

Product Data Sheet

Anti-HER2 Antibody

Catalog # Source Reactivity Applications

CPA7162 Rabbit H, M, R, D WB, IH

Description Rabbit polyclonal antibody to HER2

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

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 Polyclonal

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)

Gene Symbol ERBB2

Alternative Names HER2; MLN19; NEU; NGL; Receptor tyrosine-protein kinase erbB-2; Metastatic lymph

node gene 19 protein; MLN 19; Proto-oncogene Neu; Proto-oncogene c-ErbB-2;

Tyrosine kinase-type cell surface receptor HER2; p185erbB2; CD340

Entrez Gene 2064 (Human); 13866 (Mouse)

SwissProt P04626 (Human); P70424 (Mouse); P06494 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

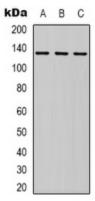
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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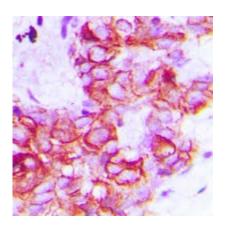




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Western blot analysis of HER2 expression in PC3 (A), MCF7 (B), NIH3T3 (C) whole cell lysates.



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 conjugacompact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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