

GluN 1

Cat.No. 114 011; Monoclonal mouse antibody, 100 µg purified IgG (lyophilized)

Data Sheet

Reconstitution/ Storage	100 µg purified IgG, lyophilized. For reconstitution add 100 µl H ₂ O to get a 1mg/ml solution in PBS. Then aliquot and store at -20°C until use.
Applications	WB: 1 : 1000 up to 1 : 10000 (AP staining) IP: yes (see remarks) ICC: 1 : 1000 (see remarks) IHC: not tested yet IHC-P/FFPE: not tested yet ELISA: yes (see remarks)
Clone	M68
Subtype	IgG2b (κ light chain)
Immunogen	Recombinant protein corresponding to AA 660 to 811 from rat GluN1 (UniProt Id: P35439)
Epitop	Epitop: AA 660 to 811 from rat GluN1 (UniProt Id: P35439)
Reactivity	Reacts with: human (Q05586), rat (P35439), mouse (P35438), zebrafish. Other species not tested yet.
Specificity	Specific for GluN 1.
Remarks	IP: For most effective IP use the solubilization protocol described in this ELISA protocol. Consider that protein-protein interaction may be affected. ICC: For proteins of the post-synaptic density (PSD) para-formaldehyde fixation is not recommended. A methanol fixation is more suitable and gives better results; recommended protocol. This antibody is suitable for the surface staining of living cells. After washing cells with bound antibodies, they can be fixed and visualized with secondary reagents. ELISA: Suitable as capture antibody for sandwich-ELISA with cat. no. 114 003 as detector antibody (protocol for sandwich-ELISA).

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

GluNs (NMDA-receptors) represent a class of glutamate receptors that are of central importance in synaptic plasticity. Multiple NMDA receptor subtypes exist: **GluN 1** and GluN 2 A-D. GluN 1 is the most important as it is required for activity. NMDA-receptors allow Ca²⁺ influx and are thought to trigger Ca²⁺ dependent postsynaptic processes involved in long term potentiation and depression.

Selected References SYSY Antibodies

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- Fusion Competent Synaptic Vesicles Persist upon Active Zone Disruption and Loss of Vesicle Docking.
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 Aguayo FI, Tejos-Bravo M, Díaz-Vélez G, Pacheco A, García-Rojo G, Corrales W, Olave FA, Aliaga E, Ulloa JL, Avalos AM, Román-Albasini L, et al. Frontiers in molecular neuroscience (2018) 11: 283. **WB; tested species: rat**
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 McKenzie C, Spanova M, Johnson A, Kainrath S, Zheden V, Sitte HH, Janovjak H. Journal of neuroscience methods (2018) : . **WB; tested species: rat**
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