Phospho-4E-BP1 (Thr36) Ab

Cat.#: AF3431 Concn.: 1mg/ml Mol.Wt.: 18kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200, IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-4E-BP1 (Thr36) Ab detects endogenous levels of 4E-

BP1 only when phosphorylated at Threonine 36

Immunogen: A synthesized peptide derived from human 4E-BP1 around

the phosphorylation site of Threonine 36

Uniprot: Q13541

Description: 4E-BP1 binds to eIF4E, preventing its assembly into the

EIF4F complex and inhibiting cap-dependent translation. Phosphorylation of 4E-BP1 disrupts this binding, activating

cap-dependent translation.

Similarity: The TOS motif mediates interaction with RPTOR, leading to

promote phosphorylation by mTORC1 complex.Belongs to

the eIF4E-binding protein family.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt

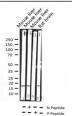


Western blot analysis of 4E-BP1 phosphorylation expression in EGF treated MDA-MB-435 whole cell lysates, The lane on the

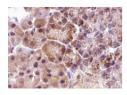
left is treated with the antigen-specific peptide.



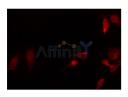
Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com



Western blot analysis of Phospho-4E-BP1 (Thr36) expression in various lysates



AF3431 at 1/100 staining human pancreas tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat antirabbit Ab was used as the secondary.



AF3431 staining Hela by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

IMPORTANT: For western blot, incubate membrane with diluted Ab in 5% w/v milk , 1X TBS, 0.1% Tween@20 at 4° C with gentle shaking, overnight.

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