## Overview

<table>
<thead>
<tr>
<th><strong>Product name</strong></th>
<th>Anti-ErbB 2 antibody [3B5]</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Description</strong></td>
<td>Mouse monoclonal [3B5] to ErbB 2</td>
</tr>
<tr>
<td><strong>Host species</strong></td>
<td>Mouse</td>
</tr>
<tr>
<td><strong>Tested applications</strong></td>
<td>Suitable for: IHC-Fr, IHC-P, WB, IP, Flow Cyt</td>
</tr>
<tr>
<td><strong>Species reactivity</strong></td>
<td>Reacts with: Mouse, Rat, Human, Monkey</td>
</tr>
<tr>
<td><strong>Predicted to work with</strong></td>
<td>Dog</td>
</tr>
<tr>
<td><strong>Immunogen</strong></td>
<td>Synthetic peptide corresponding to Human ErbB 2 aa 1242-1255 (C terminal). Sequence: TAENPEYLGLDVPV</td>
</tr>
<tr>
<td><strong>Database link</strong></td>
<td>P04626</td>
</tr>
</tbody>
</table>

### Epitope
Within amino acids 1242 - 1255 of human c-neu (TAENPEYLGLDVPV)

### Positive control
SK-BR-3 cells or breast carcinoma tissue

### General notes
For paraffin sections, we recommend pretreating with a pressure cooker, but trypsin and heat will also work well. Including 0.05% saponin during staining of paraffin sections may help reduce background. In one study ductal carcinomas in situ displaying large-cell, comedo growth type, were stained while none of 16 ductal carcinomas in situ of small-cell, papillary, or cribriform growth type were stained by this antibody. MO 01/02/07: update consignment level to 0 following settlement of consignment stock in January 07.

## Properties

<table>
<thead>
<tr>
<th><strong>Form</strong></th>
<th>Liquid</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Storage instructions</strong></td>
<td>Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.</td>
</tr>
<tr>
<td><strong>Storage buffer</strong></td>
<td>Preservative: 0.09% Sodium azide</td>
</tr>
<tr>
<td><strong>Purity</strong></td>
<td>Protein G purified</td>
</tr>
<tr>
<td><strong>Clonality</strong></td>
<td>Monoclonal</td>
</tr>
<tr>
<td><strong>Clone number</strong></td>
<td>3B5</td>
</tr>
</tbody>
</table>
Isotype

IgG1

Applications

Our Abpromise guarantee covers the use of ab16901 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<table>
<thead>
<tr>
<th>Application</th>
<th>Abreviews</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>IHC-Fr</td>
<td></td>
<td>Use a concentration of 2.5 µg/ml.</td>
</tr>
<tr>
<td>IHC-P</td>
<td>⭐⭐⭐⭐</td>
<td>Use at an assay dependent concentration.</td>
</tr>
<tr>
<td>WB</td>
<td>⭐⭐⭐⭐</td>
<td>Use a concentration of 2.5 µg/ml. Detects a band of approximately 190 kDa (predicted molecular weight: 138 kDa).</td>
</tr>
<tr>
<td>IP</td>
<td></td>
<td>Use 1µg for 10^6 cells.</td>
</tr>
<tr>
<td>Flow Cyt</td>
<td>⭐⭐⭐⭐</td>
<td>1/50 - 1/100. ab170190 - Mouse monoclonal IgG1, is suitable for use as an isotype control with this antibody.</td>
</tr>
</tbody>
</table>

Target

Function
Protein tyrosine kinase that is part of several cell surface receptor complexes, but that apparently needs a coreceptor for ligand binding. Essential component of a neuregulin-receptor complex, although neuregulins do not interact with it alone. GP30 is a potential ligand for this receptor. Regulates outgrowth and stabilization of peripheral microtubules (MTs). Upon ERBB2 activation, the MEMO1-RHOA-DIAPH1 signaling pathway elicits the phosphorylation and thus the inhibition of GSK3B at cell membrane. This prevents the phosphorylation of APC and CLASP2, allowing its association with the cell membrane. In turn, membrane-bound APC allows the localization of MACF1 to the cell membrane, which is required for microtubule capture and stabilization. In the nucleus is involved in transcriptional regulation. Associates with the 5'-TCAAATTC-3' sequence in the PTGS2/COX-2 promoter and activates its transcription. Implicated in transcriptional activation of CDKN1A; the function involves STAT3 and SRC. Involved in the transcription of rRNA genes by RNA Pol I and enhances protein synthesis and cell growth.

Tissue specificity
Expressed in a variety of tumor tissues including primary breast tumors and tumors from small bowel, esophagus, kidney and mouth.

Involvement in disease
Hereditary diffuse gastric cancer
Glioma
Ovarian cancer
Lung cancer
Gastric cancer
Chromosomal aberrations involving ERBB2 may be a cause gastric cancer. Deletions within 17q12 region producing fusion transcripts with CDK12, leading to CDK12-ERBB2 fusion leading to truncated CDK12 protein not in-frame with ERBB2.

Sequence similarities
Belongs to the protein kinase superfamily. Tyr protein kinase family. EGF receptor subfamily. Contains 1 protein kinase domain.
Post-translational modifications


Cellular localization


Images

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast cancer tissue sections labeling ErbB 2 with ab16901 at 1/400 dilution. the tissue was fixed with formaldehyde; heat mediated antigen retrieval was performed using a acetate buffer pH6. An undiluted goat anti-mouse HRP conjugated seconday antibody was used.

Anti-ErbB 2 antibody [3B5] (ab16901) at 1/250 dilution + Human Huh7 whole cell lysate at 20 µg

Secondary

Goat anti-mouse HRP conjugate at 1/5000 dilution

Predicted band size: 138 kDa

Observed band size: 138 kDa

Additional bands at: 48 kDa (possible non-specific binding)

Exposure time: 5 minutes

Blocking: 5% milk for 1 hour at 25°C.
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human Mamma tumor Her2Neu tissue labeling ErbB 2 with ab16901.

Western blot - Anti-ErbB 2 antibody [3B5] (ab16901) at 1 µg/ml + Whole cell lysate HEK293

**Predicted band size:** 138 kDa

Overlay histogram showing LoVo cells stained with ab16901 (red line). The cells were fixed with methanol (5 min) and then permeabilized with 0.1% PBS-Tween for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab16901, 1/100 dilution) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse IgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG1 [ICIGG1] (ab91353, 2µg/1x10^6 cells) used under the same conditions. Acquisition of >5,000 events was performed. This antibody gave a positive signal in LoVo cells fixed with 4% paraformaldehyde (10 min) permeabilized with 0.1% PBS-Tween used under the same conditions.
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human breast carcinoma labeling ErbB 2 with ab16901.

Please note: All products are “FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES”

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