# **Product Name: Niemann Pick C1 Rabbit Monoclonal**

**Antibody** 

Catalog #: AMRe86256



## **Summary**

**Production Name** Niemann Pick C1 Rabbit Monoclonal Antibody

**Description** Recombinant rabbit monoclonal antibody

**Host** Rabbit

**Application** WB, IHC-P, ICC/IF, FC **Reactivity** Human,Mouse,Rat

#### **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

Clonality Monoclonal

Form Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw  $\bf Storage$ 

cycles.

Supplied in 50mM Tris-Glycine(pH 7.4), 0.15M NaCl, 40% Glycerol, 0.01% sodium

azide and 0.05% protective protein. Stable for 12 months from date of receipt.

**Purification** Affinity Purification

## **Immunogen**

Gene Name Niemann Pick C1

Alternative Names NPC
Gene ID 4864
SwissProt ID 015118.

# **Application**

**Dilution Ratio** WB: 1:1000-1:5000 IHC-P: 1:50-1:100 ICC/IF: 1:50-1:100 FC: 1:100-1:200

Molecular Weight Calculated MW:142 kDa; Observed MW:160-180 kDa

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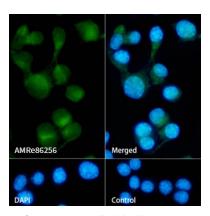


## **Background**

This gene encodes a large protein that resides in the limiting membrane of endosomes and lysosomes and mediates intracellular cholesterol trafficking via binding of cholesterol to its N-terminal domain. It is predicted to have a cytoplasmic C-terminus, 13 transmembrane domains, and 3 large loops in the lumen of the endosome - the last loop being at the N-terminus. This protein transports low-density lipoproteins to late endosomal/lysosomal compartments where they are hydrolized and released as free cholesterol. Defects in this gene cause Niemann-Pick type C disease, a rare autosomal recessive neurodegenerative disorder characterized by over accumulation of cholesterol and glycosphingolipids in late endosomal/lysosomal compartments.[provided by RefSeq, Aug 2009]

### **Research Area**

### **Image Data**



Immunofluorescence analysis of Neuro-2a cells labelling Niemann Pick C1 with AMRe86256.

#### Note

For research use only.

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