abcam

Product datasheet

Anti-DLK-1 antibody [MM0514-9D8] ab89908

2 References 3 图像

概述

产**品名称** Anti-DLK-1抗体[MM0514-9D8]

抽述 小鼠单克隆抗体[MM0514-9D8] to DLK-1

宿主 Mouse

经测试应用 适用于: IHC-P, Flow Cyt, ICC/IF

种属反应性 与反应: Human

免疫原 Recombinant fragment corresponding to Human DLK-1.

Database link: P80370

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. Upon delivery aliquot and store at -20°C or -80°C. Avoid repeated freeze / thaw

cycles.

存储溶液 Constituent: PBS

纯度 Protein G purified

纯**化**说明 The IgG fraction of culture supernatant was purified by Protein G affinity chromatography and 0.2

µm filtered.

克隆 单克隆

克隆编号 MM0514-9D8

同种型 lqG2b

应用

1

The Abpromise guarantee

Abpromise™承诺保证使用ab89908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
IHC-P		Use a concentration of 1 µg/ml. Perform heat mediated antigen retrieval before commencing with IHC staining protocol.
Flow Cyt		Use 2µg for 10 ⁶ cells. We recommend to fix with 4% formaldehyde for 10min at room temperature.
ICC/IF		Use a concentration of 5 µg/ml.

靶标

功能 May have a role in neuroendocrine differentiation.

组织**特异性** Found within the stromal cells in close contact to the vascular structure of placental villi, yolk sac,

fetal liver, adrenal cortex and pancreas and in the beta cells of the islets of Langerhans in the adult

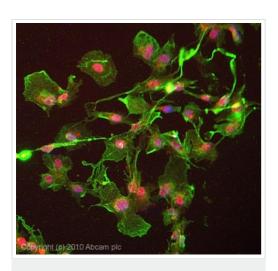
pancreas. Found also in some forms of neuroendocrine lung tumor tissue.

序列相似性 Contains 6 EGF-like domains.

翻译后修饰 N- and O-glycosylated.

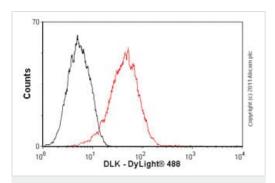
细胞定位 Membrane.

图片



Immunocytochemistry/ Immunofluorescence - Anti-DLK-1 antibody [MM0514-9D8] (ab89908)

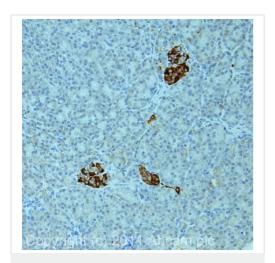
ICC/IF image of ab89908 stained HepG2 cells. The cells were 4% formaldehyde fixed (10 min) and then incubated in 1%BSA / 10% normal goat serum / 0.3M glycine in 0.1% PBS-Tween for 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab89908, 5µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 goat anti-mouse IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. DAPI was used to stain the cell nuclei (blue) at a concentration of 1.43µM.



Flow Cytometry - Anti-DLK-1 antibody [MM0514-9D8] (ab89908)

Overlay histogram showing HepG2 cells stained with ab89908 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab89908, 2µg/1x10 6 cells) for 30 min at 22 $^\circ$ C. The secondary antibody used was DyLight® 488 goat antimouse lgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22 $^\circ$ C. Isotype control antibody (black line) was mouse lgG2b [PLPV219] (ab91366, 2µg/1x10 6 cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive signal in HepG2 cells fixed with 80% methanol (5 min) used under the same conditions.

Please note that Abcam do not have any data for use of this antibody on non-fixed cells. We welcome any customer feedback.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DLK-1 antibody
[MM0514-9D8] (ab89908)

IHC image of ab89908 staining in human pancreas formalin fixed paraffin embedded tissue section, performed on a Leica BondTM system using the standard protocol F. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 20 mins. The section was then incubated with ab89908, 1µg/ml, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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