

## **Product Data Sheet**

## **Anti-HER2** Antibody

Catalog #	Source	Reactivity	Applications		
CPA9813	Mouse	Н	WB, IH		
Description		Mouse monoclonal antibody to HER2			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within human HER2.			
	-	The exact sequence is prop	rietary.		
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of HER2 protein.			
Clonality		Monoclonal			
Conjugation					
Form		Mouse IgG1. Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium			
	i	azide.			
Dilution		WB (1/500 - 1/1000), IH (1/100 - 1/300)			
Gene Symbol	ĺ	ERBB2			
Alternative Names		HER2; MLN19; NEU; NGL; Receptor tyrosine-protein kinase erbB-2; Metastatic lymph			
	1	node gene 19 protein; MLN	19; Proto-oncogene Neu; Proto-oncogene c-ErbB-2;		
		0	rface receptor HER2; p185erbB2; CD340		
Entrez Gene		2064 (Human)			
SwissProt					
Storage/Stabi			ry 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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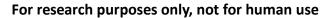
250

180 130

> 95 72

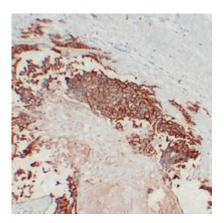
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Western blot analysis of HER2 expression in MDAMB231 (A), Hela (B) whole cell lysates. (Predicted band size: 137 kD; Observed band size: 180 kD)



Immunohistochemical analysis of HER2 staining in human breast carcinoma 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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