## GSK-3α (D80E6) Rabbit mAb





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Applications: W, IP	<b>Reactivity:</b> H M R Hm Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 51	Source/Isotype: Rabbit IgG	<b>UniProt ID:</b> #P49840	Entrez-Gene Id: 2931		
Product Usage Information	e	<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.						
Specificity/Sensitivity		GSK-3α (D80E6) Rabbit mAb detects endogenous levels of total GSK-3α protein. The antibody does not cross-react with GSK-3β.						
Species predicted to react based on 100% sequence homology		Pig						
Source / Purifi	ication	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the sequence of human GSK-3α protein.						
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). GSK-3α regulates the production of amyloid-β peptides, a major component of the plaques that accumulate with progression of Alzheimer disease. Administration of therapeutic concentrations of lithium, a GSK-3 inhibitor, attenuates amyloid-β production by specifically inhibiting the cleavage of amyloid precursor protein (APP) by γ-secretase, blocking accumulation of amyloid-β peptides in the brains of mice that overproduce APP (6).						
Background R	eferences	<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>Phiel, C.J. et al. (2003) <i>Nature</i> 423, 435-439.</li> </ol>						
Species React	ivity	Species reactivity is det	ermined by testing	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.				ר 5% w/v BSA, 1X		
Applications <b>k</b>	(ey	W: Western Blotting IP: Immunoprecipitation						
Cross-Reactivity Key H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey								
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