Phospho-NF kappaB p105/p50 (Ser907) Ab

Cat.#: AF3221 Concn.: 1mg/ml Mol.Wt.: 105kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200 IP, IF/ICC 1:100-1:500

Reactivity: Human

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-NF- kappaB p105/p50 (Ser907) Ab detects

endogenous levels of NF- kappaB p105/p50 only when

phosphorylated at Serine 907

Immunogen: A synthesized peptide derived from human NF- kappaB

p105/p50 around the phosphorylation site of Serine 907

Uniprot: P19838

Description: NFkB-p105 a transcription factor of the nuclear factor-

kappaB (NFkB) group. Undergoes cotranslational processing by the 26S proteasome to produce a 50 kD protein. The 105 kD protein is a Rel protein-specific transcription inhibitor and the 50 kD protein is a DNA binding subunit of NFkB. NFkB is a transcription regulator that is activated by various intraand extra-cellular stimuli such as cytokines, oxidant-free radicals, ultraviolet irradiation, and bacterial or viral

products.

Subcellular Location: Nucleus. Cytoplasm. Nuclear, but also found in the

cytoplasm in an inactive form complexed to an inhibitor.

Tissue Specificity: By phorbol ester and TNF.

Similarity: The C-terminus of p105 might be involved in cytoplasmic

retention, inhibition of DNA-binding, and transcription activation.Glycine-rich region (GRR) appears to be a critical

element in the generation of p50.

Storage Condition and

Buffer:

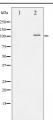
Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt

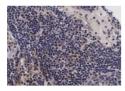


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Western blot analysis of NF kappaB p105/p50 phosphorylation expression in TNF- α treated HeLa whole cell lysates,The lane on the left is treated with the antigen-specific peptide.



AF3221 at 1/200 staining human lymph nodes tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22° C. An HRP conjugated goat antirabbit Ab was used as the secondary.



AF3221 staining HeLa by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

IMPORTANT: For western blot, incubate membrane with diluted Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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