

## Anti-ATM (Ser-1981), Phosphospecific Antibody

Catalog # AN1647

## **Product Information**

Application WB, ICC, IP
Primary Accession Q13315
Host Mouse

**Clonality** Mouse Monoclonal

IsotypeIgG1Clone NamesM366Calculated MW350687

## **Additional Information**

Gene ID 472

Other Names ataxia telangiectasia mutated, AT1 ATDC TEL1 TELO1

**Target/Specificity** Ataxia telangiectasia mutated kinase (ATM) is a serine/threonine kinase that

regulates cell cycle checkpoints and DNA repair. Mutations of ATM cause a

spectrum of defects ranging from neurodegeneration to cancer

predisposition. Activation of ATM after DNA damage involves Cdk5 mediated phosphorylation of Ser-794 followed by autophosphorylation at Ser-1891. Active ATM kinase regulates a number of proteins involved in cell cycle checkpoint control, apoptosis and DNA repair. The Cdk5–ATM pathway regulates phosphorylation and function of the ATM targets p53 and H2AX in postmitotic neurons. Other known substrates of ATM include Chk2, Chk1, CtIP, 4E-BP1, BRCA1, RPA3, SMC1, FANCD2, Rad17, Artemis, Nbs1, and the I-2 regulatory subunit of PP1. Thus, activation of Cdk5 by DNA damage may be an important initiator of ATM-dependent regulation of cell cycle checkpoints.

**Dilution** WB~~1:1000 ICC~~N/A IP~~N/A

Format Protein A Purified

**Storage** Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** Anti-ATM (Ser-1981), Phosphospecific Antibody is for research use only and

not for use in diagnostic or therapeutic procedures.

Shipping Blue Ice

## **Background**

Ataxia telangiectasia mutated kinase (ATM) is a serine/threonine kinase that regulates cell cycle checkpoints and DNA repair. Mutations of ATM cause a spectrum of defects ranging from neurodegeneration to cancer predisposition. Activation of ATM after DNA damage involves Cdk5 mediated phosphorylation of Ser-794

followed by autophosphorylation at Ser-1891. Active ATM kinase regulates a number of proteins involved in cell cycle checkpoint control, apoptosis and DNA repair. The Cdk5–ATM pathway regulates phosphorylation and function of the ATM targets p53 and H2AX in postmitotic neurons. Other known substrates of ATM include Chk2, Chk1, CtIP, 4E-BP1, BRCA1, RPA3, SMC1, FANCD2, Rad17, Artemis, Nbs1, and the I-2 regulatory subunit of PP1. Thus, activation of Cdk5 by DNA damage may be an important initiator of ATM-dependent regulation of cell cycle checkpoints.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.