## IκBβ (7B4) Mouse mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 48	<b>Source/Isotype:</b> Mouse IgG1	<b>UniProt ID:</b> #Q15653	Entrez-Gene Id: 4793
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		ІкВ $\beta$ (7B4) Mouse mAb recognizes endogenous levels of total ІкВ $\beta$ protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a recombinant protein specific to a carboxy terminal fragment of human $I\kappa B\beta$ protein.				
Background		The NF-κB/Rel transcription factors are present in the cytosol in an inactive state complexed with the inhibitory IκB proteins (1-3). Activation occurs via phosphorylation of IκBα at Ser32 and Ser36 followed by proteasome-mediated degradation that results in the release and nuclear translocation of active NF-κB (3-7). IκBα phosphorylation and resulting Rel-dependent transcription are activated by a highly diverse group of extracellular signals including inflammatory cytokines, growth factors, and chemokines. Kinases that phosphorylate IκB at these activating sites have been identified (8). The regulation of IκBβ and IκBε is similar to that of IκBα. However, the phosphorylation and ubiquitin-mediated degradation of these proteins occurs with much slower kinetics (9). IKK phosphorylation of IκBβ occurs at Ser19 and Ser23, while IκΒε can be phosphorylated at Ser18 and Ser22 (10).				
Background References		<ol> <li>Baeuerle, P.A. and Baltimore, D. (1988) Science 242, 540-6.</li> <li>Beg, A.A. and Baldwin, A.S. (1993) Genes Dev 7, 2064-70.</li> <li>Finco, T.S. et al. (1994) Proc Natl Acad Sci USA 91, 11884-8.</li> <li>Brown, K. et al. (1995) Science 267, 1485-8.</li> <li>Brockman, J.A. et al. (1995) Mol Cell Biol 15, 2809-18.</li> <li>Traenckner, E.B. et al. (1995) EMBO J 14, 2876-83.</li> <li>Chen, Z.J. et al. (1996) Cell 84, 853-62.</li> <li>Karin, M. and Ben-Neriah, Y. (2000) Annu Rev Immunol 18, 621-63.</li> <li>Hoffmann, A. et al. (2002) Science 298, 1241-5.</li> <li>Shirane, M. et al. (1999)   Biol Chem 274, 28169-74.</li> </ol>				
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**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 W: Western Blotting

Cross-Reactivity Key H: Human

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