

## 3808

## Phospho-Rictor (Thr1135) (D30A3) Rabbit mAh



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

Applications: W	Reactivity: H M Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 200	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q6R327	Entrez-Gene Id: 253260
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		Phospho-Rictor (Thr1135) (D30A3) Rabbit mAb detects endogenous levels of rictor protein only when phosphorylated at Thr1135.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to the sequence surrounding Thr1135 of human Rictor protein.				
Background		Cell growth is a fundamental biological process whereby cells accumulate mass and increase in size. The mammalian TOR (mTOR) pathway regulates growth by coordinating energy and nutrient signals with growth factor-derived signals (1). mTOR is a large protein kinase with two different complexes. One complex contains mTOR, GβL and raptor, which is a target of rapamycin. The other complex, insensitive to rapamycin, includes mTOR, GβL, Sin1, and rictor (1). The mTOR-rictor complex phosphorylates Ser473 of Akt/PKB <i>in vitro</i> (2). This phosphorylation is essential for full Akt/PKB activation. Furthermore, an siRNA knockdown of rictor inhibits Ser473 phosphorylation in 3T3-L1 adipocytes (3). This complex has also been shown to phosphorylate the rapamycin-resistant mutants of S6K1, another effector of mTOR (4). Phosphorylation of Thr1135 on rictor was identified at Cell Signaling Technology (CST) using PhosphoScan®, CST's LC-MS/MS platform for phosphorylation site discovery (5). Additional research indicates that rictor is phosphorylated at Thr1135 by p70 S6K, which negatively regulates mTORC2 protein complex as part of a negative feedback mechanism controlling Akt activity (6-8).				
Background References		<ol> <li>Sarbassov, D.D. et al. (2004) Curr. Biol. 14, 1296-1302.</li> <li>Sarbassov, D.D. et al. (2005) Science 307, 1098-1101.</li> <li>Hresko, R.C. and Mueckler, M. (2005) J. Biol. Chem. 280, 40406-40416.</li> <li>Ali, S.M. and Sabatini, D.M. (2005) J. Biol. Chem. 280, 19445-19448.</li> <li>Rush, J. et al. (2005) Nat Biotechnol 23, 94-101.</li> <li>Dibble, C.C. et al. (2009) Mol Cell Biol 29, 5657-70.</li> <li>Julien, L.A. et al. (2010) Mol Cell Biol 30, 908-21.</li> <li>Treins, C. et al. (2010) Oncogene 29, 1003-16.</li> </ol>				
Species Reactiv	rity	Species reactivity is do	etermined by testir	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				

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TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

**Applications Key** W: Western Blotting

Cross-Reactivity Key H: Human M: Mouse Mk: Monkey

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