

Anti-CD147 antibody [MEM-M6/6] - Low endotoxin, Azide free ab119114

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概述

产品名称	Anti-CD147抗体[MEM-M6/6] - Low endotoxin, Azide free
描述	小鼠单克隆抗体[MEM-M6/6] to CD147 - Low endotoxin, Azide free
宿主	Mouse
特异性	ab119114 recognizes Ig domain D2 (membrane proximal) of CD147 (Neurothelin).
经测试应用	适用于: Flow Cyt, ICC/IF
种属反应性	与反应: Human, Simian Virus 40
免疫原	Protein A-CR purified soluble recombinant Human CD147 (consisting of the CDNA encoding the hinge region, CH2-and CH3 domain of Human IgG1).
阳性对照	Flow Cyt: Human peripheral blood lymphocytes ICC/IF: SV40 cells.
常规说明	<p>Endotoxin level is less than 10 EU/mg of the protein, as determined by the LAL test.</p> <p>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.</p> <p>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</p>

性能

形式	Liquid
存放说明	Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.
存储溶液	pH: 7.40 Constituent: 99% PBS
无载体	Azide-free PBS 是
纯度	Protein A purified
纯化说明	ab119114 is Purified from hybridoma culture supernatant by protein-A affinity chromatography.

Purity is > 95% (by SDS-PAGE), 0.2 µm filter sterilized.

克隆 单克隆
克隆编号 MEM-M6/6
同种型 IgG1

应用

The Abpromise guarantee **Abpromise™** 承诺保证使用ab119114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. ab170190 - Mouse monoclonal IgG1, is suitable for use as an isotype control with this antibody.
ICC/IF		Use a concentration of 10 µg/ml.

靶标

功能 Plays pivotal roles in spermatogenesis, embryo implantation, neural network formation and tumor progression. Stimulates adjacent fibroblasts to produce matrix metalloproteinases (MMPS). May target monocarboxylate transporters SLC16A1, SLC16A3 and SLC16A8 to plasma membranes of retinal pigment epithelium and neural retina. Seems to be a receptor for oligomannosidic glycans. In vitro, promotes outgrowth of astrocytic processes.

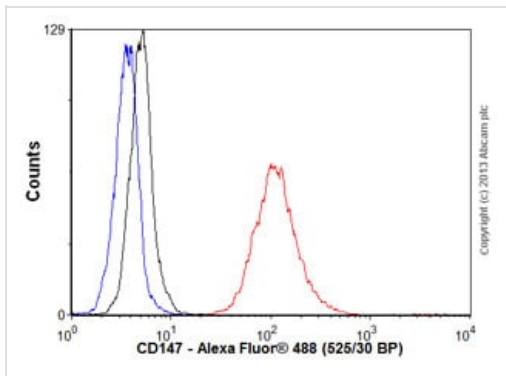
组织特异性 Present only in vascular endothelium in non-neoplastic regions of the brain, whereas it is present in tumor cells but not in proliferating blood vessels in malignant gliomas.

序列相似性 Contains 1 Ig-like C2-type (immunoglobulin-like) domain.
Contains 1 Ig-like V-type (immunoglobulin-like) domain.

翻译后修饰 N-glycosylated.

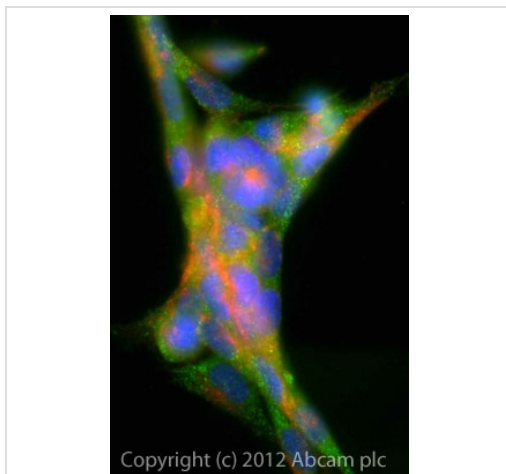
细胞定位 Cell membrane. Melanosome. Colocalizes with SLC16A1 and SLC16A8 (By similarity). Identified by mass spectrometry in melanosome fractions from stage I to stage IV.

图片



Flow Cytometry - Anti-CD147 antibody [MEM-M6/6]
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Human peripheral blood lymphocytes stained with ab119114 (red line). Human whole blood was processed using a modified protocol based on Chow *et al*, 2005 (PMID: 16080188). In brief, human whole blood was fixed in 4% formaldehyde (methanol-free) for 10 min at 22°C. Red blood cells were then lysed by the addition of Triton X-100 (final concentration - 0.1%) for 15 min at 37°C. For experimentation, cells were treated with 50% methanol (-20°C) for 15 min at 4°C. Cells were then incubated with the antibody (ab119114, 1 µg/1x10⁶ cells) for 30 min at 4°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse IgG (H&L) (**ab150113**) at 1/2000 dilution for 30 min at 4°C. Isotype control antibody (black line) was mouse IgG1 [ICIGG1] (**ab91353**, 1 µg/1x10⁶ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >30,000 total events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter. Gating strategy - peripheral blood lymphocytes.



Immunocytochemistry/ Immunofluorescence - Anti-CD147 antibody [MEM-M6/6] - Low endotoxin, Azide free (ab119114)

ICC/IF image of ab119114 stained SV40 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 (ab119114, 10 µg/ml) overnight at +4°C. The secondary antibody (green) was **ab96879**, DyLight® 488 goat anti-mouse IgG (H+L) used at a 1/250 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 1.43 µM.

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