

# PRODUCT DATA SHEET

Bioworld Technology,Inc.

# PKR Rabbit monoclonal antibody

Catalog: MB66277 Host: Rabbit Reactivity: Mouse, Rat

## **BackGround:**

An interferon-inducible, RNA-dependent protein serine/threonine kinase (PKR) has been described. PKR in earlier literature is variously known as DAI, dsJ, PI kinase, p65, p67 or TIK for the mouse kinase; and p68 or p69 for the human kinase. The PKR kinase substrate is the a subunit of protein synthesis initiation factor eIF-2. Phosphorylation of eIF-2a on Serine-51 results in inhibition of translation. Molecular cDNA clones have been isolated from both human and mouse cells. The serine/threonine kinase catalytic domains map to the car boxy terminal half of the protein while the RNA-binding domains are located in the amino terminal region. Three kinds of regulation of PKR enzymatic activity have been described. These include transcriptional regulation in response to interferon, an autoregulatory mechanism controlling PKR expres sion at the level of translation and post-translational regulation by RNA mediated autophosphorylation.

### **Product:**

Liquid in 50mM Tris-Glycine (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide and 0.05% BSA.

# **Molecular Weight:**

~ 75 kDa

## **Swiss-Prot:**

Q03963/Q63184

# **Purification&Purity:**

The antibody was purified by immunogen affinity chromatography.

# **Applications:**

WB (1/500 - 1/1000), IF/ICC (1/50 - 1/100), IP (1/10 - 1/50)

## Storage&Stability:

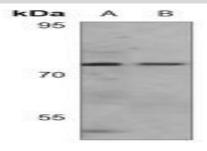
Store at  $4 \,\mathrm{C}$  short term. Aliquot and store at  $-20 \,\mathrm{C}$  long

term. Avoid freeze-thaw cycles.

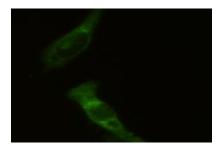
## **Specificity:**

Recognizes endogenous levels of PKR protein.

## **DATA:**



Western blot analysis of PKR expression in C6 (A), NIH3T3 (B) whole cell lysates.



Immunofluorescent analysis of PKR staining in NIH3T3 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 AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

#### Note:

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

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