ATM Antibody

Goat Polyclonal

Antigen Affinity Purified Protein ID NP_000042.2

Catalog No. A300-135A GenelD 472

Lot No. A300-135A-2

APPLICATIONS WB, IP

SPECIES REACTIVITY Human

AMOUNT 100 μl

CONCENTRATION 1000 μg/ml

STORAGE/SHELF LIFE 2 – 8° C / 1 year from date of receipt

PHYSICAL STATE Liquid

BUFFER Tris-citrate/phosphate buffer, pH 7 to 8 containing 0.09% Sodium Azide

ISOTYPE IgG
ORIGIN USA

PRODUCTION PROCEDURES

Antibody was affinity purified using an epitope specific to ATM immobilized on solid support.

The epitope recognized by A300-135A maps to a region between residues 1750 and 1800 of human Ataxia Telangiectasia Mutated using the numbering given in entry NP_000042.2 (GeneID

472).

Immunoglobulin concentration was determined by extinction coefficient: absorbance at 280 nm

of 1.4 equals 1.0 mg of IgG.

APPLICATIONS Centrifuge tube to remove product from lid. Optimal working dilutions should be determined

experimentally by the investigator. Prepare working dilution immediately before use.

Western Blot 1:1,000 - 1:5,000

Immunoprecipitation 2 – 5 µg/mg lysate

APPLICATION NOTES Western blot of lysates performed using standard western blot reagents and 3–8% SDS-PAGE.

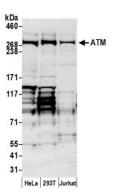
ADDITIONAL INFO https://www.bethyl.com/product/A300-135A

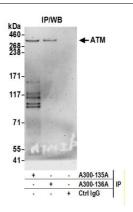
Use the link above to view SDS, a current list of citations, and other product specific information.

This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc. Eric McIntush, PhD | Chief Scientific Officer Date: June 21, 2019



ATM Antibody A300-135A





Detection of human ATM by western blot. Samples: Whole cell lysate (50 μg) from HeLa, HEK293T, and Jurkat cells prepared using NETN lysis buffer. Antibody: Affinity purified goat anti-ATM antibody A300-135A (lot A300-135A-2) used for WB at 0.4 μg/ml. Detection: Chemiluminescence with an exposure time of 75 seconds.

Detection of human ATM by western blot of immunoprecipitates. Samples: Whole cell lysate (1.0 mg per IP reaction; 20% of IP loaded) from HeLa cells prepared using NETN lysis buffer. Antibodies: Affinity purified goat anti–ATM antibody A300–135A (lot A300–135A–2) used for IP at 3 μg per reaction. ATM was also immunoprecipitated by goat anti–ATM antibody A300–136A. For blotting immunoprecipitated ATM, A300–135A was used at 1 μg/ml. Detection: Chemiluminescence with an exposure time of 3 minutes.