abcam

Product datasheet

Recombinant Human KAT4 / TBP Associated Factor 1 protein ab135218

1 Image

Description

Product name Recombinant Human KAT4 / TBP Associated Factor 1 protein

Purity > 70 % SDS-PAGE.

Expression system Baculovirus infected Sf9 cells

Accession P21675

Protein length Protein fragment

Animal free No

Nature Recombinant

Species Human

Sequence MGPGCDLLLRTAATITAAAIMSDTDSDEDSAGGGPFSLAG

FLFGNINGAG

QLEGESVLDDECKKHLAGLGALGLGSLITELTANEELTGT

DGALVNDEGW

VRSTEDAVDYSDINEVAEDESRRYQQTMGSLQPLCHSDY

DEDDYDADCED

 ${\tt IDCKLMPPPPPPGPMKKDKDQDSITGEKVDFSSSSDSE}$

SEMGPQEATQA

ESEDGKLTLPLAGIMQHDATKLLPSVTELFPEFRPGKVLR

FLRLFGPGKN

VPSVWRSARRKRKKKHRELIQEEQIQEVECSVESEVSQK

SLWNYDYAPPP

PPEQCLSDDEITMMAPVESKFSQSTGDIDKVTDTKPRVA

EWRYGPARLWY

DMLGVPEDGSGFDYGFKLRKTEHEPVIKSRMIEEFRKLEE

NNGTDLLADE

NFLMVTQLHWEDDIIWDGEDVKHKGTKPQRASLAGWLPS

SMTRNAMAYNV

QQGFAATLDDDKPWYSIFPIDNEDLVYGRWEDNIIWDAQA

MPRLLEPPVL

TLDPNDENLILEIPDEKEEATSNSPSKESKKESSLKKSRILL

GKTGVIKE EPQQNMSQPEV

Predicted molecular weight

130 kDa including tags

1

Amino acids 1 to 561

Specifications

Our Abpromise guarantee covers the use of ab135218 in the following tested applications.

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

Applications Western blot

SDS-PAGE

Form Liquid

Preparation and Storage

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

pH: 7.50

Constituents: 0.31% Glutathione, 0.002% PMSF, 0.004% DTT, 0.79% Tris HCI, 0.003% EDTA,

25% Glycerol (glycerin, glycerine), 0.88% Sodium chloride

General Info

Function Largest component and core scaffold of the TFIID basal transcription factor complex. Contains

novel N- and C-terminal Ser/Thr kinase domains which can autophosphorylate or

transphosphorylate other transcription factors. Phosphorylates TP53 on 'Thr-55' which leads to MDM2-mediated degradation of TP53. Phosphorylates GTF2A1 and GTF2F1 on Ser residues. Possesses DNA-binding activity. Essential for progression of the G1 phase of the cell cycle.

Involvement in disease Defects in TAF1 are the cause of dystonia type 3 (DYT3) [MIM:314250]; also called X-linked

dystonia-parkinsonism (XDP). DYT3 is a X-linked dystonia-parkinsonism disorder. Dystonia is defined by the presence of sustained involuntary muscle contractions, often leading to abnormal

postures. DYT3 is characterized by severe progressive torsion dystonia followed by

parkinsonism. Its prevalence is high in the Philippines. DYT3 has a well-defined pathology of extensive neuronal loss and mosaic gliosis in the striatum (caudate nucleus and putamen) which

appears to resemble that in Huntington disease.

Sequence similaritiesBelongs to the TAF1 family.

Contains 2 bromo domains.

Contains 1 HMG box DNA-binding domain.

Contains 2 protein kinase domains.

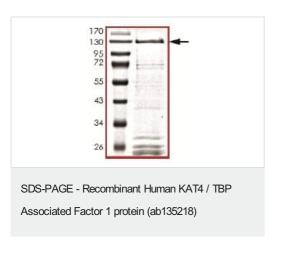
Post-translational

modifications

Phosphorylated by casein kinase II in vitro.

Cellular localization Nucleus.

Images



SDS-PAGE analysis of ab135218.

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