## Estrogen Receptor α (D6R2W) Rabbit mAb (Alexa Fluor® 647 Conjugate)



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

<b>Applications:</b> IF-IC, FC-FP	Reactivity:	<b>Sensitivity:</b> Endogenous	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P03372	Entrez-Gene Id: 2099
Product Usage Information		<b>Application</b> Immunofluorescence (Ir Flow Cytometry (Fixed/P			<b>Dilution</b> 1:50 - 1:200 1:50
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at $4^{\circ}$ C. Do not aliquot the antibody. Protect from light. Do not freeze.			
Specificity/Sensitivity		Estrogen Receptor $\alpha$ (D6R2W) Rabbit mAb (Alexa Fluor® 647 Conjugate) recognizes endogenous levels of total estrogen receptor $\alpha$ protein.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the amino terminus of human estrogen receptor $\alpha$ protein.			
Description		This Cell Signaling Technology antibody is conjugated to Alexa Fluor $^{\circledR}$ 647 fluorescent dye and tested in-house for direct flow cytometric and immunofluorescent analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated Estrogen Receptor $\alpha$ (D6R2W) Rabbit mAb #13258.			
Background		Estrogen receptor $\alpha$ (ER $\alpha$ ), a member of the steroid receptor superfamily, contains highly conserved DNA binding and ligand binding domains (1). Through its estrogen-independent and estrogen-dependent activation domains (AF-1 and AF-2, respectively), ER $\alpha$ regulates transcription by recruiting coactivator proteins and interacting with general transcriptional machinery (2). Phosphorylation at multiple sites provides an important mechanism to regulate ER $\alpha$ activity (3-5). Ser104, 106, 118, and 167 are located in the amino-terminal transcription activation function domain AF-1, and phosphorylation of these serine residues plays an important role in regulating ER $\alpha$ activity. Ser118 may be the substrate of the transcription regulatory kinase CDK7 (5). Ser167 may be phosphorylated by p90RSK and Akt (4,6). According to the research literature, phosphorylation at Ser167 may confer tamoxifen resistance in breast cancer patients (4).			
Background References		<ol> <li>Mangelsdorf, D.J. et al. (1995) Cell 83, 835-9.</li> <li>Glass, C.K. and Rosenfeld, M.G. (2000) Genes Dev 14, 121-41.</li> <li>Chen, D. et al. (1999) Mol Cell Biol 19, 1002-15.</li> <li>Campbell, R.A. et al. (2001) J Biol Chem 276, 9817-24.</li> <li>Chen, D. et al. (2000) Mol Cell 6, 127-37.</li> <li>Joel, P.B. et al. (1998) Mol Cell Biol 18, 1978-84.</li> </ol>			

**Species Reactivity** 

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

**Applications Key** 

IF-IC: Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** 

**H:** Human

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