GSK-3α (D80D1) Rabbit mAb



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Applications: W, IP, IF-IC	Reactivity: H M R Hm Mk	Sensitivity: Endogenous	MW (kDa): 51	Source/Isotype: Rabbit IgG	UniProt ID: #P49840	Entrez-Gene Id: 2931
Product Usage Information		Application Western Blotting Immunoprecipitation Immunofluorescence	(Immunocytochem	istry)	1 1	ilution :1000 :200 :50 - 1:100
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		GSK-3α (D80D1) Rabbit mAb detects endogenous levels of total GSK-3α protein. The antibody does not cross-react with GSK-3β.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to the sequence of human GSK-3 α .				
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-3 α and Ser9 of GSK-3 β (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-3 α 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 Ro	eferences	2. Srivastava, A.K. and 3. Cross, D.A. et al. (19 4. Nusse, R. (1997) <i>Cel</i> . 5. Diehl, J.A. et al. (199	Nelsh, G.I. et al. (1996) <i>Trends Cell Biol</i> 6, 274-9. Grivastava, A.K. and Pandey, S.K. (1998) <i>Mol Cell Biochem</i> 182, 135-41. Cross, D.A. et al. (1995) <i>Nature</i> 378, 785-9. Nusse, R. (1997) <i>Cell</i> 89, 321-3. Diehl, J.A. et al. (1998) <i>Genes Dev</i> 12, 3499-511. Phiel, C.J. et al. (2003) <i>Nature</i> 423, 435-9.			
Species Reacti	vity	Species reactivity is de	termined by testin	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.

Applications Key

W: Western Blotting IP: Immunoprecipitation IF-IC: Immunofluorescence (Immunocytochemistry)

Cross-Reactivity Key

H: Human M: Mouse R: Rat Hm: Hamster Mk: Monkey

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