Revision 3		
CaMKII-α Antibody	Ce T E	CHNOLOGY*
Store	Orders:	877-616-CELL (2355) orders@cellsignal.com
	Support:	877-678-TECH (8324)
#3357	Web:	info@cellsignal.com cellsignal.com
#	3 Trask Lane   Danvers   Mass	achusetts   01923   USA

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Applications: W	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 50	<b>Source/Isotype:</b> Rabbit	UniProt ID: #Q9UQM7	Entrez-Gene Id 815		
Product Usage Information	!	<b>Application</b> Western Blotting			Dilution 1:1000			
Storage		Supplied in 10 mM sc 20°C. Do not aliquot t		5), 150 mM NaCl, 100 µg.	/ml BSA and 50% gl	ycerol. Store at –		
Specificity/Sensitivity		CaMKII-α Antibody detects endogenous levels of total CaMKII-α. The peptide sequence used as the antigen is not conserved in CaMKII-β, -γ and -δ.						
Source / Purifi	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Tyr436 of human CaMKII-α Antibodies are purified by protein A and peptide affinity chromatography.						
Background		CaMKII is an important member of the calcium/calmodulin-activated protein kinase family, functioning in neural synaptic stimulation and T cell receptor signaling (1,2). CaMKII has catalytic and regulatory domains. Ca <sup>2+</sup> /calmodulin binding to the CaMKII regulatory domain relieves autoinhibition and activates the kinase (3). The activated CaMKII further autophosphorylates at Thr286 to render the kinase constitutively active (3). The threonine phosphorylation state of CaMKII can be regulated through PP1/PKA. PP1 (protein phosphatase 1) dephosphorylates phospho-CaMKII at Thr286. PKA (protein kinase A) prevents phospho-CaMKII (Thr286) dephosphorylation through an inhibitory effect on PP1 (4).						
Background Re	eferences	1. Hughes, K. et al. (2001) <i>J Biol Chem</i> 276, 36008-13. 2. Barria, A. et al. (1997) <i>Science</i> 276, 2042-5. 3. Barkai, U. et al. (2000) <i>Mol Endocrinol</i> 14, 554-63. 4. Makhinson, M. et al. (1999) <i>J Neurosci</i> 19, 2500-10.						
Species Reactiv	vity	Species reactivity is d	etermined by testir	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot B	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 K	ey	W: Western Blotting						
Cross-Reactivit	ty Key	H: Human M: Mouse	R: Rat					
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