# Aldolase A/B/C Recombinant Antibody [PSH01-81] - Rat IgG1 (Chimeric)

# HA601512

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| **Product Type:** | Recombinant Chimeric Antibody, primary antibodies |
| **Species reactivity:** | Human, Mouse, Rat |
| **Applications:** | IHC-Fr, IHC-P, WB |
| **Molecular Wt:** | Predicted band size: 40 kDa |
| **Clone number:** | PSH01-81 |

**Description:** ALDOC is a member of the class I fructose-biphosphate aldolase gene family. Expressed specifically in the hippocampus and Purkinje cells of the brain, Aldolase C is a glycolytic enzyme that catalyzes the reversible aldol cleavage of fructose-1,6-biphosphate and fructose 1-phosphate to dihydroxyacetone phosphate and either glyceraldehyde-3-phosphate or glyceraldehyde, respectively.

**Immunogen:** Synthetic peptide within Human Aldolase C aa 50-100.

**Positive control:** Mouse brain tissue, rat brain tissue, U-87 MG cell lysate, RAW264.7 cell lysate, C6 cell lysate, Mouse brain tissue lysate, Rat brain tissue lysate.

**Subcellular location:** cytoskeleton, cytosol, extracellular exosome, extracellular region

**Database links:** SwissProt: P09972 Human | P05063 Mouse | P09117 Rat

**Recommended Dilutions:**

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| **IHC-Fr** | 1:500 |
| **IHC-P** | 1:2,000 |
| **WB** | 1:5,000 |

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

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

**Purity:** Protein A affinity purified.

## Images

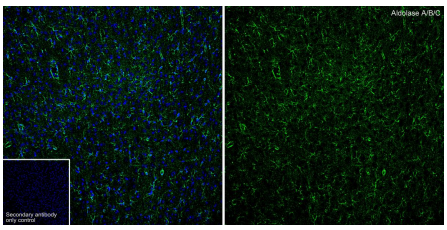


**Fig1:** Application: IHC-Fr

Species: Mouse

Site: brain

Sample: Frozen section

Antibody concentration: 1/500

Antigen retrieval: Not required

**Fig2:** Immunohistochemical analysis of paraffin-embedded mouse brain tissue with Rat anti-Aldolase A/B/C antibody (HA601512) at 1/2,000 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 $ddH_2O$ and PBS, and then probed with the primary antibody (HA601512) at 1/2,000 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.

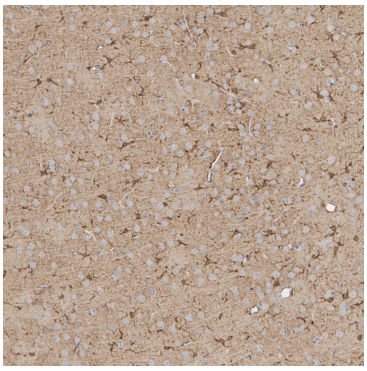

**Fig3:** Immunohistochemical analysis of paraffin-embedded rat brain tissue with Rat anti-Aldolase A/B/C antibody (HA601512) at 1/2,000 dilution.

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 $ddH_2O$ and PBS, and then probed with the primary antibody (HA601512) at 1/2,000 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.

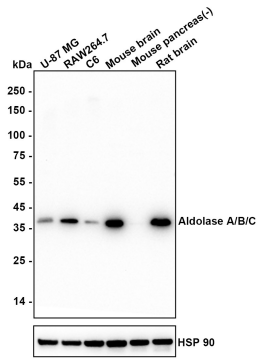


**Fig4:** Western blot analysis of Aldolase A/B/C on different lysates with Rat anti-Aldolase A/B/C antibody (HA601512) at 1/5,000 dilution.

Lane 1: U-87 MG cell lysate
Lane 2: RAW264.7 cell lysate
Lane 3: C6 cell lysate
Lane 4: Mouse brain tissue lysate
Lane 5: Mouse pancreas tissue lysate (negative)
Lane 6: Rat brain tissue lysate

Lysates/proteins at 20 µg/Lane.

Predicted band size: 40 kDa
Observed band size: 40 kDa

Exposure time: 4 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 (HA601512) at 1/5,000 dilution was used in primary antibody dilution (K1803) at 4℃ overnight. Goat Anti-Rat IgG H&L - HRP Secondary Antibody (HA1023) at 1/50,000 dilution was used for 1 hour at room temperature.



## Background References

1. Chen L, Zeng Y, Ren B, et al. ALDOC regulated the biological function and immune infiltration of gastric cancer cells. Int J Biochem Cell Biol. 2023 May; 158:106407.
2. Fan K, Wang J, Sun W, et al. MUC16 C-terminal binding with ALDOC disrupts the ability of ALDOC to sense glucose and promotes gallbladder carcinoma growth. Exp Cell Res. 2020 Sep 1;394(1):112118.