## PLK3 (D14F12) Rabbit mAb



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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

Applications: W	Reactivity: H M	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 80	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q9H4B4	Entrez-Gene Id 1263
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		PLK3 (D14F12) Rabbit mAb detects endogenous levels of total PLK3 protein.				
Species predicted to react based on 100% sequence homology		Rat, Monkey				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Cys625 of human PLK3 protein.				
Background		At least 4 distinct polo-like kinases exist in mammalian cells: PLK1, PLK2, PLK3 and PLK4/SAK (1). Like the other PLK family members, PLK3 contains an amino-terminal catalytic domain and a conserved carboxy-terminal domain termed the Polo box (2). PLK3, also called proliferation-related kinase (Prk) (3), was originally described as a fibroblast growth factor (FGF)-inducible kinase (Fnk) and identified as an immediate-early response gene responsive to FGF-1 and other mitogens (4). PLK3 is a cytokine-inducible serine/threonine kinase whose protein expression is cell cycle regulated. Though its expression is found primarily in G1 phase of the cell cycle, PLK3 is detected in G0 and in late telophase prior to cytokinesis (5). Like the other PLK family members, PLK3 functions mainly as a regulator of the cell cycle. Specifically, PLK3 is required for entry into S phase and is a critical regulator of G1 events, as indicated by RNAi-induced PLK3-depleted cells (2). PLK3 is also involved in the regulation of DNA damage response via phosphorylation of p53 on Ser20 (6). PLK3 may act as a tumor suppressor as Plk3-deficient mice develop spontaneous tumors in various organs (7). Unlike PLK1, PLK3 expression is down regulated in cancers including lung (3), head and neck (8), and colon (9).				
Background References		2. Zimmerman, W.C. a 3. Li, B. et al. (1996) <0 4. Donohue, P.J. et al. 5. Zimmerman, W.C. a 6. Xie, S. et al. (2001) < 7. Yang, Y. et al. (2008 8. Dai, W. et al. (2000)	1998) <cite>Curr Opin Cell Biol</cite> 10, 776-83.  In, W.C. and Erikson, R.L. (2007) <cite>Proc Natl Acad Sci USA</cite> 104, 1847-52.  (1996) <cite>J Biol Chem</cite> 271, 19402-8.  P.J. et al. (1995) <cite>J Biol Chem</cite> 270, 10351-7.  In, W.C. and Erikson, R.L. (2007) <cite>Cell Cycle</cite> 6, 1314-8.  (2001) <cite>J Biol Chem</cite> 276, 43305-12.  al. (2008) <cite>Cancer Res</cite> 68, 4077-85.  al. (2000) <cite>Genes Chromosomes Cancer</cite> 27, 332-6.  al. (2002) <cite>Int J Oncol</cite> 20, 121-6.			
Species Reactiv	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).
W		MARORIANIT E	11.			50/ / BGA ***

**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 M: Mouse

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