# abcam

## Product datasheet

## APC Anti-Niemann Pick Cl antibody [EPR5209] ab223985

Recombinant RabMAb

## 2 Images

#### Overview

**Product name** APC Anti-Niemann Pick C1 antibody [EPR5209]

**Description** APC Rabbit monoclonal [EPR5209] to Niemann Pick C1

**Host species** Rabbit

Conjugation APC. Ex: 645nm, Em: 660nm **Tested applications** Suitable for: Flow Cyt (Intra)

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. **Immunogen** 

Positive control Flow Cyt (intra): HepG2 cells

**General notes** 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**.

### **Properties**

**Form** Liquid

Shipped at 4°C. Upon delivery aliquot. Store at +4°C. Do Not Freeze. Store In the Dark. Storage instructions

Preservative: 0.02% Sodium azide Storage buffer

Constituents: PBS, 1% BSA

**Purity** Protein A purified

Clonality Monoclonal Clone number **EPR5209** 

**Isotype** IgG

## **Applications**

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab223985 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
Flow Cyt (Intra)		1/500. The cellular localisation of this product has been verified in ICC/IF.

Target	
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**Function** 

Involved in the intracellular trafficking of cholesterol. May play a role in vesicular trafficking in glia, a process that may be crucial for maintaining the structural and functional integrity of nerve

terminals.

Involvement in disease

Defects in NPC1 are the cause of Niemann-Pick disease type C1 (NPDC1) [MIM:257220]. A lysosomal storage disorder that affects the viscera and the central nervous system. It is due to defective intracellular processing and transport of low-density lipoprotein derived cholesterol. It causes accumulation of cholesterol in lysosomes, with delayed induction of cholesterol homeostatic reactions. Niemann-Pick disease type C1 has a highly variable clinical phenotype. Clinical features include variable hepatosplenomegaly and severe progressive neurological

dysfunction such as ataxia, dystonia and dementia. The age of onset can vary from infancy to late adulthood. An allelic variant of Niemann-Pick disease type C1 is found in people with Nova Scotia

ancestry. Patients with the Nova Scotian clinical variant are less severely affected.

Sequence similarities

Belongs to the patched family.

Contains 1 SSD (sterol-sensing) domain.

Domain

A cysteine-rich N-terminal domain and a C-terminal domain containing a di-leucine motif necessary for lysosomal targeting are critical for mobilization of cholesterol from lysosomes.

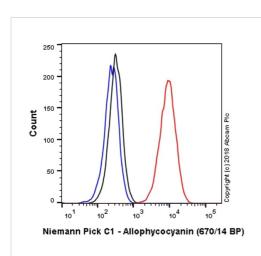
Post-translational modifications

Glycosylated.

**Cellular localization** 

Late endosome membrane. Lysosome membrane.

## **Images**

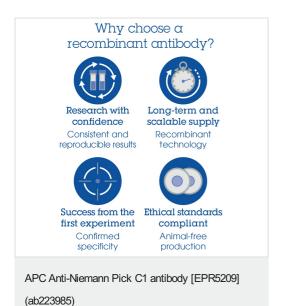


Flow Cytometry (Intracellular) - APC Anti-Niemann Pick C1 antibody [EPR5209] (ab223985) Overlay histogram showing HepG2 cells stained with ab223985 (red line). The cells were fixed with 4% formaldehyde (10 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab223985, 1/500 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was Rabbit IgG (monoclonal) Allophycocyanin (<u>ab232814</u>) used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 40 mW Red laser (640nm) and 670/14 bandpass filter.

This antibody gave a positive signal in HepG2 cells fixed with 80% methanol (5 min)/permeabilized with 0.1% PBS-Triton X-100 for 15 min used under the same conditions.



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