# abcam

# Product datasheet

# Anti-SOX10 antibody [EPR4007] ab155279



★★★★★ <u>7 Abreviews</u> <u>78 References</u> 6 Images

### Overview

**Product name** Anti-SOX10 antibody [EPR4007]

Rabbit monoclonal [EPR4007] to SOX10 **Description** 

**Host species** Rabbit

**Tested applications** Suitable for: IHC-Fr, ICC/IF, Flow Cyt (Intra), WB

Species reactivity Reacts with: Mouse, Rat, Human

**Immunogen** Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: Human brain, SH-SY5Y, A375, Mouse brain, Neuro-2a, and Rat brain lysates; ICC/IF: C6

cells. Flow Cyt (intra): A-375 cells. IHC-Fr: Mouse cerebellum

**General notes** This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

### **Properties**

**Form** Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

**Purity** Protein A purified

Clonality Monoclonal Clone number **EPR4007** 

Isotype lgG

### **Applications**

### The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab155279 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-Fr		1/50. Perform heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20).
ICC/IF	****(3)	1/500. For unpurified use at 1/250 - 1/500.
Flow Cyt (Intra)		1/200.
WB	<b>★★★★★ (1)</b>	1/1000 - 1/10000. Predicted molecular weight: 49 kDa.

### **Target**

### **Function**

Transcription factor that seems to function synergistically with the POU domain protein TST-1/OCT6/SCIP. Could confer cell specificity to the function of other transcription factors in developing and mature glia.

# Tissue specificity Involvement in disease

Expressed in fetal brain and in adult brain, heart, small intestine and colon.

Defects in SOX10 are the cause of Waardenburg syndrome type 2E (WS2E) [MIM:611584]. WS2 is a genetically heterogeneous, autosomal dominant disorder characterized by sensorineural deafness, pigmentary disturbances, and absence of dystopia canthorum. The frequency of deafness is higher in WS2 than in WS1.

Defects in SOX10 are a cause of Waardenburg syndrome type 4C (WS4C) [MIM:613266]; also known as Waardenburg-Shah syndrome. WS4C is characterized by the association of Waardenburg features (depigmentation and deafness) and the absence of enteric ganglia in the distal part of the intestine (Hirschsprung disease).

Defects in SOX10 are a cause of Yemenite deaf-blind hypopigmentation syndrome (YDBHS) [MIM:601706]. YDBHS consists of cutaneous hypopigmented and hyperpigmented spots and patches, microcornea, coloboma and severe hearing loss. Another case observed in a girl with similar skin symptoms and hearing loss but without microcornea or coloboma is reported as a mild form of this syndrome.

Defects in SOX10 are the cause of peripheral demyelinating neuropathy, central dysmyelinating leukodystrophy, Waardenburg syndrome, and Hirschsprung disease (PCWH) [MIM:609136]; also called neurologic variant of Waardenburg-Shah syndrome. PCWH is a rare, complex and more severe neurocristopathy that includes features of 4 distinct syndromes: peripheral demyelinating neuropathy, central dysmyelinating leukodystrophy, Waardenburg syndrome, and Hirschsprung disease.

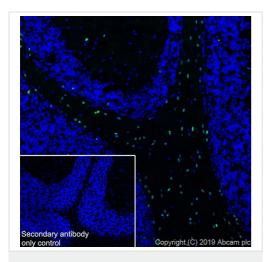
### Sequence similarities

Contains 1 HMG box DNA-binding domain.

### **Cellular localization**

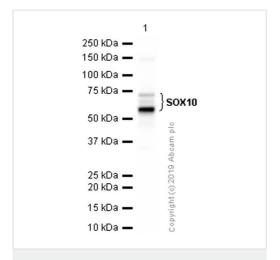
Cytoplasm. Nucleus.

## **Images**



Immunohistochemistry (Frozen sections) - Anti-SOX10 antibody [EPR4007] (ab155279)

Immunohistochemistry (Frozen) analysis of mouse cerebellum tissue sections labeling SOX10 with purified ab155279 at 1/50 (2.2  $\mu$ g/ml). Goat anti rabbit lgG (Alexa Fluor 488, ab150077) at 1/1000 (2  $\mu$ g/ml) was used as the secondary antibody. Sections were fixed with 4% paraformaldehyde and permeabilised with 0.2% Triton X-100. Negative control: PBS instead of the primary antibody. DAPI (blue) was used as nuclear counterstain. Heat mediated antigen retrieval using sodium citrate buffer (10mM citrate pH 6.0 + 0.05% Tween-20) was performed.



Western blot - Anti-SOX10 antibody [EPR4007] (ab155279)

Anti-SOX10 antibody [EPR4007] (ab155279) at 1/2000 dilution + A375 (Human malignant melanoma epithelial cell) whole cell lysate at 15  $\mu g$ 

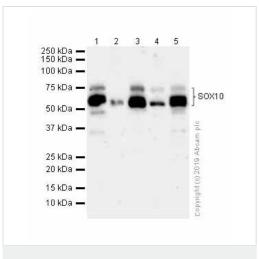
# **Secondary**

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 49 kDa Observed band size: 56-75 kDa

Exposure time: 3 seconds

Blocking and diluting buffer: 5% NFDM/TBST



Western blot - Anti-SOX10 antibody [EPR4007] (ab155279)

**All lanes :** Anti-SOX10 antibody [EPR4007] (ab155279) at 1/1000 dilution (Purified)

Lane 1: Human brain lysates

Lane 2: SH-SY5Y (Human neuroblastoma epithelial cell) whole cell

lysates

Lane 3: Mouse brain lysates

Lane 4: Neuro-2a (Mouse neuroblastoma neuroblast) whole cell

lysates

Lane 5: Rat brain lysates

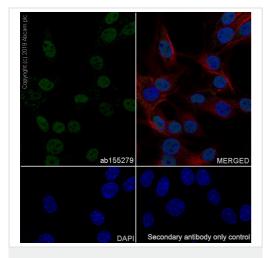
Lysates/proteins at 20 µg per lane.

### **Secondary**

**All lanes :** Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

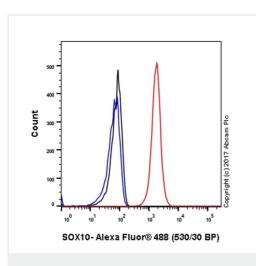
**Predicted band size:** 49 kDa **Observed band size:** 56-75 kDa

The bands observed are consistent with what have been described in PMID: 21423190



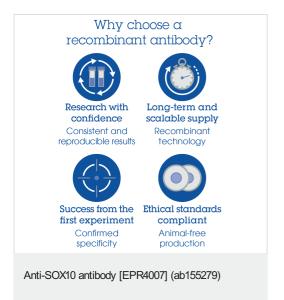
Immunocytochemistry/ Immunofluorescence - Anti-SOX10 antibody [EPR4007] (ab155279)

Immunocytochemistry/ Immunofluorescence analysis of C6 (Rat glial tumor glial cell) cells labeling SOX10 with purified ab155279 at 1:50 dilution (3.6 μg/ml). Cells were fixed in 4% Paraformaldehyde and permeabilized with 0.1% tritonX-100. Cells were counterstained with Ab195889 Anti-alpha Tubulin antibody [DM1A] - Microtubule Marker (Alexa Fluor® 594) 1:200 (2.5 μg/ml). Goat anti rabbit lgG (Alexa Fluor® 488, **ab150077**) was used as the secondary antibody at 1:1000 (2 μg/ml) dilution. DAPI (blue) was used as nuclear counterstain. PBS instead of the primary antibody was used as the secondary antibody only control.



Flow Cytometry (Intracellular) - Anti-SOX10 antibody [EPR4007] (ab155279)

Intracellular Flow Cytometry analysis ofA-375 (human malignant melanoma cell line) cells labeling with purified ab155279 at 1/200 dilution (1ug/ml) (Red). Cells were fixed with4% paraformaldehydeand permeabilised with 90% methanol. A Goat anti rabbit lgG (Alexa Fluor® 488) (ab150077) (1/2000 dilution) was used as the secondary antibody.Rabbit monoclonal lgG (Black) (ab172730) was used as a isotype control.Cell without incubation with primary antibody and secondary antibody (Blue) were used as unlabeled control.



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