

## PKA beta (catalytic subunit) (phospho Ser338) antibody

**Cat. No. GTX25816**

<b>Host</b>	Rabbit
<b>Clonality</b>	Polyclonal
<b>Isotype</b>	IgG
<b>Applications</b>	WB, ChIP assay
<b>Reactivity</b>	Human, Mouse

**Package**

50 µl

## Applications

**Application Note**

\*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	Assay dependent
ChIP assay	1:100

Not tested in other applications.

**Calculated MW** 41 kDa. ( [Note](#) )**Product Note**

Peptide competition data indicate that this antibody cross-reacts with the PKA nu subunit (64% homologous) and partially with the alpha subunit (82% homologous).

## Properties

<b>Form</b>	Liquid
<b>Buffer</b>	PBS, 0.1% BSA, 50% Glycerol
<b>Preservative</b>	0.05% Sodium azide
<b>Storage</b>	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
<b>Concentration</b>	Batch dependent (Please refer to the vial label for the specific concentration.)
<b>Immunogen</b>	The antiserum was produced against a chemically synthesized phosphopeptide derived from a region of human PKA catalytic b subunit that contains serine 338. The sequence is conserved in cow and pig.
<b>Purification</b>	Purified by antigen-affinity chromatography
<b>Conjugation</b>	Unconjugated



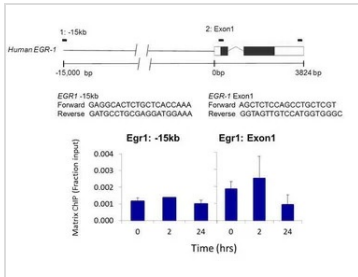
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**Note**

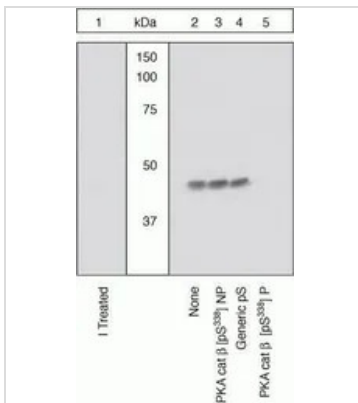
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**DATA IMAGES**

**GTx25816 ChIP assay Image**

ChIP analysis of human 5A8 J-Lat T lymphocytes culture treated with 10 µg/mL PHA (phytohemagglutinin) for 0, 2, and 24 hours using GTx25816 PKA beta (catalytic subunit) (phospho Ser338) antibody.

Immunoprecipitation was performed using a multiplex microplate Matrix ChIP assay (refer to PMID: 22098709). The precipitated DNA was detected by PCR with primer set targeting to -15kb upstream and exon-1 of the EGR1 gene.

IP reaction : 1.0ul antibody / 100ul sample


**GTx25816 WB Image**

WB (peptide competition) analysis of 3T3-L1 cells using GTx25816 PKA beta (catalytic subunit) (phospho Ser338) antibody prior incubated with the non-phosphopeptide corresponding to the immunogen (Lane 3), a generic phosphoserine-containing peptide (Lane 4), or the phosphopeptide immunogen (Lane 5) control. The data show that only the immunogen phosphopeptide blocks the signal, demonstrating the specificity of the antibody. The membrane treated with phosphatase (Lane 1) eliminates the signal further verifying that the antibody is phospho-specific.



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