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# **Parvalbumin**

Cat.No. 195 011; Monoclonal mouse antibody, 100 µg purified IgG (lyophilized)

### **Data Sheet**

Reconstitution/ Storage	100 μg purified IgG, lyophilized. Albumin and azide were added for stabilization. For <b>reconstitution</b> add 100 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: yes (see remarks) IP: yes ICC: 1:500 IHC: 1:500 up to 1:1000 IHC-P: 1:500
Clone	58E1
Subtype	IgG1 (κ light chain)
Immunogen	Full-length recombinant rat Parvalbumin (UniProt Id: P02625)
Reactivity	Reacts with: rat (P02625), mouse (P32848). No signal: zebrafish. Other species not tested yet.
Matching control	195-0P
Remarks	<b>WB</b> : The rabbit polyclonal antiserum (cat. no. 195 002) is recommended for westernblotting. Due to its small size, a tricine gel should be used.

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

### Background

**Parvalbumin** is a small, acidic calcium binding protein and belongs to the family of EF hand proteins. The protein is found in skeletal muscle and the brain of vertebrates where it locates to a specific population of GABAergic interneurons. This subset of neurons may contribute to maintaining the balance between excitation and inhibition in the cortex and the hippocampus.

### Selected References for 195 011

Isolated P/Q Calcium Channel Deletion in Layer VI Corticothalamic Neurons Generates Absence Epilepsy. Bomben VC. Aiba I. Oian J. Mark MD. Herlitze S. Noebels JL

The Journal of neuroscience: the official journal of the Society for Neuroscience (2016) 362: 405-18. . **IHC; tested species:** mouse

Adolescent intermittent ethanol exposure decreases perineuronal nets in the hippocampus in a sex dependent manner: Modulation through pharmacological inhibition of RPTPB/ζ.

Galán-Llario M, Gramage E, García-Guerra A, Torregrosa AB, Gasparyan A, Navarro D, Navarrete F, García-Gutiérrez MS, Manzanares J, Herradón G

Neuropharmacology (2024) 247: 109850. . IHC; tested species: mouse

Investigations on the Ability of the Insular Cortex to Process Peripheral Immunosuppression.

Bihorac J, Salem Y, Lückemann L, Schedlowski M, Doenlen R, Engler H, Mark MD, Dombrowski K, Spoida K, Hadamitzky M Journal of neuroimmune pharmacology: the official journal of the Society on NeuroImmune Pharmacology (2024) 191: 40.

#### IHC; tested species: rat

Sexually dimorphic role for insular perineuronal nets in aversion-resistant alcohol consumption.

Martins de Carvalho L, Chen H, Sutter M, Lasek AW

Frontiers in psychiatry (2023) 14: 1122423. . IHC; tested species: mouse

An enriched environment ameliorates the reduction of parvalbumin-positive interneurons in the medial prefrontal cortex caused by maternal separation early in life.

Irie K, Ohta KI, Ujihara H, Araki C, Honda K, Suzuki S, Warita K, Otabi H, Kumei H, Nakamura S, Koyano K, et al.

Frontiers in neuroscience (2023) 17: 1308368. . **IHC; tested species: mouse** 

The role of  $\alpha$ -tubulin tyrosination in controlling the structure and function of hippocampal neurons.

Hosseini S, van Ham M, Erck C, Korte M, Michaelsen-Preusse K

Frontiers in molecular neuroscience (2022) 15: 931859. . IHC; tested species: mouse

Activity-dependent reconnection of adult-born dentate granule cells in a mouse model of frontotemporal dementia.

Terreros-Roncal J, Flor-García M, Moreno-Jiménez EP, Pallas-Bazarra N, Rábano A, Sah N, van Praag H, Giacomini D, Schinder AF, Ávila J, Llorens-Martín M, et al.

The Journal of neuroscience: the official journal of the Society for Neuroscience (2019):.. IHC; tested species: mouse

The transgenic mouse line Igsf9-eGFP allows targeted stimulation of inferior olive efferents.

Pätz C, Brachtendorf S, Eilers J

Journal of neuroscience methods (2018) 296: 84-92. . IHC; tested species: mouse

Mechanisms of Functional Hypoconnectivity in the Medial Prefrontal Cortex of Mecp2 Null Mice.

Sceniak MP, Lang M, Enomoto AC, James Howell C, Hermes DJ, Katz DM

Cerebral cortex (New York, N.Y.: 1991) (2016) 265: 1938-1956. . IHC; tested species: mouse

### **Selected General References**

Quantitative analysis of parvalbumin-immunoreactive cells in the human epileptic hippocampus. Andrioli A et al. Neuroscience (2007) PubMed:17850980

Expression patterns of calretinin, calbindin and parvalbumin and their colocalization in neurons during development of Macaca monkey retina.

Hendrickson A et al. Exp. Eye Res. (2007) PubMed:17845803

Access the online factsheet including applicable protocols at <a href="https://sysy.com/product/195011">https://sysy.com/product/195011</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.