

Produktinformation



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Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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SMARCA2/BRM Recombinant Monoclonal Antibody [BLR301M]



Rabbit Recombinant Monoclonal

Purified RefSeq ID NP_003061.3

Catalog No. A700-301 Uniprot ID P51531 Lot No. 1 GeneID 6595

APPLICATIONS WB, IP, Flow Cyt

SPECIES REACTIVITY Human

AMOUNT 100 μ l (50+ tests)

CONCENTRATION 250 μg/ml

STORAGE/SHELF LIFE 2 - 8°C / 1 year from date of receipt

PHYSICAL STATE Liquid

BUFFER Phosphate Buffered Saline (PBS) with 0.1% BSA and 0.09% Sodium Azide

ISOTYPE IgG

CLONE # BLR301M

ORIGIN USA

PRODUCTION PROCEDURES

Recombinant antibody was purified from cell culture supernatant.

Immunogen was a peptide representing a region between residue 1325 and 1375 of human SMARCA2/BRM using the numbering given in entry NP_003061.3 (Gene ID 6595).

SMARCA2/BRM using the numbering given in entry NP_003061.3 (Gene ID 6595).

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 1:1000

Immunoprecipitation 20 µl/mg lysate

Flow Cytometry Fixed in 4% formaldehyde and permeabilized with 90% methanol. 1 µl

per 1 x 10^6 cells.

APPLICATION NOTES All western blot analysis is performed using 5% Milk-TBST for blocking and as antibody diluent.

Primary antibody is incubated overnight.

Western blots of cell lysates are performed using Goat anti-Rabbit IgG Heavy and Light Chain

Antibody (Cat. No. A120-101P).

Western blots of immunoprecipitates are performed using Goat anti-Rabbit Light Chain HRP

Conjugate (Cat. No. A120-113P) with 5% Normal Pig Serum (Cat. No. S100-020) added to the

blocking buffer.

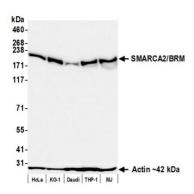
ADDITIONAL INFO https://www.fortislife.com/p/A700-301

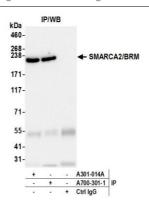
Use the link above to view SDS, a current list of citations, and other product specific information.

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

Michael Spencer, PhD

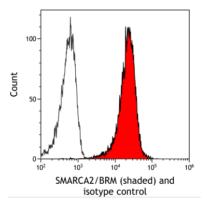
Date: December 26, 2023





Detection of human SMARCA2/BRM by western blot. Samples: Whole cell lysate (50 μg) from HeLa, KG-1, Daudi, THP-1, and MJ cells prepared using NETN lysis buffer. Antibody: Rabbit anti-SMARCA2/BRM recombinant monoclonal antibody [BLR301M] (A700-301 lot 1) used at 1:1000. Secondary: HRP-conjugated goat anti-rabbit IgG (A120-101P). Detection: Chemiluminescence with an exposure time of 10 seconds. Lower Panel: Rabbit anti-Actin recombinant monoclonal antibody [BLR057F] (A700-057).

Detection of human SMARCA2/BRM 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:* Rabbit anti–SMARCA2/BRM recombinant monoclonal antibody [BLR301M] (A700–301 lot 1) used for IP at 20 μl/mg lysate. SMARCA2/BRM was also immunoprecipitated by a second antibody against a different epitope of SMARCA2/BRM (A301–014A). For blotting immunoprecipitated SMARCA2/BRM, A700–301 was used at 1:1000. *Detection:* Chemiluminescence with an exposure time of 30 seconds.



Detection of human SMARCA2/BRM (shaded) in HeLa cells by flow cytometry. *Antibody:* Rabbit anti–SMARCA2/BRM recombinant monoclonal antibody [BLR301M] (A700–301) or isotype control (unshaded). *Secondary:* DyLight® 650–conjugated goat anti–rabbit IgG (A120–101D5).