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Neuroligin1 cytoplasmic domain

Cat.No. 129 013; Polyclonal rabbit antibody, 50 µg specific antibody (lyophilized)

Data Sheet

Reconstitution/ Storage	50 μg specific antibody, lyophilized. Affinity purified with the immunogen. Albumin and azide were added for stabilization. For reconstitution add 50 μl H ₂ O to get a 1mg/ml solution in PBS. Then aliquot and store at -20°C to -80°C until use. Antibodies should be stored at +4°C when still lyophilized. Do not freeze! For detailed information, see back of the data sheet.
Applications	WB: 1: 1000 (AP staining) IP: yes ICC: not tested yet IHC: not tested yet IHC_P: not tested yet
Immunogen	Synthetic peptide corresponding to AA 737 to 754 from mouse Neuroligin1 (UniProt Id: Q99K10)
Reactivity	Reacts with: rat (Q62765), mouse (Q99K10). Other species not tested yet.
Specificity	K.O.

TO BE USED IN VITRO / FOR RESEARCH ONLY NOT TOXIC, NOT HAZARDOUS, NOT INFECTIOUS, NOT CONTAGIOUS

Background

Neuroligins form a family of postsynaptic cell surface molecules that interact with β -neurexins. They are 110-120 kDa polypeptides with homology to acetylcholine esterase. **Neuroligin 1** and neuroligin 3 are specifically localized to post-synaptic densities of excitatory synapses whereas neuroligin 2 is found exclusively on inhibitory synapses.

Mutations in neuroligin 3 and neuroligin 4 have been implicated with a rare, heritable form of autism.

Selected References for 129 013

Optogenetic control of excitatory post-synaptic differentiation through neuroligin-1 tyrosine phosphorylation. Letellier M, Lagardère M, Tessier B, Janovjak H, Thoumine O eLife (2020) 9:.. WB. IP: tested species: mouse

Cortical reorganization of the glutamate synapse in the activity-based anorexia rat model: Impact on cognition. Mottarlini F. Targa G. Bottan G. Tarenzi B. Fumagalli F. Caffino L

Journal of neurochemistry (2022) 1614: 350-365. . WB; tested species: rat

Responsivity of serotonin transporter knockout rats to short and long access to cocaine: Modulation of the glutamate signalling in the nucleus accumbens shell.

Caffino L, Mottarlini F, Targa G, Verheij MMM, Fumagalli F, Homberg JR

British journal of pharmacology (2022) 17914: 3727-3739. . WB; tested species: rat

Hevin-calcyon interaction promotes synaptic reorganization after brain injury.

Kim JH, Jung HG, Kim A, Shim HS, Hyeon SJ, Lee YS, Han J, Jung JH, Lee J, Ryu H, Park JY, et al.

Cell death and differentiation (2021) 289: 2571-2588. . WB; tested species: mouse

A sex difference in the response of the rodent postsynaptic density to synGAP haploinsufficiency.

Mastro TL, Preza A, Basu S, Chattarji S, Till SM, Kind PC, Kennedy MB

eLife (2020) 9: . . WB; tested species: rat

Prefrontal Nectin3 Reduction Mediates Adolescent Stress-Induced Deficits of Social Memory, Spatial Working Memory, and Dendritic Structure in Mice.

Wang HL, Li JT, Wang H, Sun YX, Liu R, Wang XD, Su YA, Si TM

Neuroscience bulletin (2020):.. WB; tested species: mouse

TSPAN5 Enriched Microdomains Provide a Platform for Dendritic Spine Maturation through Neuroligin-1 Clustering. Moretto E, Longatti A, Murru L, Chamma I, Sessa A, Zapata J, Hosy E, Sainlos M, Saint-Pol J, Rubinstein E, Choquet D, et al. Cell reports (2019) 295: 1130-1146.e8. . WB; tested species: mouse

Neuroligin 1 regulates spines and synaptic plasticity via LIMK1/cofilin-mediated actin reorganization.

Liu A, Zhou Z, Dang R, Zhu Y, Qi J, He G, Leung C, Pak D, Jia Z, Xie W

The Journal of cell biology (2016) 2124: 449-63. . WB; tested species: mouse

Synaptic Contacts Enhance Cell-to-Cell Tau Pathology Propagation.

Calafate S, Buist A, Miskiewicz K, Vijayan V, Daneels G, de Strooper B, de Wit J, Verstreken P, Moechars D

Cell reports (2015) 118: 1176-83. . WB; KD verified; tested species: rat

 $Promoter-like\ sequences\ regulating\ transcriptional\ activity\ in\ neurexin\ and\ neuroligin\ genes.$

Runkel F, Rohlmann A, Reissner C, Brand SM, Missler M

Journal of neurochemistry (2013) 1271: 36-47. . WB

Selected General References

Neuroligin 1 is a postsynaptic cell-adhesion molecule of excitatory synapses. Song JY, Ichtchenko K, Südhof TC, Brose N

Proceedings of the National Academy of Sciences of the United States of America (1999) 963: 1100-5.

Access the online factsheet including applicable protocols at https://sysy.com/product/129013 or scan the QR-code.



FAQ - How should I store my antibody?

Shipping Conditions

 All our antibodies and control proteins / peptides are shipped lyophilized (vacuum freezedried) and are stable in this form without loss of quality at ambient temperatures for several weeks.

Storage of Sealed Vials after Delivery

- Unlabeled and biotin-labeled antibodies and control proteins should be stored at 4°C before reconstitution. They must not be stored in the freezer when still lyophilized!
 Temperatures below zero may cause loss of performance.
- Fluorescence-labeled antibodies should be reconstituted immediately upon receipt. Long term storage (several months) may lead to aggregation.
- **Control peptides** should be kept at -20°C before reconstitution.

Long Term Storage after Reconstitution (General Considerations)

- The storage freezer must not be of the frost-free variety ("no-frost freezer"). This cycle
 between freezing and thawing (to reduce frost-build-up), which is exactly what should be
 avoided. For the same reason, antibody vials should be placed in an area of the freezer that
 has minimal temperature fluctuations, for instance towards the back rather than on a door
 shelf.
- Aliquot the antibody and store frozen (-20°C to -80°C). Avoid very small aliquots (below 20 µl)
 and use the smallest storage vial or tube possible. The smaller the aliquot, the more the stock
 concentration is affected by evaporation and adsorption of the antibody to the surface of the
 storage vial or tube. Adsorption of the antibody to the surface leads to a substantial loss of
 activity.
- The addition of glycerol to a final concentration of 50% lowers the freezing point of your stock and keeps your antibody at -20°C in liquid state. This efficiently avoids freeze and thaw cycles.

Product Specific Hints for Storage

Control proteins / peptides

• Store at -20°C to -80°C.

Monoclonal Antibodies

- Ascites and hybridoma supernatant should be stored at -20°C up to -80°C. Prolonged storage at 4°C is not recommended! Unlike serum, ascites may contain proteases that will degrade the antibodies.
- **Purified IgG** should be stored at -20°C up to -80°C. Adding a carrier protein like BSA will increase long term stability. Many of our antibodies already contain carrier proteins. Please refer to the data-sheet for detailed information.

Polyclonal Antibodies

- Crude antisera: With anti-microbials added, they may be stored at 4°C. However, frozen storage (-20°C up to -80°C) is preferable.
- Affinity purified antibodies: Less robust than antisera. Storage at -20°C up to -80°C is
 recommended. Adding a carrier protein like BSA will increase long term stability. Most of our
 antibodies already contain carrier proteins. Please refer to the data-sheet for detailed
 information.

Fluorescence-labeled Antibodies

• Store as a liquid with 1:1 (v/v) glycerol at -20°C. Protect these antibodies from light exposure.

Avoid repeated freeze-thaw cycles for all antibodies!

FAQ - How should I reconstitute my antibody?

Reconstitution

- All our purified antibodies are lyophilized from PBS. To reconstitute the antibody in PBS, add
 the amount of deionized water given in the respective datasheet. If higher volumes are
 preferred, add water as mentioned above and then the desired amount of PBS and a
 stabilizing carrier protein (e.g. BSA) to a final concentration of 2%. Some of our antibodies
 already contain albumin. Take this into account when adding more carrier protein.
 For complete reconstitution, carefully remove the lid. After adding water, briefly vortex the
 solution. You can spin down the liquid by placing the vial into a 50 ml centrifugation tube filled
 with paper.
- If desired, add small amounts of azide or thimerosal to prevent microbial growth. This is especially recommended if you want to keep an aliquot a 4°C.
- After reconstitution of fluorescence-labeled antibodies, add 1:1 (v/v) glycerol to a final
 concentration of 50%. This lowers the freezing point of your stock and keeps your antibody in
 liquid state at -20°C.
- Glycerol may also be added to unlabeled primary antibodies. It is a suitable way to avoid freezethaw cycles.
- Please refer to our tips and hints for subsequent storage of reconstituted antibodies and control peptides and proteins.