33

Phospho-Cyclin B1 (Ser133) (9E3) Rabbit mAb



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Applications: W, IP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 55	Source/Isotype: Rabbit IgG	UniProt ID: #P14635	Entrez-Gene Id: 891	
Product Usage Information	2	Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:100		
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/Ser	nsitivity	Phospho-Cyclin B1 (Ser133) (9E3) Rabbit mAb detects endogenous levels of cyclin B1 protein only w phosphorylated at Ser133.				protein only when	
Source / Purifi	cation	Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser133 of human cyclin B1.					
Background		Cyclins are a family of proteins that activate specific cyclin-dependent kinases required for progression through the cell cycle. The entry of all eukaryotic cells into mitosis is regulated by activation of cdc2/cdk1 at the G2/M transition. This activation is a multi-step process that begins with the binding of the regulatory subunit, cyclin B1, to cdc2/cdk1 to form the mitosis-promoting factor (MPF). MPF remains in the inactive state until phosphorylation of cdc2/cdk1 at Thr161 by cdk activating kinase (CAK) (1,2) and dephosphorylation of cdc2/cdk1 at Thr14/Tyr15 by cdc25C (3-5). Five cyclin B1 phosphorylation sites (Ser116, 126, 128, 133, and 147) are located in the cytoplasmic retention signal (CRS) domain and are thought to regulate the translocation of cyclin B1 to the nucleus at the G2/M checkpoint, promoting nuclear accumulation and initiation of mitosis (6-9). While MPF itself can phosphorylate Ser126 and Ser128, polo-like kinase 1 (PLK1) phosphorylates cyclin B1 preferentially at Ser133 and possibly at Ser147 (6,10). At the end of mitosis, cyclin B1 is targeted for degradation by the anaphase-promoting complex (APC), allowing for cell cycle progression (11). Research studies have shown that cyclin B1 is overexpressed in breast, prostate, and non-small cell lung cancers (12-14).					
Background R	eferences	1. Lorca, T. et al. (1992) EMBO J 11, 2381-90. 2. Harper, J.W. and Elledge, S.J. (1998) Genes Dev 12, 285-9. 3. Norbury, C. et al. (1991) EMBO J 10, 3321-9. 4. McGowan, C.H. and Russell, P. (1993) EMBO J 12, 75-85. 5. Atherton-Fessler, S. et al. (1994) Mol Biol Cell 5, 989-1001. 6. Toyoshima-Morimoto, F. et al. (2001) Nature 410, 215-20. 7. Li, J. et al. (1997) Proc Natl Acad Sci U S A 94, 502-7. 8. Takizawa, C.G. and Morgan, D.O. (2000) Curr Opin Cell Biol 12, 658-65. 9. Santos, S.D. et al. (2012) Cell 149, 1500-13. 10. Jackman, M. et al. (2003) Nat Cell Biol 5, 143-8. 11. Gong, D. and Ferrell, J.E. (2010) Mol Biol Cell 21, 3149-61. 12. Mashal, R.D. et al. (1996) Cancer Res 56, 4159-63. 13. Kawamoto, H. et al. (2000) Cancer Res 60, 4000-4.					
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).	
Western Blot B	-	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	ley	W: Western Blotting IP: Immunoprecipitation					
Cross-Reactivi	ty Key	H: Human					
Trademarks aı	nd Patents	Cell Signaling Technology is a trademark of Cell Signaling Technology, Inc.					

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