

Product datasheet

Recombinant Human SOX2 protein ab79950

画像数 2

製品の概要

製品名	Recombinant Human SOX2 protein
タンパク質長	Full length protein

製品の詳細

由来	Recombinant
由来	Escherichia coli

アミノ酸配列

生物種	Human
配列	<p>           MYNMMETELK PPGPQQTSGG GGGNSTAAAA            GGNQKNPDR VKRPMNAFMV WSRGQRRKMA            QENPKMHNSE ISKRLGAEWK LLSETEKRPF            IDEAKRLRAL HMKEHPDYKY RPRRKTKTLM            KKDKYTLPGG LLAPGGNSMA SGVGVGAGLG            AGVNQRMSY AHMNGWSNGS YSMMQDQLGY            PQHPGLNAHG AAQMMPMHRY DVSALQYNM            TSSQTYMNGS PTYSMSYSQQ GTPGMALGSM            GSVVKSEASS SPPVVTSSSH SRAPCQAGDL            RDMISMYLPG AEVPEPAAPS RLHMSQHYQS            GPVPGTAING TLPLSHM         </p>

特性

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.

アプリケーション	<p>Sandwich ELISA</p> <p>Western blot</p> <p>SDS-PAGE</p>
エンドトキシン・レベル	< 0.100 Eu/μg
精製度	<p>&gt; 95 % SDS-PAGE.</p> <p>ab79950 is greater than 95% by SDS-PAGE gel and HPLC analyses. Endotoxin level is less than 0.1 ng per μg (1EU/μg).</p>

**製品の状態**

Lyophilised

**前処理および保存****保存方法および安定性**

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

**再構成**

Reconstituted ab79950 is stable for at least 3 months when stored in working aliquots with a carrier protein at -20°C. Avoid repeated freeze/thaw cycles.

**関連情報****機能**

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.

**関連疾患**

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 uni- or bilateral anophthalmia or microphthalmia, and esophageal atresia with trachoesophageal fistula.

**配列類似性**

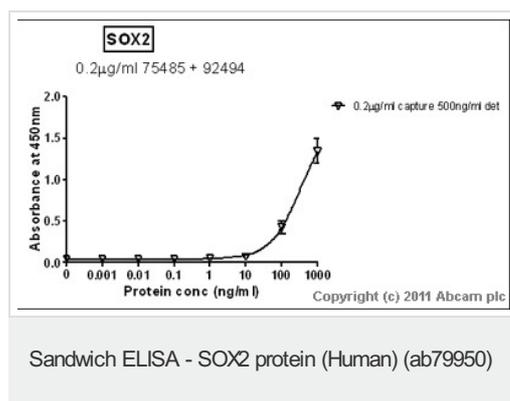
Contains 1 HMG box DNA-binding domain.

**翻訳後修飾**

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

**細胞内局在**

Nucleus.

**画像**

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.



Western blot - Recombinant Human SOX2 protein  
(ab79950)

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

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