

SANTA CRUZ BIOTECHNOLOGY, INC.

# p-Atm (Ser 1981)-R: sc-33042-R



The Power to Ouestion

## **BACKGROUND**

The phosphatidylinositol kinase (PIK) family members fall into two distinct subgroups. The first subgroup contains proteins such as the PI 3- and PI 4-kinases and the second group comprises the PIK-related kinases. The PIK-related kinases include Atm, DNA-PKCS and FRAP. These proteins have in common a region of homology at their carboxy termini that is not present in the PI 3- and PI 4-kinases. All of the members of the PIK-related kinases are also more than 270 kDa. The Atm gene is mutated in the autosomal recessive disorder ataxia telangiectasia (AT) that is characterized by cerebellar degeneration (ataxia) and the appearance of dilated blood vessels (telangiectases) in the conjunctivae of the eyes. AT cells are hypersensitive to ionizing radiation, impaired in mediating the inhibition of DNA synthesis and display delays in p53 induction. Cultured cells respond to breaks in double stranded DNA by immediate activation of ATM through autophosphorylated, inactive form of Atm.

#### REFERENCES

- Nowak, R. 1995. Discovery of AT gene sparks biomedical research bonanza. Science 268: 1700-1701.
- 2. Savitsky, K., et al. 1995. A single ataxia telangiectasia gene with a product similar to Pl-3 kinase. Science 268: 1749-1753.
- Keith, C.T., et al. 1995. PIK-related kinases: DNA repair, recombination, and cell cycle checkpoints. Science 270: 50-51.
- Hartley, K.O., et al. 1995. DNA-dependent protein kinase catalytic subunit: a relative of phosphatidylinositol 3-kinase and the ataxia telangiectasia gene product. Cell 82: 849-856.
- Hickson, I., et al. 2004. Identification and characterization of a novel and specific inhibitor of the ataxia-telangiectasia mutated kinase ATM. Cancer Res. 64: 9152-9159.
- Chen, L., et al. 2005. Ataxia-telangiectasia-mutated dependent phosphorylation of Artemis in response to DNA damage. Cancer Sci. 96: 134-141.
- 7. Liu, A., et al. 2005. Alterations of DNA damage-response genes ATM and ATR in pyothorax-associated lymphoma. Lab Invest. 85: 436-446.
- 8. Lavin, M.F., et al. 2005. Atm signaling and genomic stability in response to DNA damage. Mutat. Res. 569: 123-132.
- 9. Bartkova, J., et al. 2005. Atm activation in normal human tissues and testicular cancer. Cell Cycle 4: 838-845.

### **CHROMOSOMAL LOCATION**

Genetic locus: ATM (human) mapping to 11q23; Atm (mouse) mapping to 9 C-D.

#### **SOURCE**

p-Atm (Ser 1981)-R is a rabbit polyclonal antibody raised against a short amino acid sequence containing phosphorylated Ser 1981 of Atm of human origin.

## **PRODUCT**

Each vial contains 200  $\mu g$  lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1%  $\alpha$ 0.1% sodium azide

Blocking peptide available for competition studies, sc-33042 P, (100  $\mu$ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

#### **APPLICATIONS**

p-Atm (Ser 1981)-R is recommended for detection of Ser 1981 phosphorylated Atm of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500).

Suitable for use as control antibody for Atm siRNA (h): sc-29761 and Atm siRNA (m): sc-29762.

#### **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 B Blocking Reagent: sc-2335 (use 50 mM NaF, sc-24988, as diluent) and Western Blotting Luminol Reagent: sc-2048. 2) 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.

## **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.