Phospho-PNK1 (Ser114/Thr118) Antibody



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Applications: W	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 59	Source/Isotype: Rabbit	UniProt ID: #Q96T60	Entrez-Gene Id: 11284
Product Usage Information		Application Western Blotting		Dilution 1:1000		
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		Phospho-PNK1 (Ser114/Thr118) Antibody detects endogenous levels of PNK1 only when phosphorylated on Ser114 and Thr118.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues around Ser114 and Thr118 of human PNK1. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		PNK (polynucleotide kinase) is a DNA repair enzyme that participates in single strand break repair and non-homologous end rejoining (NHEJ) for double strand breaks. PNK possesses a 5'-DNA kinase activity and a 3'-DNA phosphatase activity (1,2). It has three domains, a C-terminal kinase domain, a central phosphatase domain, and an N-terminal forkhead associated (FHA) domain that is responsible for protein-protein interactions. Reduction in expression of PNK by RNAi sensitizes cells to ionizing radiation and topoisomerase I inhibitors (3) Phosphorylation of PNK1 at Ser114/ Thr118 was independently identified at Cell Signaling Technology (CST) using PhosphoScan®, CST's MS/MS platform for phosphorylation site discovery. Phosphorylation of PNK1 at Ser114/ Thr118 was observed in extracts of various cell lines following UV treatment. For additional information please visit PhosphoSitePlus®, CST's modification site knowledgebase, at www.phosphosite.org.				
Background References		1. Jilani, A. et al. (1999) <i>J Biol Chem</i> 274, 24176-86. 2. Dobson, C.J. and Allinson, S.L. (2006) <i>Nucleic Acids Res</i> 34, 2230-7. 3. Bernstein, N.K. et al. (2008) <i>Anticancer Agents Med Chem</i> 8, 358-67.				
Species Reacti	vity	Species reactivity is d	etermined by testin	g in at least one approve	ed 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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