abcam

Product datasheet

Recombinant Human TATA binding protein TBP ab81897

1 References 画像数 2

製品の詳細

製品名 Recombinant Human TATA binding protein TBP

生理活性 1 unit equals 1 nanogram of purified protein.

精製度 > 95 % SDS-PAGE.

発現系 Escherichia coli

アクセッション番号 <u>NM_003194</u>

タンパク質長 Full length protein

Animal free No

由来 Recombinant

生物種 Human

配列 MDQNNSLPPY AQGLASPQGA MTPGIPIFSP

PIRLEGLVLT HQQFSSYEP ELFPGLIYRM IKPRIVLLIF

VSGKVVLTGA KVRAEIYEAF ENIYPILKG FRKTT

予測される分子量39 kDa領域1 to 339

サブ His tag N-Terminus

特性

Our Abpromise guarantee covers the use of ab81897 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 Gel Supershift Assays

SDS-PAGE

Western blot

EMSA

製品の状態

Liquid

備考

1 unit equals 1 nanogram of purified protein.

前処理および保存

保存方法および安定性

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 7.9

Constituents: 0.75% Potassium chloride, 0.0154% DTT, 0.316% Tris HCl, 0.00584% EDTA, 20%

Glycerol (glycerin, glycerine)

関連情報

機能

General transcription factor that functions at the core of the DNA-binding multiprotein factor TFIID. Binding of TFIID to the TATA box is the initial transcriptional step of the pre-initiation complex (PIC), playing a role in the activation of eukaryotic genes transcribed by RNA polymerase II. Component of the transcription factor SL1/TIF-IB complex, which is involved in the assembly of the PIC (preinitiation complex) during RNA polymerase I-dependent transcription. The rate of PIC formation probably is primarily dependent on the rate of association of SL1 with the rDNA promoter. SL1 is involved in stabilization of nucleolar transcription factor 1/UBTF on rDNA.

組織特異性

関連疾患

Widely expressed, with levels highest in the testis and ovary.

Defects in TBP are the cause of spinocerebellar ataxia type 17 (SCA17) [MIM:607136]. Spinocerebellar ataxia is a clinically and genetically heterogeneous group of cerebellar disorders. Patients show progressive incoordination of gait and often poor coordination of hands, speech and eye movements, due to degeneration of the cerebellum with variable involvement of the brainstem and spinal cord. SCA17 is an autosomal dominant cerebellar ataxia (ADCA) characterized by widespread cerebral and cerebellar atrophy, dementia and extrapyramidal signs. The molecular defect in SCA17 is the expansion of a CAG repeat in the coding region of TBP. Longer expansions result in earlier onset and more severe clinical manifestations of the

disease.

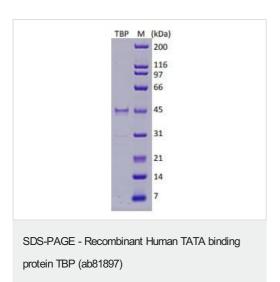
配列類似性

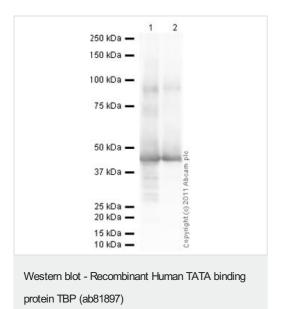
Belongs to the TBP family.

細胞内局在

Nucleus.

画像





All lanes : Anti-TATA binding protein TBP antibody - Nuclear Loading Control and ChIP Grade ($\underline{ab63766}$) at 1 $\mu g/ml$

Lane 1 : Recombinant Human TATA binding protein TBP (ab81897) at 0.1 μg

Lane 2 : Recombinant Human TATA binding protein TBP (ab81897) at 0.01 µg

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) preadsorbed (**ab97080**) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 10 seconds

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

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