abcam

Product datasheet

Alexa Fluor® 488 Anti-MiTF antibody [D5] ab201675

★★★★★ 1 Abreviews 3 References 1 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-MiTF抗体[D5]

描述 Alexa Fluor® 488荧光小鼠单克隆抗体[D5] to MiTF

宿主 Mouse

偶联物 Alexa Fluor® 488. Ex: 495nm. Em: 519nm

经测试应用 适用于: Flow Cyt (Intra)

种属反应性 与反应: Human

免疫原 Fusion protein. This information is proprietary to Abcam and/or its suppliers.

表位 N-terminal

阳性对照 Flow Cyt (Intra): MALME-3M cells.

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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

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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.

Avoid freeze / thaw cycle. Store In the Dark.

存储溶液 pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: PBS, 1% BSA, 30% Glycerol (glycerin, glycerine)

纯**度** Affinity purified

克隆 单克隆

克隆编号 D5

同种型 lgG1

轻链类型 kappa

应用

The Abpromise guarantee Abpromise™承诺保证使用ab201675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 (Intra)		1/500.

功能 Transcription factor for tyrosinase and tyrosinase-related protein 1. Binds to a symmetrical DNA

sequence (E-boxes) (5'-CACGTG-3') found in the tyrosinase promoter. Plays a critical role in the differentiation of various cell types as neural crest-derived melanocytes, mast cells, osteoclasts

and optic cup-derived retinal pigment epithelium.

组织特异性 Isoform M is exclusively expressed in melanocytes and melanoma cells. Isoform A and isoform H

are widely expressed in many cell types including melanocytes and retinal pigment epithelium (RPE). Isoform C is expressed in many cell types including RPE but not in melanocyte-lineage

cells.

疾病相关 Defects in MITF are the cause of Waardenburg syndrome type 2A (WS2A) [MIM:193510]. It is a

dominant inherited disorder characterized by sensorineural hearing loss and patches of

depigmentation. The features show variable expression and penetrance.

Defects in MITF are a cause of Waardenburg syndrome type 2 with ocular albinism (WS2-OA)

[MIM:103470]. It is an ocular albinism with sensorineural deafness.

Defects in MITF are the cause of Tietz syndrome (TIETZS) [MIM:103500]. It is an autosomal dominant disorder characterized by generalized hypopigmentation and profound, congenital,

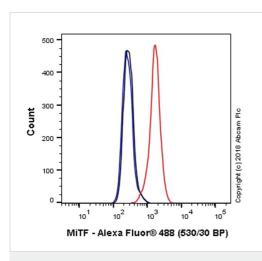
bilateral deafness. Penetrance is complete.

序列相似性 Belongs to the MiT/TFE family.

Contains 1 basic helix-loop-helix (bHLH) domain.

翻译后修饰 Phosphorylation at Ser-405 significantly enhances the ability to bind the tyrosinase promoter.

细胞定位 Nucleus.



Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-MiTF antibody [D5] (ab201675)

Overlay histogram showing MALME-3M cells stained with ab201675 (red line). The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab201675, 1/500 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was Mouse IgG1 (monoclonal) Alexa Fluor[®] 488 used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 50mW Blue laser (488nm) and 530/30 bandpass filter.

This antibody gave a positive signal in MALME-3M cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Triton X-100 for 15 min used under the same conditions.

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