

HER2 (Phospho-Thr686) Ab

Cat.#: AF8048 Mol.Wt.: 185kDa Concn.: 1mg/ml Size: 50ul.100ul.200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:1000-3000. IHC 1:50-1:200. IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: HER2 (Phospho-Thr686) Ab detects endogenous levels of

HER2 only when phosphorylated at Thr686

A synthesized peptide derived from human HER2 (Phospho-Immunogen:

Thr686)

Uniprot: P04626

Subcellular Location: Cytoplasm. Nucleus and Cell membrane. Cytoplasm,

> perinuclear region. Nucleus. Translocation to the nucleus requires endocytosis, probably endosomal sorting and is

mediated by importin beta-1/KPNB1.

Tissue Specificity: Expressed in a variety of tumor tissues including primary

breast tumors and tumors from small bowel, esophagus,

kidney and mouth.

Similarity: Belongs to the protein kinase superfamily. Tyr protein kinase

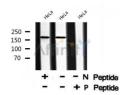
family. EGF receptor subfamily.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt



Western blot analysis HER2 (Phospho-Thr686) using HeLa whole cell lysates



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AF8048 at 1/200 staining Human breast tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF8048 at 1/200 staining Mouse kidney tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF8048 at 1/200 staining Rat kidney tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22 $^{\circ}$ C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF8048 staining MDA-MB-231 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

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