

PRODUCT DATA SHEET

Bioworld Technology, Inc.

EGFR (Phospho-Y1092) polyclonal antibody

Catalog: BS94061 Host: Rabbit Reactivity: Human, Mouse

BackGround:

Epidermal growth factor mediates its effects on cell growth through its inter-action with a cell surface glycoprotein designated the EGF receptor. Binding of EGF or TGF alpha to the EGF receptor activates tyrosine-specific protein kinase activity intrinsic to the EGF receptor. The carboxy terminal tyrosine residues on EGFR, Tyr 1068 and Tyr 1173, are the major sites of autophosphorylation, which occurs as a result of EGF binding. Once activated, EGFR mediates the binding of the phosphotyrosine binding (PTB) domain of GRB2 through direct interactions with Tyr 1068 and Tyr 1086 and through indirect interactions with Tyr 1173 in the Ras signaling pathway. Tyr 1173 of EGFR also functions as a kinase substrate. Phosphorylation of Tyr 992, Tyr 1068 and Tyr 1086 is required for conformational change in the C-terminal tail of the EGF receptor.

Product:

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

170 kDa

Swiss-Prot:

P00533(Human) Q01279(Mouse)

Purification&Purity:

ProA affinity purified

Applications:

WB:1:1,000-1:2,000 ICC:1:50-1:200 IHC:1:50-1:200

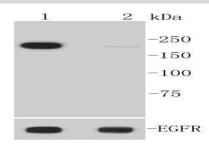
Storage&Stability:

Store at +4 $^{\circ}$ C after thawing. Aliquot store at -20 $^{\circ}$ C or -80 $^{\circ}$ C. Avoid repeated freeze / thaw cycles.

Specificity:

EGFR (Phospho-Y1092) polyclonal antibody detects endogenous levels of EGFR protein only when phosphorylated at Y1092.

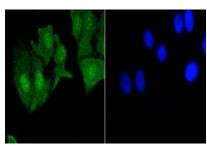
DATA:



Western blot analysis of Phospho-EGFR(Y1092) on different lysates using anti-Phospho-EGFR(Y1092) antibody at 1/1,000 dilution. Positive control:

Lane 1: A431 treated with EGF

Lane 2: Untreated A431



ICC staining Phospho-EGFR(Y1092) in BT-20 cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

Note

For research use only, not for use in diagnostic procedure.

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