

Phospho-(Ser) CDKs Substrate Antibody



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

Applications: W, IP, E-P	Sensitivity: Endogenous	Source/Isotype: Rabbit
Product Usage Information	Application	Dilution
	Western Blotting	1:1000
	Immunoprecipitation	1:100
	Peptide ELISA (DELFI A)	1:1000
Storage	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.	
Specificity/Sensitivity	Phospho-(Ser) CDKs Substrate Antibody detects phospho-serine in a (K/R)(S*)PX(K/R) motif. The antibody is phospho-specific but does not recognize phospho-serine in the absence of the CDK motif. The antibody does not cross-react with phospho-threonine- or phospho-tyrosine-containing peptides/proteins. (U.S. Patent No's.: 6,441,140; 6,982,318; 7,259,022; 7,344,714; U.S.S.N. 11,484,485; and all foreign equivalents.)	
Source / Purification	Polyclonal antibodies are produced by immunizing animals with synthetic phospho-CDK substrate peptides. Antibodies are purified by protein A and peptide affinity chromatography.	
Background	Cyclin-dependent kinases (CDKs) are a family of Ser/Thr kinases that regulate cell-cycle transitions through their association and subsequent phosphorylation of targets in a strictly ordered fashion (1). The substrates for CDKs are proline-directed. The consensus amino acid sequence for CDK substrate is (K/R)(S*)PX(K/R), where X denotes any one of the 20 amino acids (2-4) and S* is the phosphorylation site. Phospho-CDK Substrate Motif [(K/H)pSP] MultiMab™ Rabbit mAb mix recognizes phosphorylated CDK substrates at their consensus motif, providing a powerful tool for CDK target discovery and characterization, as well as HTS drug screening for potential kinase regulators.	
Background References	<ol style="list-style-type: none"> 1. Morgan, D.O. (1997) <i>Annu Rev Cell Dev Biol</i> 13, 261-91. 2. Songyang, Z. et al. (1996) <i>Mol Cell Biol</i> 16, 6486-93. 3. Songyang, Z. (1999) <i>Prog Biophys Mol Biol</i> 71, 359-72. 4. Holmes, J.K. and Solomon, M.J. (1996) <i>J Biol Chem</i> 271, 25240-6. 	
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 IP: Immunoprecipitation E-P: Peptide ELISA (DELFI A)	
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