

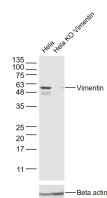
bs-0756R**[Primary Antibody]****Bioss**
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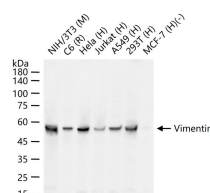
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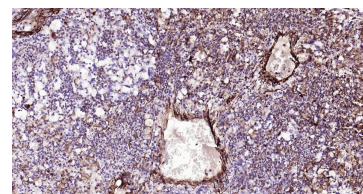
400-901-9800

Vimentin Rabbit pAb**DATASHEET****Host:** Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 7431**SWISS:** P08670**Target:** Vimentin**Immunogen:** KLH conjugated synthetic peptide derived from human Vimentin: 371-466/466.**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.**Background:** This gene encodes a member of the intermediate filament family. Intermediate filaments, along with microtubules and actin microfilaments, make up the cytoskeleton. The protein encoded by this gene is responsible for maintaining cell shape, integrity of the cytoplasm, and stabilizing cytoskeletal interactions. It is also involved in the immune response, and controls the transport of low-density lipoprotein (LDL)-derived cholesterol from a lysosome to the site of esterification. It functions as an organizer of a number of critical proteins involved in attachment, migration, and cell signaling. Mutations in this gene causes a dominant, pulverulent cataract.[provided by RefSeq, Jun 2009]**Applications:** WB (1:500-2000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (1µg/Test)**ICC/IF** (1:100)**Reactivity:** Human, Mouse, Rat
(predicted: Pig, Cow, Chicken, Goat)**Predicted MW.:** 53 kDa**Subcellular Location:** Cytoplasm**VALIDATION IMAGES**

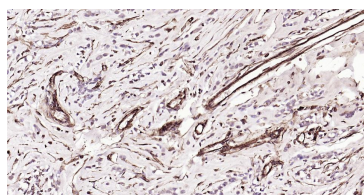
Sample: HeLa(Human) Cell Lysate at 30 ug
 KO Vimentin (Human) Cell Lysate at 30 ug
 Primary: Anti- Vimentin (bs-0756R) at 1/1000
 dilution Secondary: IRDye800CW Goat Anti-
 Rabbit IgG at 1/20000 dilution Predicted band
 size: 53 kD Observed band size: 53 kD



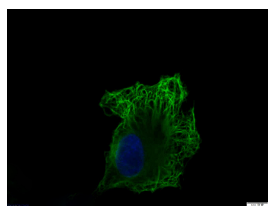
25 ug total protein per lane of various lysates
 (see on figure) probed with Vimentin polyclonal
 antibody, unconjugated (bs-0756R) at 1:1000
 dilution and 4°C overnight incubation. Followed
 by conjugated secondary antibody incubation at
 r.t. for 60 min.



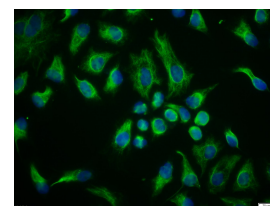
Paraformaldehyde-fixed, paraffin embedded
 Human Endometrium Cancer; Antigen retrieval
 by boiling in sodium citrate buffer (pH6.0) for 15
 min; Antibody incubation with Vimentin
 Polyclonal Antibody, Unconjugated (bs-0756R)
 at 1:200 overnight at 4°C, followed by
 conjugation to the SP Kit (Rabbit, SP-0023) and
 DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded
 Human Breast Cancer; Antigen retrieval by
 boiling in sodium citrate buffer (pH6.0) for 15
 min; Antibody incubation with Vimentin
 Polyclonal Antibody, Unconjugated (bs-0756R)
 at 1:200 overnight at 4°C, followed by



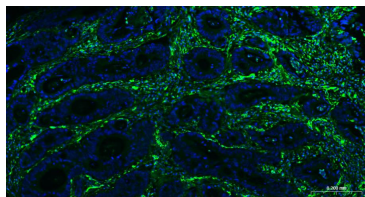
Tissue/cell: U-87MG cell; 4% Paraformaldehyde-
 fixed; Triton X-100 at room temperature for 20
 min; Blocking buffer (normal goat serum,
 C-0005) at 37°C for 20 min; Antibody incubation
 with (Vimentin) Polyclonal Antibody,
 Unconjugated (bs-0756R) 1:100, 90 minutes at



Tissue/cell: HeLa cell; 4% Paraformaldehyde-
 fixed; Triton X-100 at room temperature for 20
 min; Blocking buffer (normal goat serum,
 C-0005) at 37°C for 20 min; Antibody incubation
 with (Vimentin) Polyclonal Antibody,
 Unconjugated (bs-0756R) 1:50, 90 minutes at

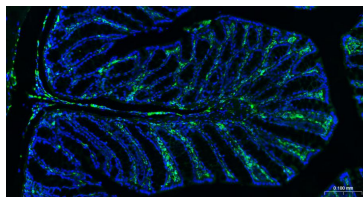
Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

conjugation to the SP Kit (Rabbit, SP-0023) and DAB (C-0010) staining.



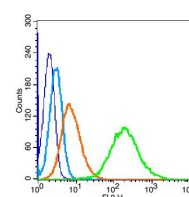
Paraformaldehyde-fixed, paraffin embedded Human Colon Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with Vimentin Polyclonal Antibody, Unconjugated (bs-0756R) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-0295G-BF488), DAPI (blue, C02-04002) was used to stain the cell nuclei.

37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody (bs-0295G-FITC) at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.

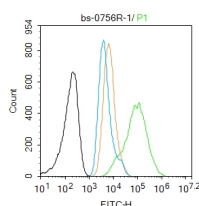


Paraformaldehyde-fixed, paraffin embedded Mouse Colon; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with Vimentin Polyclonal Antibody, Unconjugated (bs-0756R) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-0295G-BF488), DAPI (blue, C02-04002) was used to stain the cell nuclei.

37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody (bs-0295G-FITC) at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Blank control: Jurkat cells(blue). Primary Antibody:Rabbit Anti-Vimentin antibody antibody(bs-0756R), Dilution: 1µg in 100 µL 1X PBS containing 0.5% BSA; Isotype Control Antibody: Rabbit IgG(orange), used under the same conditions; Secondary Antibody: Goat anti-rabbit IgG-PE(white blue), Dilution: 1:200 in 1 X PBS containing 0.5% BSA. Protocol The cells were fixed with 2% paraformaldehyde (10 min), then permeabilized with 90% ice-cold methanol for 30 min on ice. Primary antibody (bs-0756R, 1µg /1x10⁶ cells) were incubated for 30 min on the ice, followed by 1 X PBS containing 0.5% BSA + 1 0% goat serum (15 min) to block non-specific protein-protein interactions. Then the Goat Anti-rabbit IgG/PE antibody was added into the blocking buffer mentioned above to react with the primary antibody at 1/200 dilution for 30 min on ice. Acquisition of 20,000 events was performed.



Blank control:A549. Primary Antibody (green line): Rabbit Anti-Vimentin antibody (bs-0756R) Dilution: 1µg /10⁶ cells; Isotype Control Antibody (orange line): Rabbit IgG. Secondary Antibody : Goat anti-rabbit IgG-AF488 Dilution: 1µg /test. Protocol The cells were fixed with 4% PFA (10min at room temperature)and then permeabilized with 90% ice-cold methanol for 20 min at -20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

— SELECTED CITATIONS —

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