# **Anti-IL22 antibody**

Catalog Number: 175137



#### **Product name**

Anti-IL22 antibody

## **Specificity**

Human, Mouse, Rat

## **Antibody description**

Rabbit polyclonal antibody to IL22

## Preparation

This antigen of this antibody was recombinant protein within human il-22 aa 34-179.

#### **Formulation**

Liquid, 1\*PBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide.

## **Storage**

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

## **Clonality**

Polyclonal

#### Ig Type

Rabbit IgG

### **Applications**

ELISA, WB, IHC-P

### **Dilutions**

ELISA:1:5000-1:10,000

WB:1:500

IHC-P:1:50-1:200

# Validations

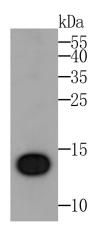
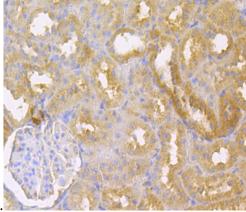


Fig1: Western blot analysis of IL-22 on IL22 recombinant protein. Proteins were transferred to a PVDF membrane and blocked with 5% BSA in PBS for 1 hour at room temperature. The primary antibody was used at a 1:500 dilution in 5% BSA at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:5,000 dilution was used for 1 hour at room



temperature.

Fig2: Immunohistochemical analysis of paraffinembedded rat kidney tissue using anti-IL-22 antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) 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 1/200 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.

# **Anti-IL22 antibody**

Catalog Number: 175137



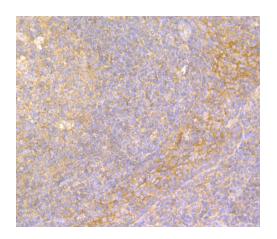


Fig3: Immunohistochemical analysis of paraffinembedded human tonsil tissue using anti-IL-22 antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the antibody at 1/200 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.

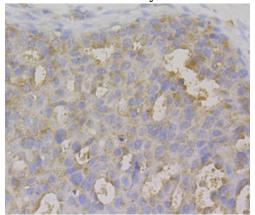


Fig4: Immunohistochemical analysis of paraffinembedded human breasr cancer tissue using anti-IL-22 antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the antibody at 1/200 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.

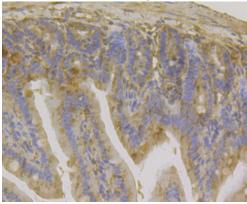


Fig5: Immunohistochemical analysis of paraffinembedded mouse colon tissue using anti-IL-22 antibody. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0) for 20 minutes. The tissues were blocked in 5% BSA for 30 minutes at room temperature, washed with ddH2O and PBS, and then probed with the antibody at 1/200 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.