

Product Data Sheet

Anti-EGFR (Phospho-Y1173) Antibody

Catalog #	Source	e Reactivity	Applications	
CQA8170	Rabbit	: H	WB, IH, IF/IC, IP	
Description		Rabbit monoclonal antibo	ody to EGFR (Phospho-Y1173)	
Immunogen		KLH-conjugated synthetic	phosphopeptide corresponding to residues surrounding	
		Y1173 of human EGFR pro	otein. The exact sequence is proprietary.	
Purification		The antibody was purified	d by immunogen affinity chromatography.	
Specificity		Recognizes endogenous levels of EGFR protein only when phosphorylated at Y1173.		
Clonality	nality Monoclonal			
Conjugation				
Form		Liquid in 50mM Tris-Glycine (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide		
		and 0.05% BSA.		
Dilution		WB (1/500 - 1/1000), IH (1,	/50 - 1/100), IF/IC (1/50 - 1/100), IP (1/10 - 1/50)	
Gene Symbol		EGFR		
Alternative Names		ERBB; ERBB1; HER1; Epidermal growth factor receptor; Proto-oncogene c-ErbB-1;		
		Receptor tyrosine-proteir	n kinase erbB-1	
Entrez Gene		1956 (Human)		
SwissProt		P00533 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid		
		freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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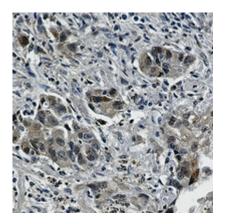
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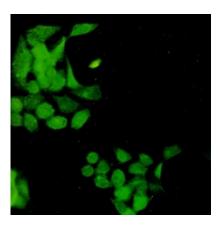
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For research purposes only, not for human use

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Western blot analysis of EGFR (Phospho-Y1173) expression in Hela (A), A549 (B) whole cell lysates. (Predicted band size: 134 kD; Observed band size: 175 kD)





Immunohistochemical analysis of EGFR (Phospho-Y1173) staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Immunofluorescent analysis of EGFR (Phospho-Y1173) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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