

Name: Jiangxi Wellanimal Gene Technology Co., Ltd.
Address: 5F, Building 2, Ganzhou National High-level Talents
Science and Innovation Park(Phase II), 1 Wudang Mountain Road,
Zhanggong District High-tech zone, Ganzhou City, Jiangxi Province.

Telephone: 400-6723-898 E-mail: biolight_info@163.com Website: http://www.wellanimal.com

Anti-HSP90AA1 Antibody, Mouse Monoclonal, G1F

HSP90A/HSPC1/HSPCA/HSP 86/HSP86/LAP-2/LPS-associated protein 2

Catalog: MA01006HuM10-G1F Size:100µL

Basic Info Host Species Reactivity

Mouse Human/Mouse/Rat

Conjugate Clonality, Isotype

None

Concentration Immunogen

1 mg/mL

Physical State Purification

Liquid Immunogen affinity purified

Clone No Applications

G1F WB/IHC

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

Specificity / Sensitivity: MA01006HuM10-G1F recognizes endogenous levels of

human HSP90AA1.

Applications The application notes include recommended starting dilutions;

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

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

Usage and

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

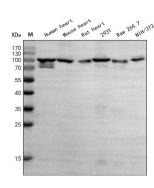


Figure 1. Application in WB

Western blot analysis of extracts of various cell lines and tissues, using HSP90AA1 antibody MAO1006HuM10-G1F at $5\mu g/mL$ Secondary antibody: HRP Mouse Anti-Rabbit IgG (H+L) at 1: 5000 dilution. Lysates/proteins: $25\mu g$ per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 1s.



QC Data



For Research Use Only!

Figure 2. Application in IHC

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human liver cancer sections labelling HSP90AA1 with purified MA01006HuM10-G1F at $10\mu g/mL$ 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.