

## **Anti-EGFR Antibody (A81016)**

## Specifications:

Name: Anti-EGFR Antibody

Description: Rabbit polyclonal antibody to EGFR.

Applications: WB, ICC/IF, IP

Recommended Dilutions: WB: 1:500-1:1,000, IP: 1:500-1:1,000, ICC/IF: 1:50-1:200

Reactivity: Human, Mouse, Rat

Immunogen: A synthetic peptide corresponding to a sequence within amino acids 1100-1200 of human

EGFR (NP\_005219.2).

Sequence: RPAGSVQNPVYHNQPLNPAPSRDPHYQDPHSTAVGNPEYLNTVQPTCVNSTFDSPAHW

AQKGSHQISLDNPDYQQDFFPKEAKPNGIFKGSTAENAEYLRV

Host: Rabbit

Clonality: Polyclonal

Isotype: IgG

Conjugate: Unconjugated

Purification: Affinity purification.

Molecular Weight: 175 kDa

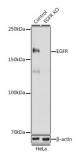
Product Form: Liquid

Formulation: Supplied in Phosphate Buffered Saline, pH 7.3, with 50% Glycerol and 0.01% Thiomersal.

Storage: Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Disclaimer: This product is for research use only. It is not intended for diagnostic or therapeutic use.

#### Images:

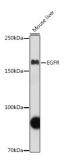


Western blot analysis of extracts from normal (control) and EGFR knockout (KO) HeLa cells, using Anti-EGFR Antibody (A81016) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 90s.

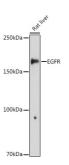


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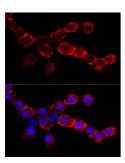
### Images continued:



Western blot analysis of extracts of Mouse liver, using Anti-EGFR Antibody (A81016) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Basic Kit. Exposure time: 90s.



Western blot analysis of extracts of Rat liver, using Anti-EGFR Antibody (A81016) at 1:1,000 dilution. The secondary antibody was Goat Anti-Rabbit IgG H&L Antibody (HRP) at 1:10,000 dilution. Lysates/proteins were present at 25µg per lane. The blocking buffer used was 3% non-fat dry milk in TBST. Detection was with a ECL Enhanced Kit (RM00021). Exposure time: 180s.



Immunofluorescence analysis of A-431 cells using Anti-EGFR Antibody (A81016) at a dilution of 1:50 (40x lens). DAPI was used to stain the cell nuclei (blue).