bsm-52360R

[Primary Antibody]

ATM Recombinant Rabbit mAb



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- DATASHEET -

Host: Rabbit Isotype: IgG
Clonality: Recombinant CloneNo.: 12A
GenelD: 472 SWISS: Q13315

Target: ATM

Purification: affinity purified by Protein A

Concentration: 1mg/ml

Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50%

Glycerol.

Shipped at 4°C. Store at -20°C for one year. Avoid repeated

freeze/thaw cycles.

Background: ATM is a 370 kDa nuclear phosphoprotein involved in the

autosomal recessive disease Ataxia Telangiectasia (AT). ATM belongs to a novel family of proteins associated with cell cycle regulation, apoptosis, and response to DNA damage repair (DNA damage caused by such things as ionizing irradiation activates ATM kinase). The C terminal region has extensive homology to the catalytic domains of Phosphatidylinositol 3 kinases (PI3 kinases).

Applications: WB (1:500-2000)

IHC-P (1:50-200) IHC-F (1:400-800) IF (1:50-200) ICC/IF (1:50)

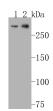
Reactivity: Human

Predicted MW.: 370 kDa

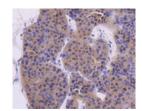
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Subcellular Cytoplasm ,Nucleus

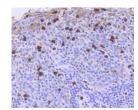
VALIDATION IMAGES



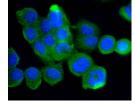
Sample: Lane 1: AGS cell lysate Lane 2: CRC cell lysate Primary: Anti-ATM (bsm-52360R) at 1:500 dilution Secondary: Goat Anti-Rabbit IgG - HRP at 1:5000 dilution Predicted band size: 370 kD Observed band size: 370 kD



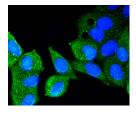
Paraformaldehyde-fixed, paraffin embedded (human liver carcinoma); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (ATM) Monoclonal Antibody, Unconjugated (bsm-52360R) at 1:50 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (human tonsil); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (ATM) Monoclonal Antibody, Unconjugated (bsm-52360R) at 1:50 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



MCF7 cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum) at 37°C for 20 min; Antibody incubation with (ATM (12A)) Monoclonal Antibody, Unconjugated (bsm-52360R) 1:50, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue) was used to stain the cell nuclei.



Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum) at 37°C for 20 min; Antibody incubation with (ATM (12A)) Monoclonal Antibody, Unconjugated (bsm-52360R) 1:50, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue) was used to stain the cell nuclei.

- SELECTED CITATIONS -

- [IF=6.814] Guo-Jian Jiang. et al. Ultraviolet B irradiation induces senescence of human corneal endothelial cells in vitro by DNA damage response and oxidative stress. J PHOTOCH PHOTOBIO B. 2022 Oct;235:112568 WB ;Human. 36137302
- [IF=5.168] Guo-Jian Jiang. et al. Carteolol triggers senescence via activation of β-arrestin–ERK–NOX4–ROS pathway in human corneal endothelial cells in vitro. CHEM-BIOL INTERACT. 2023 Apr;:110511 WB; Human. 37120125