TAF I p48 (M-19): sc-6572



The Power to Question

BACKGROUND

For gene transcription governed by RNA polymerase I, the human transcription factor SL1 (mouse TIF-IB) directs the assembly of initiation complexes at the prompter. Like TFIID, which directs transcription by RNA polymerase II, SL1/TIF-IB contains the TATA-binding protein (TBP) and a set of TBP-associated factors (TAFs). The three TAF I subunits, hTAF I p110, hTAF I p63 and hTAF p48 (or mouse TAF I p95, TAF I p68 and TAF I p48) are all integral components of SL1/TIF-IB. The mutually exclusive binding of either TAF I or TAF II subunits to TBP is believed to direct the formation of promoter and RNA polymerase-specific complexes.

CHROMOSOMAL LOCATION

Genetic locus: TAF1A (human) mapping to 1q41; Taf1a (mouse) mapping to 1.

SOURCE

TAF I p48 (M-19) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the C-terminus of TAF I p48 of mouse origin.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin. Also available as TransCruz reagent for Gel Supershift and ChIP applications, sc-6572 X, 200 μg /0.1 ml.

Blocking peptide available for competition studies, sc-6572 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

TAF I p48 (M-19) is recommended for detection of TAF I p48 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 μg per 100-500 μg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for TAF I p48 siRNA (h): sc-38486, TAF I p48 siRNA (m): sc-38487, TAF I p48 shRNA Plasmid (h): sc-38486-SH, TAF I p48 shRNA Plasmid (m): sc-38487-SH, TAF I p48 shRNA (h) Lentiviral Particles: sc-38486-V and TAF I p48 shRNA (m) Lentiviral Particles: sc-38487-V.

TAF I p48 (M-19) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of TAF I p48: 53 kDa.

Positive Controls: A-431 nuclear extract: sc-2122, Y79 cell lysate: sc-2240 or HeLa whole cell lysate: sc-2200.

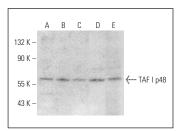
STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

DATA



TAF I p48 (M-19): sc-6572. Western blot analysis of TAF I p48 expression in A-431 nuclear extract (**A**) and Y79 (**B**), HeLa (**C**), Jurkat (**D**) and K-562 (**E**) whole cell

SELECT PRODUCT CITATIONS

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- Kenneth, N.S., et al. 2007. TRRAP and GCN5 are used by c-Myc to activate RNA polymerase III transcription. Proc. Natl. Acad. Sci. USA 104: 14917-14922.
- 9. Müller, C., et al. 2010. Nucleolar retention of a translational C/EBP α isoform stimulates rDNA transcription and cell size. EMBO J. 29: 897-909.



Try **TAF I p48 (A-10): sc-393600**, our highly recommended monoclonal alternative to TAF I p48 (M-19).