

Product Data Sheet

Anti-4EBP1 (Phospho-T46) Antibody

Catalog # Source Reactivity Applications

CQA8054 Rabbit H, M, R WB, IH, IP

Description Rabbit monoclonal antibody to 4EBP1 (Phospho-T46)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

T46 of human 4EBP1 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of 4EBP1 protein only when phosphorylated at T46.

Clonality Monoclonal

Conjugation

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

and 0.05% BSA.

Dilution WB (1/500 - 1/1000), IH (1/50 - 1/100), IP (1/10 - 1/50)

Gene Symbol EIF4EBP1

Alternative 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

Entrez Gene 1978 (Human); 13685 (Mouse); 116636 (Rat)

SwissProt Q13541 (Human); Q60876 (Mouse); Q62622 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

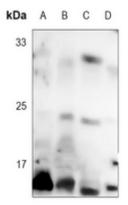
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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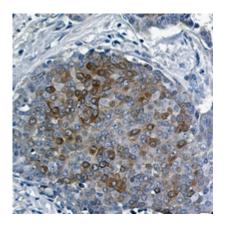
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Western blot analysis of 4EBP1 (Phospho-T46) expression in K562 (A), C6 (B), NIH3T3 (C), Hela (D) whole cell lysates. (Predicted band size: 13 kD; Observed band size: 15-20 kD)



Immunohistochemical analysis of 4EBP1 (Phospho-T46) staining in human breast carcinoma formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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