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## Anti-ACTA2 Antibody, Mouse Monoclonal, A1T

ACT-A2, ACTSA, ACTVS, Asma, GIG46

Catalog: CM00526HuM10-A1T Size:100µL

Basic Info Host Species Reactivity

Mouse Predict reacts with: Human/Mouse/Rat

Conjugate Clonality, Isotype

None

Concentration Immunogen

1.0 mg/mL A synthetic peptide of human ACTA2

Physical State Purification

Liquid Immunogen affinity purified

Clone No Applications

A1T WB/IHC

Property Form & Buffer: Supplied in PBS, 50% glycerol, PH7.4.

Specificity / Sensitivity: CM00526HuM10-A1T recognizes endogenous levels

of human ACTA2.

Applications The application notes include recommended starting dilutions;

optimal dilutions/concentrations should be determined by the end user.

WB:  $0.5 \sim 5 \mu \text{ g/mL}$  IF:  $5 \sim 20 \mu \text{ g/mL}$  IHC:  $5 \sim 20 \mu \text{ g/mL}$  ICC:  $5 \sim 20 \mu \text{ g/mL}$ 

Usage and Store at 4

Store at  $4^{\circ}C$  for frequent use.

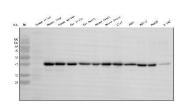
Storage

Aliquot and store at -20℃ for 12 months.

Avoid repeated freezing/thawing and violent shaking.

Please centrifuge it, before using.

QC Data



## Figure 1. Application in WB

Western blot analysis of extracts of various cells lines and tissues, using ACTA2 antibody CM00526HuM10-A1T at 1  $\mu$  g/mL. Secondary antibody: HRP Mouse Anti-Rabbit IgG (H+L) at 1: 20000 dilution. Lysates/proteins: 25ug per lane.Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 3min.



QC Data



## Figure 2. Application in IHC

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon, human lung, monkey epididymis, monkey heart, monkey kidney, monkey liver, monkey rectum, monkey stomach, monkey tongue sections labelling ACTA2 with purified CM00526HuM10-A1T at 5  $\mu$  g/mL dilution. Heat mediated antigen retrieval was performed using Heat mediated antigen retrieval using citrate buffer (PH6.0). Tissue was counterstained with Hematoxylin. Mouse specific IHC polymer detection kit HRP/DAB secondary antibody was used at 1: 2000 dilution. PBS instead of the primary antibody was used as the negative control.

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