

## Synaptopodin Recombinant Antibody [PSH10-45] - Rat IgG1 (Chimeric)

# HA601402



<b>Product Type:</b>	Recombinant Chimeric Antibody, primary antibodies
<b>Species reactivity:</b>	Human, Mouse, Rat
<b>Applications:</b>	IHC-Fr, IHC-P, WB
<b>Molecular Wt:</b>	Predicted band size: 100 kDa
<b>Clone number:</b>	PSH10-45

**Description:** Synaptopodin is a protein that in humans is encoded by the SYNPO gene. Synaptopodin is an actin-associated protein that may play a role in actin-based cell shape and motility. The name synaptopodin derives from the protein's associations with postsynaptic densities and dendritic spines and with renal podocytes.

**Immunogen:** Recombinant protein within Mouse Synaptopodin aa 1-929.

**Positive control:** Human kidney tissue, mouse brain tissue, mouse kidney tissue, rat striatum tissue, Mouse brain tissue lysate, Rat brain tissue lysate.

**Subcellular location:** Cytoplasm, cytoskeletonm Cell junction, tight junctionm Perikaryonm Cell projection, dendritic spinem Postsynaptic densitym Synapse, Cytoplasm, cytosol.

**Database links:** SwissProt: Q8N3V7 Human | Q8CC35 Mouse | Q9Z327 Rat

### Recommended Dilutions:

<b>IHC-Fr</b>	1:500
<b>IHC-P</b>	1:200-1:500
<b>WB</b>	1:2,000

**Storage Buffer:** PBS (pH7.4), 0.1% BSA, 40% Glycerol. Preservative: 0.05% Sodium Azide.

**Storage Instruction:** Shipped at 4°C. Store at +4°C short term (1-2 weeks). It is recommended to aliquot into single-use upon delivery. Store at -20°C long term.

**Purity:** Protein A affinity purified.

Hangzhou Huaan Biotechnology Co., Ltd.

Orders:0086-571-88062880

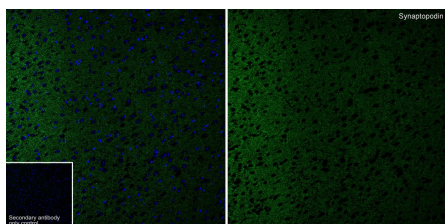
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## Images

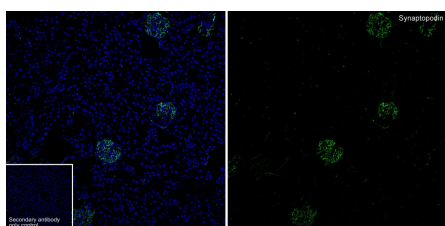
**Fig1:** Immunofluorescence analysis of frozen mouse brain tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/500 dilution.



The section was pre-treated using 1% SDS buffer (in PBS, pH 7.4) for 5 minutes at room temperature.

The tissues were blocked in 10% negative goat serum for 1 hour at room temperature, washed with PBS, and then probed with the primary antibody (HA601402, green) at 1/500 dilution overnight at 4 °C, washed with PBS. Goat Anti-Rat IgG H&L (iFluor™ 488, HA1133) was used as the secondary antibody at 1/1,000 dilution. Nuclei were counterstained with DAPI (blue).

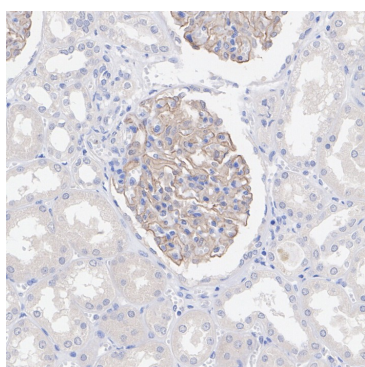
**Fig2:** Immunofluorescence analysis of frozen mouse kidney tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/500 dilution.



The section was pre-treated using 1% SDS buffer (in PBS, pH 7.4) for 5 minutes at room temperature.

The tissues were blocked in 10% negative goat serum for 1 hour at room temperature, washed with PBS, and then probed with the primary antibody (HA601402, green) at 1/500 dilution overnight at 4 °C, washed with PBS. Goat Anti-Rat IgG H&L (iFluor™ 488, HA1133) was used as the secondary antibody at 1/1,000 dilution. Nuclei were counterstained with DAPI (blue).

**Fig3:** Immunohistochemical analysis of paraffin-embedded human kidney tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/200 dilution.



The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA601402) at 1/200 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

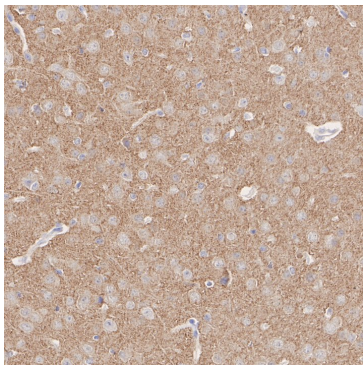
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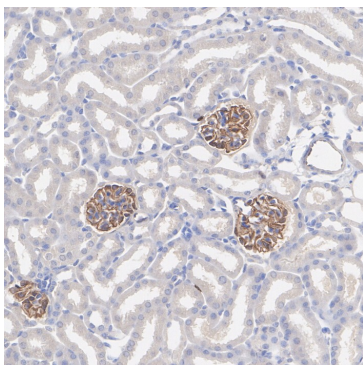
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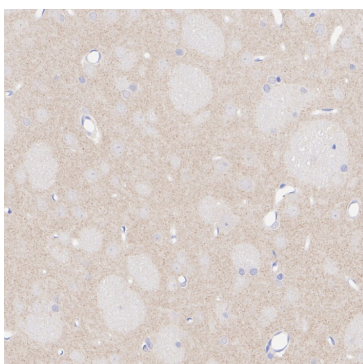
**Fig4:** Immunohistochemical analysis of paraffin-embedded mouse brain tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/500 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA601402) at 1/500 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.



**Fig5:** Immunohistochemical analysis of paraffin-embedded mouse kidney tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/200 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA601402) at 1/200 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

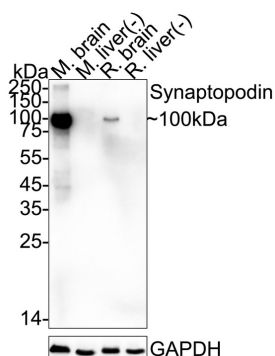


**Fig6:** Immunohistochemical analysis of paraffin-embedded rat striatum tissue with Rat anti-Synaptopodin antibody (HA601402) at 1/500 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA601402) at 1/500 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.

**Fig7:** Western blot analysis of Synaptopodin on different lysates with Rat anti-Synaptopodin antibody (HA601402) at 1/2,000 dilution.

Lane 1: Mouse brain tissue lysate  
Lane 2: Mouse liver tissue lysate (negative)  
Lane 3: Rat brain tissue lysate  
Lane 4: Rat liver tissue lysate (negative)



Lysates/proteins at 20 µg/Lane.

Predicted band size: 100 kDa  
Observed band size: 100 kDa

Exposure time: 46 seconds; ECL: K1801;  
4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (HA601402) at 1/2,000 dilution was used in primary antibody dilution (K1803) at 4 °C overnight. Goat Anti-Rat IgG H&L - HRP Secondary Antibody (HA1023) at 1/5,000 dilution was used for 1 hour at room temperature.

**Note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

## Background References

1. Wu PY et al. Synaptopodin: a key regulator of Hebbian plasticity. *Front Cell Neurosci.* 2024 Nov
2. Kruse P et al. Synaptopodin Regulates Denervation-Induced Plasticity at Hippocampal Mossy Fiber Synapses. *Cells.* 2024 Jan

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