## EGF Receptor (D1D4J) XP<sup>®</sup> Rabbit mAb (PE Conjugate) S000 S000 S000 S000 S000 S000 S000



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Applications: FC-FP, FC-L	<b>Reactivity:</b> H M	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P00533	Entrez-Gene Id: 1956		
Product Usage Information		Application Flow Cytometry (Fixed/Permeabilized) Flow Cytometry (Live)			<b>Dilution</b> 1:50 1:50		
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4°C. Do not aliquot the antibody. Protect from light. Do not freeze.					
Specificity/Sensi	tivity	EGF Receptor (D1D4J) XP <sup>®</sup> Rabbit mAb (PE Conjugate) recognizes endogenous levels of total EGFR protein.					
Source / Purifica	tion	Monoclonal antibody is produced by immunizing animals with mammalian cells expressing full length EGF receptor protein.					
Description		This Cell Signaling Technology antibody is conjugated to phycoerythrin (PE) and tested in-house for direct flow cytometry analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated EGFR (D1D4J) XP <sup>®</sup> Rabbit mAb #54359.					
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 Refe	erences	<ol> <li>Hackel, P.O. et al. (1999) <i>Curr Opin Cell Biol</i> 11, 184-9.</li> <li>Zwick, E. et al. (1999) <i>Trends Pharmacol Sci</i> 20, 408-12.</li> <li>Cooper, J.A. and Howell, B. (1993) <i>Cell</i> 73, 1051-4.</li> <li>Hubbard, S.R. et al. (1994) <i>Nature</i> 372, 746-54.</li> <li>Biscardi, J.S. et al. (1999) <i>J Biol Chem</i> 274, 8335-43.</li> <li>Emlet, D.R. et al. (1997) <i>J Biol Chem</i> 272, 4079-86.</li> <li>Levkowitz, G. et al. (1999) <i>Mol Cell</i> 4, 1029-40.</li> <li>Ettenberg, S.A. et al. (1999) <i>Oncogene</i> 18, 1855-66.</li> <li>Rojas, M. et al. (1996) <i>J Biol Chem</i> 271, 27456-61.</li> <li>Feinmesser, R.L. et al. (1999) <i>J Biol Chem</i> 274, 16168-73.</li> </ol>					
Species Reactivit	у	Species reactivity is determined by testing in at least one approved application (e.g., western blot).					
Applications Key		FC-FP: Flow Cytometry (Fixed/Permeabilized) FC-L: Flow Cytometry (Live)					
Cross-Reactivity	Key	H: Human M: Mouse					
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