

PRODUCT DATA SHEET

Bioworld Technology,Inc.

NF-kappaB p105 polyclonal antibody

Catalog: BS67441 Host: Rabbit Reactivity: Human

BackGround:

Transcription factors of the nuclear factor (NF-κB)/Rel family play a pivotal role in inflammatory and immune responses . There are five family members in mammals: RelA, c-Rel, RelB, NF-κB1 (p105/p50), and NF-kB2 (p100/p52). Both p105 and p100 are proteolytically processed by the proteasome to produce p50 and p52, respectively. Rel proteins bind p50 and p52 to form dimeric complexes that bind DNA and regulate transcription. In unstimulated cells, NF-kB is sequestered in the cytoplasm by IκB inhibitory proteins . NF-κB-activating agents can induce the phosphorylation of IkB proteins, targeting them for rapid degradation through the ubiquitin-proteasome pathway and releasing NF-kB to enter the nucleus where it regulates gene expression. NIK and IKKα (IKK1) regulate the phosphorylation and processing of NF-κB2 (p100) to produce p52, which translocates to the nucleus .Following IKK-mediated phosphorylation of p105 NF-κB at multiple sites (Ser921, 923, 927, and 932) on its carboxy-terminus, SCF/β-TrCP mediated processing produces the 50 kDa active form p50.

Product:

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Molecular Weight:

~ 120 kDa

Swiss-Prot:

P19838

Purification&Purity:

The antibody was purified by immunogen affinity chromatography.

Applications:

WB (1/500 - 1/1000), IHC (1/50 - 1/200)

Storage&Stability:

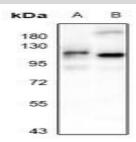
Store at 4 ℃ short term. Aliquot and store at -20 ℃ long

term. Avoid freeze-thaw cycles.

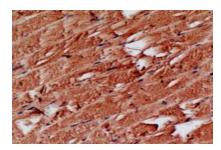
Specificity:

Recognizes endogenous levels of NF-kappaB p105 protein.

DATA:



Western blot analysis of NF-kappaB p105 expression in A375 (A), My-la2059 (B) whole cell lysates.



Immunohistochemical analysis of NF-kappaB p105 staining in human muscle 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.

Note:

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

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