# abcam

# Product datasheet

# Anti-NMDAR1 antibody [N308/48] - Neuronal Marker ab134308

12 References 4 Images

Overview

Product name Anti-NMDAR1 antibody [N308/48] - Neuronal Marker

**Description** Mouse monoclonal [N308/48] to NMDAR1 - Neuronal Marker

Host species Mouse

Tested applications

Suitable for: Flow Cyt, WB, ICC/IF

Species reactivity

Reacts with: Mouse, Rat, Human

**Immunogen** Fusion protein corresponding to Rat NMDAR1 aa 1-400 (Hinge).

Database link: P35439

Positive control WB: Human, Mouse and Rat brain membrane tissue lysate, Neuro-2a cell lysate. ICC/IF: SK-N-

BE cells. Flow Cyt: SH-SY5Y cells.

General notes The clone number has been updated from S308-48 to N308/48, both clone numbers name the

same antibody clone.

The 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. Store at -20°C.

**Storage buffer** Preservative: 0.09% Sodium azide

Constituents: 49% PBS, 50% Glycerol

**Purity** Protein G purified

Clonality Monoclonal
Clone number N308/48

**Isotype** IgG1

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# **Applications**

# The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab134308 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		Use at an assay dependent concentration.	
WB		1/1000. Predicted molecular weight: 105 kDa.	
ICC/IF		1/100.	

# **Target**

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	411	ICTI	on

NMDA receptor subtype of glutamate-gated ion channels with high calcium permeability and voltage-dependent sensitivity to magnesium. Mediated by glycine. This protein plays a key role in synaptic plasticity, synaptogenesis, excitotoxicity, memory acquisition and learning. It mediates neuronal functions in glutamate neurotransmission. Is involved in the cell surface targeting of NMDA receptors.

# Sequence similarities

Post-translational modifications

Belongs to the glutamate-gated ion channel (TC 1.A.10.1) family. NR1/GRIN1 subfamily.

 $\label{lem:nmd} \mbox{NMDA is probably regulated by $C$-terminal phosphorylation of an isoform of NR1 by PKC.}$ 

Dephosphorylated on Ser-897 probably by protein phosphatase 2A (PPP2CB). Its

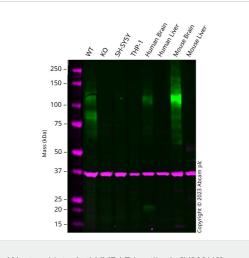
phosphorylated state is influenced by the formation of the NMDAR-PPP2CB complex and the NMDAR channel activity.

#### Cellular localization

Cell membrane. Cell junction > synapse > postsynaptic cell membrane. Cell junction > synapse > postsynaptic cell membrane > postsynaptic density. Enriched in post-synaptic plasma membrane

and post-synaptic densities.

# Images



Western blot - Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308)

**All lanes :** Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308) at 1/1000 dilution

Lane 1: Wild-type Neuro-2a cell lysate

Lane 2: GRIN1 knockout Neuro-2a cell lysate

Lane 3: SH-SY5Y UNBOILED cell lysate

Lane 4: THP-1 UNBOILED cell lysate

Lane 5: Human Brain UNBOILED cell lysate

Lane 6: Human Liver UNBOILED cell lysate

Lane 7: Mouse Brain UNBOILED cell lysate

Lane 8: Mouse Liver UNBOILED cell lysate

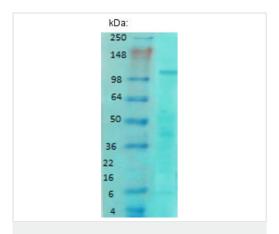
Lysates/proteins at 20 µg per lane.

Developed using the ECL technique.

Performed under reducing conditions.

**Predicted band size:** 105 kDa **Observed band size:** 120 kDa

Western blot: Anti-GRIN1 antibody [N308/48] (ab134308) staining at 1/1000 dilution, shown in green; Rabbit Anti-GAPDH antibody [EPR16891] (ab181602) loading control staining at 1/20000 dilution, shown in magenta. In Western blot, ab134308 was shown to bind specifically to GRIN1. A band was observed at 120 kDa in wild-type Neuro-2a cell lysates with no signal observed at this size in GRIN1 knockout cell line ab281960 (knockout cell lysate ab282987). To generate this image, unboiled wild-type and GRIN1 knockout Neuro-2a cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3% milk in TBS-0.1% Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4°C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Mouse IgG H&L 800CW and Goat anti-Rabbit IgG H&L 680RD at 1/20000 dilution.



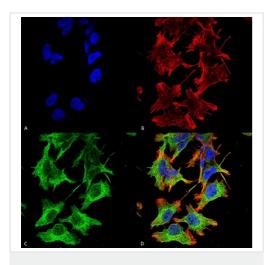
Western blot - Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308)

Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308) at 1/1000 dilution + Rat brain membrane tissue lysate

#### Secondary

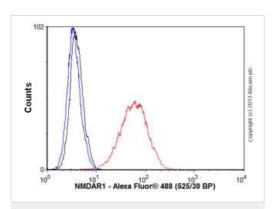
Goat anti-mouse IgG HRP

Predicted band size: 105 kDa



Immunocytochemistry/ Immunofluorescence - Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308)

SK-N-BE cells labeling NMDAR1 using ab134308 at 1/100 dilution in ICC/IF. Cells were fixed using 4% formaldehyde for 15 minutes at room temperature. Incubated with primary antibody for 1 hour at room temperature. Secondary antibody was a goat anti-mouse ATTO 488 (green) at 1/100 dilution for 1 hour at room temperature. Counterstained with Phalloidin Texas Red F-actin stain. Nuclei were stained with DAPI (blue).



Flow Cytometry - Anti-NMDAR1 antibody [N308/48] - Neuronal Marker (ab134308)

Overlay histogram showing SH-SY5Y cells stained with ab134308 (red line). The cells were fixed with 4% paraformaldehyde (10 min) and incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab134308,  $1\mu g/1x10^6$  cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse lgG (H&L) (ab150113) at 1/2000 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse lgG1 [ICIGG1] (ab91353,  $1\mu g/1x10^6$  cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control. Acquisition of >5,000 events were collected using a 20mW

Argon ion laser (488nm) and 525/30 bandpass filter. This antibody gave a positive result in 80% methanol (5 min) fixed SH-SY5Y cells used under the same conditions. Please note that Abcam do not have any data for use of this antibody on non-fixed cells. We welcome any customer feedback.

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

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