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Phospho-Estrogen Receptor α (Ser167) (D5W3Z) Rabbit mAb



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

Applications Key

WB: Western Blotting IF-IC: Immunofluorescence (Immunocytochemistry)

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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