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# Synaptopodin

Cat.No. 163 004; Polyclonal Guinea pig antibody, 100 µl antiserum (lyophilized)

## **Data Sheet**

Reconstitution/ Storage	100 μl antiserum, lyophilized. For <b>reconstitution</b> add 100 μl H <sub>2</sub> O, then aliquot and store at -20°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 : 500 up to 1 : 1000 (AP staining) (see remarks) IP: not tested yet ICC: 1 : 500 IHC: 1 : 200 IHC-P: 1 : 200 up to 1 : 500 ExM: external data
Immunogen	Recombinant protein corresponding to residues near the central region of mouse Synaptopodin. (UniProt Id: Q8CC35)
Reactivity	Reacts with: rat (Q9Z327), mouse (Q8CC35), human (Q8N3V7). Other species not tested yet.
Specificity	This antibody detects the renal Synpo-long and the neuronal Synpo-short isofoms but is negativ for the T-variant. K.O. validated PubMed: <u>28922860</u>
Matching control	163-0P
Remarks	<b>WB</b> : The antibody is less sensitive than the rabbit antiserum (cat. no. <u>163 002</u> ).

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

### Background

Synaptopodin is the founding member of a class of proline-rich actin-associated proteins and has been originally identified in podocytes of renal glomeruli (1, 2). In brain it localizes to the post-synaptic density (PSD) and the spine aparatus (1). In humans, three isoforms of synaptopodin with a distinct expression profile have been identified, neuronal Synpo-short, renal Synpo-long and Synpo-T. All three isoforms interact with alpha-actinin and induce alpha-actinin-induced actin filaments (3). Synaptopodin deficient mice lack the dendritic spine apparatus and exhibit impaired activity-dependent long-term synaptic plasticity (4).

## Selected References for 163 004

Expansion-enhanced super-resolution radial fluctuations enable nanoscale molecular profiling of pathology specimens. Kylies D, Zimmermann M, Haas F, Schwerk M, Kuehl M, Brehler M, Czogalla J, Hernandez LC, Konczalla L, Okabayashi Y, Menzel J, et al.

Nature nanotechnology (2023) : . . IHC-P, EXM; tested species: human,mouse

Smad4 promotes diabetic nephropathy by modulating glycolysis and OXPHOS. Li J, Sun YBY, Chen W, Fan J, Li S, Qu X, Chen Q, Chen R, Zhu D, Zhang J, Wu Z, et al. EMBO reports (2020) : e48781. . **WB, IHC; tested species: mouse** 

CPT1A Protects Podocytes From Lipotoxicity and Apoptosis In Vitro and Alleviates Diabetic Nephropathy In Vivo. Xie Y, Yuan Q, Tang B, Xie Y, Cao Y, Qiu Y, Zeng J, Wang Z, Su H, Zhang C Diabetes (2024) 736: 879-895. . **IHC-P, ICC; tested species: mouse,human** 

Pathology-oriented multiplexing enables integrative disease mapping. Kuehl M, Okabayashi Y, Wong MN, Gernhold L, Gut G, Kaiser N, Schwerk M, Gräfe SK, Ma FY, Tanevski J, Schäfer PSL, et al. Nature (2025) : . . **IHC-P; tested species: mouse** 

DOT1L protects against podocyte injury in diabetic kidney disease through phospholipase C-like 1. Hu Y, Ye S, Kong J, Zhou Q, Wang Z, Zhang Y, Yan H, Wang Y, Li T, Xie Y, Chen B, et al. Cell communication and signaling : CCS (2024) 221: 519. . **IHC; tested species: mouse** 

Maintenance of Lognormal-Like Skewed Dendritic Spine Size Distributions in Dentate Granule Cells of TNF, TNF-R1, TNF-R2, and TNF-R1/2-Deficient Mice.

Rößler N, Smilovic D, Vuksic M, Jedlicka P, Deller T The Journal of comparative neurology (2024) 5327: e25645. . **IHC; tested species: mouse** 

The proteasome modulates endocytosis specifically in glomerular cells to promote kidney filtration. Sachs W, Blume L, Loreth D, Schebsdat L, Hatje F, Koehler S, Wedekind U, Sachs M, Zieliniski S, Brand J, Conze C, et al. Nature communications (2024) 151: 1897. **IHC; tested species: mouse** 

Loss of tumor necrosis factor (TNF)-receptor 1 and TNF-receptor 2 partially replicate effects of TNF deficiency on dendritic spines of granule cells in mouse dentate gyrus. Smilovic D, Rietsche M, Fellenz M, Drakew A, Vuksic M, Deller T The Journal of comparative neurology (2023) 5312: 281-293. . **IHC; tested species: mouse** 

The classical pathway triggers pathogenic complement activation in membranous nephropathy. Seifert L, Zahner G, Meyer-Schwesinger C, Hickstein N, Dehde S, Wulf S, Köllner SMS, Lucas R, Kylies D, Froembling S, Zielinski S, et al.

Nature communications (2023) 141: 473. . IHC-P; tested species: mouse

Asparaginyl endopeptidase protects against podocyte injury in diabetic nephropathy through cleaving cofilin-1. Lei C, Li M, Qiu Y, Xie Y, Hao Z, Yin X, Zhang Z, Su H, Yang L, Lin J, Hammes HP, et al. Cell death & disease (2022) 132: 184. . **IHC; tested species: mouse** 

An ex vivo culture model of kidney podocyte injury reveals mechanosensitive, synaptopodin-templating, sarcomere-like structures.

Jiang S, Alisafaei F, Huang YY, Hong Y, Peng X, Qu C, Puapatanakul P, Jain S, Miner JH, Genin GM, Suleiman HY, et al. Science advances (2022) 835: eabn6027. . **ICC; tested species: human,mouse** 



Access the online factsheet including applicable protocols at <u>https://sysy.com/product/163004</u> 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.