bsm-63191R

[Primary Antibody]

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IHC-P (1:50-200)

IHC-F (1:50-200)

ICC/IF (1:50-200)

IF (1:50-200) Flow-Cyt (1:50-100)

IP (1:20-50)

eIF4EBP1 Recombinant Rabbit mAb

DATASHEET -

Host: Rabbit Isotype: IgG Clonality: Recombinant CloneNo.: 9F8 **GeneID: 1978 SWISS:** Q13541

Target: eIF4EBP1

Immunogen: A synthesized peptide derived from human 4E BP1: 1-46.

Purification: affinity purified by Protein A

Storage: 10mM phosphate buffered saline(pH 7.4) with 150mM sodium chloride, 0.05% BSA, 0.02% Proclin300 and 50% glycerol.

Store at 4°C for short term. Store at -20°C for long term. Avoid

repeated freeze/thaw cycles.

Background: 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.

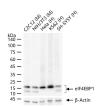
Reactivity: Human, Mouse, Rat

Applications: WB (1:500-2000)

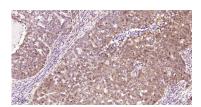
Predicted MW.: 13 kDa

Subcellular Cytoplasm ,Nucleus

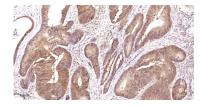
VALIDATION IMAGES -



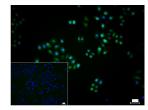
25 ug total protein per lane of various lysates (see on figure) probed with eIF4EBP1 monoclonal antibody, unconjugated (bsm-63191R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



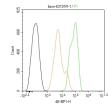
Paraformaldehyde-fixed, paraffin embedded Human Breast Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with eIF4EBP1 Monoclonal Antibody, Unconjugated (bsm-63191R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Human Colon Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with eIF4EBP1 Monoclonal Antibody, Unconjugated (bsm-63191R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.



4% Paraformaldehyde-fixed Hela (H) cell; Triton X-100 at r.t. for 20 min: Antibody incubation with (4E-BP1) monoclonal Antibody, unconjugated (bsm-63191R) 1:100, 90 min at 37°C; followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-60295G-BF488) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.



The Hela (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% icecold methanol for 20 min at -20°C, the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.). followed by secondary antibody incubation for 40 min at room temperature. Primary Antibody (green):Rabbit Anti-4E-BP1 antibody (bsm-63191R,1:100); Isotype Control (orange): Rabbit IgG (bs-0295P), Blank control (black): PBS. Acquisition of 20,000 events was performed.