abcam

Product datasheet

Anti-ATM antibody [5C2] ab2618

★★★★★ 1 Abreviews   17 References   3 Images

Overview

Product name	Anti-ATM antibody [5C2]
Description	Mouse monoclonal [5C2] to ATM
Host species	Mouse
Tested applications	**Suitable for:** WB, ICC/IF
Species reactivity	**Reacts with:** Mouse, Rat, Human, Monkey
Immunogen	Recombinant fragment, corresponding to amino acids 980-1512 of Human ATM
Positive control	Raji or Akata whole cell lysate.
General notes	ab2618 should always be kept frozen.

Properties

Form	Liquid
Storage instructions	Shipped on Dry Ice. Upon delivery aliquot. Store at -20°C or -80°C. Avoid freeze / thaw cycle. Please see notes section.
Storage buffer	Preservative: None Constituents: 10mM PBS, pH 7.4
Purification notes	Purified from ascities fluid by Protein G chromatography to at least 95% homogeneity as determined by SDS-PAGE.
Clonality	Monoclonal
Clone number	5C2
Myeloma	unknown
Isotype	IgG1
Light chain type	kappa

Applications

Our Abpromise guarantee covers the use of **ab2618** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB	★★★★★	1/1000. Predicted molecular weight: 350 kDa.
ICC/IF		1/100 - 1/1000.

Target

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. Recognizes the substrate consensus sequence [ST]-Q. Phosphorylates 'Ser-139' of histone variant H2AX/H2AFX at double strand breaks (DSBs), thereby regulating DNA damage response mechanism. 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 p53/TP53, FANCD2, NFKBIA, BRCA1, CTIP, nibrin (NBN), TERF1, RAD9 and DCLRE1C. 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.
Tissue specificity	Found in pancreas, kidney, skeletal muscle, liver, lung, placenta, brain, heart, spleen, thymus, testis, ovary, small intestine, colon and leukocytes.
Involvement in disease	Defects in ATM are the cause of ataxia telangiectasia (AT) [MIM:208900]; also known as Louis-Bar syndrome, which includes four complementation groups: A, C, D and E. This rare recessive disorder is characterized by progressive cerebellar ataxia, dilation of the blood vessels in the conjunctiva and eyeballs, immunodeficiency, growth retardation and sexual immaturity. AT patients have a strong predisposition to cancer; about 30% of patients develop tumors, particularly lymphomas and leukemias. Cells from affected individuals are highly sensitive to damage by ionizing radiation and resistant to inhibition of DNA synthesis following irradiation. Note=Defects in ATM contribute to T-cell acute lymphoblastic leukemia (TALL) and T-prolymphocytic leukemia (TPLL). TPLL is characterized by a high white blood cell count, with a predominance of prolymphocytes, marked splenomegaly, lymphadenopathy, skin lesions and serous effusion. The clinical course is highly aggressive, with poor response to chemotherapy and short survival time. TPLL occurs both in adults as a sporadic disease and in younger AT patients. Note=Defects in ATM contribute to B-cell non-Hodgkin lymphomas (BNHL), including mantle cell lymphoma (MCL). Note=Defects in ATM contribute to B-cell chronic lymphocytic leukemia (BCLL). BCLL is the commonest form of leukemia in the elderly. It is characterized by the accumulation of mature CD5+ B lymphocytes, lymphadenopathy, immunodeficiency and bone marrow failure.
Sequence similarities	Belongs to the PI3/PI4-kinase family. ATM subfamily. Contains 1 FAT domain. Contains 1 FATC domain.

|	Contains 1 PI3K/PI4K domain.
Domain	The FATC domain is required for interaction with KAT5.
Post-translational modifications	Phosphorylated by NUAK1/ARK5. Autophosphorylation on Ser-367, Ser-1893, Ser-1981 correlates with DNA damage-mediated activation of the kinase. Acetylation, on DNA damage, is required for activation of the kinase activity, dimer-monomer transition, and subsequent autophosphorylation on Ser-1981. Acetylated in vitro by KAT5/TIP60.
Cellular localization	Nucleus. Cytoplasmic vesicle. Primarily nuclear. Found also in endocytic vesicles in association with beta-adaptin.

Images
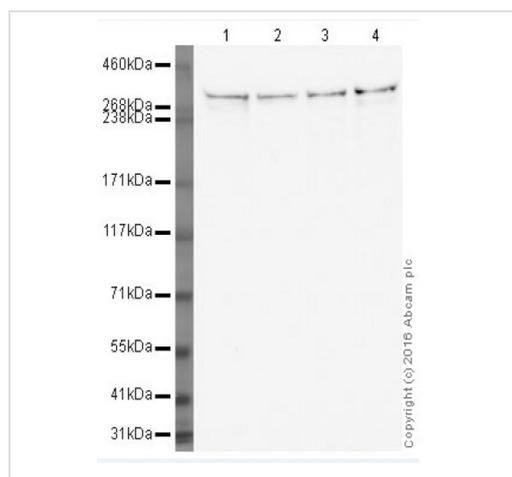
Western blot - Anti-ATM antibody [5C2] (ab2618)

All lanes : Anti-ATM antibody [5C2] (ab2618) at 1/1000 dilution

Lane 1 : HeLa whole cell lysate
Lane 2 : HeLa nuclear lysate
Lane 3 : SK-N-SH whole cell lysate
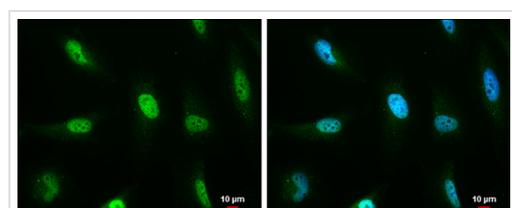**Lane 4 :** Raji whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary
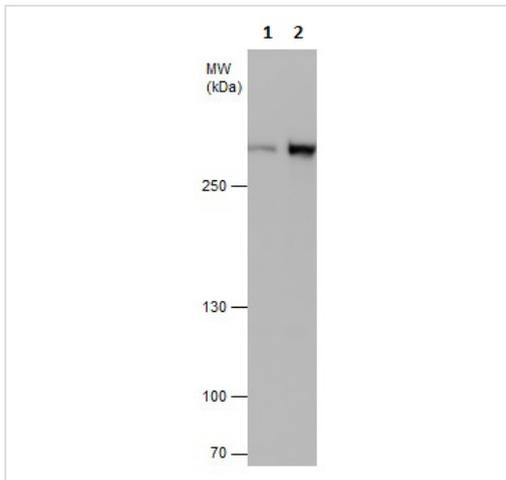**All lanes :** Goat polyclonal to Mouse IgG - H&L - Pre-Adsorbed (HRP) at 1/5000 dilution

Predicted band size: 350 kDa

Exposure time: 12 minutes

Immunocytochemistry/ Immunofluorescence - Anti-ATM antibody [5C2] (ab2618)

Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling ATM (green) with ab2618 at a dilution of 1/200. Cells were fixed with 4% paraformaldehyde at room temperature for 15 minutes. Blue: Hoechst 33342 staining.

All lanes : Anti-ATM antibody [5C2] (ab2618)
at 1/1000 dilution

Lane 1 : HeLa whole cell extract

Lane 2 : HeLa nuclear cell extract

Lysates/proteins at 30 µg per lane.

Predicted band size: 350 kDa

Western blot - Anti-ATM antibody [5C2] (ab2618)

Separated by 5% SDS-PAGE.

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