

## KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody

Rabbit monoclonal antibody Catalog # AGI1815

### **Specification**

# KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody - Product Information

Application WB, FC, ICC Primary Accession 013541

Reactivity Rat, Human, Mouse

Clonality Monoclonal Isotype Rabbit IgG

Calculated MW Predicted, 13 kDa, observed, 15-20 kDa

**KDa** 

Gene Name EIF4EBP1

Aliases EIF4EBP1; Eukaryotic Translation Initiation

Factor 4E Binding Protein 1; PHAS-I; 4E-BP1; Phosphorylated Heat- And

Acid-Stable Protein Regulated By Insulin 1; Eukaryotic Translation Initiation Factor 4E-Binding Protein 1; EIF4E-Binding

Protein 1; 4EBP1; BP-1

Immunogen A synthesized peptide derived from human

Phospho-4E BP1 (Thr46)

# KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody - Additional Information

Gene ID 1978

**Other Names** 

Eukaryotic translation initiation factor 4E-binding protein 1, 4E-BP1, eIF4E-binding protein 1, Phosphorylated heat- and acid-stable protein regulated by insulin 1, PHAS-I, EIF4EBP1

# KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody - Protein Information

### Name EIF4EBP1

### **Function**

Repressor of translation initiation that regulates EIF4E activity by preventing its assembly into the eIF4F complex: hypophosphorylated form competes with EIF4G1/EIF4G3 and strongly binds to EIF4E, leading to repress translation. In contrast, hyperphosphorylated form dissociates from EIF4E, allowing interaction between EIF4G1/EIF4G3 and EIF4E, leading to initiation of translation. Mediates the regulation of protein translation by hormones, growth factors and other stimuli that signal through the MAP kinase and mTORC1 pathways.

#### **Cellular Location**

Cytoplasm. Nucleus. Note=Localization to the nucleus is unaffected by phosphorylation status.



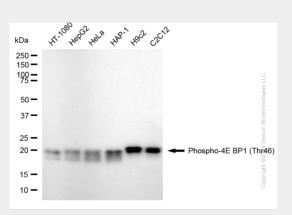
{ECO:0000250|UniProtKB:Q60876}

## KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody - Protocols

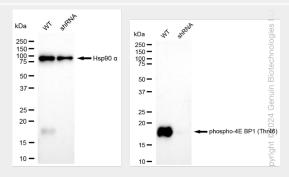
Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

## KD-Validated Anti-Phospho-4E BP1 (Thr46) Rabbit Monoclonal Antibody - Images

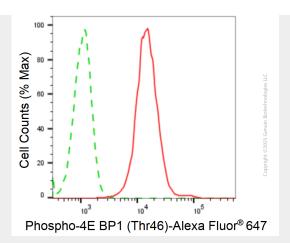


Western blotting analysis using anti-phospho-4E BP1 (Thr46) antibody (Cat#AGI1815). Total cell lysates (30  $\mu$ g) from various cell lines were loaded and separated by SDS-PAGE. The blot was incubated with anti-phospho-4E BP1 (Thr46) antibody (Cat#AGI1815, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody respectively.

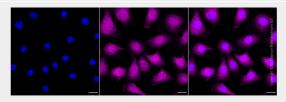


Western blotting analysis using anti-phospho-4E BP1 (Thr46) antibody (Cat#AGI1815). Phospho-4E BP1 (Thr46) expression in wild-type (WT) and EIF4EBP1 shRNA knockdown (KD) HeLa cells with 20  $\mu$ g of total cell lysates.  $\beta$ -Tubulin serves as a loading control. The blot was incubated with anti-phospho-4E BP1 (Thr46) antibody (Cat#AGI1815, 1:5,000) and HRP-conjugated goat anti-rabbit secondary antibody respectively.





Flow cytometric analysis of phospho-4E BP1 (Thr46) expression in C2C12 cells using anti-phospho-4E BP1 (Thr46) antibody (Cat#AGI1815, 1:2,000). Green, isotype control; red, phospho-4E BP1 (Thr46).



Immunocytochemical staining of C2C12 cells with anti-Phospho-4E BP1 (Thr46) antibody (Cat#AGI1815, 1:1,000). Nuclei were stained blue with DAPI; Phospho-4E BP1 (Thr46) was stained magenta with Alexa Fluor® 647. Images were taken using Leica stellaris 5. Protein abundance based on laser Intensity and smart gain: Medium. Scale bar, 20 µm.