## **EGFR Recombinant Antibody**



## **RACO0280**

## **Product Information**

Size: **Protein Background:** 

50ul Receptor tyrosine kinase binding ligands of the EGF family and activating several signaling cascades to convert extracellular cues into appropriate cellular responses.

Reactivity: Known ligands include EGF, TGFA/TGF-alpha, amphiregulin, epigen/EPGN,

BTC/betacellulin, epiregulin/EREG and HBEGF/heparin-binding EGF. Ligand binding triggers receptor homo- and/or heterodimerization and autophosphorylation on key Source: cytoplasmic residues. The phosphorylated receptor recruits adapter proteins like GRB2

which in turn activates complex downstream signaling cascades. Activates at least 4 major downstream signaling cascades including the RAS-RAF-MEK-ERK, PI3 kinase-

AKT, PLCgamma-PKC and STATs modules.

Gene ID:

**EGFR** 

Uniprot ELISA, WB, IHC, IF, FC

P00533 **Recommended dilutions:** 

1:200, FC:1:20-1:200 Epidermal growth factor receptor (EC 2.7.10.1) (Proto-oncogene c-ErbB-1) (Receptor

Immunogen:

Synonyms:

A synthesized peptide derived from human EGFR (ErbB 1).

tyrosine-protein kinase erbB-1), EGFR, ERBB ERBB1 HER1

Storage:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.

Human

Homo sapiens (Human)

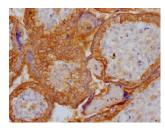
Isotype:

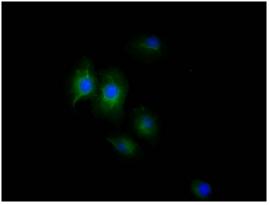
Rabbit IgG

**Applications:** 

WB:1:500-1:5000, IHC:1:50-1:200, IF:1:20-

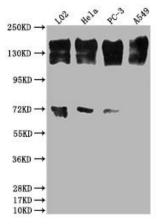
## **Product Images**





IHC image of RACO0280 diluted at 1:100 and staining in paraffinembedded human placenta tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a Goat antirabbit IgG polymer labeled by HRP and visualized using 0.05% DAB.

Immunofluorescence staining of A549 Cells with RACO0280 at 1:50, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeated by 0.2% TritonX-100, and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. Nuclear DNA was labeled in blue with DAPI. The secondary antibody was FITC-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L).



Western Blot

Positive WB detected in (L02 whole cell lysate) Hela whole cell lysate) PC-3 whole cell lysate) A549 whole cell lysate) All lanes: EGFR antibody at 1:2000

Secondary

Goat polyclonal to rabbit IgG at 1:50000 dilution

Predicted band size: 135, 45, 78, 70 kDa

Observed band size: 165 kDa