Sin1 (D7G1A) Rabbit mAb

100 μl (10 western blots)



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New 07/13

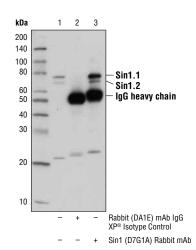
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Applications Species Cross-Reactivity* Molecular Wt. Isotype W. IP H. R. Mk 78. 74 kDa Rabbit IgG** Endogenous

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 that is a component of two different complexes. The mTOR complex 1 (mTORC1), a target of rapamycin, contains mTOR, GβL, and raptor. mTORC2, insensitive to rapamycin, includes mTOR, GBL, Sin1, and rictor (1). The mTORC2 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). mTORC2 has also been shown to phosphorylate the rapamycin-resistant mutants of S6K1, another effector of mTOR (4). In addition, phosphorylation of Sin1 at Thr86 by Akt/PKB was shown to regulate the activity of mTORC2 in adipocytes upon stimulation by growth factors (5).

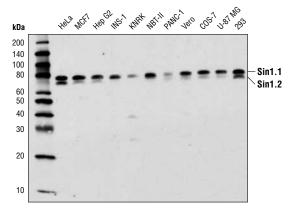
Specificity/Sensitivity: Sin1 (D7G1A) Rabbit mAb recognizes endogenous levels of total Sin1 protein, including isoforms Sin1.1 and Sin1.2.

Source/Purification: Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human Sin1 protein.



Rabbit (DA1E) mAb IgG XP® Isotype Control #3900 (lane 2) or Sin1 (D7G1A) Rabbit mAb (lane 3). Lane 1 is 10% input. Western blot analysis was performed using Sin1 (D7G1A) Rahhit mAh

Immunoprecipitation of Sin1 from Hep G2 cell extracts using



Western blot analysis of extracts from various cell lines using Sin1 (D7G1A)

Entrez Gene ID #79109 UniProt ID #Q9BPZ7

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.

- *Species cross-reactivity is determined by western blot.
- **Anti-rabbit secondary antibodies must be used to detect this antibody.

Recommended Antibody Dilutions:

1:1000 Western blotting 1:50 Immunoprecipitation

For product specific protocols please see the web page for this product at www.cellsignal.com.

Please visit www.cellsignal.com for a complete listing of recommended complementary products.

Background References:

- (1) Sarbassov, D.D. et al. (2004) Curr Biol 14, 1296-302.
- (2) Sarbassov, D.D. et al. (2005) Science 307, 1098-101.
- (3) Hresko, R.C. and Mueckler, M. (2005) J Biol Chem 280,
- (4) Ali, S.M. and Sabatini, D.M. (2005) J Biol Chem 280. 19445-8.
- (5) Humphrey, S.J. et al. (2013) Cell Metab 17, 1009-20.

IMPORTANT: For western blots, incubate membrane with diluted antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween-20 at 4°C with gentle shaking, overnight.