abcam

Product datasheet

Alexa Fluor® 647 Anti-Ndufs1 antibody [EPR11521(B)] ab198956



重组 RabMAb

2 图像

概述

产品名称 Alexa Fluor® 647荧光Anti-Ndufs1抗体[EPR11521(B)]

描述 Alexa Fluor® 647荧光兔单克隆抗体[EPR11521(B)] to Ndufs1

宿主 Rabbit

偶联物 Alexa Fluor® 647. Ex: 652nm. Em: 668nm

经测试应用 适用干: ICC/IF 种属反应性 与反应: Human

预测可用于: Mouse, Rat 📤

免疫原 Synthetic peptide within Human Ndufs1. The exact sequence is proprietary.

Database link: P28331

阳性对照 ICC/IF: HepG2 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 see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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Life Technologies Corporation, 5781 Van Allen Way, Carlsbad, CA 92008 USA or **outlicensing@thermofisher.com**.

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

纯**度** Protein A purified

克隆 单克隆

克隆编号 EPR11521(B)

同种型 lgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab198956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
ICC/IF		1/100. This product gave a positive signal in HepG2 cells fixed with 100% methanol (5 min)

靶	杤
70	PJ

功能

Core subunit of the mitochondrial membrane respiratory chain NADH dehydrogenase (Complex I) that is believed to belong to the minimal assembly required for catalysis. Complex I functions in

the transfer of electrons from NADH to the respiratory chain. The immediate electron acceptor for the enzyme is believed to be ubiquinone (By similarity). This is the largest subunit of complex I and it is a component of the iron-sulfur (IP) fragment of the enzyme. It may form part of the active site

crevice where NADH is oxidized.

疾病相关 Defects in NDUFS1 are a cause of mitochondrial complex I deficiency (MT-C1D) [MIM:252010].

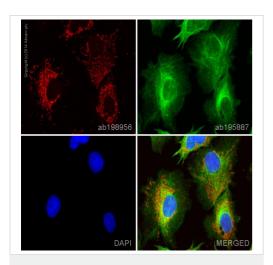
A disorder of the mitochondrial respiratory chain that causes a wide range of clinical disorders, from lethal neonatal disease to adult-onset neurodegenerative disorders. Phenotypes include macrocephaly with progressive leukodystrophy, non-specific encephalopathy, cardiomyopathy, myopathy, liver disease, Leigh syndrome, Leber hereditary optic neuropathy, and some forms of

Parkinson disease.

序列相似性 Belongs to the complex I 75 kDa subunit family.

Contains 1 2Fe-2S ferredoxin-type domain.

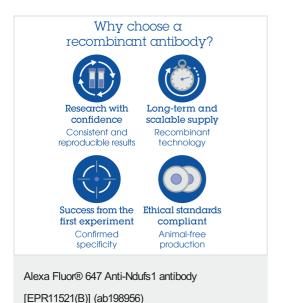
细**胞定位** Mitochondrion inner membrane.



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-Ndufs1 antibody [EPR11521(B)] (ab198956)

ab198956 staining Ndufs1 in HepG2 cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab198956 at 1/100 dilution (shown in red) and ab195887, Mouse monoclonal to alpha Tubulin (Alexa Fluor® 488), at 1/250 dilution (shown in green). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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