

# Phospho-HER4(Y1162) Antibody

Affinity Purified Rabbit Polyclonal Antibody (Pab)

Catalog # AP3122a

## Product Information

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| <b>Application</b>       | IF, WB, IHC-P, E                                |
| <b>Primary Accession</b> | <a href="#">Q15303</a>                          |
| <b>Other Accession</b>   | <a href="#">Q62956</a> , <a href="#">Q61527</a> |
| <b>Reactivity</b>        | Human, Rat, Mouse                               |
| <b>Predicted</b>         | Mouse, Rat                                      |
| <b>Host</b>              | Rabbit  |
| <b>Clonality</b>         | Polyclonal                                      |
| <b>Isotype</b>           | Rabbit IgG                                      |
| <b>Calculated MW</b>     | 146808  |

## Additional Information

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|---------------------------|--|
| <b>Gene ID</b>            | 2066   |
| <b>Other Names</b>        | Receptor tyrosine-protein kinase erbB-4, Proto-oncogene-like protein c-ErbB-4, Tyrosine kinase-type cell surface receptor HER4, p180erbB4, ERBB4 intracellular domain, 4ICD, E4ICD, s80HER4, ERBB4, HER4 |
| <b>Target/Specificity</b> | This HER4 Antibody is generated from rabbits immunized with a KLH conjugated synthetic phosphopeptide corresponding to amino acid residues surrounding Y1162 of human HER4.                              |
| <b>Dilution</b>           | IF~~1:10~50 WB~~1:1000 IHC-P~~1:100~500 E~~Use at an assay dependent concentration.  |
| <b>Format</b>             | Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.                             |
| <b>Storage</b>            | Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.  |
| <b>Precautions</b>        | Phospho-HER4(Y1162) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.   |

## Protein Information

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|-----------------|-------|
| <b>Name</b>     | ERBB4 |
| <b>Synonyms</b> | HER4  |

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|--------------------------|---|
| <b>Function</b>          | <p>Tyrosine-protein kinase that plays an essential role as cell surface receptor for neuregulins and EGF family members and regulates development of the heart, the central nervous system and the mammary gland, gene transcription, cell proliferation, differentiation, migration and apoptosis. Required for normal cardiac muscle differentiation during embryonic development, and for postnatal cardiomyocyte proliferation. Required for normal development of the embryonic central nervous system, especially for normal neural crest cell migration and normal axon guidance. Required for mammary gland differentiation, induction of milk proteins and lactation. Acts as cell-surface receptor for the neuregulins NRG1, NRG2, NRG3 and NRG4 and the EGF family members BTC, EREG and HBEGF. Ligand binding triggers receptor dimerization and autophosphorylation at specific tyrosine residues that then serve as binding sites for scaffold proteins and effectors. Ligand specificity and signaling is modulated by alternative splicing, proteolytic processing, and by the formation of heterodimers with other ERBB family members, thereby creating multiple combinations of intracellular phosphotyrosines that trigger ligand- and context- specific cellular responses. Mediates phosphorylation of SHC1 and activation of the MAP kinases MAPK1/ERK2 and MAPK3/ERK1. Isoform JM-A CYT-1 and isoform JM-B CYT-1 phosphorylate PIK3R1, leading to the activation of phosphatidylinositol 3-kinase and AKT1 and protect cells against apoptosis. Isoform JM-A CYT-1 and isoform JM-B CYT-1 mediate reorganization of the actin cytoskeleton and promote cell migration in response to NRG1. Isoform JM-A CYT-2 and isoform JM-B CYT-2 lack the phosphotyrosine that mediates interaction with PIK3R1, and hence do not phosphorylate PIK3R1, do not protect cells against apoptosis, and do not promote reorganization of the actin cytoskeleton and cell migration. Proteolytic processing of isoform JM-A CYT-1 and isoform JM-A CYT-2 gives rise to the corresponding soluble intracellular domains (4ICD) that translocate to the nucleus, promote nuclear import of STAT5A, activation of STAT5A, mammary epithelium differentiation, cell proliferation and activation of gene expression. The ERBB4 soluble intracellular domains (4ICD) colocalize with STAT5A at the CSN2 promoter to regulate transcription of milk proteins during lactation. The ERBB4 soluble intracellular domains can also translocate to mitochondria and promote apoptosis.</p> |
| <b>Cellular Location</b> | Cell membrane; Single-pass type I membrane protein. Note=In response to NRG1 treatment, the activated receptor is internalized  |
| <b>Tissue Location</b>   | 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.   |

## Background

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HER4/ERBB4 is a member of the type I receptor tyrosine kinase subfamily that includes EGFR, ERBB2 and ERBB3. It is a receptor for NDF/heregulin.

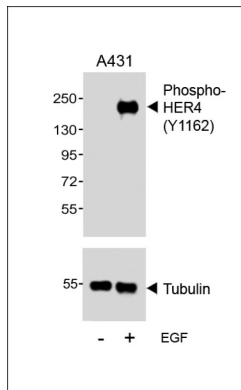
## References

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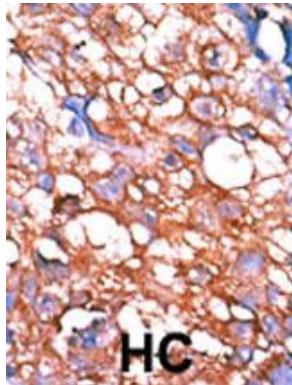
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 Hughes, D.P., et al., Cancer Res. 64(6):2047-2053 (2004).  
 Cheng, Q.C., et al., J. Biol. Chem. 278(40):38421-38427 (2003).  
 Chaudhury, A.R., et al., J. Neuropathol. Exp. Neurol. 62(1):42-54 (2003).  
 Komuro, A., et al., J. Biol. Chem. 278(35):33334-33341 (2003).

## Images

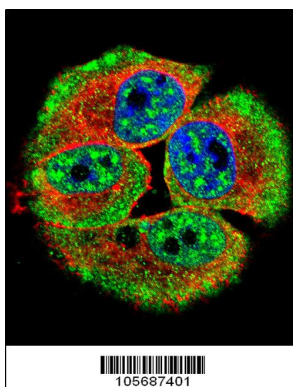
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Western blot analysis of lysates from A431 cell line, untreated or treated with EGF, 100ng/ml, using (Cat. #AP3122a)(upper) or Tubulin (lower).



Formalin-fixed and paraffin-embedded human cancer tissue reacted with the primary antibody, which was peroxidase-conjugated to the secondary antibody, followed by AEC staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated. BC = breast carcinoma; HC = hepatocarcinoma.



Confocal immunofluorescent analysis of Phospho-HER4-Y1162 Antibody(Cat#AP3122a) with MCF-7 cell followed by Alexa Fluor 488-conjugated goat anti-rabbit IgG (green). Actin filaments have been labeled with Alexa Fluor 555 phalloidin (red).DAPI was used to stain the cell nuclear (blue).

## Citations

- [Analysis of the tyrosine kinome in melanoma reveals recurrent mutations in ERBB4.](#)
- [Choice of fixative is crucial to successful immunohistochemical detection of phosphoproteins in paraffin-embedded tumor tissues.](#)

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.