

### **Product Data Sheet**

# **Anti-Glucocorticoid Receptor (pS211) Antibody**

Catalog # Source Reactivity Applications

CPA4130 Rabbit H, M, B WB, IH

**Description** Rabbit polyclonal antibody to Glucocorticoid Receptor (pS211)

Immunogen KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human Glucocorticoid Receptor (pS211). The exact sequence is

proprietary.

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

**Specificity** Recognizes endogenous levels of Glucocorticoid Receptor (pS211) protein.

**Clonality** Polyclonal

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

and 0.01% sodium azide.

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

Gene Symbol NR3C1

Alternative Names GRL; Glucocorticoid receptor; GR; Nuclear receptor subfamily 3 group C member 1

Entrez Gene 2908 (Human)

SwissProt P04150 (Human); P06537 (Mouse)

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

freeze/thaw cycles.

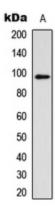
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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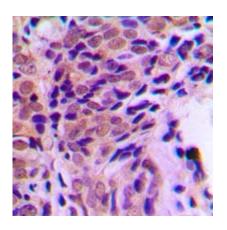
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Western blot analysis of Glucocorticoid Receptor (pS211) expression in A549 (A) whole cell lysates.



Immunohistochemical analysis of Glucocorticoid Receptor (pS211) staining in human prostate 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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