

Product Data Sheet

Anti-EIF2S2 (Phospho-S67) Antibody

Catalog # Source Reactivity Applications

CPA3585 Rabbit H, M, R, Mk WB, IH

Description Rabbit polyclonal antibody to EIF2S2 (Phospho-S67)

Immunogen KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding

S67 of human EIF2S2 protein. The exact sequence is proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of EIF2S2 protein only when phosphorylated at S67.

Clonality Polyclonal

Conjugation

Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,

and 0.01% sodium azide.

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

Gene Symbol EIF2S2

Alternative Names EIF2B; Eukaryotic translation initiation factor 2 subunit 2; Eukaryotic translation

initiation factor 2 subunit beta; eIF-2-beta

Entrez Gene 8894 (Human); 67204 (Mouse)

SwissProt P20042 (Human); Q99L45 (Mouse)

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

freeze/thaw cycles.

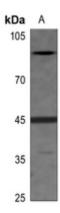
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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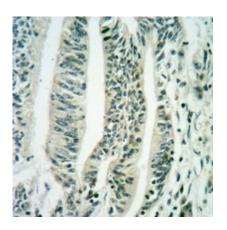
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Western blot analysis of EIF2S2 (Phospho-S67) expression in DLD (A) whole cell lysates. (Predicted band size: 38 kD; Observed band size: 45 kD)



Immunohistochemical analysis of EIF2S2 (Phospho-S67) staining in human colorectal cancer 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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