abcam

Product datasheet

Anti-NIPAl antibody ab121744

2 Images

Overview

Product name Anti-NIPA1 antibody

Description Rabbit polyclonal to NIPA1

Host species Rabbit

Tested applications Suitable for: ICC/IF, IHC-P

Species reactivity Reacts with: Human

Immunogen Recombinant Protein Epitope Signature Tag, corresponding to amino acids 183-226

(HGPTNIMVYI SICSLLGSFT VPSTKGIGLA AQDILHNNPS SQRA) of Human NIPA1.

Run BLAST with EXPASY IN Run BLAST with S NCBI

Positive control Human urinary bladder tissue.

General notesThe Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7.20

Preservative: 0.02% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 59% PBS

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab121744 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
ICC/IF		Use a concentration of 1 - 4 µg/ml. Recommend PFA Fixation and Triton X-100 treatment
IHC-P		1/20 - 1/50. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Target

Function Acts as a Mg(2+) transporter. Can also transport other divalent cations such as Fe(2+), Sr(2+),

Ba(2+), Mn(2+) and Co(2+) but to a much less extent than Mg(2+).

Tissue specificity Widely expressed with highest levels in neuronal tissues.

Involvement in disease Defects in NIPA1 are the cause of spastic paraplegia autosomal dominant type 6 (SPG6)

[MIM:600363]. Spastic paraplegia is a degenerative spinal cord disorder characterized by a slow,

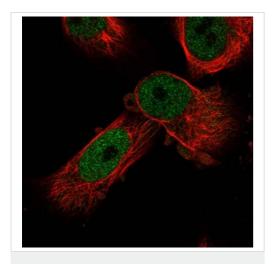
gradual, progressive weakness and spasticity of the lower limbs.

Sequence similarities Belongs to the NIPA family.

Cellular localization Cell membrane. Early endosome. Recruited to the cell membrane in response to low extracellular

magnesium.

Images



Immunocytochemistry/ Immunofluorescence - Anti-NIPA1 antibody (ab121744)

Immunofluorescent staining of Human cell line U-251MG shows positivity in nuclei but not nucleoli. Recommended concentration of ab121744 1-4 μ g/ml. Cells treated with PFA/Triton X-100.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-NIPA1 antibody (ab121744)

ab121744, at a 1/10 dilution, staining NIPA1 in paraffin-embedded Human urinary bladder tissue by Immunohistochemistry.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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