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## Phospho-IKKα/β (Ser176/180) Antibody II 2007 707



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Applications: W	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 85 IKK-alpha 87 IKK-beta	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #O14920, #O15111	<b>Entrez-Gene Id:</b> 3551, 1147		
Product UsageApplicationInformationWestern Blotting		<b>Dilution</b> 1:1000						
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				erol. Store at –		
Specificity/Sensitivity		Phospho-ΙΚΚα/β (Ser176/180) Antibody II detects endogenous levels of ΙΚΚα and ΙΚΚβ only when phosphorylated at Ser176 and Ser180 or Ser177 and Ser181, respectively.						
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a phosphopeptide corresponding to a region surrounding Ser177/181 of ΙΚΚβ. Antibodies are purified by protein A and peptide affinity chromatography.						
Background		The NF-κB/Rel transcription factors are present in the cytosol in an inactive state, complexed with the inhibitory IκB proteins (1-3). Most agents that activate NF-κB do so through a common pathway based on phosphorylation-induced, proteasome-mediated degradation of IκB (3-7). The key regulatory step in this pathway involves activation of a high molecular weight IκB kinase (IKK) complex whose catalysis is generally carried out by three tightly associated IKK subunits. IKKα and IKKβ serve as the catalytic subunits of the kinase and IKKγ serves as the regulatory subunit (8,9). Activation of IKK depends upon phosphorylation at Ser177 and Ser181 in the activation loop of IKKβ (Ser176 and Ser180 in IKKα), which causes conformational changes, resulting in kinase activation (10-13).						
Background R	eferences	<ol> <li>Baeuerle, P.A. and Baltimore, D. (1988) <i>Science</i> 242, 540-6.</li> <li>Beg, A.A. and Baldwin, A.S. (1993) <i>Genes Dev</i> 7, 2064-70.</li> <li>Finco, T.S. et al. (1994) <i>Proc Natl Acad Sci USA</i> 91, 11884-8.</li> <li>Brown, K. et al. (1995) <i>Science</i> 267, 1485-8.</li> <li>Brockman, J.A. et al. (1995) <i>Mol Cell Biol</i> 15, 2809-18.</li> <li>Traenckner, E.B. et al. (1995) <i>EMBO J</i> 14, 2876-83.</li> <li>Chen, Z.J. et al. (1996) <i>Cell</i> 84, 853-62.</li> <li>Zandi, E. et al. (1997) <i>Cell</i> 91, 243-52.</li> <li>Karin, M. (1999) <i>Oncogene</i> 18, 6867-74.</li> <li>DiDonato, J.A. et al. (1997) <i>Nature</i> 388, 548-54.</li> <li>Mercurio, F. et al. (1997) <i>Science</i> 278, 860-6.</li> <li>Johnson, L.N. et al. (1996) <i>Cell</i> 85, 149-58.</li> <li>Delhase, M. et al. (1999) <i>Science</i> 284, 309-13.</li> </ol>						
Species Reacti	vitv	Species reactivity is o	determined by testing	in at least one appro	ved application (e.g., w	vestern blot).		
Western Blot E	-	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.						
Applications K	ey	W: Western Blotting						
Cross-Reactivi	ty Key	H: Human M: Mouse R: Rat Mk: Monkey						
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