

Anti-vascular Amyloid 1-42 antibody [mOC31] - Conformation-Specific ab201059

重组 RabMAb

★★★★★ **2 Abreviews** **6 References** **5 图像**

概述

产品名称	Anti-vascular Amyloid 1-42抗体[mOC31] - Conformation-Specific
描述	兔单克隆抗体[mOC31] to vascular Amyloid 1-42 - Conformation-Specific
宿主	Rabbit
经测试应用	适用于: Dot blot, IHC-P, IHC-FrFI
种属反应性	与反应: Mouse, Human
免疫原	The details of the immunogen for this antibody are not available.
阳性对照	beta Amyloid (Aβ) 1-42 IHC-P: FFPE human brain Alzheimer
常规说明	<p>This antibody was developed as part of a collaboration between Abcam and Professor Charles Glabe, UC Irvine.</p> <p>ab201059 recognizes a conformation-specific discontinuous epitope of beta Amyloid that maps to segments EFGRHSGY and ED (Hatami et al. 2014). It specifically recognizes a subset of vascular amyloid that does not colocalize with thioflavin S (Hatami et al. 2014, McLean et al. 2013).</p> <p>For further information on the immunogen, please refer to Hatami et al. 2014 and Kayed et al. 2007.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

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 long

	term. Avoid freeze / thaw cycle.
存储溶液	pH: 7.2 Preservative: 0.01% Sodium azide Constituents: 59% PBS, 40% Glycerol, 0.05% BSA
纯度	Protein A purified
克隆	单克隆
克隆编号	mOC31
同种型	IgG

应用

The Abpromise guarantee

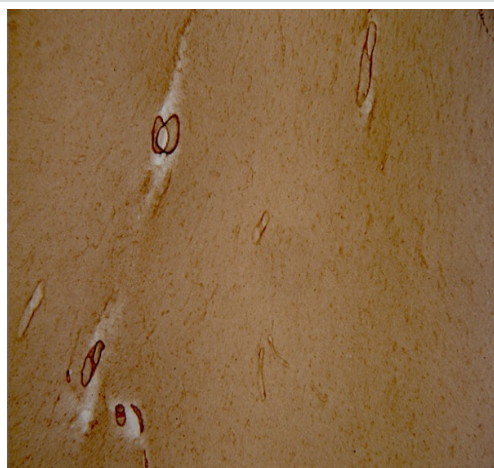
Abpromise™ 承诺保证使用ab201059于以下的经测试应用

“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
Dot blot		1/100.
IHC-P		1/250 - 1/500. Do not perform antigen retrieval.
IHC-FrFI		Use at an assay dependent concentration.

靶标

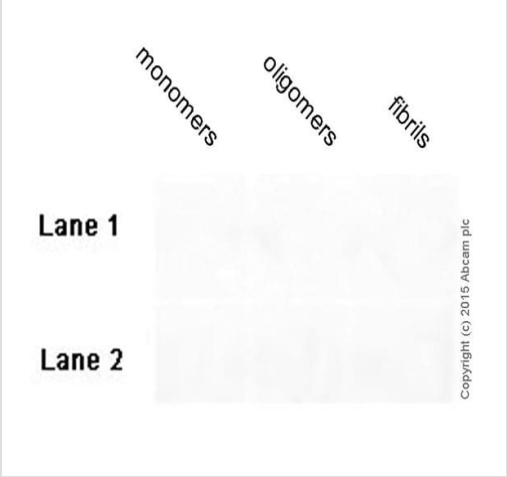
图片



Immunohistochemistry - Free Floating - Anti-vascular Amyloid 1-42 antibody [mOC31] - Conformation-Specific (ab201059)

Image courtesy of Professor Charles Glabe, UC Irvine

Immunohistochemical staining of human brain tissue from a patient with a diagnosis of Alzheimers disease, male, 81 years, 5 hour post mortem index, tangle stage 5, plaque stage B, mini mental status exam score 12. Sections were cut using a vibratome. No antigen retrieval was performed. Free floating sections were stained using ab201059 at a dilution of 50 ng/mL. The secondary antibody used was a biotinylated goat anti-rabbit at a dilution of 1/225, which was blocked with normal goat serum. The sample was visualized using ABC solution (1 hour incubation) followed by 1-4 minutes of DAB. The sample was mounted and allowed to dry overnight, followed by dehydration in increasingly concentrated ethanol solutions.



Dot Blot - Anti-vascular Amyloid 1-42 antibody
[mOC31] - Conformation-Specific (ab201059)

Negative control (secondary ab only):

Lane 1: beta Amyloid (Aβ) 1-40.

Lane 2: beta Amyloid (Aβ) 1-42.

Primary antibody omitted.

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated ([ab97051](#)) at 1/5000 dilution was used as secondary antibody.

Blocking and diluting buffer: 5% NFDM/TBST.

Exposure time: 30 seconds.

Why choose a recombinant antibody?

Research with confidence
Consistent and reproducible results

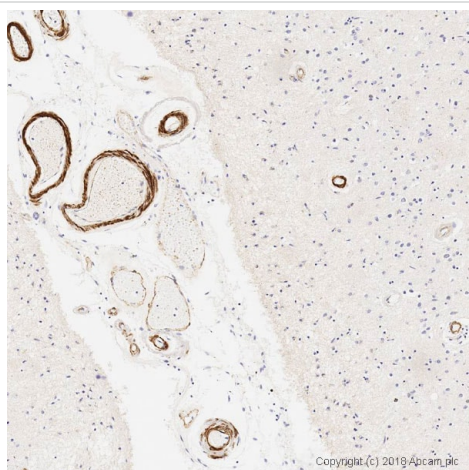
Long-term and scalable supply
Recombinant technology

Success from the first experiment
Confirmed specificity

Ethical standards compliant
Animal-free production

Anti-vascular Amyloid 1-42 antibody [mOC31] -
Conformation-Specific (ab201059)

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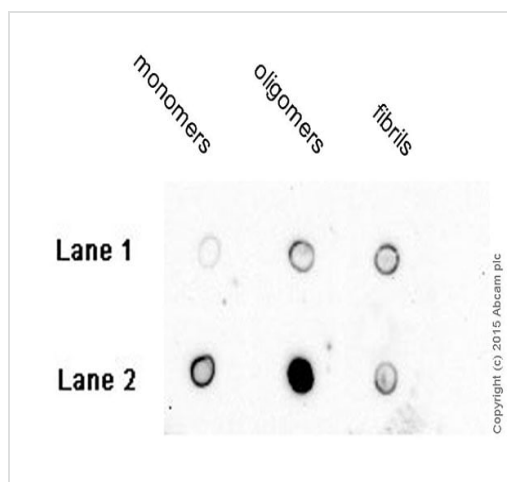


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-vascular Amyloid 1-42 antibody [mOC31] - Conformation-Specific (ab201059)

IHC image of beta Amyloid staining in a formalin fixed paraffin embedded human Alzheimer brain tissue section, performed on a Leica Bond™ system using the standard protocol F. No antigen retrieval was performed prior to staining. The section was incubated with **ab201090** at 1/500 dilution for 15 mins at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

Image showing staining of vascular Amyloid as described in Hatami et al. 2014



Dot Blot - Anti-vascular Amyloid 1-42 antibody [mOC31] - Conformation-Specific (ab201059)

Dot blot analysis of beta Amyloid 1-42 labeled with ab201059 at 1/100 dilution.

Lane 1: beta Amyloid (A β) 1-40.

Lane 2: beta Amyloid (A β) 1-42.

Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated (**ab97051**) at 1/5000 dilution was used as secondary antibody.

Blocking and diluting buffer: 5% NFDm/TBST.

Exposure time: 1 minute.

Antibody reactivity was assessed using a dot blot, which is a non-quantitative method that maintains the native conformation of beta Amyloid. Beta Amyloid 1-40 and 1-42 peptides underwent the following aggregation conditions before being spotted onto a nitrocellulose membrane and detected using ab201059:

Monomers: 0.3 mg of beta Amyloid peptide was dissolved in 30 μ l 100 mM NaOH and incubated at room temperature for 10 minutes. It was then diluted with 970 μ l of 1% SDS and boiled for five minutes.

Oligomers: 0.3 mg of beta Amyloid peptide was dissolved in 30 μ l 100 mM NaOH and incubated at room temperature for 10 minutes. It was then diluted with 970 μ l of 10 mM phosphate buffer pH 7.4 containing 0.02% sodium azide and incubated at room temperature for four days.

Fibrils: 0.3 mg of beta Amyloid peptide was dissolved in 1 ml 50% hexafluoroisopropanol (HFIP) with 0.02% sodium azide. It was then stirred constantly for nine days; the first seven with a cap on and the final two with the cap removed to allow evaporation of the HFIP. Fibrils were then sedimented at 20,000 rpm in a microcentrifuge for

20 minutes and resuspended in 1 ml of PBS + 0.02% sodium azide.

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