

## Progesterone Receptor A/B Antibody



Orders:

877-616-CELL (2355) orders@cellsignal.com

Support:

877-678-TECH (8324)

Web:

info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

## For Research Use Only. Not for Use in Diagnostic Procedures.

| Applications:<br>WB          | Reactivity:<br>H                               | Sensitivity:<br>Endogenous   | <b>MW (kDa):</b><br>90 (PR-A), 118<br>(PR-B) | <b>Source:</b><br>Rabbit | <b>UniProt ID:</b><br>#P06401 | Entrez-Gene Id:<br>5241 |  |
|------------------------------|--|--|--|--------------------------|-------------------------------|-------------------------|--|
| Product Usage<br>Information | Application                                    |  |  | Dilution                 |                               |                         |  |
|                              | We   | estern Blotting  |  |                          | 1:1000                        |                         |  |
| Storage                      | •  | Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.   |  |                          |                               |                         |  |
| Specificity / Sensitiv       |  | Progesterone Receptor A/B Antibody detects endogenous levels of total progesterone receptor A and B proteins. This antibody does not cross-react with other PR family members.   |  |                          |                               |                         |  |
| Source / Purification        | -  | Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Tyr541 of human progesterone receptor.   |  |                          |                               |                         |  |
| Background                   | A. P<br>but (<br>pho:<br>sites<br>Ser3<br>Ser2 | Human progesterone receptor (PR) is expressed as two forms: the full length PR-B and the short form PR-A. PR-A lacks the first 164 amino acid residues of PR-B (1,2). Both PR-A and PR-B are ligand activated, but differ in their relative ability to activate target gene transcription (3,4). The activity of PR is regulated by phosphorylation; at least seven serine residues are phosphorylated in its amino-terminal domain. Three sites (Ser81, Ser102, and Ser162) are unique to full length PR-B, while other sites (Ser190, Ser294, Ser345, and Ser400) are shared by both isoforms (5). Phosphorylation of PR-B at Ser190 (equivalent to Ser26 of PR-A) is catalyzed by CDK2 (6). Mutation of Ser190 results in decreased activity of PR (7), suggesting that the phosphorylation at Ser190 may be critical to its biological function. |  |                          |                               |                         |  |
| Background Referer           | 2. K<br>3. G<br>4. W<br>5. C<br>6. Zl          | <ol> <li>Evans, R.M. (1988) Science 240, 889-895.</li> <li>Kastner, P. et al. (1990) EMBO J. 112, 1603-1614.</li> <li>Giangrande, P.H. et al. (2000) Mol. Cell. Biol. 20, 3102-3115.</li> <li>Wen, D.X. et al. (1994) Mol. Cell. Biol. 14, 8356-8364.</li> <li>Clemm, D.L. et al. (2000) Mol. Endocrinol. 14, 52-65.</li> <li>Zhang, Y. et al. (1997) Mol. Endocrinol. 11, 823-832.</li> <li>Takimoto, G.S. et al. (1996) J. Biol. Chem. 271, 13308-13316.</li> </ol>  |  |                          |                               |                         |  |

**Species Reactivity** 

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

**Western Blot Buffer** 

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** 

WB: Western Blotting

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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