abcam

Product datasheet

Alexa Fluor® 488 Anti-Huntingtin antibody [EPR5526] ab197506



重组 RabMAb

2 图像

概述

产品名称 Alexa Fluor® 488荧光Anti-Huntingtin抗体[EPR5526]

描述 Alexa Fluor® 488荧光兔单克隆抗体[EPR5526] to Huntingtin

宿主 Rabbit

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

适用于: ICC/IF 经测试应用

不适用于: Flow Cyt (Intra)

种属反应性 与反应: Mouse

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ICC/IF: Neuro-2a cells. 阳性对照

This product is a recombinant monoclonal antibody, which offers several advantages including: 常规说明

- 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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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), PBS, 1% BSA

纯**度** Protein A purified

 克隆
 单克隆

 克隆编号
 EPR5526

同种型 IgG

应用

The Abpromise guarantee Abpromise™承诺保证使用ab197506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.

应用说明 Is unsuitable for Flow Cyt (Intra).

靶标

功能 May play a role in microtubule-mediated transport or vesicle function.

组织特异性 Expressed in the brain cortex (at protein level). Widely expressed with the highest level of

expression in the brain (nerve fibers, varicosities, and nerve endings). In the brain, the regions where it can be mainly found are the cerebellar cortex, the neocortex, the striatum, and the

hippocampal formation.

疾病相关 Defects in HTT are the cause of Huntington disease (HD) [MIM:143100]. HD is an autosomal

dominant neurodegenerative disorder characterized by involuntary movements (chorea), general motor impairment, psychiatric disorders and dementia. Onset of the disease occurs usually in the third or fourth decade of life and symptoms progressively worsen leading to death in 10 to 20 years. Onset and clinical course depend on the degree of poly-Gln repeat expansion, longer expansions resulting in earlier onset and more severe clinical manifestations. HD affects 1 in 10,000 individuals of European origin. Neuropathology of Huntington disease displays a distinctive pattern with loss of neurons, especially in the caudate and putamen (striatum).

序列相似性 Belongs to the huntingtin family.

Contains 10 HEAT repeats.

结构域 The N-terminal Gln-rich and Pro-rich domain has great conformational flexibility and is likely to

exist in a fluctuating equilibrium of alpha-helical, random coil, and extended conformations.

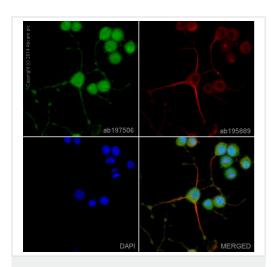
翻译后修饰 Cleaved by apopain downstream of the polyglutamine stretch. The resulting N-terminal fragment is

cytotoxic and provokes apoptosis.

Forms with expanded polyglutamine expansion are specifically ubiquitinated by SYVN1, which

Cytoplasm. Nucleus. The mutant Huntingtin protein colocalizes with AKAP8L in the nuclear matrix of Huntington's disease neurons.

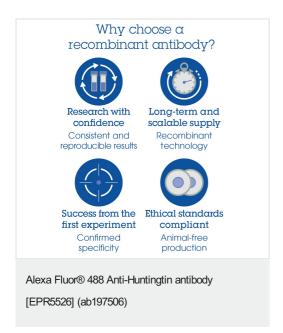
图片



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-Huntingtin antibody [EPR5526] (ab197506)

ab197506 staining Huntingtin in Neuro-2a differentiated cells. The cells were fixed with 100% methanol (5 min), permeabilised in 0.1% Triton X-100 for 5 minutes and then blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated with ab197506 at 1/100 dilution (shown in green) and ab195889, Mouse monoclonal [DM1A] to alpha Tubulin (Alexa Fluor® 594, shown in red) at 1/167 dilution overnight at +4°C. Nuclear DNA was labelled in blue with DAPI.

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



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