

TFIIF RAP 30 (H-148): sc-292295

BACKGROUND

In eukaryotic systems, initiation of transcription from protein-coding genes is a complex process requiring RNA polymerase II and broad families of auxiliary transcription factors. Such factors can be divided into two major functional classes: the basal factors that are required for transcription of all Pol II genes, including TFIIA, TFIIB, TFIID, TFIIE, TFIIF and TFIIH; and sequence-specific factors that regulate gene expression. The basal transcription factors and Pol II form a specific multiprotein complex near the transcription start site by interacting with core promoter elements such as the TATA box generally located 25-30 base pairs upstream of the transcription start site. TFIIF, a heteromer composed of a small (RAP 30) and a large (RAP 74) subunit, is required for RNA polymerase II to assemble into a preinitiation complex formed by promoter DNA and the general factors TFIID, TFIIA and TFIIB. In addition, TFIIF stimulates transcription elongation by RNA polymerase II.

CHROMOSOMAL LOCATION

Genetic locus: GTF2F2 (human) mapping to 13q14.12; Gtf2f2 (mouse) mapping to 14 D3.

SOURCE

TFIIF RAP 30 (H-148) is a rabbit polyclonal antibody raised against amino acids 102-249 mapping near the C-terminus of TFIIF RAP 30 of human origin.

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Available as TransCruz reagent for Gel Supershift and ChIP applications, sc-292295 X, 200 µg/0.1 ml.

APPLICATIONS

TFIIF RAP 30 (H-148) is recommended for detection of TFIIF RAP 30 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).

TFIIF RAP 30 (H-148) is also recommended for detection of TFIIF RAP 30 in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for TFIIF RAP 30 siRNA (h): sc-38521, TFIIF RAP 30 siRNA (m): sc-38522, TFIIF RAP 30 shRNA Plasmid (h): sc-38521-SH, TFIIF RAP 30 shRNA Plasmid (m): sc-38522-SH, TFIIF RAP 30 shRNA (h) Lentiviral Particles: sc-38521-V and TFIIF RAP 30 shRNA (m) Lentiviral Particles: sc-38522-V.

TFIIF RAP 30 (H-148) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

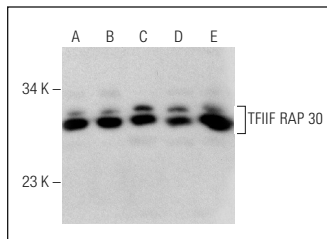
Molecular Weight of TFIIF RAP 30: 30 kDa.

Positive Controls: K-562 nuclear extract: sc-2130, Jurkat nuclear extract: sc-2132 or HeLa nuclear extract: sc-2120.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit IgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit IgG-HRP: sc-2030 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunoprecipitation: use Protein A/G PLUS-Agarose: sc-2003 (0.5 ml agarose/2.0 ml). 3) Immunofluorescence: use goat anti-rabbit IgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit IgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



TFIIF RAP 30 (H-148): sc-292295. Western blot analysis of TFIIF RAP 30 expression in K-562 (A), Jurkat (B), A-431 (C), HeLa (D) and U-937 (E) nuclear extracts.

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.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.



Try **TFIIF RAP 30 (15): sc-136408** or **TFIIF RAP 30 (G-3): sc-374304**, our highly recommended monoclonal alternatives to TFIIF RAP 30 (H-148).