36 Store at -20C Phospho-Progesterone Receptor (Ser294) Antibody **Cell Signaling** TECHNOLOGY® 877-616-CELL (2355) Orders: 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. UniProt ID: Applications: **Reactivity:** Sensitivity: MW (kDa): Source: Entrez-Gene Id: 90 (PR-A), 118 WB Н Endogenous Rabbit #P06401 5241 (PR-B)

(PR-B)		
Product Usage Information	Application Western Blotting	Dilution 1:1000
Storage	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.	
Specificity / Sensitivity	Phospho-Progesterone Receptor (Ser294) Antibody recognizes endogenous levels of progesterone receptor B (PR-B) and progesterone receptor A (PR-A) proteins only when phosphorylated at Ser294 and Ser130, respectively. This antibody does not cross-react with other progesterone receptor family members.	
Species predicted to react based on 100% sequence homology:	Mouse, Rat, Pig, Horse	
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser294 of human progesterone receptor B (PR-B) protein. Antibodies are purified by protein A and peptide affinity chromatography.	
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 (Ser19, 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. Progesterone receptor Ser294 is as an important hormone-dependent phospho-acceptor site that serves as an extracellular signaling "sensor." Research studies indicate that p44/p42 MAP kinases are responsible for phosphorylation of progesterone receptor at Ser294 following hormone binding. This phosphorylation event promotes nuclear retention of the receptor, enhanced transcriptional activity, and ubiquitin-dependent proteasomal degradation (6,8,9).	
Background References	 Evans, R.M. (1988) Science 240, 889 Kastner, P. et al. (1990) EMBO J. 112 Giangrande, P.H. et al. (2000) Mol. C Wen, D.X. et al. (1994) Mol. Cell. Bio Clemm, D.L. et al. (2000) Mol. Endocrin Takimoto, G.S. et al. (1996) J. Biol. C Lange, C.A. et al. (2000) Proc Natl A Daniel, A.R. et al. (2007) Steroids 72 	2, 1603-1614. ell. Biol. 20, 3102-3115. l. 14, 8356-8364. rinol. 14, 52-65. ol. 11, 823-832. hem. 271, 13308-13316. cad Sci U S A 97, 1032-7.
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 nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.	

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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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