

NF-kappaB p105 monoclonal antibody

Catalog: MB66917

Host: Mouse

Reactivity: Human

BackGround:

Transcription factors of the nuclear factor κВ $(NF-\kappa B)/Rel$ family play a pivotal role in inflammatory and immune responses . There are five family members in mammals: RelA, c-Rel, RelB, NF-KB1, and NF-KB2. 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 IKB inhibitory proteins . NF-kB-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 IKKa regulate the phosphorylation and processing of NF-kB2 to produce p52, which translocates to the nucleus .Following IKK-mediated phosphorylation of p105 NF-κB at multiple sites on its carboxy-terminus, SCF/β-TrCP mediated processing produces the 50 kDa active form p50.

Product:

Mouse IgG1 kappa. Liquid in PBS, pH 7.3, 30% glycerol, and 0.01% sodium azide.

Molecular Weight:

~ 50,120 kDa

Swiss-Prot:

P19838

Purification&Purity:

This antibody is purified through a protein G column.

Applications:

WB (1/500 - 1/1000), IHC (1/50 - 1/200), FC (1/10 - 1/50)

Storage&Stability:

Store at 4 ${\rm C}$ short term. Aliquot and store at -20 ${\rm C}$ long

term. Avoid freeze-thaw cycles.

Specificity:

Bioworld Technology, Inc.

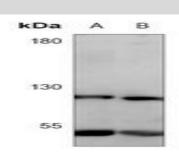
 Add:
 1660 South Highway 100, Suite 500 St. Louis Park, MN 55416,USA.

 Email:
 info@bioworlde.com

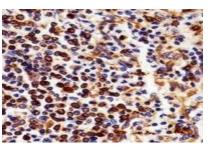
 Tel:
 6123263284

 Fax:
 6122933841
 Recognizes endogenous levels of NF-kappaB p105 protein.

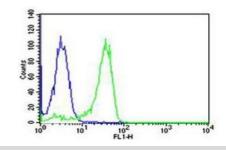
DATA:



Western blot analysis of NF-kappaB p105 expression in Daudi (A), Jurkat (B) whole cell lysates.



Immunohistochemical analysis of NF-kappaB p105 staining in human spleen 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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