# Anti-KNG1 antibody

Catalog Number: 176585



#### **Product name**

Anti-KNG1 antibody

## **Specificity**

Human, Mouse

## Antibody description

Rabbit monoclonal antibody to KNG1

## **Preparation**

This antigen of this antibody was recombinant protein within human kiningen 1 aa 500-644.

#### **Formulation**

Liquid, 1\*TBS (pH7.4), 0.05% BSA, 40% Glycerol. Preservative: 0.05% Sodium Azide.

# Storage

Store at +4°C after thawing. Aliquot store at -20°C. Avoid repeated freeze / thaw cycles.

### **Clonality**

Monoclonal

## Ig Type

Rabbit IgG

### **Applications**

WB, IHC-P

#### **Dilutions**

WB: 1:500-1:2,000

IHC-P: 1:50-1:200

#### **Validations**

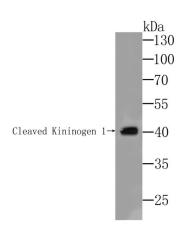
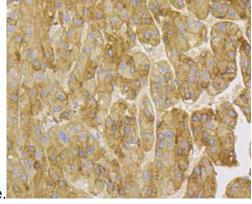


Fig1: Western blot analysis of Kininogen 1 on mouse stomach lysate. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody was used at a 1:500 dilution in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:5,000 dilution was used for 1 hour at room



temperature

Fig2: Immunohistochemical analysis of paraffinembedded human liver tissue using anti-Kininogen 1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the antibody at 1/100 dilution, for 30 minutes at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chrogen. Counter stained with hematoxylin and mounted with DPX.

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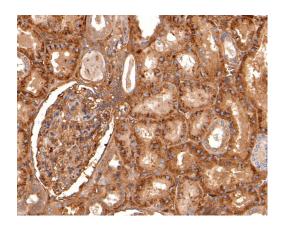


Fig3: Immunohistochemical analysis of paraffin-

embedded human kidney tissue using anti-Kininogen 1 antibody. The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 8.0-8.4) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the antibody at 1/100 dilution, for 30 minutes at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chrogen. Counter stained with hematoxylin and mounted with DPX.