

Phospho-ATR-T1989 Rabbit pAb

Catalog No.: AP1248

2 Publications

Basic Information

Observed MW

300kDa

Calculated MW

301kDa

Category

Primary antibody

Applications

ELISA, WB, IHC-P

Cross-Reactivity

Human, Mouse, Rat

Background

The protein encoded by this gene is a serine/threonine kinase and DNA damage sensor, activating cell cycle checkpoint signaling upon DNA stress. The encoded protein can phosphorylate and activate several proteins involved in the inhibition of DNA replication and mitosis, and can promote DNA repair, recombination, and apoptosis. This protein is also important for fragile site stability and centrosome duplication. Defects in this gene are a cause of Seckel syndrome 1.

Recommended Dilutions

WB 1:500 - 1:1000

IHC-P 1:50 - 1:200

Immunogen Information

Gene ID

545

Swiss Prot

Q13535

Immunogen

A synthetic phosphorylated peptide around T1989 of human ATR (NP_001175.2).

Synonyms

FRP1; MEC1; SCKL; FCTCS; SCKL1; Phospho-ATR-T1989

Contact

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Product Information

Source

Rabbit

Isotype

IgG

Purification

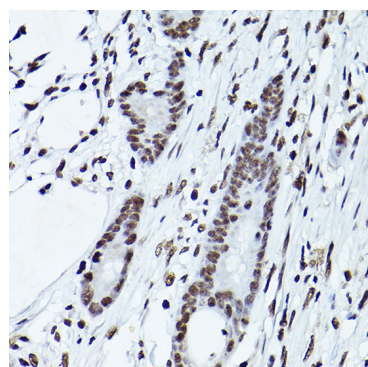
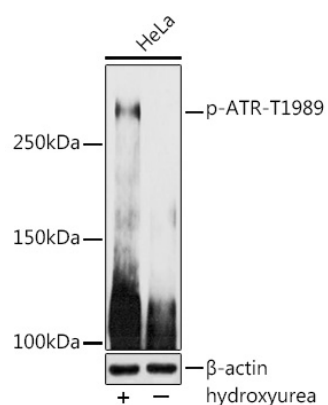
Affinity purification

Storage

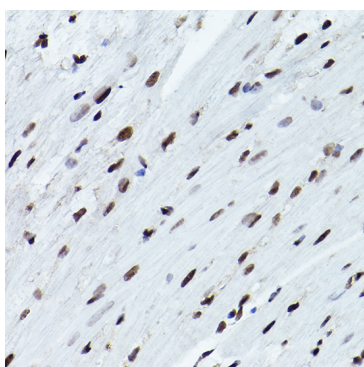
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300, 50% glycerol, pH7.3.

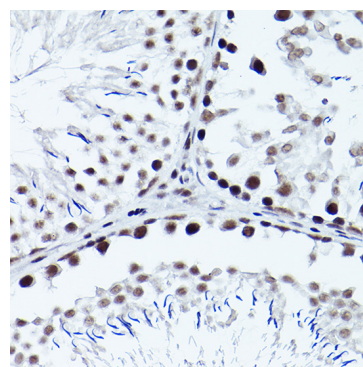
Validation Data



Immunohistochemistry analysis of Phospho-ATR-T1989 in paraffin-embedded human colon carcinoma using Phospho-ATR-T1989 Rabbit pAb (AP1248) at dilution of 1:100 (40x lens). Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.



Immunohistochemistry analysis of Phospho-ATR-T1989 in paraffin-embedded mouse heart using Phospho-ATR-T1989 Rabbit pAb (AP1248) at dilution of 1:100 (40x lens). Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.



Immunohistochemistry analysis of Phospho-ATR-T1989 in paraffin-embedded rat testis using Phospho-ATR-T1989 Rabbit pAb (AP1248) at dilution of 1:100 (40x lens). Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.