

# PRODUCT DATA SHEET

Bioworld Technology,Inc.

# **EEF2** monoclonal antibody

Catalog: MB66520 Host: Mouse Reactivity: Human

#### **BackGround:**

Eukaryotic elongation factor 2 (eEF2) catalyzes the translocation of peptidyl-tRNA from the A site to the P site on the ribosome. It has been shown that phosphorylation of eEF2 at threonine 56 by eEF2 kinase inhibits its activity. eEF2 kinase is normally dependent on Ca2+ ions and calmodulin. eEF2 kinase can also be activated by PKA in response to elevated cAMP levels, which are generally increased in stress- or starvation-related conditions. A variety of treatments known to raise intracellular Ca2+ or cAMP levels have been shown to result in increased phosphorylation of eEF2, and thus to inhibit peptide-chain elongation. The inactive phosphorylated eEF2 can be converted to its active nonphosphorylated form by a protein phosphatase, most likely a form of protein phosphatase-2A (PP-2A). Insulin, which activates protein synthesis in a wide range of cell types, induces rapid dephosphorylation of eEF2 through mTOR signaling and may involve modulation of the activity of the PP-2A or the eEF2 kinase or both.

### **Product:**

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.

# **Molecular Weight:**

~ 95 kDa

# **Swiss-Prot:**

P13639

# **Purification&Purity:**

The antibody was purified by immunogen affinity chromatography.

#### **Applications:**

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

# Storage&Stability:

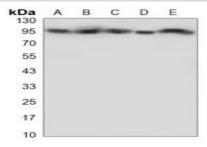
Store at 4 ℃ short term. Aliquot and store at -20 ℃ long

term. Avoid freeze-thaw cycles.

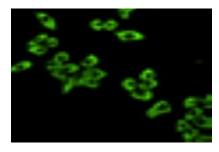
# **Specificity:**

Recognizes endogenous levels of EEF2 protein.

#### **DATA:**



Western blot analysis of EEF2 expression in HL60 (A), Jurkat (B), SHSY5Y (C), U20S (D), Hela (E) whole cell lysates.



Immunofluorescent analysis of EEF2 staining in HeLa 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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