

Phospho-Chk2 (Ser19) Antibody



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

Applications: W	Reactivity:	Sensitivity: Endogenous	MW (kDa): 62	Source/Isotype: Rabbit	UniProt ID: #O96017	Entrez-Gene Id: 11200
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM soo 20°C. Do not aliquot th		δ), 150 mM NaCl, 100 μg/	ml BSA and 50% gl	ycerol. Store at –
Specificity/Sensitivity		Phospho-Chk2 (Ser19) Antibody detects endogenous levels of Chk2 only when phosphorylated at serine 19. The antibody does not cross-react with Chk2 phosphorylated at other sites.				
Source / Purificat	tion	Polyclonal antibodies corresponding to reside peptide affinity chrom	dues surrounding S	nmunizing animals with a Ser19 of human Chk2. Ar	a synthetic phospho itibodies are purific	opeptide ed by protein A and
Background		Chk2 is the mammalian orthologue of the budding yeast Rad53 and fission yeast Cds1 checkpoint kinases (1-3). The amino-terminal domain of Chk2 contains a series of seven serine or threonine residues (Ser19, Thr26, Ser28, Ser33, Ser35, Ser50, and Thr68) each followed by glutamine (SQ or TQ motif). These are known to be preferred sites for phosphorylation by ATM/ATR kinases (4,5). After DNA damage by ionizing radiation (IR), UV irradiation, or hydroxyurea treatment, Thr68 and other sites in this region become phosphorylated by ATM/ATR (5-7). The SQ/TQ cluster domain, therefore, seems to have a regulatory function. Phosphorylation at Thr68 is a prerequisite for the subsequent activation step, which is attributable to autophosphorylation of Chk2 at residues Thr383 and Thr387 in the activation loop of the kinase domain (8).				
Background References		1. Allen, J.B. et al. (1994) <i>Genes Dev.</i> 8, 2401-2415. 2. Weinert, T.A. et al. (1994) <i>Genes Dev.</i> 8, 652-665. 3. Murakami, H. and Okayama, H. (1995) <i>Nature</i> 374, 817-819. 4. Kastan, M.B. and Lim, D.S. (2000) <i>Nat. Rev. Mol. Cell Biol.</i> 1, 179-186. 5. Matsuoka, S. et al. (2000) <i>Proc. Natl. Acad. Sci. USA</i> 97, 10389-10394. 6. Melchionna, R. et al. (2000) <i>Nat. Cell Biol.</i> 2, 762-765. 7. Ahn, J.Y. et al. (2000) <i>Cancer Res.</i> 60, 5934-5936. 8. Lee, C.H. and Chung, J.H. (2001) <i>J. Biol. Chem.</i> 276, 30537-30541.				
Species Reactivit	v	Species reactivity is de	stermined by testin	g in at least one approve	ed application (e.g.	western hlot)

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

Cross-Reactivity Key H: Human

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