

Anti-ATM (C-terminal region) Antibody

Catalog # AN1645

Product Information

Application WB, FC, ICC, IP

Primary Accession Q13315 Host Mouse

Clonality Mouse Monoclonal

IsotypeIgG2bClone NamesM361Calculated 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 FC~~1:10~50 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 (C-terminal region) 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.

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