SA2 Antibody

Goat Polyclonal

Antigen Affinity Purified Protein ID NP_006594.3

Catalog No. A300-159A GenelD 10735

Lot No. A300-159A-2

APPLICATIONS WB, IP, ICC

SPECIES REACTIVITY Human, Mouse

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 SA2 immobilized on solid support.

The epitope recognized by A300–159A maps to a region between residues 1125 and 1175 of

human Stromal Antigen 2 using the numbering given in entry NP_006594.3 (GeneID 10735).

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 - 1:10,000

Immunocytochemistry 1:500 – 1:2,000

APPLICATION NOTES Western blot of lysates performed using standard western blot reagents and 4–8% SDS-PAGE.

2 - 10 µg/mg lysate

ADDITIONAL INFO https://www.bethyl.com/product/A300-159A

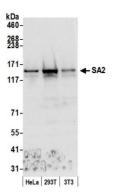
Immunoprecipitation

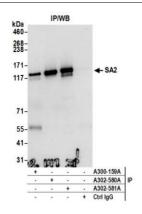
Use the link above to view SDS, a current list of citations, and other product specific information.

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



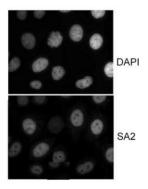
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Detection of human and mouse SA2 by western blot. Samples: Whole cell lysate (50 μ g) from HeLa, HEK293T, and mouse NIH 3T3 cells prepared using NETN lysis buffer. Antibody: Affinity purified goat anti–SA2 antibody A300–159A (lot A300–159A–2) used for WB at 0.1 μ g/ml. Detection: Chemiluminescence with an exposure time of 10 seconds.

Detection of human SA2 by western blot of immunoprecipitates. *Samples:* Whole cell lysate (0.5 or 1.0 mg per IP reaction; 20% of IP loaded) from HeLa cells prepared using NETN lysis buffer. *Antibodies:* Affinity purified goat anti–SA2 antibody A300–159A (lot A300–159A–2) used for IP at 6 μg per reaction. SA2 was also immunoprecipitated by rabbit anti–SA2 antibodies A302–581A and A301–582A. For blotting immunoprecipitated SA2, A300–159A was used at 1 μg/ml. *Detection:* Chemiluminescence with an exposure time of 10 seconds.



Localization of human SA2. *Sample:* HeLa cells that were extracted for 5 min. at 4C in 0.5% Triton in CSK buffer. *Antibody:* Affinity purified goat anti-SA2 (BL147G; Cat. No. A300-159A) used at 1 µg/ml.