

## **Product Data Sheet**

## **Anti-HER2** Antibody

Catalog #	Source	e Reactivity	Applications		
CPA7002	Rabbit	-	WB, IH, IF/IC		
	Rabbit				
Description		Rabbit polyclonal antibody to HER2			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
		region of human HER2. The exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of HER2 protein.			
Clonality Polycl		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 (1/100 - 1/200), IF/IC (1/100 - 1/500)			
Gene Symbol		ERBB2			
Alternative Names		HER2; MLN19; NEU; NGL; Receptor tyrosine-protein kinase erbB-2; Metastatic lymph			
		node gene 19 protein; MI	N 19; Proto-oncogene Neu; Proto-oncogene c-ErbB-2;		
		Tyrosine kinase-type cell	surface receptor HER2; p185erbB2; CD340		
Entrez Gene		2064 (Human)			
SwissProt		P04626 (Human)			
Storage/Stabi	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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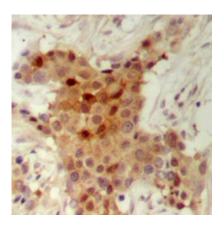
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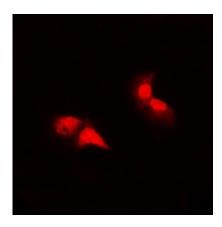
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Western blot analysis of HER2 expression in MDA (A), SKOV3 (B) whole cell lysates. (Predicted band size: 137 kD; Observed band size: 185 kD)



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.



Immunofluorescent analysis of HER2 staining in MCF 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.

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