## **Anti-SOX1 Antibody [PSH10-51]**

## **HA723215**



Product Type: Recombinant Rabbit monoclonal IgG, primary antibodies

Species reactivity: Human, Mouse, Rat, Cynomolgus monkey, Pig

Applications: WB, IHC-Fr, IHC-P

Molecular Wt: Predicted band size: 39 kDa

Clone number: PSH10-51

**Description:** Sox genes comprise a family of genes that are related to the mammalian sex determining

gene SRY. These genes similarly contain sequences that encode for the HMG-box domain, which is responsible for the sequence-specific DNA-binding activity. Sox genes encode putative transcriptional regulators implicated in the decision of cell fates during development and the control of diverse developmental processes. The highly complex group of Sox genes cluster at least 40 different loci that rapidly diverged in various animal lineages. At present, 30 Sox genes have been identified. Members of this family have been shown to be conserved during evolution and to play key roles during animal development. Some are

involved in human diseases, including sex reversal.

**Immunogen:** Recombinant protein within Human SOX1 aa 1-391.

Positive control: Human cerebellum tissue, mouse cerebellum tissue, mouse E14.5 embryonic eyeball tissue,

rat cerebellum tissue, rat E14.5 embryo tissue, NCCIT cell lysate, Mouse brain tissue lysate,

Rat brain tissue lysate.

Subcellular location: Nucleus.

Database links: SwissProt: 000570 Human | P53783 Mouse

**Recommended Dilutions:** 

WB 1:2,000 IHC-Fr 1:500 IHC-P 1:2,000

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.

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Technical:0086-571-89986345

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## **Images**

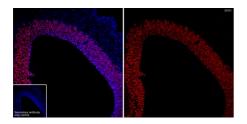


Fig1: Application: IHC-Fr

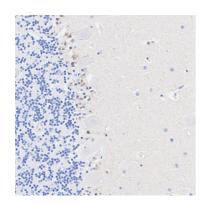
Species: Mouse

Site: E14.5 embryonic brain

Sample: Frozen section

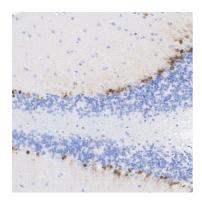
Antibody concentration: 1:500

Antigen retrieval: Not required



**Fig2:** Immunohistochemical analysis of paraffin-embedded human cerebellum tissue with Rabbit anti-SOX1 antibody (HA723215) at 1/2,000 dilution.

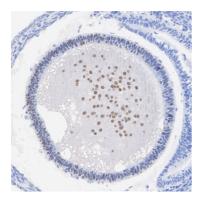
The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA723215) at 1/2,000 dilution for 1 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 mouse cerebellum tissue with Rabbit anti-SOX1 antibody (HA723215) at 1/2,000 dilution.

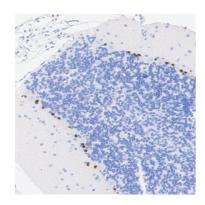
The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH<sub>2</sub>O and PBS, and then probed with the primary antibody (HA723215) at 1/2,000 dilution for 1 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.

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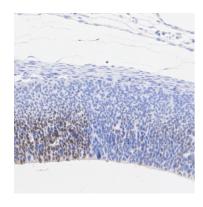
**Fig4:** Immunohistochemical analysis of paraffin-embedded mouse E14.5 embryonic eyeball tissue with Rabbit anti-SOX1 antibody (HA723215) at 1/2,000 dilution.

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<sub>2</sub>O and PBS, and then probed with the primary antibody (HA723215) 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.



**Fig5:** Immunohistochemical analysis of paraffin-embedded rat cerebellum tissue with Rabbit anti-SOX1 antibody (HA723215) 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 $_2$ O and PBS, and then probed with the primary antibody (HA723215) 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.



**Fig6:** Immunohistochemical analysis of paraffin-embedded rat E14.5 embryo tissue with Rabbit anti-SOX1 antibody (HA723215) 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<sub>2</sub>O and PBS, and then probed with the primary antibody (HA723215) 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.

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kDa 250-150-150-100-75-55-45-35-25-14**Fig7:** Western blot analysis of SOX1 on different lysates with Rabbit anti-SOX1 antibody (HA723215) at 1/2,000 dilution.

Lane 1: NCCIT cell lysate

Lane 2: Mouse brain tissue lysate Lane 3: Rat brain tissue lysate

Lysates/proteins at 20 µg/Lane.

Predicted band size: 39 kDa Observed band size: 39 kDa

Exposure time: 3 minutes; ECL: K1801;

4-20% SDS-PAGE gel.

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

## **Background References**

- 1. Sun X et al. Anti-SOX1 Antibodies in Paraneoplastic Neurological Syndrome. J Clin Neurol. 2020 Oct
- 2. Kanwore K et al. SOX1 Is a Backup Gene for Brain Neurons and Glioma Stem Cell Protection and Proliferation. Mol Neurobiol. 2021 Jun