

# Phospho-S6 Ribosomal Protein (RPS6)-S235/236 polyclonal antibody

Catalog: BS79397

Host: Rabbit

Reactivity: Human, Mouse, Rat

## BackGround:

Ribosomes, the organelles that catalyze protein synthesis, consist of a small 40S subunit and a large 60S subunit. Together these subunits are composed of 4 RNA species and approximately 80 structurally distinct proteins. This gene encodes a cytoplasmic ribosomal protein that is a component of the 40S subunit. The protein belongs to the S6E family of ribosomal proteins. It is the major substrate of protein kinases in the ribosome, with subsets of five C-terminal serine residues phosphorylated by different protein kinases. Phosphorylation is induced by a wide range of stimuli, including growth factors, tumor-promoting agents, and mitogens. Dephosphorylation occurs at growth arrest. The protein may contribute to the control of cell growth and proliferation through the selective translation of particular classes of mRNA. As is typical for genes encoding ribosomal proteins, there are multiple processed pseudogenes of this gene dispersed through the genome.

## Product:

1mg/ml in PBS with 0.02% sodium azide, 50% glycerol, pH7.2

## Molecular Weight:

32KDa

## Swiss-Prot:

P62753

## Purification&Purity:

The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen and the purity is > 95% (by SDS-PAGE).

## Applications:

WB, 1:500 - 1:2000

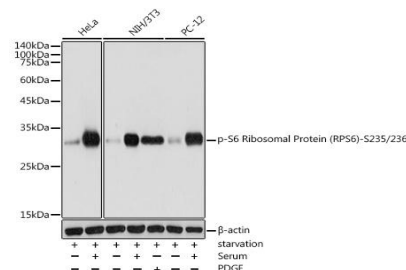
## Storage&Stability:

Store at 4 °C short term. Aliquot and store at -20 °C long term. Avoid freeze-thaw cycles.

## Modification:

Phosphorylated

## DATA:



Western blot analysis of extracts of various cell lines, using Phospho-S6 Ribosomal Protein -S235/236 antibody at 1:1000 dilution. HeLa cells NIH/3T3 cells and PC-12 cells were treated by 10% FBS at 37 °C for 30 minutes after serum-starvation overnight. NIH/3T3 cells were treated by PDGF at 37 °C for 30 minutes after serum-starvation overnight. Secondary antibody: HRP Goat Anti-Rabbit IgG at 1:10000 dilution. Lysates/proteins: 25ug per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit. Exposure time: 1s.

## Note:

For research use only, not for use in diagnostic procedure.

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