

PRODUCT DATA SHEET

Bioworld Technology,Inc.

ERα (Phospho-S118) polyclonal antibody

Catalog: BS94082 Host: Rabbit Reactivity: Human

BackGround:

Estrogen receptor alpha (ERa, ER, ESR, ESRA, Era, NR3A1, estrogen receptor 1) is a ligand-activated transcription factor composed of several domains important for hormone binding, DNA binding and activation of transcription. Alternative splicing results in several ERa mRNA transcripts, which differ primarily in their 5' untranslated regions. ERa undergoes phosphorylation in response to estradiol binding. Human ERa is predominately phosphorylated on Ser 118 and to a lesser extent on Ser 104 and Ser 106. In response to activation of the mitogen-activated protein kinase pathway, phosphorylation occurs on Ser 118 and Ser 167. These serine residues are all located within the activation function 1 region of the N-terminal domain of ERa. In contrast, activation of protein kinase A increases the phosphorylation of Ser 236, which is located in the DNA-binding domain. Src kinase-dependent Tyr 537 phosphorylation may enhance estrogen binding to ERa. Mutation of Tyr 537 of the human ERa produces receptors having a range of constitutive activity.

Product:

Rabbit IgG, 1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

Molecular Weight:

66 kDa

Swiss-Prot:

P03372(Human)

Purification&Purity:

ProA affinity purified

Applications:

WB:1:1,000 ICC:1:50-1:200 IHC:1:50-1:200

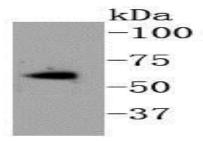
Storage&Stability:

Store at +4 $^{\circ}$ C after thawing. Aliquot store at -20 $^{\circ}$ C or -80 $^{\circ}$ C. Avoid repeated freeze / thaw cycles.

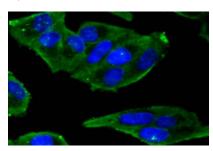
Specificity:

ER α (Phospho-S118) polyclonal antibody detects endogenous levels of ER α protein only when phosphorylated at S118.

DATA:



Western blot analysis of Estrogen Receptor alpha(phospho S118) on MCF-7 cell lysates using anti- phospho-Estrogen Receptor alpha(S118) antibody at 1/1,000 dilution.



ICC staining phospho-Estrogen Receptor alpha(S118) in Hela cells (green). The nuclear counter stain is DAPI (blue). Cells were fixed in paraformaldehyde, permeabilised with 0.25% Triton X100/PBS.

Note:

For research use only, not for use in diagnostic procedure.

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