

## **Product Data Sheet**

## Anti-HER4 (Phospho-Y1284) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3924	Rabbit	H <i>,</i> M, R	WB, IH, IF/IC		
DescriptionRabbit polyclonal antibody to HER4 (Phospho-Y1284)					
Immunogen	K	LH-conjugated synthetic ph	osphopeptide corresponding to residues surrounding		
	Y	1284 of human HER4 prote	in. The exact sequence is proprietary.		
Purification	T	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous leve	ls of HER4 protein only when phosphorylated at Y1284.		
Clonality	P	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	aı	nd 0.01% sodium azide.			
Dilution	W	VB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	E	RBB4			
Alternative Na	a <mark>mes</mark> H	ER4; Receptor tyrosine-pro	tein kinase erbB-4; Proto-oncogene-like protein		
	C-	-ErbB-4; Tyrosine kinase-ty	pe cell surface receptor HER4; p180erbB4		
Entrez Gene	20	066 (Human); 13869 (Mou	se); 59323 (Rat)		
SwissProt	Q	(15303 (Human); Q61527 (	Mouse); Q62956 (Rat)		
Storage/Stabi	lity Sl	hipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fr	eeze/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

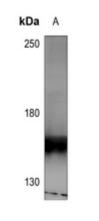
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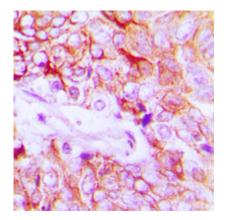
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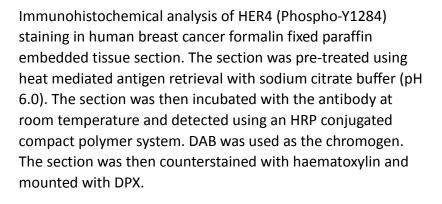
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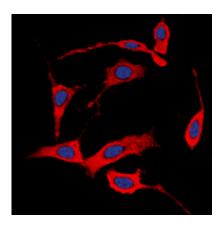
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Western blot analysis of HER4 (Phospho-Y1284) expression in rat brain (A) whole cell lysates. (Predicted band size: 146 kD; Observed band size: 150 kD)







Immunofluorescent analysis of HER4 (Phospho-Y1284) staining in MCF7 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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