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## SNAP29

Cat.No. 111 303; 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 <b>reconstitution</b> add 50 μl H <sub>2</sub> 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 IP: not tested yet ICC: 1: 500 IHC: yes IHC-P: not tested yet
Immunogen	Recombinant protein corresponding to AA 1 to 257 from rat SNAP29 (UniProt Id: Q9Z2P6)
Reactivity	Reacts with: human (O95721), rat (Q9Z2P6), mouse (Q9ERB0), hamster. Other species not tested yet.
Specificity	K.O. validated PubMed: <u>34069872</u>
Remarks	ICC: The following fixatives are possible: methanol, 4% formaldehyde/PFA

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

#### Background

**SNAP29**, also known as **GS32**, is an ubiquitously distributed relative of SNAP25 and SNAP23 that is ubiquitously distributed among intracellular membranes and that is also found in the cytosol of mammalian cells. As a Q-SNARE it forms SNARE complexes in vitro but its precise role in intracellular membrane traffic is not known.

#### Selected References for 111 303

SNAP-25 gene family members differentially support secretory vesicle fusion. Arora S, Saarloos I, Kooistra R, van de Bospoort R, Verhage M, Toonen RF Journal of cell science (2017) 13011: 1877-1889. . **WB, ICC** 

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

Lysosomal exocytosis releases pathogenic a-synuclein species from neurons in synucleinopathy models.

Xie YX, Naseri NN, Fels J, Kharel P, Na Y, Lane D, Burré J, Sharma M Nature communications (2022) 131: 4918. . **WB; tested species: mouse** 

Generation and Characterization of a CRISPR/Cas9-Mediated SNAP29 Knockout in Human Fibroblasts.

Martens MC, Edelkamp J, Seebode C, Schäfer M, Stählke S, Krohn S, Jung O, Murua Escobar H, Emmert S, Boeckmann L International journal of molecular sciences (2021) 2210: . . WB; KO verified; tested species: human

Cardiac SNARE Expression in Health and Disease.

Bowman PRT, Smith GL, Gould GW

Frontiers in endocrinology (2019) 10: 881. . WB; tested species: mouse

Dynamics of the mouse brain cortical synaptic proteome during postnatal brain development.

Gonzalez-Lozano MA, Klemmer P, Gebuis T, Hassan C, van Nierop P, van Kesteren RE, Smit AB, Li KW
Scientific reports (2016) 6: 35456. . **WB** 

The SNARE protein vti1a functions in dense-core vesicle biogenesis.

Walter AM, Kurps J, de Wit H, Schöning S, Toft-Bertelsen TL, Lauks J, Ziomkiewicz I, Weiss AN, Schulz A, Fischer von Mollard G, Verhage M, et al.

The EMBO journal (2014) 3315: 1681-97. . WB; tested species: mouse

Quantitative proteomic and genetic analyses of the schizophrenia susceptibility factor dysbindin identify novel roles of the biogenesis of lysosome-related organelles complex 1.

Gokhale A, Larimore J, Werner E, So L, Moreno-De-Luca A, Lese-Martin C, Lupashin VV, Smith Y, Faundez V The Journal of neuroscience: the official journal of the Society for Neuroscience (2012) 3211: 3697-711. . WB

#### **Selected General References**

A SNARE complex mediating fusion of late endosomes defines conserved properties of SNARE structure and function. Antonin W et al. EMBO J. (2000) PubMed:11101518

Selective interaction of complexin with the neuronal SNARE complex. Determination of the binding regions. Pabst S et al. J. Biol. Chem. (2000) PubMed:10777504

GS32, a novel Golgi SNARE of 32 kDa, interacts preferentially with syntaxin 6. Wong SH et al. Mol. Biol. Cell (1999) PubMed:9880331

Membrane fusion and exocytosis.

Jahn R et al. Annu. Rev. Biochem. (1999) PubMed:10872468

Access the online factsheet including applicable protocols at <a href="https://sysy.com/product/111303">https://sysy.com/product/111303</a> 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.