# abcam

## Product datasheet

## Recombinant Human SOX2 protein ab80520

**Description** 

Product name Recombinant Human SOX2 protein

Purity > 95 % SDS-PAGE.

Purity: Greater than 95% by SDS-PAGE gel and HPLC analyses. Endotoxin Level: Endotoxin

level is less than 0.1 ng per μg (1EU/μg).

Endotoxin level < 0.100 Eu/μg
Expression system Escherichia coli

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 TLPLSHMGGY GRKKRRORRR

**Specifications** 

Our Abpromise guarantee covers the use of ab80520 in the following tested applications.

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

**Applications** SDS-PAGE

Form Lyophilized

**Preparation and Storage** 

**Stability and Storage** Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

1

#### Reconstitution

The lyophilized protein is stable for at least 2 years from date of receipt at -20°C. Reconstituted Sox2-TAT is stable for at least 3 months when stored in working aliquots with a carrier protein at -20°C. 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.

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

## Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- · We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit <a href="https://www.abcam.com/abpromise">https://www.abcam.com/abpromise</a> or contact our technical team.

### Terms and conditions

· Guarantee only valid for products bought direct from Abcam or one of our authorized distributors