abcam

Product datasheet

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

יעלאעבע RabMAb

画像数 2

製品の概要

特記事項

製品名 Alexa Fluor® 488 Anti-Huntingtin antibody [EPR5526]

製品の詳細 Alexa Fluor® 488 Rabbit monoclonal [EPR5526] to Huntingtin

由来種 Rabbit

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

アプリケーション 適用あり: ICC/IF

適用なし: Flow Cyt (Intra)

種交差性 交差種: Mouse

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

ポジティブ・コントロール ICC/IF: Neuro-2a cells.

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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製品の特性

製品の状態 Liquid

保存方法 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.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

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

精製度 Protein A purified

ポリ/モノ モノクローナル **ウローン名** EPR5526

アイソタイプ IgG

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab197506の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		1/100.

追加情報 Is unsuitable for Flow Cyt (Intra).

ターゲット情報

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

組織特異性 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.

関連疾患 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).

配列類似性 Belongs to the huntingtin family.

Contains 10 HEAT repeats.

ドメイン 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.

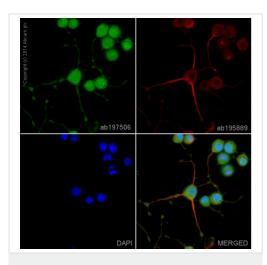
翻訳後修飾 Cleaved by apopain downstream of the polyglutamine stretch. The resulting N-terminal fragment is

cytotoxic and provokes apoptosis.

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

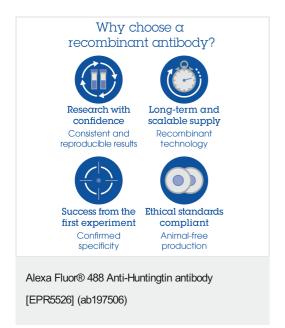
Cytoplasm. Nucleus. The mutant Huntingtin protein colocalizes with AKAP8L in the nuclear matrix of Huntington's disease neurons.

画像



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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