

Product Data Sheet

Anti-HER2 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA3215	Rabbit	H, M, R, D	WB, IH		
Description		Rabbit polyclonal antibody to	HER2		
Immunogen		KLH-conjugated synthetic per	otide encompassing a sequence within the C-term		
		region of human HER2. The e	xact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous level	s of HER2 protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
		and 0.01% sodium azide.			
Dilution		WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol		ERBB2			
Alternative Na	ames	HER2; MLN19; NEU; NGL; Red	eptor tyrosine-protein kinase erbB-2; Metastatic lymph		
		node gene 19 protein; MLN 1	9; Proto-oncogene Neu; Proto-oncogene c-ErbB-2;		
		Tyrosine kinase-type cell surf	ace receptor HER2; p185erbB2; CD340		
Entrez Gene		2064 (Human); 13866 (Mous	e)		
SwissProt		P04626 (Human); P70424 (M	ouse); P06494 (Rat)		
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

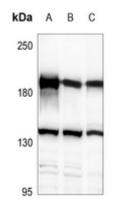
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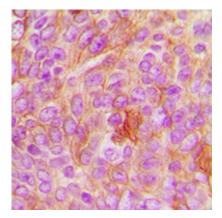


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Western blot analysis of HER2 expression in MCF7 (A), HCT116 (B), SGC7901 (C) whole cell lysates. (Predicted band size: 137 kD; Observed band size: 185; 137 kD)



Immunohistochemical analysis of HER2 staining in human breast 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.

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