abcam

Product datasheet

Alexa Fluor® 568 Anti-Dopamine Transporter antibody [EPR19695] ab303466

Recombinant

RabMAb

3 Images

Overview

Product name Alexa Fluor® 568 Anti-Dopamine Transporter antibody [EPR19695]

Description Alexa Fluor® 568 Rabbit monoclonal [EPR19695] to Dopamine Transporter

Host species Rabbit

Conjugation Alexa Fluor® 568. Ex: 578nm, Em: 603nm

Tested applications Suitable for: IHC-P

Unsuitable for: IHC-Fr

Species reactivity Reacts with: Mouse, Rat

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Mouse and rat striatum FFPE tissue setions.

General notesThis 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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outlicensing@thermofisher.com.

Properties

Form Liquid

Storage instructions 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. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 68% PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

Purity Protein A purified

ClonalityMonoclonalClone numberEPR19695

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab303466 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Application notes Is unsuitable for IHC-Fr.

Target

Function Amine transporter. Terminates the action of dopamine by its high affinity sodium-dependent

reuptake into presynaptic terminals.

Involvement in disease Defects in SLC6A3 are the cause of dystonia-parkinsonism infantile (DYTPRI) [MIM:613135]. It is

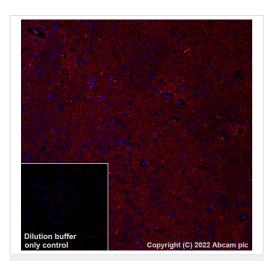
a neurodegenerative disorder characterized by infantile onset of parkinsonism and dystonia. Other neurologic features include global developmental delay, bradikinesia and pyramidal tract

signs.

Sequence similarities Belongs to the sodium:neurotransmitter symporter (SNF) (TC 2.A.22) family. SLC6A3 subfamily.

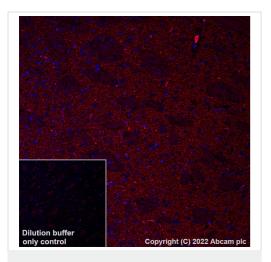
Cellular localization Membrane.

Images



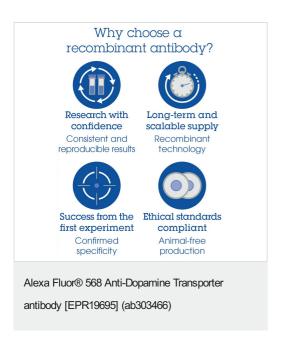
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 568 Anti-Dopamine Transporter antibody [EPR19695] (ab303466) Immunohistochemical analysis of paraffin-embedded rat striatum tissue labeling Dopamine Transporter with ab303466 at 1/100 dilution (5.0 μ g/mL). Positive staining is observed on rat striatum. The section was incubated with ab303466 for 60 mins at room temperature (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®]RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 40 mins was used.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 568 Anti-Dopamine Transporter antibody [EPR19695] (ab303466) Immunohistochemical analysis of paraffin-embedded mouse striatum tissue labeling Dopamine Transporter with ab303466 at 1/100 dilution (5.0 μ g/mL). Positive staining is observed on mouse striatum. The section was incubated with ab303466 for 60 mins at room temperature (shown in red). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®]RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

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