

bsm-33187M**[Primary Antibody]****alpha smooth muscle Actin Mouse mAb****BioSS**
ANTIBODIES

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— DATASHEET —**Host:** Mouse**Isotype:** IgG**Clonality:** Monoclonal**CloneNo.:** 3F9**GeneID:** 59**SWISS:** P62736**Target:** alpha smooth muscle Actin**Purification:** affinity purified by Protein G**Concentration:** 1mg/ml

Storage: Size : 50ul/100ul/200ul
0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.
Size : 200ug (PBS only)
0.01M PBS
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.

Background: All eukaryotic cells express Actin, which often constitutes as much as 50% of total cellular protein. Actin filaments can form both stable and labile structures and are crucial components of microvilli and the contractile apparatus of muscle cells. While lower eukaryotes, such as yeast, have only one Actin gene, higher eukaryotes have several isoforms encoded by a family of genes. At least six types of Actin are present in mammalian tissues and fall into three classes. alpha-Actin expression is limited to various types of muscle, whereas beta- and gamma-Actin are the principle constituents of filaments in other tissues. Members of the small GTPase family regulate the organization of the Actin cytoskeleton. Rho controls the assembly of Actin stress fibers and focal adhesion. Rac regulates Actin filament accumulation at the plasma membrane. Cdc42 stimulates formation of filopodia.

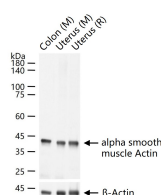
Applications: WB (1:1000-5000)**IHC-P** (1:200-500)**IHC-F** (1:200-500)**IF** (1:200-500)**Flow-Cyt** (1ug/Test)**ICC/IF** (1:100-200)

Reactivity: Human, Mouse, Rat
(predicted: Rabbit, Chicken)

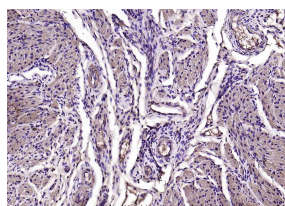
Predicted
MW.: 42 kDa

Observed
MW.: 42 kDa

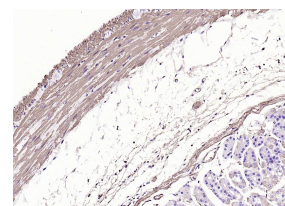
Subcellular
Location: Cytoplasm

— VALIDATION IMAGES —

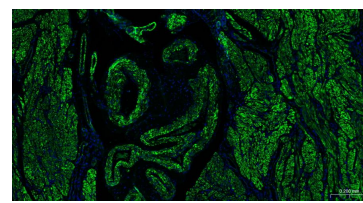
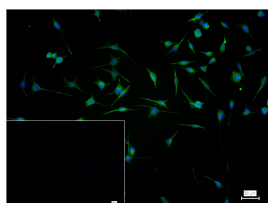
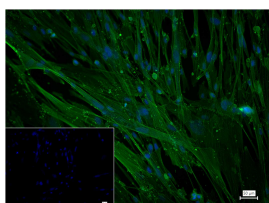
25 ug total protein per lane of various lysates (see on figure) probed with alpha smooth muscle Actin monoclonal antibody, unconjugated (bsm-33187M) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Paraformaldehyde-fixed, paraffin embedded (rat uterus); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (alpha smooth muscle Actin) Monoclonal Antibody, Unconjugated (bsm-33187M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (mouse stomach); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (alpha smooth muscle Actin) Monoclonal Antibody, Unconjugated (bsm-33187M) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Mouse)(sp-0024) instructions and DAB staining.

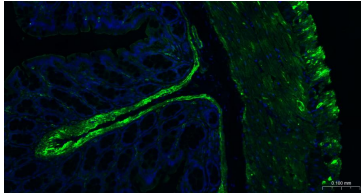


Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

4% Paraformaldehyde-fixed WI-38 (treated with 5 ng/mL TGF- β 1 for 72h) (H) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (alpha smooth muscle Actin) monoclonal Antibody, unconjugated (bsm-33187M) 1:100, 90 min at 37°C; followed by BF488 conjugated Goat Anti-Mouse IgG antibody (green, bs-60296G-BF488) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.

4% Paraformaldehyde-fixed NIH/3T3 treated with 5 ng/mL TGF- β 1 for 72h (M) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (alpha smooth muscle Actin) monoclonal Antibody, unconjugated (bsm-33187M) 1:100, 90 min at 37°C; followed by BF488 conjugated Goat Anti-Mouse IgG antibody (green, bs-60296G-BF488) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.

Paraformaldehyde-fixed, paraffin embedded Human Uterus; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with alpha smooth muscle Actin Monoclonal Antibody, Unconjugated (bsm-33187M) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Mouse IgG antibody (green, bs-0296G-FITC), DAPI (blue, C02-04002) was used to stain the cell nuclei.



Paraformaldehyde-fixed, paraffin embedded Rat Colon; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with alpha smooth muscle Actin Monoclonal Antibody, Unconjugated (bsm-33187M) at 1:200 overnight at 4°C. Followed by conjugated Goat Anti-Mouse IgG antibody (green, bs-0296G-FITC), DAPI (blue, C02-04002) was used to stain the cell nuclei.

— SELECTED CITATIONS —

- **[IF=18]** Zetao Wang. et al. Nano-vibration exciter: Hypoxia-inducible factor 1 signaling pathway-mediated extracellular vesicles as bioactive glass substitutes for bone regeneration. BIOACT MATER. 2024 Oct;40:460 IF ;Rat. 10.1016/j.bioactmat.2024.06.023
- **[IF=16.6]** Li Yin. et al. UPP1 promotes lung adenocarcinoma progression through the induction of an immunosuppressive microenvironment. NAT COMMUN. 2024 Feb;15(1):1-23 IF ;Human. 38331898
- **[IF=15.7]** Li Xiao. et al. Low-intensity focused ultrasound-activated piezoelectric gel bandage for diabetic wound repair and neuropathic pain relief. NAT COMMUN. 2026 Mar;; IF ;Rat. 41857055
- **[IF=13.6]** Juan Yan. et al. Engineered exosomes reprogram Gli1+ cells in vivo to prevent calcification of vascular grafts and autologous pathological vessels. SCI ADV. 2023 Jul;9(29) WB ;Human. 37478186
- **[IF=11.5]** Ning Yang. et al. Nintedanib-amplified effects of HSP47 siRNA on liver fibrosis therapy by inhibiting collagen secretion. J CONTROL RELEASE. 2025 Nov;;114378 IF ;MOUSE. 41187814