RPA16 (H-83): sc-292363



The Power to Question

BACKGROUND

RNA polymerases transcribe nuclear genes for ribosomal RNA, thus representing ribosomal biogenesis. RNA polymerase I (Pol I) is located in the nucleolus and transcribes class I genes, which code for large ribosomal RNA. Different subunits of the Pol I transcription machinery are targets of various physiological stimuli, which suggests that multiple signaling pathways are involved in carrying out Pol I transcription. RPA16, RPA40 and RPA135 are subunits of Pol I that associate with each other at an early stage of RNA Pol I assembly. RPA40 is essential for the function and integrity of the complex and is also an essential subunit of RNA polymerase III (Pol III).

REFERENCES

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- Seither, P., et al. 1997. Molecular cloning and characterization of the cDNA encoding the largest subunit of mouse RNA polymerase I. Mol. Gen. Genet. 255: 180-186.
- Hoeger, H., et al. 1998. Deficient transcription of subunit RPA 40 of RNA polymerase I and III in heart of rats with neonatal asphyxia. Life Sci. 62: 275-282.
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- Chen, H.K., et al. 1999. Human Nopp140, which interacts with RNA polymerase I: implications for rRNA gene transcription and nucleolar structural organization. Mol. Cell Biol. 19: 8536-8546.
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CHROMOSOMAL LOCATION

Genetic locus: POLR1D (human) mapping to 13q12.2; Polr1d (mouse) mapping to $5\ G3$.

SOURCE

RPA16 (H-83) is a rabbit polyclonal antibody raised against amino acids 51-133 mapping at the C-terminus of RPA16 of human origin.

PRODUCT

Each vial contains 200 μg lgG 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-292363 X, 200 $\mu g/0.1$ ml.

RESEARCH USE

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

APPLICATIONS

RPA16 (H-83) is recommended for detection of RPA16 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).

RPA16 (H-83) is also recommended for detection of RPA16 in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for RPA16 siRNA (h): sc-38240, RPA16 siRNA (m): sc-38241, RPA16 shRNA Plasmid (h): sc-38240-SH, RPA16 shRNA Plasmid (m): sc-38241-SH, RPA16 shRNA (h) Lentiviral Particles: sc-38240-V and RPA16 shRNA (m) Lentiviral Particles: sc-38241-V.

RPA16 (H-83) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of RPA16: 15 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use goat anti-rabbit lgG-HRP: sc-2004 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible goat anti-rabbit lgG-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 lgG-FITC: sc-2012 (dilution range: 1:100-1:400) or goat anti-rabbit lgG-TR: sc-2780 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

STORAGE

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

PROTOCOLS

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



Try **RPA16 (2774C3a): sc-81636**, our highly recommended monoclonal alternative to RPA16 (H-83).

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