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Phospho-EGF Receptor (Tyr1068) Antibody



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Applications: W, IHC-P	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 175	Source/Isotype: Rabbit	UniProt ID: #P00533	Entrez-Gene Id: 1956		
Product Usage Information		Application Western Blotting Immunohistochemistry (Paraffin)			Dilution 1:1000 1:100 - 1:400			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at - 20°C. Do not aliquot the antibody.				/cerol. Store at –		
Specificity/Sensitivity		Phospho-EGF Receptor (Tyr1068) Antibody detects endogenous levels of EGF receptor only when phosphorylated at tyrosine 1068. The antibody may cross-react with other activated EGF receptor family members (e.g. ErbB2), and cross-reacts slightly with activated PDGF receptor .						
Source / Purific	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Tyr1068 of human EGF receptor. Antibodies are purified by protein A and peptide affinity chromatography.						
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 PLCy binds at phospho-Tyr992, resulting in activation of PLCy-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 Re	eferences	 Hackel, P.O. et al. (1999) <i>Curr Opin Cell Biol</i> 11, 184-9. Zwick, E. et al. (1999) <i>Trends Pharmacol Sci</i> 20, 408-12. Cooper, J.A. and Howell, B. (1993) <i>Cell</i> 73, 1051-4. Hubbard, S.R. et al. (1994) <i>Nature</i> 372, 746-54. Biscardi, J.S. et al. (1999) <i>J Biol Chem</i> 274, 8335-43. Emlet, D.R. et al. (1997) <i>J Biol Chem</i> 272, 4079-86. Levkowitz, G. et al. (1999) <i>Mol Cell</i> 4, 1029-40. Ettenberg, S.A. et al. (1999) <i>Oncogene</i> 18, 1855-66. Rojas, M. et al. (1996) <i>J Biol Chem</i> 271, 27456-61. Feinmesser, R.L. et al. (1999) <i>J Biol Chem</i> 274, 16168-73. 						
Species Reactiv	vity	Species reactivity is do	etermined by testing	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot B	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.				1 5% w/v BSA, 1X		
Applications K	ey	W: Western Blotting IHC-P: Immunohistochemistry (Paraffin)						
Cross-Reactivit	ту Кеу	H: Human M: Mouse R: Rat						
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