# Phospho-EGFR-T669 Rabbit pAb

Catalog No.: AP0025 1 Publications



## **Basic Information**

#### **Observed MW**

200kDa

## **Calculated MW**

134kDa

### Category

Primary antibody

## **Applications**

WB,IP,ELISA

#### **Cross-Reactivity**

Human

# **Background**

The protein encoded by this gene is a transmembrane glycoprotein that is a member of the protein kinase superfamily. This protein is a receptor for members of the epidermal growth factor family. EGFR is a cell surface protein that binds to epidermal growth factor, thus inducing receptor dimerization and tyrosine autophosphorylation leading to cell proliferation. Mutations in this gene are associated with lung cancer. EGFR is a component of the cytokine storm which contributes to a severe form of Coronavirus Disease 2019 (COVID-19) resulting from infection with severe acute respiratory syndrome coronavirus-2 (SARS-CoV-2).

## **Recommended Dilutions**

**WB** 1:500 - 1:2000

IP 0.5μg-4μg antibody for 200μg-400μg extracts of

whole cells

**ELISA** Recommended starting

concentration is 1 µg/mL.

Please optimize the
concentration based on
your specific assay
requirements.

# Immunogen Information

**Gene ID**Swiss Prot
1956
P00533

#### **Immunogen**

Synthetic peptide. This information is considered to be commercially sensitive.

# **Synonyms**

ERBB; ERRP; HER1; mENA; ERBB1; PIG61; NISBD2; Phospho-EGFR-T669

## **Contact**

<b>a</b>		400-999-6126
$\bowtie$		cn.market@abclonal.com.cn
$\odot$	Ī	www.abclonal.com.cn

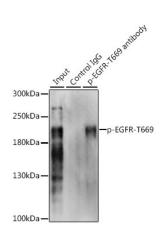
## **Product Information**

SourceIsotypePurificationRabbitIgGAffinity purification

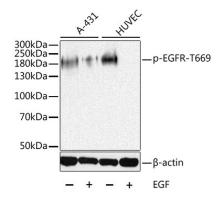
#### Storage

Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.02% sodium azide,50% glycerol,pH7.3.



Immunoprecipitation analysis of 200  $\mu$ g extracts of A-431 cells, using 3  $\mu$ g Phospho-EGFR-T669 pAb (AP0025). Western blot was performed from the immunoprecipitate using Phospho-EGFR-T669 pAb (AP0025) at a dilution of 1:1000. A-431 cells were treated by EGF (100 ng/mL) at 37°C for 30 minutes after serum-starvation overnight.



Western blot analysis of lysates from A-431 and HUVEC cells, using Phospho-EGFR-T669 Rabbit pAb (AP0025) at 1:1000 dilution. A431 cells were treated by EGF (100ng/mL) for 30 minutes after serum-starvation overnight. HUVEC cells were treated by EGF.

Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% BSA.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.