

**Phospho-Estrogen Receptor α (Ser167)
(D5W3Z) Rabbit mAb****Orders:** 877-616-CELL (2355)
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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IF-IC	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 66	Source/Isotype: Rabbit IgG	UniProt ID: #P03372	Entrez-Gene Id: 2099
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**Product Usage
Information****Application**Western Blotting
Immunofluorescence (Immunocytochemistry)**Dilution**1:1000
1:800**Storage**Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C . Do not aliquot the antibody.**Specificity/Sensitivity**Phospho-Estrogen Receptor α (Ser167) (D5W3Z) Rabbit mAb recognizes endogenous levels of ER α protein only when phosphorylated at Ser167.**Source / Purification**Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser167 of human ER α protein.**Background**

Estrogen receptor α (ER α), 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 α regulates transcription by recruiting coactivator proteins and interacting with general transcriptional machinery (2). Phosphorylation at multiple sites provides an important mechanism to regulate ER α 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 α 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). ER α can be phosphorylated at Ser167 by various kinases such as S6K1, RSK, and Aurora A (7-9). Phosphorylation on Ser167 promotes ER α -dependent transcription and cellular proliferation, and is attributed to increased resistance to tamoxifen treatment (6, 9, 10). Various studies have shown that increased Ser167 phosphorylation correlates with poor prognosis in different cancer types (11, 12)

Background References

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3. Chen, D. et al. (1999) *Mol Cell Biol* 19, 1002-15.
4. Campbell, R.A. et al. (2001) *J Biol Chem* 276, 9817-24.
5. Chen, D. et al. (2000) *Mol Cell* 6, 127-37.
6. Joel, P.B. et al. (1998) *Mol Cell Biol* 18, 1978-84.
7. Yamnik, R.L. et al. (2009) *J Biol Chem* 284, 6361-9.
8. Yamnik, R.L. and Holz, M.K. (2010) *FEBS Lett* 584, 124-8.
9. Zheng, X.Q. et al. (2014) *Oncogene* 33, 4985-96.
10. Wang, Y. et al. (2015) *J Mol Endocrinol* 54, 351-61.
11. López-Calderero, I. et al. (2014) *Hum Pathol* 45, 2437-46.
12. Kato, E. et al. (2014) *Cancer Sci* 105, 1307-12.

Species Reactivity

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

Western Blot BufferIMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween $\text{\textcircled{R}}$ 20 at 4°C with gentle shaking, overnight.**Applications Key****W:** Western Blotting **IF-IC:** Immunofluorescence (Immunocytochemistry)**Cross-Reactivity Key****H:** Human**Trademarks and Patents**

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