

Kit Includes	Quantity	Applications	Reactivity	MW (kDa)	Isotype
Phospho-IKKα (Ser176)/IKKβ (Ser177) (C84E11) Rabbit mAb #2078	40 µl	W	H M (R) (Mk) (B)	85 (IKKα), 87 (IKKβ)	Rabbit IgG
Phospho-IKKα/β (Ser176/180) (16A6) Rabbit mAb #2697	40 µl	W IHC-P IHC-F	H M R Mk (B)	85 IKK-α 87 IKK-β	Rabbit IgG
Phospho-IKKα/β (Ser176/180) Antibody II #2694	40 µl	W	H M R Mk	85 IKK-α 87 IKK-β	Rabbit
Anti-rabbit IgG, HRP-linked Antibody #7074	100 µl				Goat

Applications Key: W=Western Blotting IHC-P=Immunohistochemistry (Paraffin) IHC-F=Immunohistochemistry (Frozen)

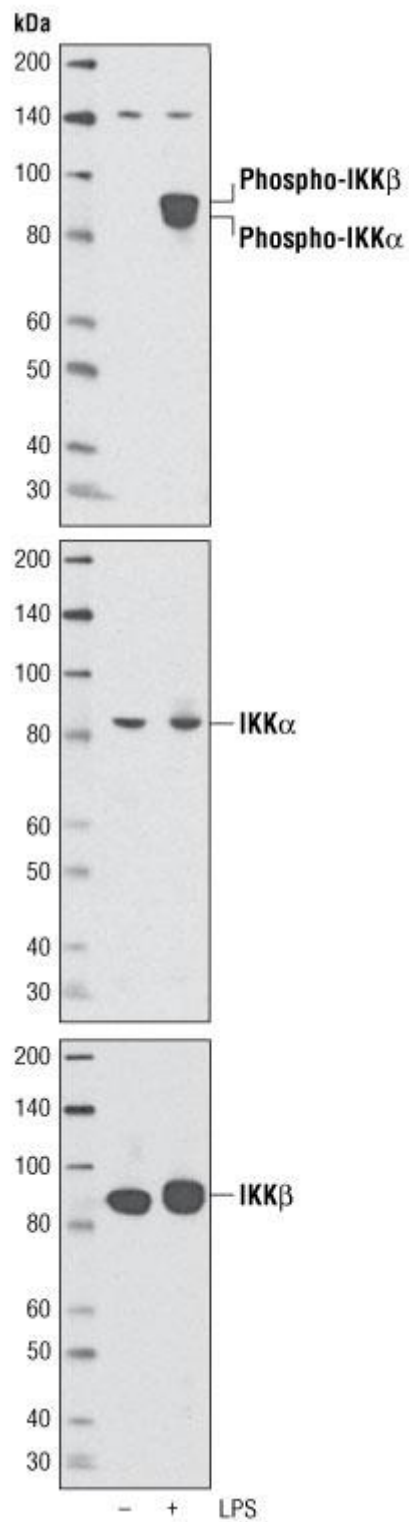
Reactivity Key: H=Human M=Mouse R=Rat Mk=Monkey B=Bovine

Species enclosed in parentheses are predicted to react based on 100% sequence homology.

Specificity / Sensitivity

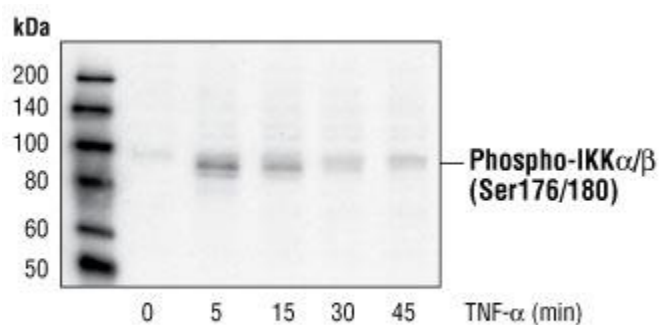
Phospho-IKKα/β (Ser176/180) Antibody, Phospho-IKKα/β (Ser176/180) Antibody II, and Phospho-IKKα/β (Ser176/180) (16A6) Rabbit mAb detect IKKα only when phosphorylated at Ser176/180 and IKKβ only when phosphorylated at Ser177/181.

Western Blotting



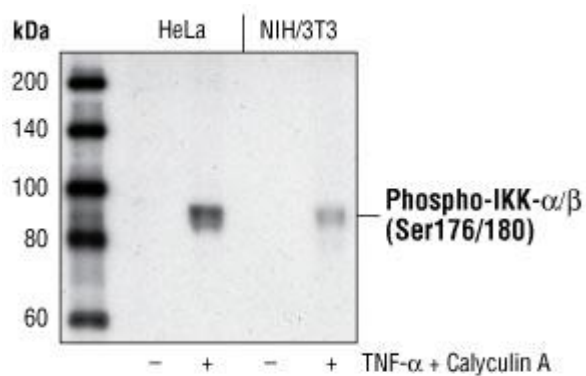
Western blot analysis of extracts from differentiated THP-1 cells, untreated or LPS-treated for 15 minutes, using Phospho-IKK α (Ser176)/IKK β (Ser177) (C84E11) Rabbit mAb #2078 (upper), IKK α Antibody #2682 (middle) and IKK β (2C8) Rabbit mAb #2370 (lower).

Western Blotting



Western blot analysis of extracts from HeLa cells, treated with TNF- α (20ng/ml) for the indicated times, using Phospho-IKK α/β (Ser176/180) Antibody II #2694.

Western Blotting



Western blot analysis of extracts from HeLa and NIH/3T3 cells, treated with TNF- α (20 ng/ml) and calyculin A #9902 (50 nM), using Phospho-IKK α/β (Ser176/180) (16A6) Rabbit mAb #2697.

Description

The Phospho-IKK α/β (Ser176/180) Antibody Sampler Kit contains reagents to examine protein levels of IKK α when phosphorylated at Ser176/180 and IKK β when phosphorylated at Ser177/181. The kit contains primary and secondary antibodies to perform four Western blots with each antibody.

Source / Purification

Polyclonal antibodies #2687 and #2694 are produced by immunizing rabbits with a synthetic phosphopeptide corresponding to residues surrounding Ser176/180 of human IKK α and Ser177/181 of IKK β , respectively, and are purified by protein A and peptide affinity chromatography. Monoclonal antibody #2697 is produced by immunizing rabbits with a synthetic phosphopeptide corresponding to residues surrounding Ser176/180 of human IKK α .

Background

The NF- κ B/Rel transcription factors are present in the cytosol in an inactive state, complexed with the inhibitory I κ B proteins (1-3). Most agents that activate NF- κ B do so through a common pathway based on phosphorylation-induced, proteasome-mediated degradation of I κ B (3-7). The key regulatory step in this pathway involves activation of a high molecular weight I κ B kinase (IKK) complex whose catalysis is generally carried out by three tightly associated IKK subunits. IKK α and IKK β serve as the catalytic subunits of the kinase and IKK γ serves as the regulatory subunit (8,9). Activation of IKK depends upon phosphorylation at Ser177 and Ser181 in the activation loop of IKK β (Ser176 and Ser180 in IKK α), which causes conformational changes, resulting in kinase activation (10-13).

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4. [Brown, K. et al. \(1995\) *Science* 267, 1485-8.](#)
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