## **ATR Antibody**

Rabbit Polyclonal

Antigen Affinity Purified Protein ID NP\_001175.1

Catalog No. A300-137A GeneID 545

Lot No. A300-137A-3

**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** Antibody was affinity purified using an epitope specific to ATR immobilized on solid

**PROCEDURES** support.

The epitope recognized by A300-137A maps to a region between residues 400 and 450 of

human Ataxia Telangiectasia and Rad3-related using the numbering given in entry

NP\_001175.1 (GeneID 545).

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 1:2,000 - 1:10,000

Immunoprecipitation 2 – 10 µg/mg lysate

ADDITIONAL INFO https://www.bethyl.com/product/A300-137A

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

IP-western blot protocol: https://www.bethyl.com/content/protocol\_IP\_WB

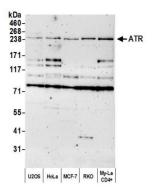
This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc.

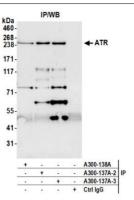
Brian McWilliams, PhD

Date: May 25, 2021



ATR Antibody A300-137A





Detection of human ATR by western blot. Samples: Whole cell lysate (10  $\mu$ g) from U2OS, HeLa, MCF-7, RKO, and My-La CD4+ cells prepared using NETN lysis buffer. Antibody: Affinity purified rabbit anti-ATR antibody (A300-137A lot 3) used for WB at 0.1  $\mu$ g/ml. Detection:

Chemiluminescence with an exposure time of 3 minutes.

Detection of human ATR 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 rabbit anti–ATR antibody (A300–137A lot 3) used for IP at 6 μg per reaction. ATR was also immunoprecipitated by a previous lot of this antibody (A300–137A lot 2) and a second antibody against a different epitope of rabbit ATR (A300–138A). For blotting immunoprecipitated ATR, A300–137A was used at 0.1 μg/ml. Detection: Chemiluminescence with an exposure time of 10 seconds.