Anti-Phospho-HER2 / ErbB2 (Y1221 + Y1222) Antibody [JE44-12]

HA721433



Product Type: Recombinant Rabbit monoclonal IgG, primary antibodies

Species reactivity: Human
Applications: WB

Molecular Wt: Predicted band size: 138 kDa

Clone number: JE44-12

Description: HER-2 (also called HER-2/neu, c-erbB2, ERBB2 or neu) is a transmembrane receptor tyrosine kinase. HER-2

is a proto-oncogene, i.e. its activation causes malignant transformation and increases the malignant potential (cell proliferation, invasiveness etc.) of the cells. Amplification of HER-2 gene invariably leads to over-expression of its protein product. The magnitude of over- expression is usually 10-100 folds or even more. Over-expressed HER-2 protein disturbs the HER-receptor family signalling networks, i.e. signalling mediated via EGFR receptor, HER-3 and HER-4. In tumours, HER-2 is over-expressed in 15-25% of primary breast cancers. Metastases usually have the same amplification status as the primary tumours. HER-2 amplification and over-expression are typical features of hormone receptor negative, rapidly growing histologic grade 2-3 tumours. Of the histologic types, Paget's disease is almost invariably HER-2 positive, whereas only a small minority of lobular and tubular carcinomas shows HER-2 amplification. HER-2 amplification and over-expression can also be found in intestinal type gastric and gastroesophageal carcinomas, ovarian carcinomas, high grade endometrial carcinomas and some salivary duct tumours. Low-level copy number increases have been found also in rare cases of lung tumours. Because of its central importance in breast cancer therapy selection, standardization of HER-2 IHC

assays and slide interpretation are of outmost clinical and economical importance.

Immunogen: Synthetic phosphopeptide corresponding to residues surrounding tyrosines 1221/1222 of human ErbB2 protein

Positive control: SK-Br-3 whole cell lysate.

Subcellular location: Cell membrane, Nucleus, Cytoplasm.

Database links: SwissProt P04626 Human

Recommended Dilutions:

WB 1:1.000

Storage Buffer: PBS (pH7.4), 0.1% BSA, 40% Glycerol. Preservative: 0.05% SodiumAzide.

Storage Instruction: Store at +4 $^{\circ}$ C after thawing. Aliquot store at -20 $^{\circ}$ C. Avoid repeated freeze / thaw cycles.

Purity: Protein A affinity purified.

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Images

kDa 250-250-150-75-50-37-25-20-15-10-■ GAPDH - + λpp **Fig1:** Western blot analysis of Phospho-HER2 / ErbB2 (Y1221 + Y1222) on different lysates with Rabbit anti-Phospho-HER2 / ErbB2 (Y1221 + Y1222) antibody (HA721433) at 1/1,000 dilution.

Lane 1: SK-Br-3 whole cell lysate

Lane 2: SK-Br-3 treated with App for 1 hour whole cell lysate

Lysates/proteins at 20 µg/Lane.

Predicted band size: 138 kDa Observed band size: 250 kDa

Exposure time: 3 minutes;

4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (HA721433) at 1/1,000 dilution was used in 5% NFDM/TBST at room temperature for 2 hours. Goat Anti-Rabbit IgG - HRP Secondary Antibody (HA1001) at 1:100,000 dilution was used for 1 hour at room temperature.

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- Ellis CM, Dyson MJ, Stephenson TJ, Maltby EL. HER2 amplification status in breast cancer: a comparison between immunohistochemical staining and fluorescence in situ hybridisation using manual and automated quantitative image analysis scoring techniques. J Clin Pathol. 2005 Jul;58(7):710-4.
- 2. Gancberg D, Jarvinen T, di Leo A, Rouas G, Cardoso F, Paesmans M, Verhest A, Piccart MJ, Isola J, Larsimont D. Evaluation of HER-2/NEU protein expression in breast cancer by immunohistochemistry: an interlaboratory study assessing the reproducibility of HER-2/NEU testing. Breast Cancer Res Treat. 2002 Jul;74(2):113-20.