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-20°C

#57295

# PhosphoPlus® Rictor (Thr1135) Antibody Duet



Cell Signaling  
TECHNOLOGY®

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**Entrez-Gene ID** #253260  
**UniProt ID** #Q6R327

New 05/18

**For Research Use Only. Not For Use In Diagnostic Procedures.**

Products Included	Product #	Quantity	Mol. Wt.	Isotype
Phospho-Rictor (Thr1135) (D30A3) Rabbit mAb	3806	100 µl	200 kDa	Rabbit IgG
Rictor (53A2) Rabbit mAb	2114	100 µl	200 kDa	Rabbit IgG

See [www.cellsignal.com](http://www.cellsignal.com) for individual component applications, species cross-reactivity, dilutions and additional application protocols.

**Description:** PhosphoPlus® Duets from Cell Signaling Technology (CST) provide a means to assess protein activation status. Each Duet contains an activation-state and total protein antibody to your target of interest. These antibodies have been selected from CST's product offering based upon superior performance in specified applications.

**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 *in vitro* (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).

**Specificity/Sensitivity:** Rictor (53A2) Rabbit mAb detects endogenous levels of total Rictor protein. Phospho-Rictor (Thr1135) (D30A3) Rabbit mAb detects endogenous levels of Rictor protein only when phosphorylated at Thr1135.

**Source/Purification:** Monoclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the sequence surrounding Gln1681 of human Rictor and a synthetic phosphopeptide corresponding to the sequence surrounding Thr1135 of human Rictor protein.

**Storage:** Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. *Do not aliquot the antibody.*

#### Background References:

- (1) Sarbassov, D.D. et al. (2004) *Curr. Biol.* 14, 1296-1302.
- (2) Sarbassov, D.D. et al. (2005) *Science* 307, 1098-1101.
- (3) Hresko, R.C. and Mueckler, M. (2005) *J. Biol. Chem.* 280, 40406-40416.
- (4) Ali, S.M. and Sabatini, D.M. (2005) *J. Biol. Chem.* 280, 19445-19448.
- (5) Rush, J. et al. (2005) *Nat Biotechnol* 23, 94-101.
- (6) Dibble, C.C. et al. (2009) *Mol Cell Biol* 29, 5657-70.
- (7) Julien, L.A. et al. (2010) *Mol Cell Biol* 30, 908-21.
- (8) Treins, C. et al. (2010) *Oncogene* 29, 1003-16.

U.S. Patent No. 7,429,487, foreign equivalents, and child patents deriving therefrom.

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**Applications:** W—Western IP—Immunoprecipitation IHC—Immunohistochemistry ChIP—Chromatin Immunoprecipitation IF—Immunofluorescence F—Flow cytometry E-P—ELISA-Peptide **Species Cross-Reactivity:** H—human M—mouse R—rat Hm—hamster Mk—monkey Mi—mink C—chicken Dm—D. melanogaster X—Xenopus Z—zebrafish B—bovine Dg—dog Pg—pig Sc—S. cerevisiae Ce—C. elegans Hr—Horse All—all species expected **Species** enclosed in parentheses are predicted to react based on 100% homology.