

Rudolf-Wissell-Str. 28a 37079 Göttingen, Germany

Phone: +49 551-50556-0
Fax: +49 551-50556-384
E-mail: sales@sysy.com
Web: www.sysy.com

Synapsin1

Cat.No. 106 008; Recombinant rabbit antibody, 50 µg recombinant IgG (lyophilized)

Data Sheet

50 μg purified recombinant IgG , IgO , IgO holds and a side were added for stabilization. For reconstitution add 50 μIgO 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 IgO holds and freeze! For detailed information, see back of the data sheet.
WB: 1: 1000 up to 1: 10000 (AP staining) IP: yes ICC: 1: 1000 IHC: 1: 200 up to 1: 500 IHC-P: 1: 1000 IHC-G: 1: 500 ExM: external data (see remarks) DNA-PAINT: external data (see remarks)
Rb46.1
IgG1 (κ light chain)
full-length recombinant rat Synapsin1 (UniProt Id: P09951)
AA 435 to 475 from rat Synapsin1 (UniProt Id: P09951)
Reacts with: human (P17600), rat (P09951), mouse (O88935), mammals. Weaker signal: chicken, zebrafish, other vertebrates. Other species not tested yet.
Specific for synapsin 1a and 1b independent of phosphorylation state. K.O. validated
This antibody is a chimeric antibody based on the well known monoclonal mouse antibody clone 46.1. The constant regions of the heavy and light chains have been replaced by rabbit specific sequences. Therefore, the antibody can be used with standard anti-rabbit secondary reagents. The antibody has been expressed in mammalian cells. IHC-G: 9% glyoxal fixation is recommended. ExM: This antibody has been successfully used for the epitope-preserving magnified analysis of the proteome (eMAP) expansion microscopy method (Park et al. 2021. PMID: 34767453). DNA-PAINT: This antibody has been successfully used for DNA-PAINT application (see Unterauer et al., 2024; PMID: 38552614).

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

Background

Synapsins are neuron-specific phosphoproteins that play a fundamental role in synaptic vesicle trafficking and neurotransmitter release. They are exclusively associated with small synaptic vesicles in presynaptic terminals, with little or no expression in non-neuronal tissues including neuroendocrine cells (1–4). In mammals, three distinct genes—SYN1, SYN2, and SYN3—encode more than eight isoforms through alternative splicing. Synapsin1 is one of the most specific markers of synapses throughout both the central and peripheral nervous systems. In addition to presynaptic terminals, it is localized to sensory nerve endings and peripheral innervation of the gastrointestinal tract, including the small intestine, where it contributes to neurotransmitter release in enteric and extrinsic nerves (2,3). Two splice variants, synapsin1a and synapsin1b, interact with synaptic vesicle membranes and the cytoskeletal proteins actin and spectrin (1). Synapsin2, also expressed in the nervous system, exists in at least two splice variants, whereas synapsin3 displays a more restricted distribution, being enriched in hippocampal neurons and developing neural circuits (4).

Synapsins are major neuronal phosphoproteins and substrates of several kinases, including PKA, CaMK I, and CaMK II, with synapsin1 serving as a reference substrate for calmodulin-dependent protein kinases (1,4). Beyond their established neuronal role, recent studies have implicated synapsins in glioblastoma biology. In particular, synapsin3 has been shown to promote neuronal-like differentiation of glioblastoma stem cells by antagonizing Notch signaling, thereby reducing tumor stemness and progression (5). Moreover, glioblastoma cells can exploit synaptic communication pathways, underscoring a broader role for synaptic proteins in tumor growth and plasticity (6).

Selected References for 106 008

Spatial proteomics in neurons at single-protein resolution.

Unterauer EM, Shetab Boushehri S, Jevdokimenko K, Masullo LA, Ganji M, Sograte-Idrissi S, Kowalewski R, Strauss S, Reinhardt SCM, Perovic A, Marr C, et al.

Cell (2024) 1877: 1785-1800.e16. . DNA PAINT; tested species: rat

Methods for shipping live primary cortical and hippocampal neuron cultures from postnatal mice.

Sammoura FM, Popova D, Morris A, Hart RP, Richardson JR

Current research in neurobiology (2023) 4: 100069. . ICC; tested species: mouse

Selected General References

A phospho-switch controls the dynamic association of synapsins with synaptic vesicles. Hosaka M et al. Neuron (1999) PubMed:10571231

Integrated proteogenomic characterization of glioblastoma evolution. Kim KH et al. Cancer Cell (2024) PubMed:38215747

Essential functions of synapsins I and II in synaptic vesicle regulation. Rosahl TW et al. Nature (1995) PubMed:7777057

The synaptic vesicle cycle: a cascade of protein-protein interactions. Südhof TC et al. Nature (1995) PubMed:7791897

Synaptic vesicles and exocytosis.

Jahn R et al. Annu. Rev. Neurosci. (1994) PubMed:8210174

et al. () PubMed:39994412

Access the online factsheet including applicable protocols at https://sysy.com/product/106008 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.