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# a-Tubulin

Cat.No. 302 211; Recombinant mouse antibody, 100 µg recombinant IgG (lyophilized)

# **Data Sheet**

Reconstitution/ Storage	100 $\mu$ g purified recombinant IgG, lyophilized. Albumin and azide were added for stabilization. For <b>reconstitution</b> add 100 $\mu$ l 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 up to 1: 5000 (AP staining)  IP: yes  ICC: 1: 500 up to 1: 1000  IHC: 1: 500  IHC_P: 1: 500 up to 1: 1000  ELISA: yes (see remarks)
Clone	3A2
Subtype	IgG1 (κ light chain)
Immunogen	Synthetic peptide corresponding to residues near the carboxy terminus of human α-Tubulin 4A. (UniProt Id: P68366)
Reactivity	Reacts with: human (P68366), rat, mouse, vertebrates, invertebrates, yeast. Other species not tested yet.
Specificity	Specific for α-tubulin (glu- and tyr-α-tubulin)
Matching control	302-21P
Remarks	<b>ELISA</b> : Suitable as capture antibodies for sandwich-ELISA with cat. no. 302 203 as detector antibody.

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

### **Background**

Microtubules are involved in a wide variety of cellular activities ranging from mitosis and transport events to cell movement and the maintainance of cell shape.

Tubulin itself is a globular protein which consists of two polypeptides,  $\mathbf{a}$ -tubulin and  $\beta$ -tubulin.  $\alpha$ - and  $\beta$ -tubulin dimers are assembled to 13 protofilaments that form a microtubule of 22 nm diameter. Assembled microtubules can be detyrosinated by a carboxypeptidaseS called vasohibins / SVBPs. Detyrosinated  $\alpha$ -tubulin is referred to as  $\mathbf{Glu}$ - $\mathbf{a}$ -tubulin and occurs for exemple in neurons. This reaction can be reverted by Tubulin tyrosine ligase (TTL) that ads a C-terminal tyrosin to Glu  $\alpha$ -tubulin. Another post-translational modification of  $\alpha$ -tubulin is C-terminal polyglutamylation which is also characteristic for microtubules in neuronal cells and the mitotic spindle. A third variant of detyrosinated  $\alpha$ -tubulin is  $\Delta 2$ -tubulin which lacks the C-terminal glutamic acid. It cannot be tyrosinated by TTL and is one of the dominant  $\alpha$ -tubulin isoforms in neurons.

#### Selected References for 302 211

Heat denaturation enables multicolor X10-STED microscopy. Saal KA, Shaib AH, Mougios N, Crzan D, Opazo F, Rizzoli SO Scientific reports (2023) 131: 5366. EXM; tested species: rat

Regulated Dynamic Trafficking of Neurexins Inside and Outside of Synaptic Terminals.

Neupert C, Schneider R, Klatt O, Reissner C, Repetto D, Biermann B, Niesmann K, Missler M, Heine M

The Journal of neuroscience: the official journal of the Society for Neuroscience (2015) 3540: 13629-47.. ICC

Liprin-α2 promotes the presynaptic recruitment and turnover of RIM1/CASK to facilitate synaptic transmission. Spangler SA, Schmitz SK, Kevenaar JT, de Graaff E, de Wit H, Demmers J, Toonen RF, Hoogenraad CC The Journal of cell biology (2013) 2016: 915-28. • WB; tested species: rat

Reduced synaptic depression in human neurons carrying homozygous disease-causing STXBP1 variant L446F. Öttl M. Toonen RF. Verhage M

Human molecular genetics (2024):.. WB; tested species: stem cells

Tomosyn affects dense core vesicle composition but not exocytosis in mammalian neurons. Subkhangulova A, Gonzalez-Lozano MA, Groffen AJA, van Weering JRT, Smit AB, Toonen RF, Verhage M eLife (2023) 12: . . WB; tested species: mouse

A Versatile Synaptotagmin-1 Nanobody Provides Perturbation-Free Live Synaptic Imaging And Low Linkage-Error in Super-Resolution Microscopy.

Queiroz Zetune Villa Real K, Mougios N, Rehm R, Sograte-Idrissi S, Albert L, Rahimi AM, Maidorn M, Hentze J, Martínez-Carranza M, Hosseini H, Saal KA, et al.

Small methods (2023): e2300218. . ICC; tested species: rat

Reduced dynamin-1 levels in neurons lacking MUNC18-1.
Lammertse HCA, Moro A, Saarloos I, Toonen RF, Verhage M
Journal of cell science (2022) 13522:.. WB; tested species: mouse

The molecular chaperone Cysteine String Protein is required for monomeric SNARE proteins to assemble in trans complexes during human sperm acrosomal exocytosis.

Montero KF, Berberián MV, Mayorga LS, Tomes CN, Ruete MC

Biology of reproduction (2022):.. WB; tested species: human

Plasticity-Related Gene 5 Is Expressed in a Late Phase of Neurodifferentiation After Neuronal Cell-Fate Determination. Gross I, Brandt N, Vonk D, Köper F, Wöhlbrand L, Rabus R, Witt M, Heep A, Plösch T, Hipp MS, Bräuer AU, et al. Frontiers in cellular neuroscience (2022) 16: 797588. **WB; tested species: mouse** 

Access the online factsheet including applicable protocols at <a href="https://sysy.com/product/302211">https://sysy.com/product/302211</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.