SAMTOR is an S-adenosylmethionine sensor for the mTORC1 pathway


mTOR complex 1 (mTORC1) regulates cell growth and metabolism in response to multiple environmental cues. Nutrients signal via the Rag guanosine triphosphatases (GTPases) to promote the localization of mTORC1 to the lysosomal surface, its site of activation. We identified SAMTOR, a previously uncharacterized protein, which inhibits mTORC1 signaling by interacting with GATOR1, the GTPase activating protein (GAP) for RagA/B. We found that the methyl donor S-adenosylmethionine (SAM) disrupts the SAMTOR-GATOR1 complex by binding directly to SAMTOR with a dissociation constant of approximately 7 μM. In cells, methionine starvation reduces SAM levels below this dissociation constant and promotes the association of SAMTOR with GATOR1, thereby inhibiting mTORC1 signaling in a SAMTOR-dependent fashion. Methionine-induced activation of mTORC1 requires the SAM binding capacity of SAMTOR. Thus, SAMTOR is a SAM sensor that links methionine and one-carbon metabolism to mTORC1 signaling.

The mechanistic target of rapamycin complex 1 (mTORC1) protein kinase is the central component of a pathway that regulates anabolic and catabolic processes in response to environmental signals, including growth factors and nutrients (1–3). Amino acids promote the translocation of mTORC1 to the lysosomal surface, where its activator RhoB resides. This localization depends on the heterodimeric Rag GTPases, which consist of RagA or RagB bound to RagD, and also has nucleotide exchange activity (4, 5). The KICSTOR complex binds GATOR1 and recruits it to the lysosome, and, like GATOR1, is necessary for amino acid starvation to inhibit mTORC1 signaling (6, 7, 8, 9). The KICSTOR complex binds GATOR1 and/or stably expressing a KICSTOR component (Flag-GATOR1), we verified that SAMTOR coimmunoprecipitated with GATOR1 and KICSTOR, and both complexes are negative regulators of mTORC1 signaling.

To search for proteins that bind to GATOR1 or KICSTOR, we mined the BioPlex protein-protein interaction database generated by immunoprecipitation followed by mass spectrometry of more than 5000 proteins stably expressed in human embryonic kidney (HEK)-293T cells (20). This analysis revealed C7orf60, a previously unstudied protein, as a putative interaction partner of all known components of GATOR1 (Depdc5, Nprl3, Nprl2) and KICSTOR (Kaptn, ITFG2, C1orf86, SZT2). For reasons described below, we renamed C7orf60 as SAMTOR. SAMTOR binds SAM with a dissociation constant of approximately 7 μM and is a SAM sensor that links methionine and one-carbon metabolism to mTORC1 signaling. Using an antibody against SAMTOR to probe anti-FLAG immunoprecipitates prepared from cells having endogenously Flag-tagged components of GATOR1 (Depdc5) or GATOR2 (WDR59) or stably expressing a KICSTOR component (Flag-Kaptn), we verified that SAMTOR coimmunoprecipitated with GATOR1 and KICSTOR, but not GATOR2 (Fig. 1A). Moreover, transiently expressed SAMTOR coimmunoprecipitated with endogenous GATOR1 and KICSTOR, as detected by the presence of their Nprl3 and SZT2 components, respectively. Loss of a component of GATOR1 or KICSTOR, but not of GATOR2, severely reduced the interaction of SAMTOR with KICSTOR or GATOR1, respectively (Fig. 1B). Furthermore, overexpressed GATOR2 coimmunoprecipitated with SAMTOR only when KICSTOR was coexpressed (fig. S1A).

Thus, SAMTOR binds to the supercomplex of GATOR1 and KICSTOR, and both complexes are required for the interaction to occur (Fig. 1C).

Orthologs of SAMTOR are encoded in the genomes of vertebrates and some invertebrates, such as Drosophila melanogaster. We could not identify SAMTOR orthologs in Caenorhabditis elegans or Saccharomyces cerevisiae (Fig. 1D).

To determine whether SAMTOR regulates mTORC1 signaling, we overexpressed SAMTOR in HEK-293T cells and monitored the phosphorylation of Thr389 of S6 kinase 1 (S6K1), a canonical mTORC1 substrate. SAMTOR expression suppressed mTORC1 signaling in a dose-dependent fashion (Fig. 2A), establishing SAMTOR as a negative regulator of the pathway. Amino acids activate mTORC1 by promoting its localization to the lysosomal surface (4, 8). Consistent with SAMTOR inhibiting the amino acid sensing pathway upstream of mTORC1, overexpression of green fluorescent protein (GFP)-tagged SAMTOR displaced mTOR from lysosomes to an extent similar to that seen with GFP-Sestrin2, an inhibitor of GATOR2 (21, 22) (Fig. 2B).

To position the SAMTOR function within the mTORC1 pathway, we performed epistasis experiments with established mTORC1 regulators. Overexpression of SAMTOR inhibited mTORC1 signaling when coexpressed with the wild-type RagA and RagC heterodimer, but not with the constitutively active mutant heterodimer (RagA Q66L and RagC S75N) that bypasses the requirement for amino acids for maintaining mTORC1 activity (Fig. 2C) (4, 5). In addition, SAMTOR did not inhibit mTORC1 signaling in cells lacking either a GATOR1 or KICSTOR component. Thus, SAMTOR acts upstream of the Rag GTPases and requires GATOR1 and KICSTOR to inhibit mTORC1 signaling (Fig. 2D). In combination with the interaction data, these results are consistent with SAMTOR promoting the function of GATOR1 and/or KICSTOR, which are both negative regulators of mTORC1 signaling.

Sequence analyses predict that SAMTOR contains a class I Rossmann fold methyltransferase domain (Fig. 3A and fig. S2) (23). These domains are known to bind S-adenosylmethionine (SAM) and exist in methyltransferases in bacteria, archaee, and eukaryotes (24). To determine whether SAMTOR binds SAM, we developed an equilibrium binding assay based on one we used to detect the binding of leucine to Sestrin2 (16) and determined that SAMTOR binds SAM with a dissociation constant of approximately 7 μM (Fig. 3B). A competition binding assay revealed that, as with other SAM-binding proteins, SAMTOR can also bind S-adenosylhomocysteine (SAH), the demethylated form of SAM (Fig. 3B).

Given these findings, we asked whether SAM and SAH regulate the interaction of SAMTOR and GATOR1-KICSTOR. Indeed, SAM and SAH, but not methionine, homocysteine, adenosine, 5-methylthioadenosine, leucine, or isoleucine, disrupted the interaction when added directly to the immunopurified complex kept at 4°C (Fig. 3, C and D). Thus, SAM disrupts the interaction between SAMTOR and GATOR1-KICSTOR analogously to how leucine and arginine induce the

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release of Sestrin2 and CASTOR1 from GATOR2, respectively (16, 19). Given that SAH has the same effect, it is unlikely that a methylation event is required for SAM to dissociate SAMTOR from GATOR1-KICSTOR.

Mutagenesis of highly conserved residues in human SAMTOR yielded two mutants, Gly172 → Ala (G172A) and Asp190 → Ala (D190A), that no longer bound SAM (Fig. 3E and fig. S2, A and B). These mutants coimmunoprecipitated greater amounts of endogenous GATOR1 and KICSTOR. In contrast, SAMTOR does not have the same effect, it is unlikely that a methylation event is required for SAM to dissociate SAMTOR from GATOR1-KICSTOR. Consistent with the effects of SAM on the interaction between SAMTOR and GATOR1-KICSTOR, we hypothesized that methionine starvation promotes the dissociation of SAMTOR from GATOR1-KICSTOR, and the interaction requires both GATOR1 and KICSTOR, but not SAMTOR. FLAG immunoprecipitates were prepared from wild-type, Nprl3-deficient, and WDR24-deficient HEK-293T cells transiently expressing the indicated cDNAs. FLAG immunoprecipitates and lysates were analyzed as in (A). (C) Model showing how SAMTOR interacts with GATOR1 and KICSTOR. (D) Presence or absence of gene orthologs of SAMTOR in several model organisms.
hibitory effects of methionine restriction on mTORC1 signaling (Fig. 4H), indicating that it of SAM to methionine-starved cells reactivated its capacity to bind SAM. Moreover, the addition for the mTORC1 pathway to detect changes in Because MAT2A is essential in human cells, that the loss of methionine adenosyltransferase (MAT2A) would prevent mTORC1 from sensing methionine while leaving its activation by SAM largely intact (Fig. 4I).

Several properties of SAMTOR suggest that it functions as a SAM sensor that signals methionine deficiency to mTORC1 (Fig. 4I): (i) SAMTOR binds SAM with an affinity that is compatible with the drop in intracellular SAM concentrations caused by methionine starvation, (ii) SAMTOR is required for methionine starvation to inhibit mTORC1 signaling, and (iii) SAMTOR mutants that do not bind SAM cannot signal methionine deficiency to mTORC1. Because SAM levels can be affected by the availability of folate, betaine, and vitamin B_{12}, SAMTOR may also link mTORC1 signaling to the availability of these metabolites.

The Rag GTPase pathway senses and integrates the presence of multiple amino acids upstream of mTORC1 (4, 8). Sestrin1 and Sestrin2 detect leucine, whereas CASTOR1 and SLC38A9 sense cytosolic and lysosomal arginine, respectively (16, 19). In contrast to the Sestrins and CASTOR1, which bind to GATOR2, SAMTOR interacts with GATOR1-KICSTOR. Our genetic data suggest that SAMTOR potentiates GATOR1 function through an unknown mechanism that may involve disruption of the binding of GATOR1 to GATOR2. The interaction between SAMTOR and GATOR1 requires KICSTOR, which may reflect either a composite binding site or the requirement for KICSTOR to localize GATOR1 to the lysosomal surface. In addition, structural information will be needed if we are to understand how the binding of SAM to SAMTOR disrupts its inter- action with GATOR1 and KICSTOR.

Unlike leucine and arginine, which directly bind sensors upstream of mTORC1, methionine

**Fig. 2. SAMTOR is a negative regulator of mTORC1 signaling that acts upstream of the Rag GTPases, GATOR1, and KICSTOR.** (A) Transient overexpression of SAMTOR inhibits mTORC1 signaling. FLAG immunoprecipitates were prepared from HEK-293T cells transfected with 2 ng of FLAG-S6K1 cDNA along with either hemagglutinin (HA)-tagged metap2 cDNA or increasing amounts of HA-SAMTOR cDNA. FLAG immunoprecipitates and cell lysates were analyzed by immunoblotting for the phosphorylation states and levels of the indicated proteins. (B) Overexpression of GFP-SAMTOR displaces mTOR from lysosomes, similar to the effect of GFP-Sestrin2. Wild-type HEK-293T cells transiently expressing GFP-metap2, GFP-SAMTOR, or GFP-Sestrin2 were processed for immunofluorescence detection of mTOR and the lysosomal marker LAMP2. In all images, insets represent selected fields magnified 5.12× as well as their overlays. Scale bar, 10 μm. (C) SAMTOR functions upstream of the Rag GTPases to regulate the mTORC1 pathway. HEK-293T cells expressing the indicated cDNAs were starved of amino acids for 50 min or starved and restimulated with amino acids for 10 min. FLAG immunoprecipitates and cell lysates were analyzed as in (A). (D) SAMTOR functions upstream of GATOR1 and KICSTOR. FLAG immunoprecipitates and cell lysates prepared from wild-type, Nprl3-deficient, or SZT2-deficient HEK-293T cell lines expressing the indicated cDNAs were analyzed as in (A).
Fig. 3. S-adenosylmethionine binds SAMTOR to disrupt its interaction with GATOR1 and KICSTOR. (A) Schematic of the human SAMTOR protein indicating the class I Rossmann fold methyltransferase domain. Shown is an alignment of partial sequences of this domain from SAMTOR in indicated species. Amino acid positions are colored from white to blue in order of increasing sequence similarity. Orange dots denote the Gly172 and Asp190 residues of human SAMTOR. (B) SAMTOR binds SAM and SAH. Purified FLAG-SAMTOR protein was analyzed by SDS–polyacrylamide gel electrophoresis followed by Coomassie blue staining. Binding assays were performed with purified FLAG-SAMTOR incubated with the indicated concentrations of [H]SAM, unlabeled SAM, or SAH. Values for each point are means ± SD of three technical replicates from one representative experiment. The experiment was performed twice. (C) SAM and SAH disrupt the interaction of SAMTOR with GATOR1 in vitro. FLAG immunoprecipitates were prepared from endogenously FLAG-tagged Depdc5 HEK-293T cells. SAM and SAH were added directly to the immunoprecipitates at the indicated concentrations. FLAG immunoprecipitates and cell lysates were analyzed by immunoblotting for the levels of the indicated proteins. (D) The interaction between SAMTOR and GATOR1 is disrupted by 100 μM SAM or SAH, but not by 1 mM methionine, homocysteine, adenosine, 5-methylthioadenosine, leucine, or isoleucine. The experiment was performed and analyzed as in (C). (E) Wild-type HA-SAMTOR, but not HA-SAMTOR G172A or D190A, binds SAM. HA-tagged wild-type and mutant SAMTOR proteins were prepared from HEK-293T cells expressing the indicated cDNAs, and binding assays were performed as in (B). A representative experiment is shown; values are means ± SD of three technical replicates. Two-tailed t tests were used for comparisons between two groups. *P < 0.001; ns, not significant. The experiment was repeated three times. (F) HA-SAMTOR G172A and D190A coimmunoprecipitate more endogenous GATOR1 and KICSTOR than does wild-type SAMTOR, and the interactions are insensitive to SAM added in vitro. HA immunoprecipitates and cell lysates were prepared from HEK-293T cells transiently expressing wild-type HA-SAMTOR or its mutants G172A or D190A, SAM was added to the immunoprecipitates where indicated. HA immunoprecipitates and cell lysates were analyzed as in (C). (G) HA-SAMTOR G172A and D190A inhibit mTORC1 activity to similar extents as wild-type SAMTOR. FLAG immunoprecipitates were prepared from HEK-293T cells transfected with the indicated cDNAs. FLAG immunoprecipitates and cell lysates were analyzed by immunoblotting for the phosphorylation states and levels of the indicated proteins.
Fig. 4. SAMTOR senses SAM to signal methionine sufficiency to mTORC1.

(A) HEK-293T cells were incubated with or without methionine for 2 hours before sample preparation for liquid chromatography/mass spectrometry (LC/MS)–based measurements of the absolute amounts of the indicated metabolites. The dissociation constant $K_d$ of SAMTOR for SAM is indicated. (B) Methionine starvation increases the interaction between SAMTOR and GATOR1. HEK-293T cells transiently expressing HA-tagged metap2 or SAMTOR were kept in growth medium (RPMI) or starved of methionine for 2 hours (~Met) and then restimulated for 20 min with 100 mM methionine (+Met) or 1 mM SAM(+SAM). HA immunoprecipitates and cell lysates were analyzed by immunoblotting for the levels of the indicated proteins. (C) In SAMTOR-depleted cells, the mTORC1 pathway is resistant to methionine starvation. HEK-293T cells stably coexpressing Cas9 and the indicated guides were incubated in media with or without methionine for 2 hours. Cell lysates were analyzed by immunoblotting for the phosphorylation states and the levels of the indicated proteins. (D) The loss of SAMTOR does not affect the sensitivity of the mTORC1 pathway to leucine or arginine starvation. SAMTOR-deficient HEK-293T cells with or without FLAG-SAMTOR expression were starved of the indicated amino acid for 2 hours. Cell lysates were analyzed as in (C). (E) In cells without SAMTOR, mTOR colocalizes with lysosomes even upon methionine starvation. SAMTOR-deficient or control HEK-293T cells were treated as indicated for 2 hours before processing for immunofluorescence detection of mTOR and the lysosomal marker LAMP2. In all images, insets represent selected fields magnified 3.07× as well as their overlays. Scale bar, 10 μm. (F) Reexpression in SAMTOR-null cells of wild-type SAMTOR, but not the SAM-binding G172A mutant of SAMTOR, restored the capacity of the mTORC1 pathway to sense methionine sufficiency. SAMTOR-null cells were transfected with the indicated cDNAs and the cells were treated as in (C) before preparing lysates and FLAG immunoprecipitates. FLAG immunoprecipitates and cell lysates were analyzed as in (C). (G) In Drosophila S2R+ cells depleted of dSamtor or dSesn, the dTOR pathway is resistant to methionine or leucine starvation, respectively. S2R+ cells were transfected with dsRNAs targeting the indicated mRNAs and starved of the indicated amino acids for 1 hour. Cell lysates were analyzed as in (C). (H) Acute loss of MAT2A using a doxycycline-suppressible (dox-off) system attenuates the capacity of mTORC1 to sense methionine but leaves SAM signaling largely intact. MAT2A dox-off HEK-293T cells were treated with doxycycline (30 ng/ml) for 50 hours before starving them as in (C). Cell lysates were analyzed as in (C). (I) Model depicting how SAM sensing by SAMTOR signals methionine levels to mTORC1. Substrates receiving a methyl group from SAM include DNA, RNA, proteins, and phospholipids. N5-me-THF, N5-methyl-tetrahydrofolate; DMG, dimethylglycine; Pi, inorganic phosphate; PPI, pyrophosphate.
is sensed indirectly through SAM. SAM is a central metabolite required for most methylation reactions, including that of DNA (37), histones (25, 30), and phospholipids (32), and our work highlights its additional role as a signaling molecule. Whereas S. cerevisiae does not have a SAMTOR homolog, the yeast TOR pathway does sense methionine through the regulated methylation of the PP2A family of phosphatases (33).

In metazoans, the mTORC1 pathway senses multiple amino acids, which suggests that these nutrients were, at times, scarce during their evolution. Two inferences can be drawn from the existence of SAMTOR: (i) SAM can become limiting in certain nutritional states, and (ii) modulation of mTORC1 under these conditions is beneficial for maintaining organismal homeostasis. Indeed, diets low in methionine reduce tissue SAM levels, improve insulin sensitivity, and extend lifespan in mice and rats (34–38). It is intriguing to speculate that these benefits might be mediated in part via the SAMTOR-dependent inhibition of mTORC1, which is well appreciated for its impact on glucose metabolism and the aging process (7). Given that SAMTOR has a SAM-binding pocket, it may be possible to modulate SAMTOR function pharmacologically.

REFERENCES AND NOTES

15. G. Wyant et al., Cell, in press.

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SUPPLEMENTARY MATERIALS

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Materials and Methods

Supplementary Text

Figs. S1 to S4

References (39–43)

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SAMTOR is an S-adenosylmethionine sensor for the mTORC1 pathway

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SAMTOR joins the family
The amino acid methionine is widely appreciated to have interesting effects on animal physiology. Diets low in methionine increase longevity and overall health, particularly glucose homeostasis. Gu et al. describe a potential molecular link between the effects of methionine restriction and the growth controller mTOR complex 1 (mTORC1), a well-validated regulator of life span and health span in many organisms. They identify a protein that they named SAMTOR as a component of the nutrient-sensing pathway upstream of mTORC1. SAMTOR directly binds S-adenosylmethionine (SAM), a metabolite made from methionine, and is necessary for regulating mTORC1 in response to methionine.

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