

TORC1/CRTC1 Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: M	Sensitivity: Endogenous	MW (kDa): 78	Source/Isotype: Rabbit	UniProt ID: #Q6UUV9	Entrez-Gene Id: 23373
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		TORC1/CRTC1 Antibody recognizes endogenous levels of total TORC1 (CRTC1) protein. This antibody does not cross-react with TORC2 and TORC3 proteins.				
Species predicted to react based on 100% sequence homology		Human				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly296 of human TORC1 (CRTC1) protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Glucose homeostasis is regulated by hormones and cellular energy status. Elevations of blood glucose during feeding stimulate insulin release from pancreatic β -cells through a glucose sensing pathway. Feeding also stimulates release of gut hormones such as glucagon-like peptide-1 (GLP-1), which further induces insulin release, inhibits glucagon release and promotes β -cell viability. CREB-dependent transcription likely plays a role in both glucose sensing and GLP-1 signaling (1). The protein CRTC2 (CREB-regulated transcription coactivator 2)/TORC2 (transducer of regulated CREB activity 2) functions as a CREB co-activator (2,3) and is implicated in mediating the effects of these two pathways (4). In quiescent cells, CRTC2/TORC2 is phosphorylated at Ser171 and becomes sequestered in the cytoplasm via an interaction with 14-3-3 proteins. Glucose and gut hormones lead to the dephosphorylation of CRTC2/TORC2 and its dissociation from 14-3-3 proteins. Dephosphorylated CRTC2/TORC2 enters the nucleus to promote CREB-dependent transcription. CRTC2/TORC2 plays a key role in the regulation of hepatic gluconeogenic gene transcription in response to hormonal and energy signals during fasting (5).				
		CRTC1/TORC1, CRTC2/ dependent transcripti at Ser151 in mouse hy	TORC2 and CRTC3/ on of HTLV-1 long t pothalamic cells univator, CRTC1/TORC	RC1 and CRTC3/TORC3 a TORC3 associate with th erminal repeats (6,7). CF nder basal conditions (8) 1 is dephosphorylated a nce and fertility (8).	ne HTLV Tax protein RTC1/TORC1 is highl n. When these cells a	to promote Tax- y phosphorylated are exposed to
Background References		1. Hinke, S.A. et al. (2004) <i>J Physiol</i> 558, 369-80. 2. Conkright, M.D. et al. (2003) <i>Mol Cell</i> 12, 413-23. 3. Iourgenko, V. et al. (2003) <i>Proc Natl Acad Sci U S A</i> 100, 12147-52. 4. Screaton, R.A. et al. (2004) <i>Cell</i> 119, 61-74. 5. Koo, S.H. et al. (2005) <i>Nature</i> 437, 1109-11. 6. Koga, H. et al. (2004) <i>J Biol Chem</i> 279, 52978-83. 7. Siu, Y.T. et al. (2006) <i>J Virol</i> 80, 7052-9. 8. Altarejos, J.Y. et al. (2008) <i>Nat Med</i> 14, 1112-7.				

Species Reactivity

Species reactivity is determined by testing in at least one approved 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

Cross-Reactivity Key M: Mouse

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