

4407

Phospho-EGF Receptor (Tyr1173) (53A5) Rabbit mAb



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Applications: W, W-F, IP, IHC-P	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 175	Source/Isotype: Rabbit IgG	UniProt ID: #P00533	Entrez-Gene Id: 1956
Product Usage Information		Application Western Blotting Fluorescent Western Immunoprecipitation Immunohistochemistr	y (Paraffin)	Dilution 1:1000 1:1000 1:50 1:125 - 1:500		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody. For a carrier free (BSA and azide free) version of this product see product #47737.				
Specificity/Sensitivity		Phospho-EGF Receptor (Tyr1173) (53A5) Rabbit mAb detects endogenous EGF receptors only when phosphorylated at Tyr1173. This antibody may cross-react with other activated receptor tyrosine kinases.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr1173 of human EGF receptor.				
Background		The epidermal growth factor (EGF) receptor is a transmembrane tyrosine kinase that belongs to the HER/ErbB protein family. Ligand binding results in receptor dimerization, autophosphorylation, activation of downstream signaling, internalization, and lysosomal degradation (1,2). Phosphorylation of EGF receptor (EGFR) at Tyr845 in the kinase domain is implicated in stabilizing the activation loop, maintaining the active state enzyme, and providing a binding surface for substrate proteins (3,4). c-Src is involved in phosphorylation of EGFR at Tyr845 (5). The SH2 domain of PLCγ binds at phospho-Tyr992, resulting in activation of PLCγ-mediated downstream signaling (6). Phosphorylation of EGFR at Tyr1045 creates a major docking site for the adaptor protein c-Cbl, leading to receptor ubiquitination and degradation following EGFR activation (7,8). The GRB2 adaptor protein binds activated EGFR at phospho-Tyr1068 (9). A pair of phosphorylated EGFR residues (Tyr1148 and Tyr1173) provide a docking site for the Shc scaffold protein, with both sites involved in MAP kinase signaling activation (2). Phosphorylation of EGFR at specific serine and threonine residues attenuates EGFR kinase activity. EGFR carboxy-terminal residues Ser1046 and Ser1047 are phosphorylated by CaM kinase II; mutation of either of these serines results in upregulated EGFR tyrosine autophosphorylation (10).				
Background References		 Hackel, P.O. et al. (1999) Curr Opin Cell Biol 11, 184-9. Zwick, E. et al. (1999) Trends Pharmacol Sci 20, 408-12. Cooper, J.A. and Howell, B. (1993) Cell 73, 1051-4. Hubbard, S.R. et al. (1994) Nature 372, 746-54. Biscardi, J.S. et al. (1999) J Biol Chem 274, 8335-43. Emlet, D.R. et al. (1997) J Biol Chem 272, 4079-86. Levkowitz, G. et al. (1999) Mol Cell 4, 1029-40. Ettenberg, S.A. et al. (1999) Oncogene 18, 1855-66. Rojas, M. et al. (1996) J Biol Chem 271, 27456-61. Feinmesser, R.L. et al. (1999) J Biol Chem 274, 16168-73. 				

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting W-F: Fluorescent Western IP: Immunoprecipitation IHC-P: Immunohistochemistry

(Paraffin)

Cross-Reactivity Key H: Human M: Mouse R: Rat

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