Cyclin B1 Antibody



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Applications: W, IF-IC	Reactivity: H M R Hm Mk	Sensitivity: Endogenous	MW (kDa): 55	Source/Isotype: Rabbit	UniProt ID: #P14635	Entrez-Gene Id: 891
Product Usage Information	2	Application Western Blotting Immunofluorescence	(Immunocytochem	istry)		Dilution 1:1000 1:400
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Cyclin B1 Antibody detects endogenous levels of total cyclin B1.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a syntheic peptide corresponding to residues near the amino terminus of human cyclin B1. Antibodies are purified using peptide affinity chromatography.				
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 References		1. Lorca, T. et al. (1992) <i>EMBO J</i> 11, 2381-90. 2. Harper, J.W. and Elledge, S.J. (1998) <i>Genes Dev</i> 12, 285-9. 3. Norbury, C. et al. (1991) <i>EMBO J</i> 10, 3321-9. 4. McGowan, C.H. and Russell, P. (1993) <i>EMBO J</i> 12, 75-85. 5. Atherton-Fessler, S. et al. (1994) <i>Mol Biol Cell</i> 5, 989-1001. 6. Toyoshima-Morimoto, F. et al. (2001) <i>Nature</i> 410, 215-20. 7. Li, J. et al. (1997) <i>Proc Natl Acad Sci U S A</i> 94, 502-7. 8. Takizawa, C.G. and Morgan, D.O. (2000) <i>Curr Opin Cell Biol</i> 12, 658-65. 9. Santos, S.D. et al. (2012) <i>Cell</i> 149, 1500-13. 10. Jackman, M. et al. (2003) <i>Nat Cell Biol</i> 5, 143-8. 11. Gong, D. and Ferrell, J.E. (2010) <i>Mol Biol Cell</i> 21, 3149-61. 12. Mashal, R.D. et al. (1996) <i>Cancer Res</i> 56, 4159-63. 13. Kawamoto, H. et al. (1997) <i>Am J Pathol</i> 150, 15-23. 14. Soria, J.C. et al. (2000) <i>Cancer Res</i> 60, 4000-4.				

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 **IF-IC:** Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey

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