## SANTA CRUZ BIOTECHNOLOGY, INC.

# ATRIP (V-20): sc-33411



#### BACKGROUND

DNA damage or incomplete replication of DNA results in the inhibition of cell cycle progression at the G<sub>1</sub> to S or the G<sub>2</sub> to M phase transition by conserved regulatory mechanisms known as cell cycle checkpoints. Checkpoint proteins include Rad17, which is involved in regulating cell cycle progression at the G1 checkpoint, Chk1, Chk2, Rad1, Rad9 and Hus1, which are involved in regulating cell cycle arrest at the G2 checkpoint. In response to DNA damage, ATM and ATR kinases are important for cell cycle checkpoint response signalling. ATR-interacting protein (ATRIP), also designated ATM and Rad3related-interacting protein, is required for checkpoint signaling after DNA damage. It is also important for ATR, which regulates DNA replication and damage checkpoint responses, expression. ATRIP is a ubiquitously expressed protein that can form heterodimers with ATR. After dimerization they bind the RPA complex and are recruited to single stranded DNA. ATRIP is a nuclear protein that may also play a role in protein stabilization.

#### REFERENCES

- 1. Cortez, D., et al. 2001. ATR and ATRIP: partners in checkpoint signaling. Science 294: 1713-1716.
- 2. Zou, L., et al. 2003. Sensing DNA damage through ATRIP recognition of RPA-ssDNA complexes. Science 300: 1542-1548.
- 3. Ball, H.L., et al. 2005. ATRIP oligomerization is required for ATR-dependent checkpoint signaling. J. Biochem. 280: 31390-31396.
- 4. Kim, S.M., et al. 2005. Phosphorylation of Chk1 by ATM- and Rad3-related (ATR) in Xenopus egg extracts requires binding of ATRIP to ATR but not the stable DNA-binding. J. Biochem. 280: 38355-38364.

#### **CHROMOSOMAL LOCATION**

Genetic locus: ATRIP (human) mapping to 3p21.31.

#### SOURCE

ATRIP (V-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of ATR-interacting protein 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.

Blocking peptide available for competition studies, sc-33411 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

#### **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.

#### **APPLICATIONS**

ATRIP (V-20) is recommended for detection of ATRIP of 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).

ATRIP (V-20) is also recommended for detection of ATRIP in additional species, including equine, canine, bovine and porcine.

Suitable for use as control antibody for ATRIP siRNA (h): sc-44800, ATRIP shRNA Plasmid (h): sc-44800-SH and ATRIP shRNA (h) Lentiviral Particles: sc-44800-V.

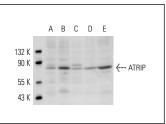
Molecular Weight of ATRIP: 86 kDa.

Positive Controls: MCF7 nuclear extract: sc-2149, U-937 nuclear extract: sc-2156 or PC-3 nuclear extract: sc-2152.

#### **RECOMMENDED SECONDARY REAGENTS**

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (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 donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz<sup>™</sup> Mounting Medium: sc-24941.

#### DATA



ATRIP (V-20): sc-33411. Western blot analysis of ATRIP expression in UV-treated HeLa (A), MCF7 (B), THP-1 (C), U-937 (D) and PC-3 (E) nuclear extracts.

#### **RESEARCH USE**

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

