1/1/24, 7:39 AM Revision 1

Phospho-GSK-3β (Thr390) Antibody					CHNOLOGY® 877-616-CELL (2355) orders@cellsignal.com
				Support:	877-678-TECH (8324)
#3548				Web:	info@cellsignal.com cellsignal.com
			3 Trask	Lane   Danvers   Ma	ssachusetts   01923   USA
For Research Use Only. Not for		edures.			
Applications: Reacti WB H		<b>MW (kDa):</b> 46	Source: Rabbit	<b>UniProt ID:</b> #P49841	Entrez-Gene Id: 2932
Product Usage	Application			Dilution	
Information	Western Blotting			1:1000	
Storage	Supplied in 10 mM sodi 20°C. Do not aliquot the		), 150 mM NaCl, 10	0 μg/ml BSA and 50%	6 glycerol. Store at –
Specificity / Sensitivity	Phospho-GSK-3β (Thr3 phosphorylated at Thr3		s endogenous level	s of human GSK-3β ι	protein only when
Source / Purification	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr390 of human GSK-3β. Antibodies are purified by peptide affinity chromatography.				
Background	Glycogen synthase kinase-3 (GSK-3) was initially identified as an enzyme that regulates glycogen synthesis in response to insulin (1). GSK-3 is a ubiquitously expressed serine/threonine protein kinase that phosphorylates and inactivates glycogen synthase. GSK-3 is a critical downstream element of the PI3K/Akt cell survival pathway whose activity can be inhibited by Akt-mediated phosphorylation at Ser21 of GSK-3α and Ser9 of GSK-3β (2,3). GSK-3 has been implicated in the regulation of cell fate in <i>Dictyostelium</i> and is a component of the Wnt signaling pathway required for <i>Drosophila, Xenopus</i> , and mammalian development (4). GSK-3 has been shown to regulate cyclin D1 proteolysis and subcellular localization (5). The phosphorylation of GSK-3β at Thr390 was found to be a possible substrate of p38 MAPK and was reported by several labs using phosphoproteomic analysis on mitotic cell extracts (6-10). Phosphorylation of this site was also identified at Cell Signaling Technology (CST) using PhosphoScan <sup>®</sup> , CST's LC-MS/MS platform for modification site discovery (11). Please visit PhosphoSitePlus <sup>®</sup> , CST's modification site knowledgebase, at www.phosphosite.org for more information.				
Background References	<ol> <li>Welsh, G.I. et al. (1996) <i>Trends Cell Biol</i> 6, 274-9.</li> <li>Srivastava, A.K. and Pandey, S.K. (1998) <i>Mol Cell Biochem</i> 182, 135-41.</li> <li>Cross, D.A. et al. (1995) <i>Nature</i> 378, 785-9.</li> <li>Nusse, R. (1997) <i>Cell</i> 89, 321-3.</li> <li>Diehl, J.A. et al. (1998) <i>Genes Dev</i> 12, 3499-511.</li> <li>Thornton, T.M. et al. (2008) <i>Science</i> 320, 667-70.</li> <li>Daub, H. et al. (2008) <i>Mol Cell</i> 31, 438-48.</li> <li>Dephoure, N. et al. (2008) <i>Proc Natl Acad Sci USA</i> 105, 10762-7.</li> <li>Lowery, D.M. et al. (2007) <i>EMBO J</i> 26, 2262-73.</li> <li>Beausoleil, S.A. et al. (2004) <i>Proc Natl Acad Sci USA</i> 101, 12130-5.</li> <li>Rush, J. et al. (2005) <i>Nat Biotechnol</i> 23, 94-101.</li> </ol>				
Species Reactivity	Species reactivity is dete	ermined by testing ir	n at least one appro	ved 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	WB: Western Blotting				
Cross-Reactivity Key	H: human M: mouse R: X: Xenopus Z: zebrafish GP: Guinea Pig Rab: ra	B: bovine Dg: dog	Pg: pig Sc: S. cere		-

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Limited Uses

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