abcam

Product datasheet

Recombinant Human SOX2 protein ab79950

2 Images

Description

Product name Recombinant Human SOX2 protein

Purity > 95 % SDS-PAGE.

ab79950 is greater than 95% by SDS-PAGE gel and HPLC analyses. Endotoxin level is less than

0.1 ng per μ g ($1EU/\mu$ g).

Endotoxin level < 0.100 Eu/μg
Expression system Escherichia coli

Accession P48431

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MYNMMETELK PPGPQQTSGG GGGNSTAAAA

GGNQKNSPDR VKRPMNAFMV WSRGQRRKMA QENPKMHNSE ISKRLGAEWK LLSETEKRPF IDEAKRLRAL HMKEHPDYKY RPRRKTKTLM KKDKYTLPGG LLAPGGNSMA SGVGVGAGLG AGVNQRMDSY AHMNGWSNGS YSMMQDQLGY PQHPGLNAHG AAQMQPMHRY DVSALQYNSM TSSQTYMNGS PTYSMSYSQQ GTPGMALGSM GSVVKSEASS SPPVVTSSSH SRAPCQAGDL

RDMISMYLPG AEVPEPAAPS RLHMSQHYQS

GPVPGTAING TLPLSHM

Amino acids 1 to 317

Specifications

Our Abpromise guarantee covers the use of ab79950 in the following tested applications.

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

Applications Sandwich ELISA

Western blot

HPLC

SDS-PAGE

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Preparation and Storage

Stability and Storage

Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Reconstitution

Reconstituted ab79950 is stable for at least 3 months when stored in working aliquots with a carrier protein at -20⁰C. Avoid repeated freeze/thaw cycles.

General Info

Function

Transcription factor that forms a trimeric complex with OCT4 on DNA and controls the expression of a number of genes involved in embryonic development such as YES1, FGF4, UTF1 and ZFP206 (By similarity). Critical for early embryogenesis and for embryonic stem cell pluripotency.

Involvement in disease

Defects in SOX2 are the cause of microphthalmia syndromic type 3 (MCOPS3) [MIM:206900]. Microphthalmia is a clinically heterogeneous disorder of eye formation, ranging from small size of a single eye to complete bilateral absence of ocular tissues (anophthalmia). In many cases, microphthalmia/anophthalmia occurs in association with syndromes that include non-ocular abnormalities. MCOPS3 is characterized by the rare association of malformations including unior bilateral anophthalmia or microphthalmia, and esophageal atresia with trachoesophageal fistula.

Sequence similarities

Contains 1 HMG box DNA-binding domain.

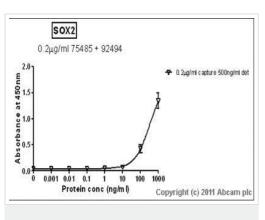
Post-translational modifications

Sumoylation inhibits binding on DNA and negatively regulates the FGF4 transactivation.

Cellular localization

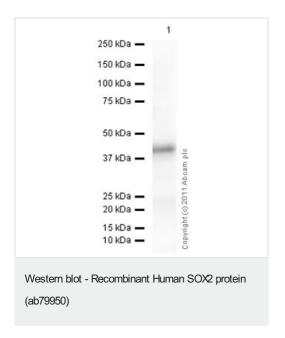
Nucleus.

Images



Sandwich ELISA - Recombinant Human SOX2 protein (ab79950)

Standard Curve for SOX2 (Analyte: SOX2 protein (Human) (ab79950)); dilution range 1pg/ml to 1µg/ml using Capture Antibody Mouse monoclonal [57CT23.3.4] to SOX2 (ab75485) at 0.2µg/ml and Detector Antibody Rabbit monoclonal [EPR3131] to SOX2 (ab92494) at 0.5µg/ml.



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

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