

Phospho-ATM(S1981) Antibody

Affinity Purified Rabbit Polyclonal Antibody (Pab)

Catalog # AP3504a

Product Information

Application	DB, IF, E
Primary Accession	Q13315
Reactivity	Human, Rat, Mouse
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Clone Names	RB8121
Calculated MW	350687

Additional Information

Gene ID	472
Other Names	Serine-protein kinase ATM, Ataxia telangiectasia mutated, A-T mutated, ATM
Target/Specificity	This Phospho-ATM-pS1981 antibody is generated from rabbits immunized with a KLH conjugated synthetic phosphopeptide corresponding to amino acid residues surrounding S1981 of human ATM.
Dilution	DB~~1:500 IF~~1:10~50 E~~Use at an assay dependent concentration.
Format	Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.
Storage	Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.
Precautions	Phospho-ATM(S1981) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

Protein Information

Name	ATM
Function	Serine/threonine protein kinase which activates checkpoint signaling upon double strand breaks (DSBs), apoptosis and genotoxic stresses such as ionizing ultraviolet A light (UVA), thereby acting as a DNA damage sensor (PubMed: 10550055 , PubMed: 10839545 , PubMed: 10910365 , PubMed: 12556884 , PubMed: 14871926 , PubMed: 15064416 , PubMed: 15448695 , PubMed: 15456891 , PubMed: 15790808 ,

PubMed:[15916964](#), PubMed:[17923702](#), PubMed:[21757780](#), PubMed:[24534091](#), PubMed:[35076389](#), PubMed:[9733514](#)). Recognizes the substrate consensus sequence [ST]-Q (PubMed:[10550055](#), PubMed:[10839545](#), PubMed:[10910365](#), PubMed:[12556884](#), PubMed:[14871926](#), PubMed:[15448695](#), PubMed:[15456891](#), PubMed:[15916964](#), PubMed:[17923702](#), PubMed:[24534091](#), PubMed:[9733514](#)). Phosphorylates 'Ser-139' of histone variant H2AX at double strand breaks (DSBs), thereby regulating DNA damage response mechanism (By similarity). Also plays a role in pre-B cell allelic exclusion, a process leading to expression of a single immunoglobulin heavy chain allele to enforce clonality and monospecific recognition by the B-cell antigen receptor (BCR) expressed on individual B-lymphocytes. After the introduction of DNA breaks by the RAG complex on one immunoglobulin allele, acts by mediating a repositioning of the second allele to pericentromeric heterochromatin, preventing accessibility to the RAG complex and recombination of the second allele. Also involved in signal transduction and cell cycle control. May function as a tumor suppressor. Necessary for activation of ABL1 and SAPK. Phosphorylates DYRK2, CHEK2, p53/TP53, FBXW7, FANCD2, NFKBIA, BRCA1, CREBBP/CBP, RBBP8/CTIP, FBXO46, MRE11, nibrin (NBN), RAD50, RAD17, PELI1, TERF1, UFL1, RAD9, UBQLN4 and DCLRE1C (PubMed:[10550055](#), PubMed:[10766245](#), PubMed:[10802669](#), PubMed:[10839545](#), PubMed:[10910365](#), PubMed:[10973490](#), PubMed:[11375976](#), PubMed:[12086603](#), PubMed:[15456891](#), PubMed:[19965871](#), PubMed:[21757780](#), PubMed:[24534091](#), PubMed:[26240375](#), PubMed:[26774286](#), PubMed:[30171069](#), PubMed:[30612738](#), PubMed:[30886146](#), PubMed:[30952868](#), PubMed:[38128537](#), PubMed:[9733515](#), PubMed:[9843217](#)). May play a role in vesicle and/or protein transport. Could play a role in T-cell development, gonad and neurological function. Plays a role in replication-dependent histone mRNA degradation. Binds DNA ends. Phosphorylation of DYRK2 in nucleus in response to genotoxic stress prevents its MDM2-mediated ubiquitination and subsequent proteasome degradation (PubMed:[19965871](#)). Phosphorylates ATF2 which stimulates its function in DNA damage response (PubMed:[15916964](#)). Phosphorylates ERCC6 which is essential for its chromatin remodeling activity at DNA double-strand breaks (PubMed:[29203878](#)). Phosphorylates TTC5/STRAP at 'Ser-203' in the cytoplasm in response to DNA damage, which promotes TTC5/STRAP nuclear localization (PubMed:[15448695](#)). Also involved in pexophagy by mediating phosphorylation of PEX5: translocated to peroxisomes in response to reactive oxygen species (ROS), and catalyzes phosphorylation of PEX5, promoting PEX5 ubiquitination and induction of pexophagy (PubMed:[26344566](#)).

Cellular Location

Nucleus. Cytoplasmic vesicle. Cytoplasm, cytoskeleton, microtubule organizing center, centrosome {ECO:0000250|UniProtKB:Q62388}. Peroxisome matrix. Note=Primarily nuclear (PubMed:9050866, PubMed:9150358). Found also in endocytic vesicles in association with beta-adaptin (PubMed:9707615). Translocated to peroxisomes in response to reactive oxygen species (ROS) by PEX5 (PubMed:26344566)

Tissue Location

Found in pancreas, kidney, skeletal muscle, liver, lung, placenta, brain, heart, spleen, thymus, testis, ovary, small intestine, colon and leukocytes

Background

ATM belongs to the PI3/PI4-kinase family. This protein is an important cell cycle checkpoint kinase that phosphorylates; thus, it functions as a regulator of a wide variety of downstream proteins, including tumor suppressor proteins p53 and BRCA1, checkpoint kinase CHK2, checkpoint proteins RAD17 and RAD9, and DNA repair protein NBS1. ATM and the closely related kinase ATR are thought to be master controllers of cell cycle checkpoint signaling pathways that are required for cell response to DNA damage and for genome stability. Mutations in the gene encoding ATM are associated with ataxia telangiectasia, an autosomal

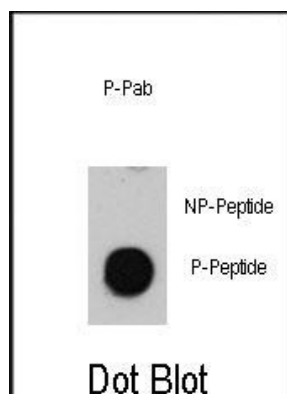
recessive disorder.

References

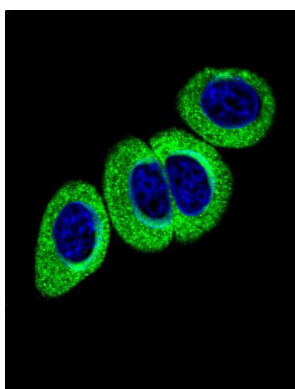
Brunet,J.,Clin. Genet. 73 (5), 465-473 (2008)

Tsai,W.B., Nat. Cell Biol. 10 (4), 460-467 (2008)

Images



Dot blot analysis of anti-Phospho-ATM-pS1981 Antibody (Cat.#AP3504a) on nitrocellulose membrane. 50ng of Phospho-peptide or Non Phospho-peptide per dot were adsorbed. Antibody working concentrations are 0.5ug per ml.



Confocal immunofluorescent analysis of Phospho-ATM-pS1981 Antibody (Cat.#AP3504a) with Hela cell followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). DAPI was used to stain the cell nuclear (blue).

Citations

- [Hepatitis B virus X stimulates redox signaling through activation of ataxia telangiectasia mutated kinase.](#)

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