

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in



ASH2 Antibody

Rabbit Polyclonal

Antigen Affinity Purified Protein ID Q9UBL3
Catalog No. A300–489A GeneID 9070

Lot No. A300-489A-2

APPLICATIONS WB, IP, IHC, ChIP, ChIP-chip

SPECIES REACTIVITY Human

PRESUMED REACTIVITY Based on 100% sequence identity, this antibody is predicted to react with 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 ASH2 immobilized on solid support.

The epitope recognized by A300-489A maps to a region between residue 575 and the C-terminus (residue 628) of human Absent, Small, or Homeotic-Like 2 using the numbering given

in Swiss-Prot entry Q9UBL3 (GeneID 9070).

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 - 1:20.000

Immunoprecipitation $2 - 5 \mu g/mg$ lysate

Immunohistochemistry 1:500 - 1:2,000. Epitope retrieval with citrate buffer pH 6.0 is

recommended for FFPE tissue sections.

ChIP 1 – 5 µg. Previous lots of this antibody have performed in this

application.

ChIP-chip 10 µg. Previous lots of this antibody have performed in this

application.

APPLICATION NOTES Western blot of immunoprecipitates performed using Normal Pig Serum (Cat. No. \$100-020),

Goat anti-Rabbit Light Chain HRP Conjugate (Cat. No. A120-113P) and 4-8% SDS-PAGE

(link to IP-western blot protocol in Additional Info section below).

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

IHC HUMAN CONTROLS Prostate Carcinoma

ADDITIONAL INFO https://www.bethyl.com/product/A300-489A

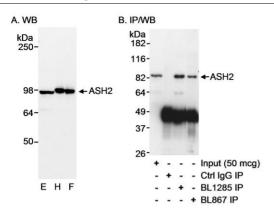
Use the link above to view SDS, a current list of citations, and other product specific information.

IP-western blot protocol: https://www.bethyl.com/content/protocol_IP_WB

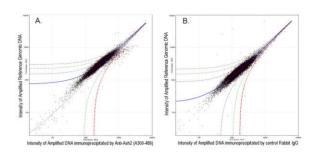
This document certifies that this product has met all of the quality control standards defined by Bethyl Laboratories, Inc. Eric McIntush, PhD | Chief Scientific Officer

Date: June 21, 2019

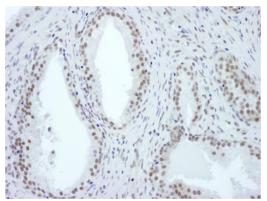
ASH2 Antibody A300-489A



Detection of human ASH2 by western blot and immunoprecipitation. Samples: A. Whole cell lysate from HEK293T cells that were mock transfected (E, 50 μg) or transfected with ASH2 expression constructs containing HA-tagged ASH2 (H, 25 μg) or Flag-tagged ASH2 (F, 25 μg). B. Whole cell lysate from one 10cm plate of normal 293T cells (~1 mg protein; 1/2 of IP loaded/lane). Antibodies: Affinity purified rabbit anti-ASH2 antibody A300-489A used at 1 μg/ml for WB (A and B) and at 5 μg/plate for IP. ASH2 was also immunoprecipitated with rabbit anti-ASH2 antibody A300-107A using 5 μg/plate. Detection: Chemiluminescence with an exposure time of 1 second (A and B).



ChIP-chip scatter plot of anti-Ash2 enriched DNA binding sites versus input reference DNA. A. 10 µg of A300-489A was used to immunoprecipitate chromatin from K-562 cells according to Ren et al (Genes Dev. 2002 16: 245-256). immunoprecipitatesd DNA and reference DNA were amplified via ligation-mediated PCR and the products labeled with fluorescent dNTPs. The labeled ChIP and reference DNA were pooled, hybridized to a DNA microarray, and analyzed. Data points below the +3 SD curve (red line) represent significantly enriched binding sites. B. As a control, a similar experiment was performed using normal rabbit IgG. Compared to the anti-Ash2 ChIP, normal rabbit IgG showed little enrichment.



Detection of human ASH2 by immunohistochemistry. *Sample:* FFPE section of human prostate carcinoma. *Antibody:* Affinity purified rabbit anti- ASH2 (Cat. No. A300-489A Lot2) used at a dilution of 1:1,000 (1µg/ml). *Detection:* DAB