

Product datasheet

anti-alpha-Smooth Muscle Actin mouse monoclonal, 1A4/ASM-1, liquid, purified, sample

Short overview

Cat. No. 690001S

 Quantity
 200 μl (100 μg/ml)

 Concentration
 100 μg/ml (20 μg)

Product description

HostMouseAntibody TypeMonoclonalIsotypeIgG2aClone1A4/ASM-1

Immunogen Synthetic N-terminus decapeptide of alpha-smooth-muscle isoform of actin

Formulation PBS, pH 7.4 with 0.09% sodium azide and 0.5% BSA

UniprotIDP62739 (Bovine), P08023 (Chicken), P62736 (Human), P62737 (Mouse), P62738 (Rat)SynomymActin, aortic smooth muscle, Alpha-actin-2, Cell growth-inhibiting gene 46 protein [Cleaved into:

Actin, aortic smooth muscle, intermediate form], ACTA2, ACTSA, ACTVS, GIG46

Conjugate Unconjugated

Purification Affinity chromatography

Storage Short term at 2-8°C; long term storage in aliguots at -20°C; avoid freeze/thaw cycles

Intended use Research use only Application ICC/IF, IHC, WB

Reactivity Bovine, Chicken, Horse, Human, Mouse, Rat

Applications

Immunocytochemistry (ICC) Assay dependent

Immunohistochemistry (IHC) - frozen 1:200-1:1,000 (100-500 ng/ml)

Immunohistochemistry (IHC) - paraffin 1:200-1:1,000 (100-500 ng/ml, protease treatment and/or microwave

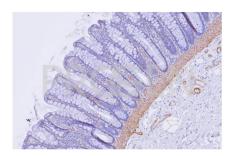
treatment recommended)

Western Blot (WB) 1:1,000 (100 ng/ml)

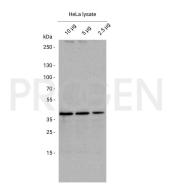
Background

1A4/ASM-1 represents an excellent marker for myogenic soft tissue tumors and smooth muscle differentiation. Polypeptide reacting: specific for alpha-smooth-muscle isoform of actin (43 kDa). Tumors specifically detected: leiomyosarcoma, leiomyoma, certain stromal cells surrounding infiltrating ductal carcinoma of breast. Tested cultured cell lines: Stress fibers of smooth muscle-derived cells and some smooth muscle subtype fibroblasts

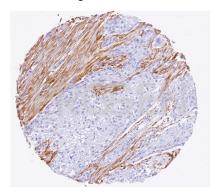
Product images



IHC analysis of human colon using anti-alpha-Smooth Muscle Actin antibody. IHC was performed on formalin fixed paraffin embedded sections. The samples were deparaffinized with xylol and ethanol followed by heat induced antigen retrieval with 10 mM citrate buffer. After preparation the tissue was blocked with normal serum for 20 min at RT. The primary antibody anti-alpha-Smooth Muscle Actin mouse monoclonal, 1A4/ASM-1 (Cat. No. 690001) was diluted in PBS (antibody concentration 400 ng/ml) and incubated at 4°C over-night. The secondary antibody ImmPRESS HRP anti-mouse IgG was incubated for 20 min at RT. Slides were incubated with DAB solution until a brown staining is visable and with Haemalaun for a few minutes. The 10x picture was acquired using microscopy (courtesy of J.Hess, University Hospital Heidelberg).



Western blot analysis of human HeLa cell lysate with anti-alpha-Smooth Muscle Actin antibody. Western blot analysis was performed on either $10~\mu g$, $5~\mu g$ or $2.5~\mu g$ of HeLa lysate. Cells were lysed in PBS by homogenization. The PVDF membrane was blocked with 5% dry milk in PBST for 1~h at RT. The primary antibody anti-alpha-Smooth Muscle Actin mouse monoclonal, 1A4/ASM-1 (Cat. No. 690001) was diluted in blocking buffer (antibody concentration $0.1~\mu g/ml$) and incubated for 1~h at RT. The secondary antibody goat anti-mouse IgG polyclonal, HRP conjugate was also diluted in blocking buffer (antibody concentration $0.2~\mu g/ml$) and incubated for 1~h at RT. The bands were visualized by chemiluminescent detection using PierceTM ECL Western Blotting Substrate.



IHC analysis of head and neck squamous cell carcinoma using anti-alpha-Smooth Muscle Actin antibody. IHC was performed on formalin fixed paraffin embedded sections. The samples were deparaffinized with xylol and ethanol followed by heat induced antigen retrieval with 10 mM citrate buffer. After preparation the tissue was blocked with normal serum for 20 min at RT. The primary antibody anti-alpha-Smooth Muscle Actin mouse monoclonal, 1A4/ASM-1 (Cat. No. 690001) was diluted in PBS (antibody concentration 200 ng/ml) and incubated at 4°C over-night. The secondary antibody ImmPRESS HRP anti-mouse IgG was incubated for 20 min at RT. Slides were incubated with DAB solution until a brown staining is visable and with Haemalaun for a few minutes. The 10x picture was acquired using microscopy (courtesy of J.Hess, University Hospital Heidelberg).

References

Publication	Species	Application
Poosti, F. et al. Inhibition of renal fibrosis with a human	Human, Mouse	ICC-IF, IHC-P-IF
CXCL9-derived glycosaminoglycan-binding peptide. Clin.		
<u>Transl. Immunol. 11, 1–18 (2022).</u>		
Jiang, D. et al. MSCs rescue impaired wound healing in a	human,mouse	WB,IHC-IF (paraffin),ICC-IF
murine LAD1 model by adaptive responses to low TGF-Î21		
levels. EMBO.Rep. 21, e49115 (2020)		
Schwinghammer, U. et al. α2-Adrenergic Receptor in Liver	human	IHC-IF,ICC-IF
Fibrosis: Implications for the Adrenoblocker Mesedin. Cells. 9,	_	
(2020)		
Buniatian, G. et al. Antifibrotic Effects of Amyloid-Beta and Its	mouse	IHC-IF,IHC (paraffin)
Loss in Cirrhotic Liver. Cells. 9, (2020)		
Munir, S. et al. TLR4-dependent shaping of the wound site by	mouse	IHC (frozen)/IF
MSCs accelerates wound healing. EMBO Rep. 21,		
e48777(2020).		