

## **Product Data Sheet**

# Anti-Estrogen Receptor alpha (Phospho-S106) Antibody

Reactivity **Applications** Catalog # **Source** CPA3929 Rabbit H, M, R, Mk, Rb WB, IH, IF/IC

Rabbit polyclonal antibody to Estrogen Receptor alpha (Phospho-S106)

KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding **Immunogen** 

S106 of human Estrogen Receptor alpha protein. The exact sequence is proprietary.

The antibody was purified by immunogen affinity chromatography. **Purification** 

**Specificity** Recognizes endogenous levels of Estrogen Receptor alpha protein only when

phosphorylated at S106.

**Clonality** Polyclonal

Conjugation

Description

**Form** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

WB (1/500 - 1/1000), IH (1/100 - 1/200), IF/IC (1/100 - 1/500) Dilution

**Gene Symbol** ESR1

**Alternative Names** ESR; NR3A1; Estrogen receptor; ER; ER-alpha; Estradiol receptor; Nuclear receptor

subfamily 3 group A member 1

**Entrez Gene** 2099 (Human); 13982 (Mouse); 24890 (Rat)

**SwissProt** P03372 (Human); P19785 (Mouse); P06211 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

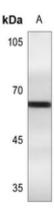
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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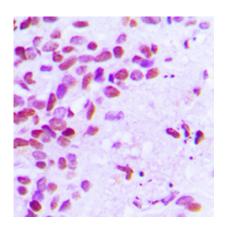
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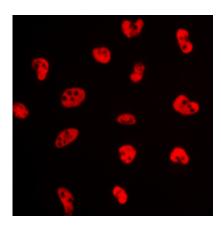
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Western blot analysis of Estrogen Receptor alpha (Phospho-S106) expression in rat lung (A) whole cell lysates. (Predicted band size: 66 kD; Observed band size: 66 kD)



Immunohistochemical analysis of Estrogen Receptor alpha (Phospho-S106) staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of Estrogen Receptor alpha (Phospho-S106) staining in MCF7 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 hidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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