

## Phospho-HER4 (Tyr1056) Ab

Cat.#: AF8370 Concn.: 1mg/ml Mol.Wt.: 180kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:1000-3000, 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: HER4 (Phospho-Tyr1056) Ab detects endogenous levels of

HER4 only when phosphorylated at Tyr1056.

Immunogen: A synthesized peptide derived from human HER4 (Phospho-

Tyr1056).

Uniprot: Q15303

Subcellular Location: Membrane and Nucleus. Following proteolytical processing

E4ICD (E4ICD1 or E4ICD2 generated from the respective isoforms) is translocated to the nucleus. Significantly more E4ICD2 than E4ICD1 is found in the nucleus. E4ICD2

colocalizes with YAP1 in the nucleus.

Tissue Specificity: Expressed at highest levels in brain, heart, kidney, in

addition to skeletal muscle, parathyroid, cerebellum, pituitary, spleen, testis and breast. Lower levels in thymus, lung, salivary gland, and pancreas. Isoform JM-A CYT-1 and isoform JM-B CYT-1 are expressed in cerebellum, but only

the isoform JM-B is expressed in the heart.

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

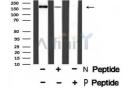
family. EGF receptor subfamily.

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 of HER4 (Phospho-Tyr1056) using Insulin treated HeLa whole cell lysates.

-/+ means absence or presence of N peptide□non-phospho peptide) and P peptide(phospho peptide).



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AF8370 staining HuvEc 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.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween020 at  $4^{\circ}$ C with gentle shaking, overnight.

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