




Product datasheet

Anti-SOX10 antibody ab180228

★☆☆☆☆ **1 Abreviews** [2 Images](#)

Overview

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|----------------------------|---|
| Product name | Anti-SOX10 antibody |
| Description | Goat polyclonal to SOX10 |
| Host species | Goat |
| Tested applications | Suitable for: WB |
| Species reactivity | Reacts with: Mouse, Human Predicted to work with: Rat, Cow, Dog, Pig  |
| Immunogen | Synthetic peptide corresponding to Human SOX10 aa 351-364 (internal sequence) (Cysteine residue). (NP_008872.1). Sequence: DAKAQVKTETAGPQ Database link: P56693 <div>  Run BLAST with  Run BLAST with </div> |
| Positive control | Human brain cerebellum lysate. Mouse brain lysate. |
| General notes | <p>The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.</p> <p>If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As</p> |

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 | pH: 7.30 Preservative: 0.02% Sodium azide Constituents: 99% Tris buffered saline, 0.5% BSA |
| Purity | Immunogen affinity purified |

| | |
|---------------------------|--|
| Purification notes | ab180228 was purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide. |
| Clonality | Polyclonal |
| Isotype | IgG |

Applications

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab180228 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 |
|-------------|-----------|---|
| WB | | Use a concentration of 0.03 - 0.1 µg/ml. Detects a band of approximately 50-52 kDa (predicted molecular weight: 49 kDa). 1 hour primary incubation is recommended for this product. |

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 | Expressed in fetal brain and in adult brain, heart, small intestine and colon. |
| Involvement in disease | <p>Defects in SOX10 are the cause of Waardenburg syndrome type 2E (WS2E) [MIM:611584]. WS2E 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.</p> <p>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).</p> <p>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.</p> <p>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.</p> |
| Sequence similarities | Contains 1 HMG box DNA-binding domain. |
| Cellular localization | Cytoplasm. Nucleus. |

Images



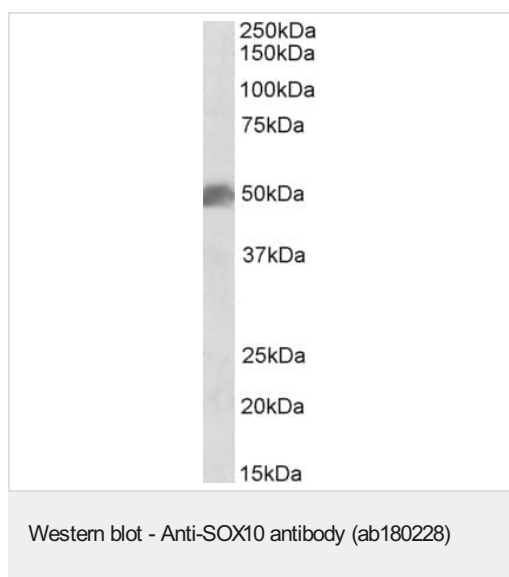
Anti-SOX10 antibody (ab180228) at 0.1 µg/ml + Mouse brain lysate in RIPA buffer at 35 µg

Developed using the ECL technique.

Predicted band size: 49 kDa

Observed band size: 52 kDa

Primary incubation 1 hour at room temperature.



Anti-SOX10 antibody (ab180228) at 0.1 µg/ml + Human brain cerebellum lysate (in RIPA buffer) at 35 µg

Developed using the ECL technique.

Predicted band size: 49 kDa

Observed band size: 48 kDa

Primary incubation was 1 hour.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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