

Product Data Sheet

Anti-HER3 Antibody

Catalog # Source Reactivity Applications

CPA6088 Rabbit H, M, R, B, Mk, Z WB, IH

Description Rabbit polyclonal antibody to HER3

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the C-term

region of human HER3. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of HER3 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 - 1/200)

Gene Symbol ERBB3

Alternative Names HER3; Receptor tyrosine-protein kinase erbB-3; Proto-oncogene-like protein

c-ErbB-3; Tyrosine kinase-type cell surface receptor HER3

Entrez Gene 2065 (Human); 13867 (Mouse); 29496 (Rat)

SwissProt P21860 (Human); Q61526 (Mouse); Q62799 (Rat)

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

freeze/thaw cycles.

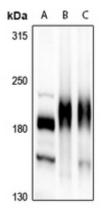
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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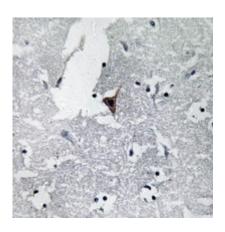
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Western blot analysis of HER3 expression in MCF7 (A), C6 (B), A375 (C) whole cell lysates. (Predicted band size: 148 kD; Observed band size: 185 kD)



Immunohistochemical analysis of HER3 staining in human brain 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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