

Anti-ATRIP Antibody
Rabbit polyclonal antibody to ATRIP
Catalog # AP60121**Specification**

Anti-ATRIP Antibody - Product Information

Application	WB, FC, IF/IC, IHC
Primary Accession	Q8WXE1
Other Accession	Q8BMG1
Reactivity	Human, Mouse, Rat
Host	Rabbit
Clonality	Polyclonal
Calculated MW	85838

Anti-ATRIP Antibody - Additional Information**Gene ID** 84126**Other Names**

AGS1; ATR-interacting protein; ATM and Rad3-related-interacting protein

Target/Specificity

KLH-conjugated synthetic peptide encompassing a sequence within the N-term region of human ATRIP. The exact sequence is proprietary.

Dilution

WB~~WB (1/500 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500), FC (1/100 - 1/200)

FC~~1:10~50

IF/IC~~N/A

IHC~~1:100~500

Format

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.

Storage

Store at -20 °C. Stable for 12 months from date of receipt

Anti-ATRIP Antibody - Protein Information**Name** ATRIP**Synonyms** AGS1**Function**

Required for checkpoint signaling after DNA damage. Required for ATR expression, possibly by stabilizing the protein.

Cellular Location

Nucleus. Note=Redistributes to discrete nuclear foci upon DNA damage

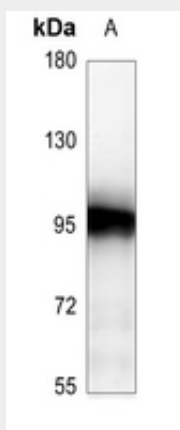
Tissue Location

Ubiquitous..

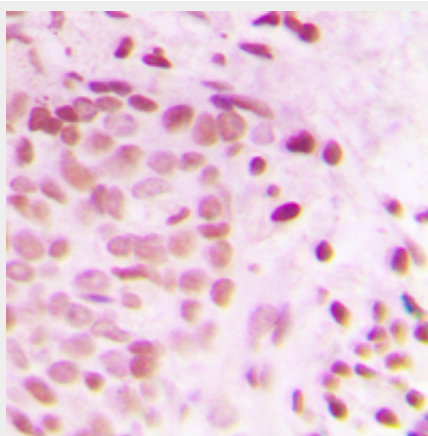
Anti-ATRIP Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

Anti-ATRIP Antibody - Images

Western blot analysis of ATRIP expression in Jurkat (A) whole cell lysates.



Immunohistochemical analysis of ATRIP staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room

temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of ATRIP staining in MCF7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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