## Atherin Antibody

Rabbit Polyclonal

| Antigen Affinity Purified | Protein ID | NP_612361.1 |  |
| :--- | :--- | :--- | :--- |
| Catalog No. | A303-578A | GeneID | 90378 |

Lot No.
A303-578A-1

APPLICATIONS
WB, IP

SPECIES REACTIVITY
AMOUNT
CONCENTRATION
STORAGE/SHELF LIFE
PHYSICAL STATE
BUFFER
ISOTYPE
ORIGIN
PRODUCTION PROCEDURES

## APPLICATIONS

## APPLICATION NOTES

ADDITIONAL INFO

Human
$100 \mu \mathrm{l}$
$1000 \mu \mathrm{~g} / \mathrm{ml}$
$2-8^{\circ} \mathrm{C} / 1$ year from date of receipt
Liquid
Tris-citrate/phosphate buffer, pH 7 to 8 containing 0.09\% Sodium Azide
IgG
USA
Antibody was affinity purified using an epitope specific to Atherin immobilized on solid support.
The epitope recognized by A303-578A maps to a region between residue 488 and 538 of human Sterile Alpha Motif Domain Containing 1 using the numbering given in entry NP_612361.1 (GeneID 90378).

Antibody concentration was determined by extinction coefficient: absorbance at 280 nm of 1.4 equals 1.0 mg of IgG .

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 | $2-10 \mu \mathrm{~g} / \mathrm{mg}$ lysate |

Western blot of immunoprecipitates performed using Normal Pig Serum (Cat. No. S100-020), Goat anti-Rabbit Light Chain HRP Conjugate (Cat. No. A120-113P) and 4-20\% SDS-PAGE (link to IP-western blot protocol in Additional Info section below).

Western blot of lysates performed using standard western blot reagents and 4-20\% SDS-PAGE. https://www.bethyl.com/product/A303-578A
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. Eric McIntush, PhD \| Chief Scientific Officer

Date: June 21, 2019


Detection of human Atherin by western blot and immunoprecipitation. Samples: Whole cell lysate from HeLa (15 and $50 \mu \mathrm{~g}$ for WB; 1 mg for IP, 20\% of IP loaded), HEK293T (T; 50 $\mu \mathrm{g}$ ) and Jurkat (J; $50 \mu \mathrm{~g}$ ) cells. Antibodies: Affinity purified rabbit anti-Atherin antibody A303-578A used for WB at $0.1 \mu \mathrm{~g} / \mathrm{ml}(\mathrm{A})$ and $1 \mu \mathrm{~g} / \mathrm{ml}$ (B) and used for IP at $6 \mu \mathrm{~g} / \mathrm{mg}$ lysate. Atherin was also immunoprecipitated by rabbit anti-Atherin antibody BL12736, which recognizes an upstream epitope. Detection: Chemiluminescence with exposure times of 3 minutes (A) and 30 seconds (B).

