Phospho-Tuberin/TSC2 (Ser939) Antibody



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Applications: W	Reactivity: H M	Sensitivity: Endogenous	MW (kDa): 200	Source/Isotype: Rabbit	UniProt ID: #P49815	Entrez-Gene Id: 7249		
Product Usage Information		Application Western Blotting		Dilution 1:1000				
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Stor 20°C. Do not aliquot the antibody.				ycerol. Store at –		
Specificity/Sensitivity		Phospho-Tuberin/TSC2 (Ser939) Antibody detects endogenous levels of tuberin only when phosphorylated at serine 939. This antibody does not cross-react with tuberin phosphorylated at other sites.						
Source / Purifi	PurificationPolyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues around Ser939 of human tuberin. Antibodies are purified by protein A and peptide affinity chromatography.							
Background		Tuberin is a product of the TSC2 tumor suppressor gene and an important regulator of cell proliferation and tumor development (1). Mutations in either <i>TSC2</i> or the related <i>TSC1</i> (hamartin) gene cause tuberous sclerosis complex (TSC), an autosomal dominant disorder characterized by development of multiple, widespread non-malignant tumors (2). Tuberin is directly phosphorylated at Thr1462 by Akt/PKB (3). Phosphorylation at Thr1462 and Tyr1571 regulates tuberin-hamartin complexes and tuberin activity (3-5). In addition, tuberin inhibits the mammalian target of rapamycin (mTOR), which promotes inhibition of p70 S6 kinase, activation of eukaryotic initiation factor 4E binding protein 1 (4E-BP1, an inhibitor of translation initiation), and eventual inhibition of translation (3,6,7). Tuberin is phosphorylated on Ser939 and Thr1462 in response to PI3K activation and that the human TSC complex is a direct biochemical target of the PI3K/Akt pathway (3). This data complements Drosophila genetics studies suggesting the possible involvement of the tuberin-hamartin complex in the PI3K/Akt mediated insulin pathway (8-10).						
Background R	eferences	 Soucek, T. et al. (1998) <i>Proc Natl Acad Sci U S A</i> 95, 15653-8. Sparagana, S.P. and Roach, E.S. (2000) <i>Curr Opin Neurol</i> 13, 115-9. Manning, B.D. et al. (2002) <i>Mol Cell</i> 10, 151-62. Aicher, L.D. et al. (2001) <i>J Biol Chem</i> 276, 21017-21. Dan, H.C. et al. (2002) <i>J Biol Chem</i> 277, 35364-70. Goncharova, E.A. et al. (2002) <i>J Biol Chem</i> 277, 30958-67. Inoki, K. et al. (2002) <i>Nat Cell Biol</i> 4, 648-57. Gao, X. and Pan, D. (2001) <i>Genes Dev</i> 15, 1383-92. Potter, C.J. et al. (2001) <i>Cell</i> 105, 357-68. Tapon, N. et al. (2001) <i>Cell</i> 105, 345-55. 						
Species Reacti	vity	Species reactivity is de	etermined by testing	g in at least one approve	ed application (e.g.,	western blot).		
Western Blot E	Buffer	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.				n 5% w/v nonfat		
Applications K	ey	W: Western Blotting						
Cross-Reactivi	ty Key	H: Human M: Mouse						
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