell membrane in a PODXL-dependent manner (By similarity). Colocalizes with actin in microvilli-rich apical regions of the syncytiotrophoblast. Found in microvilli, ruffling membrane and filopodia of HeLa cells. Present in lipid rafts of T-cells.

图片



Western blot - Anti-EBP50/NHERF-1 antibody [EBP-10] (ab9526)

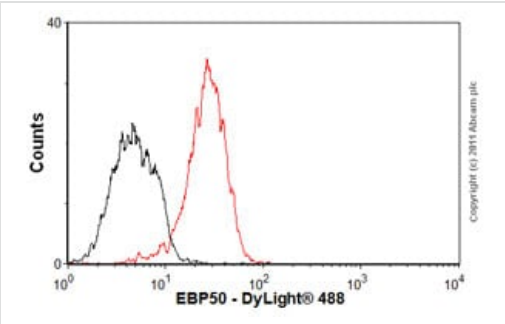
All lanes : Anti-EBP50/NHERF-1 antibody [EBP-10] (ab9526) at 2 µg/ml

Lane 1 : HepG2 (Human hepatocellular liver carcinoma cell line) Whole Cell Lysate
Lane 2 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate
Lane 3 : Caco 2 (Human colonic carcinoma cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

Secondary
All lanes : Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Observed band size: 48 kDa



Flow Cytometry - Anti-EBP50/NHERF-1 antibody [EBP-10] (ab9526)

Overlay histogram showing HepG2 cells stained with ab9526 (red line). The cells were fixed with 80% methanol (5 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 protein-protein interactions followed by the antibody (ab9526, 1µg/1x10^6 cells) for 30 min at 22ºC. The secondary antibody used was DyLight® 488 goat anti-mouse IgG (H+L) (**ab96879**) at 1/500 dilution for 30 min at 22ºC. Isotype control antibody (black line) was mouse IgG2b [PLPV219] (**ab91366**, 2µg/1x10^6 cells) used under the same conditions. Acquisition of >5,000 events was performed.

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