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## Progesterone Receptor B Antibody



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

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 118	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P06401	<b>Entrez-Gene Id:</b> 5241		
Product Usage Information		<b>Application</b> Western Blotting	<b>Dilution</b> 1:1000					
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				ycerol. Store at –		
Specificity/Sensitivity		Progesterone Receptor B Antibody detects endogenous levels of total progesterone receptor B protein. This antibody does not cross-react with other PR family members.						
Source / Purifi	cation		lyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to idues surrounding Thr73 of human progesterone receptor.					
Background		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 R	eferences	<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 Reacti	vity	Species reactivity is de	termined by testin	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	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.				1 5% w/v BSA, 1X		
Applications K	ey	W: Western Blotting						
Cross-Reactivi	ty Key	H: Human						
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