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## Phospho-cdc2 (Thr14) Antibody

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

<b>Applications:</b> W	<b>Reactivity:</b> H Hm Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 34	<b>Source/Isotype:</b> Rabbit	<b>UniProt ID:</b> #P06493	<b>Entrez-Gene Id:</b> 983
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<b>Product Usage Information</b>	<b>Application</b> Western Blotting	<b>Dilution</b> 1:1000
<b>Storage</b>	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.	
<b>Specificity/Sensitivity</b>	Phospho-cdc2 (Thr14) Antibody detects endogenous levels of cdc2 (CDK1) only when phosphorylated at Thr14. Based on sequence similarity, the antibody may cross-react with CDK2 and CDK3.	
<b>Species predicted to react based on 100% sequence homology</b>	Mouse, Rat, D. melanogaster, Xenopus, Bovine	
<b>Source / Purification</b>	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr14 of human cdc2. Antibodies are purified by protein A and peptide affinity chromatography.	
<b>Background</b>	The entry of eukaryotic cells into mitosis is regulated by cdc2 kinase activation, a process controlled at several steps including cyclin binding and phosphorylation of cdc2 at Thr161 (1). However, the critical regulatory step in activating cdc2 during progression into mitosis appears to be dephosphorylation of cdc2 at Thr14 and Tyr15 (2). Phosphorylation at Thr14 and Tyr15, resulting in inhibition of cdc2, can be carried out by Wee1 and Myt1 protein kinases (3,4). The cdc25 phosphatase may be responsible for removal of phosphates at Thr14 and Tyr15 and subsequent activation of cdc2 (1,5).	
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Atherton-Fessler, S. et al. (1994) <i>Mol Biol Cell</i> 5, 989-1001.</li> <li>2. Norbury, C. et al. (1991) <i>EMBO J</i> 10, 3321-9.</li> <li>3. McGowan, C.H. and Russell, P. (1993) <i>EMBO J</i> 12, 75-85.</li> <li>4. Wells, N.J. et al. (1999) <i>J Cell Sci</i> 112 ( Pt 19), 3361-71.</li> <li>5. Hunter, T. (1995) <i>Cell</i> 80, 225-36.</li> </ol>	

<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).
<b>Western Blot Buffer</b>	<b>IMPORTANT:</b> 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.
<b>Applications Key</b>	<b>W:</b> Western Blotting
<b>Cross-Reactivity Key</b>	<b>H:</b> Human <b>Hm:</b> Hamster <b>Mk:</b> Monkey
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