## p16INK4a Recombinant Monoclonal Antibody [BLR318M]



Rabbit Recombinant Monoclonal

Purified RefSeq ID NP\_000068.1

Catalog No. A700-318 Uniprot ID P42771 Lot No. 1 GeneID 1029

**APPLICATIONS** WB, IP, IHC, ICC, Flow Cyt

SPECIES REACTIVITY Human

**AMOUNT** 100 μl (50+ tests)

CONCENTRATION 500 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 # BLR318M

**ORIGIN** USA

PRODUCTION PROCEDURES

Recombinant antibody was purified from cell culture supernatant.

Immunogen was a peptide representing a region between residue 106 and the C-terminus of human Cyclin-dependent kinase inhibitor 2A isoform p16INK4a using the numbering

given in entry NP\_000068.1 (Gene ID 1029).

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

Immunoprecipitation 12 µl/mg lysate

Immunohistochemistry 1:100 to 1:500. Epitope retrieval with citrate buffer pH 6.0 is

recommended for FFPE tissue sections.

Immunocytochemistry 1:100 to 1:500. Epitope retrieval with citrate buffer pH 6.0 is

recommended for FFPE cell sections.

Flow Cytometry Fixed in 4% formaldehyde and permeabilized with 90% methanol. 0.5

ul 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 (A120-101P).

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

Conjugate (A120-113P) with 5% Normal Pig Serum (S100-020) added to the blocking buffer.

IHC HUMAN CONTROLS Breast Carcinoma, Ovarian Carcinoma, A-2058 Cells, HEK293T Cells, HeLa Cells, OVCAR-8 Cells,

SUP-T1 Cells

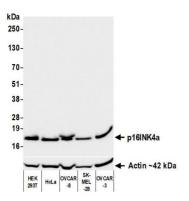
ADDITIONAL INFO https://www.fortislife.com/p/A700-318

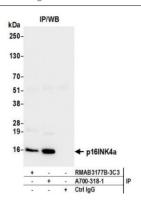
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Michael Spencer, PhD

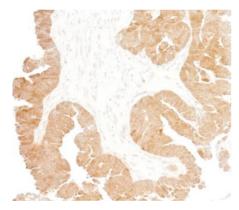
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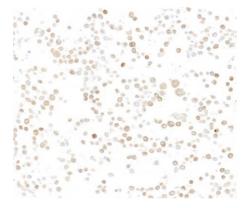




Detection of human p16lNK4a by western blot. Samples: Whole cell lysate (50 μg) from HEK293T, HeLa, OVCAR-8, SK-MEL-28, and OVCAR-3 cells prepared using NETN lysis buffer. Antibody: Rabbit anti-p16lNK4a recombinant monoclonal antibody [BLR318M] (A700-318 lot 1) used at 1:1000. Secondary: HRP-conjugated goat anti-rabbit IgG (A120-101P). Detection: Chemiluminescence with an exposure time of 3 seconds.

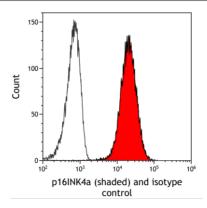
Detection of human p16INK4a by western blot of immunoprecipitates. Samples: Whole cell lysate (1.0 mg per IP reaction; 20% of IP loaded) from HEK293T cells prepared using NETN lysis buffer. Antibodies: Rabbit anti-p16INK4a recombinant monoclonal antibody [BLR318M] (A700–318 lot 1) used for IP at 12 μl/mg lysate. p16INK4a was also immunoprecipitated by a second antibody against a different epitope of p16INK4a (RMAB3177B–3C3). For blotting immunoprecipitated p16INK4a, A700–318 was used at 1:1000. Detection: Chemiluminescence with an exposure time of 10 seconds.





**Detection of human p16INK4a by immunohistochemistry.** *Sample:* FFPE section of ovarian carcinoma. *Antibody:* Rabbit anti-p16INK4a recombinant monoclonal antibody [BLR318M] (A700-318). *Secondary:* HRP-conjugated goat anti-rabbit IgG (A120-501P).

**Detection of human p16INK4a by immunocytochemistry.** *Sample:* FFPE section of HEK293T cells. *Antibody:* Rabbit anti-p16INK4a recombinant monoclonal antibody [BLR318M] (A700-318). *Secondary:* HRP-conjugated goat anti-rabbit IgG (A120-501P).



Detection of human p16INK4a (shaded) in HeLa cells by flow cytometry. *Antibody:* Rabbit anti-p16INK4a recombinant monoclonal antibody [BLR318M] (A700-318) or isotype control (unshaded). *Secondary:* DyLight® 650-conjugated goat anti-rabbit IgG (A120-101D5).