

Anti-Progesterone Receptor (Phospho-S294) Antibody

Catalog #	Source	Reactivity	Applications
CPA1882	Rabbit	H, M, R	WB, IF/IC
Description	Rabbit polyclonal antibody to Progesterone Receptor (Phospho-S294)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding S294 of human Progesterone Receptor protein. The exact sequence is proprietary.		
Purification	The antibody was purified by immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of Progesterone Receptor protein only when phosphorylated at S294.		
Clonality	Polyclonal		
Conjugation			
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.		
Dilution	WB (1/500 - 1/1000), IF/IC (1/50 - 1/200)		
Gene Symbol	PGR		
Alternative Names	NR3C3; Progesterone receptor; PR; Nuclear receptor subfamily 3 group C member 3		
Entrez Gene	5241 (Human); 25154 (Rat)		
SwissProt	P06401 (Human); Q00175 (Mouse); Q63449 (Rat)		
Storage/Stability	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, CHIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference

Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb- Rabbit, S- Sheep, Z- Zebrafish

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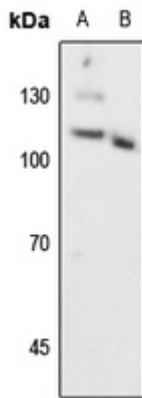
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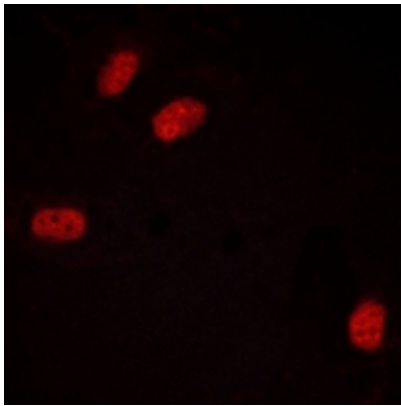
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Product Data Sheet



Western blot analysis of Progesterone Receptor (Phospho-S294) expression in HEK293T (A), HeLa (B) whole cell lysates. (Predicted band size: 98 kD; Observed band size: 118 kD)



Immunofluorescent analysis of Progesterone Receptor (Phospho-S294) staining in HeLa cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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