

Phospho-NF kappaB p100/p52 (Ser866) Antibody

Cat.#: AF3373 Concn.: 1mg/ml Mol.Wt.: 100kDa 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,

ELISA(peptide) 1:20000-1:40000

Reactivity: Human, Mouse, Rat

Purification: The antibody is from purified rabbit serum by affinity

purification via sequential chromatography on phosphopeptide and non-phospho-peptide affinity columns.

Specificity: Phospho-NF kappaB p100/p52 (Ser866) Antibody detects

endogenous levels of NF kappaB p100/p52 only when

phosphorylated at Serine 866.

Immunogen: A synthesized peptide derived from human NF- kappaB

p100/p52 around the phosphorylation site of Ser866.

Uniprot: Q00653

Description: NFkB-p100 a transcription factor of the nuclear factor-

kappaB (NFkB) group. Precursor of the p52 subunit of the nuclear factor NF-kappa-B, which binds to the kappa-B consensus sequence 5'-GGRNNYYCC-3', located in the enhancer region of genes involved in immune response and

acute phase reactions.

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.



Western blot analysis of extracts from LPS treated RAW264.7 cells, using Phospho-NF kappaB p100/p52 (Ser866) Antibody. The lane on the left was treated with blocking peptide.

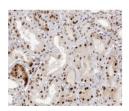


Affinity Biosciences

website:www.affbiotech.com order:order@affbiotech.com



Western blot analysis of extracts from mouse spleen/rat lung, using Phospho-NF kappaB p100/p52 (Ser866) Antibody.
-/+ means absence or presence of N peptide(non-phospho peptide) and P peptide(phospho peptide)



AF3373 at 1/200 staining human kidney 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 antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary antibody.



AF3373 staining RAW264.7 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 antibody 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 antibody.

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

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