## VE-Cadherin Recombinant Monoclonal Antibody [BLR239K]



Rabbit Recombinant Monoclonal

Purified RefSeq ID NP\_033998.2

Catalog No. A700-239 Uniprot ID P55284 Lot No. 1 GeneID 12562

**APPLICATIONS** WB, IP, IHC, ICC

SPECIES REACTIVITY Mouse

**AMOUNT** 100  $\mu$ l (50+ tests)

CONCENTRATION 500 μg/ml

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

PHYSICAL STATE Liquid

BUFFER Borate Buffered Saline (BBS) pH 8.2 with 0.1% BSA and 0.09% Sodium Azide

**ISOTYPE** IgG

CLONE # BLR239K
ORIGIN USA

PRODUCTION

Recombinant antibody was purified from cell culture supernatant.

**PROCEDURES** 

Immunogen was a recombinant protein representing the extracellular domain of mouse

Cadherin-5 (aka VE-Cadherin, residues 25-595, NP\_033998.2).

**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 µI/1 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.

**IHC MOUSE CONTROLS** 

Gut, Lung, Uterus, bEnd.3 Cells

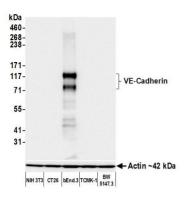
ADDITIONAL INFO https://www.fortislife.com/p/A700-239

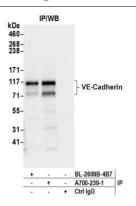
Use the link above to view SDS, a current list of citations, and other product specific information.

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Michael Spencer, PhD

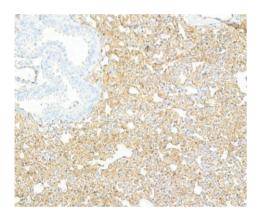
Date: April 14, 2023





Detection of mouse VE–Cadherin by western blot. *Samples:* Whole cell lysate (2 μg) from NIH 3T3, CT26, bEnd.3, TCMK–1, and BW5147.3 cells prepared using NETN lysis buffer. *Antibody:* Rabbit anti–VE–Cadherin recombinant monoclonal antibody [BLR239K] (A700–239 lot 1) used at 1:1000. *Secondary:* HRP–conjugated goat anti–rabbit IgG (A120–101P). *Detection:* Chemiluminescence with an exposure time of 3 seconds. Lower Panel: Rabbit anti–Actin recombinant monoclonal antibody [BLR057F] (A700–057).

Detection of mouse VE-Cadherin by western blot of immunoprecipitates. *Samples:* Whole cell lysate (1.0 mg per IP reaction; 5% of IP loaded) from bEnd.3 cells prepared using NETN lysis buffer. *Antibodies:* Rabbit anti-VE-Cadherin recombinant monoclonal antibody [BLR239K] (A700–239 lot 1) used for IP at 12 µl/mg lysate. VE-Cadherin was also immunoprecipitated by a second antibody against a different epitope of VE-Cadherin (BL-2609B-4B7). For blotting immunoprecipitated VE-Cadherin, A700–239 was used at 1:1000. *Detection:* Chemiluminescence with an exposure time of 1 second.



**Detection of mouse VE–Cadherin by immunohistochemistry.** *Sample:* FFPE section of mouse lung. *Antibody:* Rabbit anti–mVE–Cadherin recombinant monoclonal antibody [BLR239K] (A700–239). *Secondary:* HRP–conjugated goat anti–rabbit IgG (A120–501P).

**Detection of mouse VE–Cadherin by immunohistochemistry.** *Sample:* FFPE section of mouse uterus. *Antibody:* Rabbit anti–mVE–Cadherin recombinant monoclonal antibody [BLR239K] (A700–239). *Secondary:* HRP–conjugated goat anti–rabbit IgG (A120–501P).



**Detection of mouse VE-Cadherin by immunocytochemistry.** *Sample:* FFPE section of bEnd cells. *Antibody:* Rabbit anti-mVE-Cadherin recombinant monoclonal antibody [BLR239K] (A700-239). *Secondary:* HRP-conjugated goat anti-rabbit IgG (A120-501P).