## 0286

## IKKβ (2C8) Rabbit mAb



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<b>Applications:</b> W, W-S	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 87	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #O14920	Entrez-Gene Id 3551
Product Usage Information		<b>Application</b> Western Blotting Simple Western™		<b>Dilution</b> 1:1000 1:10 - 1:50		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		IKKβ (2C8) Rabbit mAb detects endogenous levels of total IKKβ protein. The antibody does not cross-react with IKKα or IKKγ.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues at the carboxy terminus of human IKK $\beta$ protein.				
Background		The NF- $\kappa$ B/Rel transcription factors are present in the cytosol in an inactive state, complexed with the inhibitory I $\kappa$ B proteins (1-3). Most agents that activate NF- $\kappa$ B do so through a common pathway based on phosphorylation-induced, proteasome-mediated degradation of I $\kappa$ B (3-7). The key regulatory step in this pathway involves activation of a high molecular weight I $\kappa$ B kinase (IKK) complex whose catalysis is generally carried out by three tightly associated IKK subunits. IKK $\alpha$ and IKK $\beta$ serve as the catalytic subunits of the kinase and IKK $\gamma$ serves as the regulatory subunit (8,9). Activation of IKK depends upon phosphorylation at Ser177 and Ser181 in the activation loop of IKK $\beta$ (Ser176 and Ser180 in IKK $\alpha$ ), which causes conformational changes, resulting in kinase activation (10-13).				
Background References		1. Baeuerle, P.A. and Baltimore, D. (1988) <i>Science</i> 242, 540-6. 2. Beg, A.A. and Baldwin, A.S. (1993) <i>Genes Dev</i> 7, 2064-70. 3. Finco, T.S. et al. (1994) <i>Proc Natl Acad Sci USA</i> 91, 11884-8. 4. Brown, K. et al. (1995) <i>Science</i> 267, 1485-8. 5. Brockman, J.A. et al. (1995) <i>Mol Cell Biol</i> 15, 2809-18. 6. Traenckner, E.B. et al. (1995) <i>EMBO J</i> 14, 2876-83. 7. Chen, Z.J. et al. (1996) <i>Cell</i> 84, 853-62. 8. Zandi, E. et al. (1997) <i>Cell</i> 91, 243-52. 9. Karin, M. (1999) <i>Oncogene</i> 18, 6867-74. 10. DiDonato, J.A. et al. (1997) <i>Nature</i> 388, 548-54. 11. Mercurio, F. et al. (1997) <i>Science</i> 278, 860-6. 12. Johnson, L.N. et al. (1999) <i>Science</i> 284, 309-13.				
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).				
Western Blot Buffer		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 Key		<b>W</b> : Western Blotting <b>W-S</b> : Simple Western™				
Cross-Reactivity Key		H: Human M: Mouse R: Rat Mk: Monkey				

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