1618

Phospho-Chk1 (Ser317) (D7H2) Rabbit mAh



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Applications: W	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 56	Source/Isotype: Rabbit IgG	UniProt ID: #O14757	Entrez-Gene Id: 1111
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage				5), 150 mM NaCl, 100 μg. oot aliquot the antibody.	/ml BSA, 50% glycei	rol and less than
Specificity/Sensitivity		Phospho-Chk1 (Ser317) (D7H2) Rabbit mAb recognizes endogenous levels of Chk1 protein only when phosphorylated at Ser317.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser317 of human Chk1 protein.				
Background		Chk1 kinase acts downstream of ATM/ATR kinase and plays an important role in DNA damage checkpoint control, embryonic development, and tumor suppression (1). Activation of Chk1 involves phosphorylation at Ser317 and Ser345 by ATM/ATR, followed by autophosphorylation of Ser296. Activation occurs in response to blocked DNA replication and certain forms of genotoxic stress (2). While phosphorylation at Ser345 serves to localize Chk1 to the nucleus following checkpoint activation (3), phosphorylation at Ser317 along with site-specific phosphorylation of PTEN allows for re-entry into the cell cycle following stalled DNA replication (4). Chk1 exerts its checkpoint mechanism on the cell cycle, in part, by regulating the cdc25 family of phosphatases. Chk1 phosphorylation of cdc25A targets it for proteolysis and inhibits its activity through 14-3-3 binding (5). Activated Chk1 can inactivate cdc25C via phosphorylation at Ser216, blocking the activation of cdc2 and transition into mitosis (6). Centrosomal Chk1 has been shown to phosphorylate cdc25B and inhibit its activation of CDK1-cyclin B1, thereby abrogating mitotic spindle formation and chromatin condensation (7). Furthermore, Chk1 plays a role in spindle checkpoint function through regulation of aurora B and BubR1 (8). Research studies have implicated Chk1 as a drug target for cancer therapy as its inhibition leads to cell death in many cancer cell lines (9).				
Background References		 Liu, Q. et al. (2000) Genes Dev 14, 1448-59. Zhao, H. and Piwnica-Worms, H. (2001) Mol Cell Biol 21, 4129-39. Jiang, K. et al. (2003) J Biol Chem 278, 25207-17. Martin, S.A. and Ouchi, T. (2008) Mol Cancer Ther 7, 2509-16. Chen, M.S. et al. (2003) Mol Cell Biol 23, 7488-97. Zeng, Y. et al. (1998) Nature 395, 507-10. Löffler, H. et al. (2006) Cell Cycle 5, 2543-7. Zachos, G. et al. (2007) Dev Cell 12, 247-60. Garber, K. (2005) J Natl Cancer Inst 97, 1026-8. 				
Species Reactivit	:y	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).
Western Blot Buf	ffer	IMPORTANT: For west	ern blots, incubate	membrane with diluted	primary antibody i	n 5% w/v nonfat

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 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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