

Regulation of TORC1 in Response to Amino Acid Starvation via Lysosomal Recruitment of TSC2

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SUMMARY

TOR complex 1 (TORC1) is a potent anabolic regulator of cellular growth and metabolism. When cells have sufficient amino acids, TORC1 is active due to its lysosomal localization mediated via the Rag GTPases. Upon amino acid removal, the Rag GTPases release TORC1, causing it to become cytoplasmic and inactive. We show here that, upon amino acid removal, the Rag GTPases also recruit TSC2 to the lysosome, where it can act on Rheb. Only when both the Rag GTPases and Rheb are inactive is TORC1 fully released from the lysosome. Upon amino acid withdrawal, cells lacking TSC2 fail to completely release TORC1 from the lysosome, fail to completely inactivate TORC1, and fail to adjust physiologically to amino acid starvation. These data suggest that regulation of TSC2 subcellular localization may be a general mechanism to control its activity and place TSC2 in the amino-acid-sensing pathway to TORC1.

INTRODUCTION

Small GTPases act as molecular switches that alternate between GTP-bound and GDP-bound states, thereby regulating a vast array of cellular parameters, including mitochondrial activity, cell growth, cell metabolism, and cell morphology. Small GTPases are activated or inactivated by guanine nucleotide exchange factors (GEFs) and GTPase-activating proteins (GAPs), respectively, which regulate the GDP/GTP load of the GTPase. Hence, understanding how GEFs and GAPs are regulated is an important aspect of understanding GTPase function. Compared to the regulation of GEFs, relatively little is known about how the activity of GAPs is regulated (Cherfils and Zeghouf, 2013). GAPs acting on members of the Rho and Arf GTPase superfamilies are activated via membrane recruitment, causing rearrangements in the structure of the GAPs upon membrane binding (Cherfils and Zeghouf, 2013). Regulation of GAPs acting on members of the Ras superfamily of GTPases is less well understood. One such GAP is composed of the TSC1/TSC2/TBC1D7 trimeric tumor suppressor complex (Dibble

et al., 2012), which acts on the small GTPase Rheb. Although it is known that activity of this complex is regulated by phosphorylation on multiple sites, it is not yet clear how these phosphorylations affect TSC1/2 activity at the molecular level.

The kinase TOR complex 1 (TORC1) is a potent anabolic regulator of cellular growth and metabolism that is often hyperactivated in human cancers (Guertin and Sabatini, 2007; Heitman et al., 1991; Huang and Manning, 2008; Laplante and Sabatini, 2012; Proud, 2011). To be active, TORC1 needs to bind a molecule of Rheb in the active, GTP-bound state (Inoki et al., 2003; Tee et al., 2003). By inactivating Rheb, the TSC1/2 complex is therefore a critical upstream inhibitor of TORC1. The TSC1/2 complex acts as a central point of integration of almost all known inputs regulating TORC1, including cellular stresses such as low oxygen or low ATP, and various growth-promoting signals, such as PI3K, Ras, TNF, and Wnt signaling (reviewed in Huang and Manning, 2008). The importance of the tuberous sclerosis complex (TSC) on TORC1 signaling and growth is highlighted by the fact that TSC2-inactivating mutations have been found in various human growth-related diseases (Huang and Manning, 2008). One other important input regulating TORC1 activity is the availability of amino acids (Blommaert et al., 1995; Efeyan et al., 2012; Hara et al., 1998; Jewell et al., 2013). Whether TSC2 is also involved in regulating TORC1 in response to amino acids, however, is unclear because various studies have come to differing conclusions (Gao et al., 2002; Roccio et al., 2006; Smith et al., 2005).

Unlike all the other inputs that regulate TORC1 via TSC1/2 and Rheb, amino acids regulate TORC1 via a separate set of small GTPases, the Rag GTPases (Kim et al., 2008; Sancak et al., 2008). The Rag GTPases form heterodimeric complexes consisting of RagA or RagB bound to RagC or RagD. These complexes are stably anchored to lysosomal membranes via the LAMTOR/Ragulator complex (Sancak et al., 2010). In the presence of amino acids, the Rag dimers are in an “active” conformation with RagA or RagB bound to GTP and RagC or RagD bound to GDP. The active Rag dimers recruit TORC1 to the lysosomal surface, where it binds Rheb to form an active holoenzyme. In the absence of amino acids, the Rag GAP complex termed GATOR1 causes the Rag dimers to switch into an inactive conformation containing GDP-bound RagA/B, thereby releasing TORC1 from the lysosomal surface (Bar-Peled et al., 2013; Panchaud et al., 2013). This causes TORC1 to become inactive, presumably because it no longer binds active Rheb on the

lysosomal surface. Hence, TORC1 activation can currently be viewed as consisting of two aspects—the activation of Rheb in response to a plethora of regulatory inputs and the localization of TORC1 to lysosomal membranes in response to amino acids, which allows it to meet Rheb.

In this study, we uncover subcellular localization as a mechanism regulating activity of the TSC1/2 GAP complex. We find that, upon amino acid removal, TSC1/2 is recruited to lysosomes via binding to the Rag proteins, thereby bringing TSC1/2 in close proximity to its target, Rheb. This suggests that relocalization of GAPs to the vicinity of their substrates is one mechanism for their regulation. Unexpectedly, we find that regulation of Rheb by TSC1/2 upon amino acid starvation is required for TORC1 to be released from lysosomal membranes. This suggests a “dual anchoring” mechanism of TORC1 at the lysosome, perhaps with the Rag proteins playing a crucial role in recruiting TORC1 to the lysosomal membrane and Rheb helping to retain it there. We find that cells lacking TSC2 are impaired in their response to amino acid starvation, failing to efficiently turn off TORC1. As a result, cells lacking TSC2 are very sensitive to amino acid starvation and die under conditions that control cells can cope with. In sum, our data indicate that the TSC1/2 complex is responsive to amino acid starvation and participates in amino acid signaling to TORC1. Hence, the TSC1/2 complex appears to play a role in regulating TORC1 in response to all regulatory inputs known to date.

RESULTS

Rag GTPases Bind TSC2

While studying regulation of TORC1 by the Rag GTPases, we noticed that *Drosophila* cells with reduced Rag protein levels are unable to completely shut off TORC1 in the absence of amino acids. When treated with medium lacking amino acids, control S2 cells shut off TORC1 activity, assayed via S6 kinase (S6K) phosphorylation, to roughly 7% the level of fully fed cells (Figure 1A). In contrast, S2 cells with RagA or RagC knockdown are significantly impaired in their response to amino acid removal, retaining circa 50% the TORC1 activity levels of fully fed cells (Figure 1A). In this and all subsequent experiments, amino acid removal is performed in the presence of dialyzed serum, thereby specifically removing amino acids, but not growth factors, from cell culture media. Furthermore, immunoblot quantifications are performed on a LI-COR imaging system, providing a means to quantitatively study TORC1 activity (see [Experimental Procedures](#)). This phenotype was reiterated with independent dsRNAs targeting nonoverlapping regions of RagA and RagC (Figure S1A available online), proving specificity of the phenotype. Furthermore, knockdown of LAMTOR3 also led to similar, impaired TORC1 inactivation upon amino acid withdrawal (data not shown). Similar effects can also be observed in human HEK293FT cells (Figure S1C) and in previous reports (see Figure 3H in [Sancak et al., 2008](#)). As previously shown ([Kim et al., 2008](#); [Sancak et al., 2008](#)), S2 cells and HEK293FT cells with Rag protein knockdown also show severely compromised reactivation of TORC1 upon amino acid readdition (Figures S1B and S1D). Together, these data indicate that the Rag GTPases not only activate TORC1 in the presence of amino

acids but also actively repress TORC1 in the absence of amino acids. Since the Rag proteins “let go” of TORC1 in the absence of amino acids ([Sancak et al., 2008](#)), it is not easy to explain how they could also be actively repressing TORC1. Thus, the Rag GTPases appear to have an additional activity besides their ability to reversibly bind TORC1.

We hypothesized that the Rag proteins might be recruiting inhibitory factors to the lysosome upon amino acid starvation. To investigate this, we immunoprecipitated FLAG-RagA and FLAG-RagC from *Drosophila* S2 cells and performed shotgun mass spectrometry analysis to identify interacting partners. Among the identified proteins, as expected, were known components of the Regulator complex ([Sancak et al., 2010](#)) such as p14, p18, and MP1, as well as TOR and Raptor (Figure 1B). This analysis also found significant amounts of Tsc2 as a Rag-binding protein (Figure 1B). One possibility could be that binding of Tsc2 to the Rag GTPases is indirect due to the Rag GTPases binding TORC1, which binds Rheb, which binds Tsc2. However, as shown below, binding between Tsc2 and the Rag GTPases increases when the Rag GTPases are in the inactive state and, hence, bind less TORC1, arguing against this possible explanation. Because Tsc2 is a negative component of the TOR signaling pathway, we decided to study this interaction in more detail.

We first aimed to confirm the interaction between Tsc2 and the Rag GTPases by coimmunoprecipitation (coIP). Indeed, FLAG-tagged *Drosophila* RagA and RagC were able to coIP epitope-tagged Tsc2, but not an unrelated protein, Medea (Figure 1C). Likewise, epitope-tagged human TSC2 was able to coIP human Rag GTPases in HEK293FT cells (Figure 1D). As discussed below, a complex consisting of human RagA and RagC can also coIP endogenous TSC2 (Figure 2E). In sum, the interaction between the Rag GTPases and TSC2 appears to be specific and evolutionarily conserved from flies to humans.

Delineation of Interacting Regions of TSC2 and the Rag Proteins

We next aimed to characterize in more detail the binding between TSC1/2 and the Rag complex. We first asked which component of the Rag complex is binding TSC1/2. Immunoprecipitation of tagged TSC2 showed that it binds RagA significantly more strongly than the other Rag proteins (Figure 1D), suggesting that RagA is a likely binding partner for the TSC1/2 complex. To study the residues in RagA involved in TSC2 binding, we exploited the fact that RagA has significantly stronger binding to TSC2 compared to RagB despite the two proteins being almost identical. Compared to RagB, RagA is lacking an N-terminal extension of 33 amino acids and has 5 amino acid substitutions at the C terminus of the protein (Figure 2A). We asked which of these differences is important for RagA binding to TSC2. Either removal of the N-terminal extension of RagB or introduction of the five C-terminal amino acid changes into RagB caused improved binding to TSC2 (Figures 2A and 2B), indicating that both the N-terminal and the C-terminal residues affect TSC2 binding. We then introduced the five RagA-specific amino acid substitutions individually into RagB and found that each of them improves the binding of RagB to TSC2 (Figures 2A and 2C). Hence, each of these five amino acids of RagA contributes toward TSC2 binding, and, indeed, these five amino

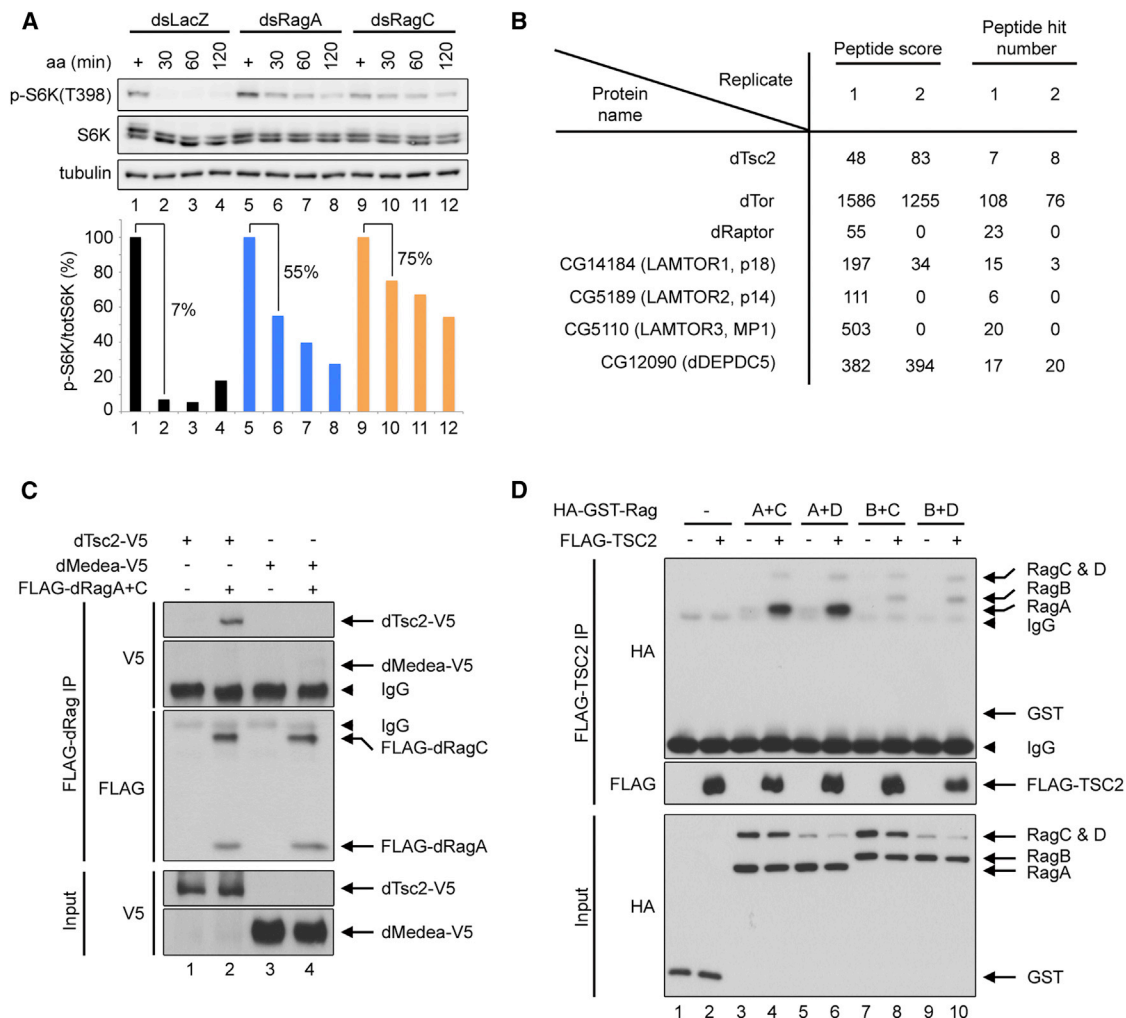


Figure 1. Conserved Binding of TSC2 to the Rag GTPase Complex

(A) *Drosophila* Rag proteins are required for complete TORC1 inactivation upon amino acid withdrawal. Immunoblots of S2 cells treated with dsRNAs against LacZ (control), RagA, or RagC incubated with media lacking only amino acids in the presence of dialyzed serum for the indicated times.

(B) Shotgun mass spectrometry analysis identifies Tsc2 as a Rag-binding protein. Two statistics of peptide abundance are shown for Tsc2 and control proteins from two independent FLAG-RagA + FLAG-RagC immunoprecipitates (IP) from S2 cells.

(C) Confirmation by colP that Rag proteins bind Tsc2, but not an unrelated protein, Medea. Immunoblots of anti-FLAG IPs from *Drosophila* S2 cells expressing FLAG-tagged RagA + RagC and V5-tagged Tsc2 or Medea, as indicated.

(D) Human TSC2 binds human Rag dimeric complexes. Immunoblots of anti-FLAG IPs from HEK293FT cells expressing FLAG-tagged TSC2 and HA-GST-tagged Rag proteins or an HA-GST control, as indicated.

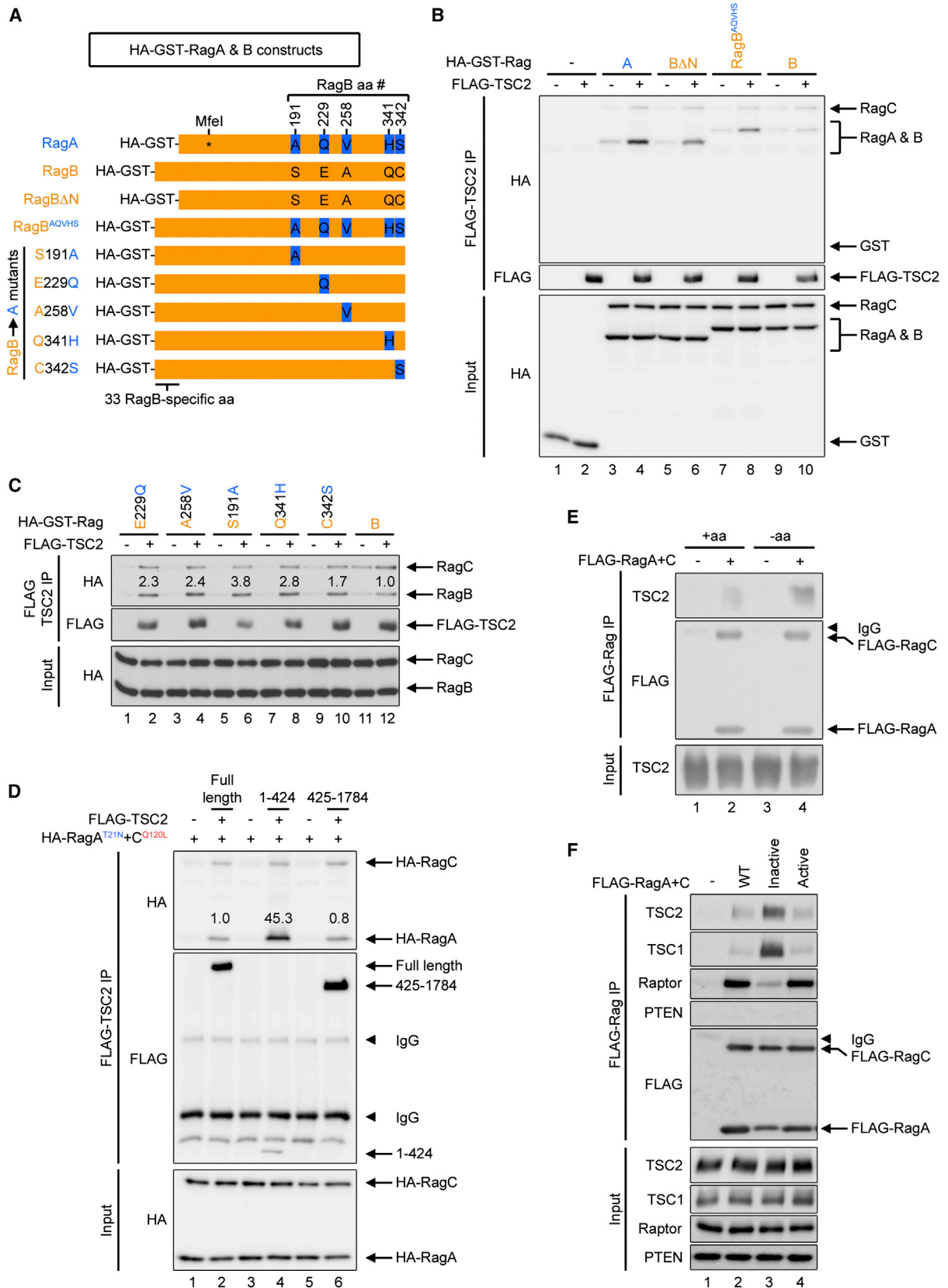
acid differences between RagA and RagB are evolutionarily conserved.

To identify the binding partner in the TSC1/2 complex, we colP ed epitope-tagged TSC1 or TSC2 individually with RagA+C, revealing that TSC2, but not TSC1, binds the Rag complex (Figure S2A). To study which residues in TSC2 are responsible for Rag binding, we generated a series of C-terminal truncations of TSC2 and tested them by colP (Figure S2B). This analysis revealed that the N-terminal 424 amino acids of TSC2 bind very strongly the Rag proteins but that, nonetheless, the remaining amino acids 425–1,784 also have Rag binding capacity (Figure 2D). In sum, binding between the two complexes appears to be mediated via TSC2 binding to RagA. Although the interaction

surfaces are complex, binding depends strongly on five individual amino acids in RagA identified in this analysis.

Binding between Rag Proteins and TSC2 Increases upon Amino Acid Removal

We asked whether binding between the Rag proteins and TSC2 depends on cellular amino acid signaling because our initial findings suggested that the Rag complex might be recruiting a TOR inhibitor upon amino acid removal. Indeed, a complex consisting of human RagA and RagC ("RagA+C") was able to colP endogenous TSC2 (lanes 1 and 2, Figure 2E), and this interaction became stronger upon amino acid removal (lanes 2 versus 4, Figure 2E). One mechanistic explanation could be



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that binding between TSC2 and the Rag GTPases depends on the state of activation of the Rag proteins, which switch to the “inactive” RagA^{low-nucleotide}/RagC^{GTP} conformation upon amino acid removal. To test this, we assayed binding between endogenous TSC2 and mutant variants of the RagA/RagC complex, locked into either the active (RagA^{GTP}/RagC^{low-nucleotide}) or inactive (RagA^{low-nucleotide}/RagC^{GTP}) states (Sancak et al., 2008), and found that, indeed, TSC2 binds the Rag proteins much more strongly when they are in the inactive state (Figure 2F). Binding of endogenous TSC1 to the Rag proteins closely paralleled binding of TSC2 (Figure 2F), indicating that TSC2 is mediating binding of the entire TSC1/2 complex because TSC1 cannot bind the Rag proteins by itself (Figure S2A). As expected (Sancak et al., 2008), Raptor binding was strongest when the Rag proteins were in the active state (Figure 2F), therefore anticorrelating with TSC2 binding. As a negative control, endogenous PTEN did not show any binding to the Rag proteins (Figure 2F). This preference for binding the inactive Rag complex cannot be ascribed to the state of activation of any one of the two Rags but rather appears to depend on the conformation of both RagA and RagC because states of intermediate activation (RagA^{GTP}/RagC^{GTP} or RagA^{low-nucleotide}/RagC^{low-nucleotide}) also showed intermediate levels of TSC2 binding (Figure S2C). In sum, binding of TSC2 to the Rag complex appears to anticorrelate with binding of mTORC1 to the Rag complex, with mTORC1 binding predominantly when the Rag complex is active in the presence of amino acids and TSC2 binding predominantly when the Rag complex is inactive in the absence of amino acids.

TSC2 Is Recruited to the Lysosome in a Rag-Dependent Manner upon Amino Acid Removal

Preferential binding of TSC2 for the inactive Rag complex, which is localized on the lysosome, raised the possibility that TSC2 might be recruited to lysosomes upon amino acid removal. Using an antibody that specifically recognizes endogenous TSC2 (Dibble et al., 2012 and Figure S3A), we found that, in the presence of amino acids, TSC2 is diffusely cytoplasmic in HEK293FT cells in a pattern that is not concentrated on lysosomes/late endosomes (marked by LAMP2, henceforward referred to as “lysosomes”) relative to elsewhere in the cytoplasm (+aa, Figure 3A). Upon removal of amino acids, however, TSC2 quickly accumulates on lysosomes (15 min –aa, Figure 3A) and remains associated with lysosomes for hours (240 min –aa, Figure 3A). Upon

readdition of amino acids, TSC2 quickly becomes diffusely cytoplasmic again (Figure 3B). Similar relocalization of TSC2 to lysosomes could also be observed in mouse embryonic fibroblasts (MEFs) (Figure 3C), and it was confirmed at the ultrastructural level by anti-TSC2 immunoelectron microscopy (Figure S3B). Furthermore, TSC1 also relocalizes to lysosomes upon amino acid removal, suggesting that the whole TSC1/2 complex relocalizes (Figure S3C).

In agreement with the biochemical data presented above, lysosomal accumulation of TSC2 upon amino acid withdrawal is mediated via the Rag proteins because it is strongly blunted upon amino acid starvation in MEFs lacking *p14/LAMTOR2* in which the Rag proteins are not lysosomally localized (Sancak et al., 2010) (Figures 4A and 4A'), in MEFs or HEK293FT cells transiently lacking expression of LAMTOR components due to siRNA-mediated knockdown (data not shown), or in MEFs in which the Rag proteins have been knocked down (Figure 4B). As expected (Sancak et al., 2008), Rag knockdown caused reduced TORC1 activation upon amino acid readdition (Figure S4A), as well as reduced TOR lysosomal localization (Figure S4B), confirming successful knockdown of the Rag proteins. In agreement with the biochemical data showing preferential binding of TSC2 for the inactive Rag complex, recruitment of TSC2 to lysosomes in the absence of amino acids was also abrogated upon knockdown of the Rag GAP GATOR1 complex components DEPDC5, NPRL2, and NPRL3 (Figures 4C and S4E). In this setup, the Rag complex is present on lysosomes but remains in the activated state (Bar-Peled et al., 2013; Panchoaud et al., 2013), leading, as previously shown, to increased TORC1 activity and lysosomal localization in the absence of amino acids (Figures S4C and S4D). These data suggest that recruitment of TSC2 to the lysosome upon amino acid withdrawal is due to the Rag proteins changing to the inactive conformation.

TSC2 Is Required for Complete Inactivation of TORC1 upon Amino Acid Removal

Having seen that TSC2 is recruited to lysosomes upon amino acid removal, we asked whether this has a functional consequence in terms of TORC1 activity. We tested whether cells lacking TSC2 are impaired in their ability to turn off TORC1 upon amino acid removal. Unlike control *Drosophila* S2 cells, which shut off TORC1 almost completely upon amino acid starvation, S2 cells with decreased Tsc2 levels show only a partial, initial drop in TORC1 activity, after which they retain a

Figure 2. TSC2 Binding to the Rag Proteins Depends on Cellular Amino Acid Signaling

- (A) Schematic diagram of RagA, RagB, and various mutants used in (B)–(D) to delineate the binding interfaces between TSC2 and RagA.
 (B) Both N-terminal and C-terminal regions of RagA are involved in TSC2 binding. CoIP of FLAG-TSC2 with RagA, RagB, and hybrid versions of the two proteins as depicted in (A), showing that both the N-terminal 33 amino acids, as well as the 5 C-terminal amino acids specific for RagA, impact TSC2 binding.
 (C) All five C-terminal amino acids specific for RagA, as shown in (A), contribute toward TSC2 binding, as each one improves the binding of RagB to TSC2 when introduced singly. CoIP experiments were performed as in (B). In (B) and (C), HA-GST-RagC was coexpressed with the RagA/B variants.
 (D) The N-terminal 424 amino acids of TSC2 bind strongly to Rag proteins, although the remaining amino acids 425–1,784 also contribute toward Rag binding. CoIP in HEK293FT cells of truncated versions of FLAG-TSC2 with HA-tagged RagA and RagC. Inactive-locked RagA^{low-nucleotide} and RagC^{GTP} were used, as this complex binds TSC2 most strongly, as in (F).
 (E) Amino acid starvation increases binding of endogenous TSC2 to the Rag GTPases. Immunoblots of anti-FLAG IPs from HEK293FT cells expressing FLAG-tagged RagA + RagC, treated with medium containing or lacking amino acids in the presence of dialyzed FBS for 1 hr.
 (F) Endogenous TSC2 and TSC1 bind most strongly to Rag dimers in the inactive state, which antiparallels Raptor binding. Immunoblots of anti-FLAG IPs from HEK293FT cells expressing FLAG-tagged WT RagA + RagC and mutants locked into the active RagA^{GTP}/RagC^{low-nucleotide} or inactive RagA^{low-nucleotide}/RagC^{GTP} states.

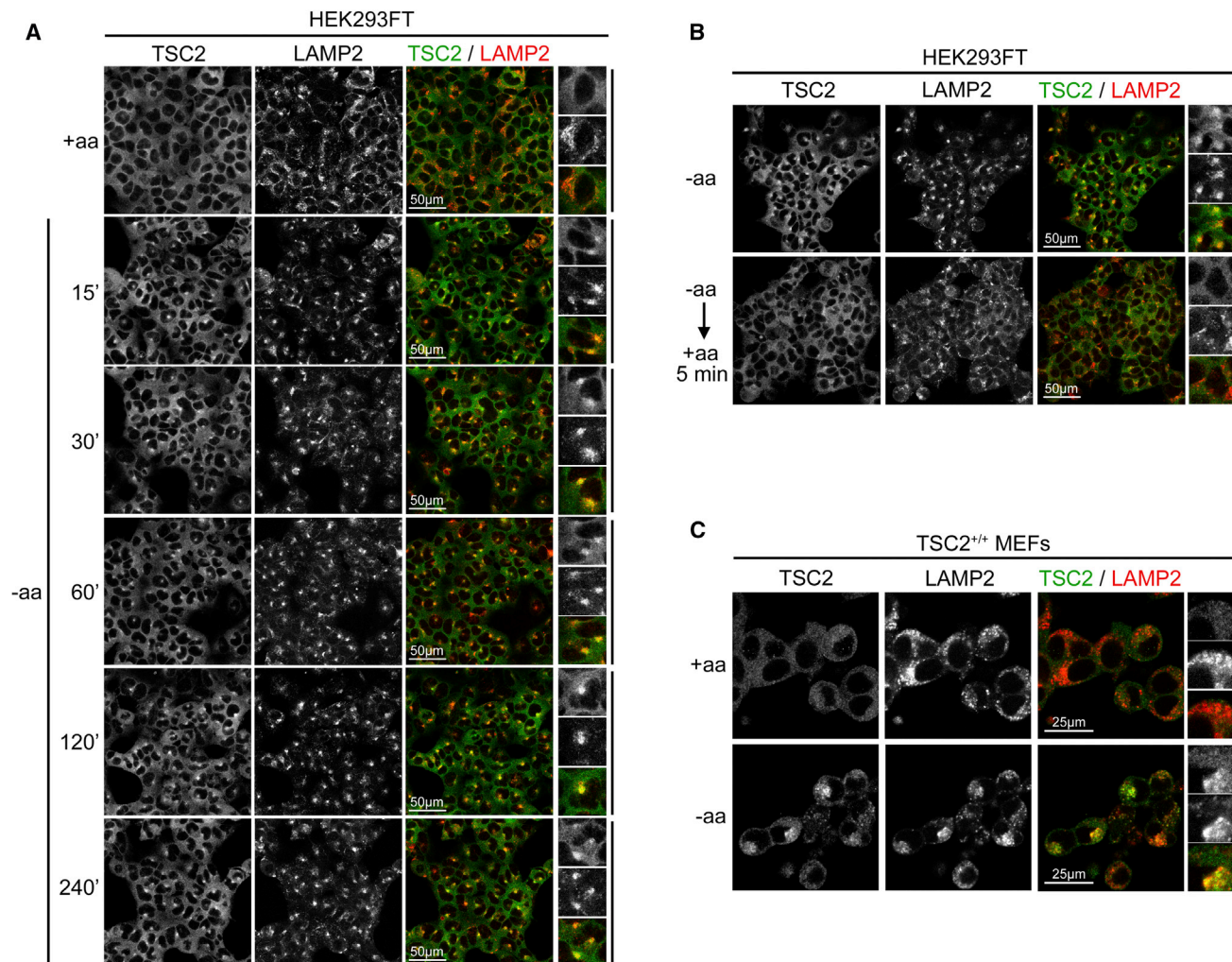


Figure 3. Amino Acid Starvation Causes TSC2 Lysosomal Localization in a Rag-Dependent Manner

(A and C) TSC2 accumulates on lysosomes upon amino acid deprivation. Confocal micrograph of HEK293FT cells (A) or MEFs (C) treated with medium containing or lacking all amino acids in the presence of dialyzed FBS for the indicated times, stained for TSC2 (green) and the lysosomal marker LAMP2 (red). LAMP2 aggregates do not show TSC2 accumulation in the +aa condition.

(B) TSC2 quickly relocates away from lysosomes upon amino acid readdition. HEK293FT cells were starved for amino acids in the presence of dialyzed serum for 4 hr and then resupplied with amino-acid-containing media for 5 min. Cells were stained and imaged as in (A).

significantly elevated amount of TORC1 activity stably for up to 2 hr (Figure 5A). This cannot be explained simply by elevated starting levels of TORC1 activity in the *Tsc2* knockdown cells because S2 cells lacking *PTEN* have similarly elevated starting levels of TORC1 activity but nonetheless efficiently shut off TORC1 upon amino acid removal (Figure 5B). These data are in agreement with previous observations (Gao et al., 2002). As in *Drosophila* cells, MEFs lacking TSC2 or TSC1 are also unable to completely inactivate mTORC1 upon amino acid starvation (Figures 5C and S5A). Also, in mouse cells, this cannot be explained by elevated starting levels of mTORC1 activity because *PTEN* knockout MEFs also start with elevated mTORC1 activity but efficiently shut off mTORC1 upon amino acid withdrawal (Figure 5D). Treatment of *TSC1*^{-/-} or *TSC2*^{-/-} MEFs with the TORC1-specific inhibitor rapamycin eliminated the

aberrantly elevated S6K phosphorylation, showing that it is reflecting elevated activity of TORC1 and not another kinase (Figures 5C and S5A). An alternate explanation could be that lack of the TSC1/2 complex leads to reduced phosphatase activity toward S6K. To test this, we performed a rapamycin treatment time course to compare the rate of S6K dephosphorylation in control and TSC2 knockdown conditions. In both MEFs and *Drosophila* S2 cells, however, rapamycin treatment caused rapid dephosphorylation of S6K with similar kinetics in cells containing or lacking TSC2 (Figures S5F and S5G, see quantification), indicating that phosphatase activity is unchanged. We wondered whether cells lacking TSC2 might simply have a delayed response to amino acid starvation, in which case extending the time of incubation in medium lacking amino acids would allow them to more completely shut off mTORC1. Astoundingly,

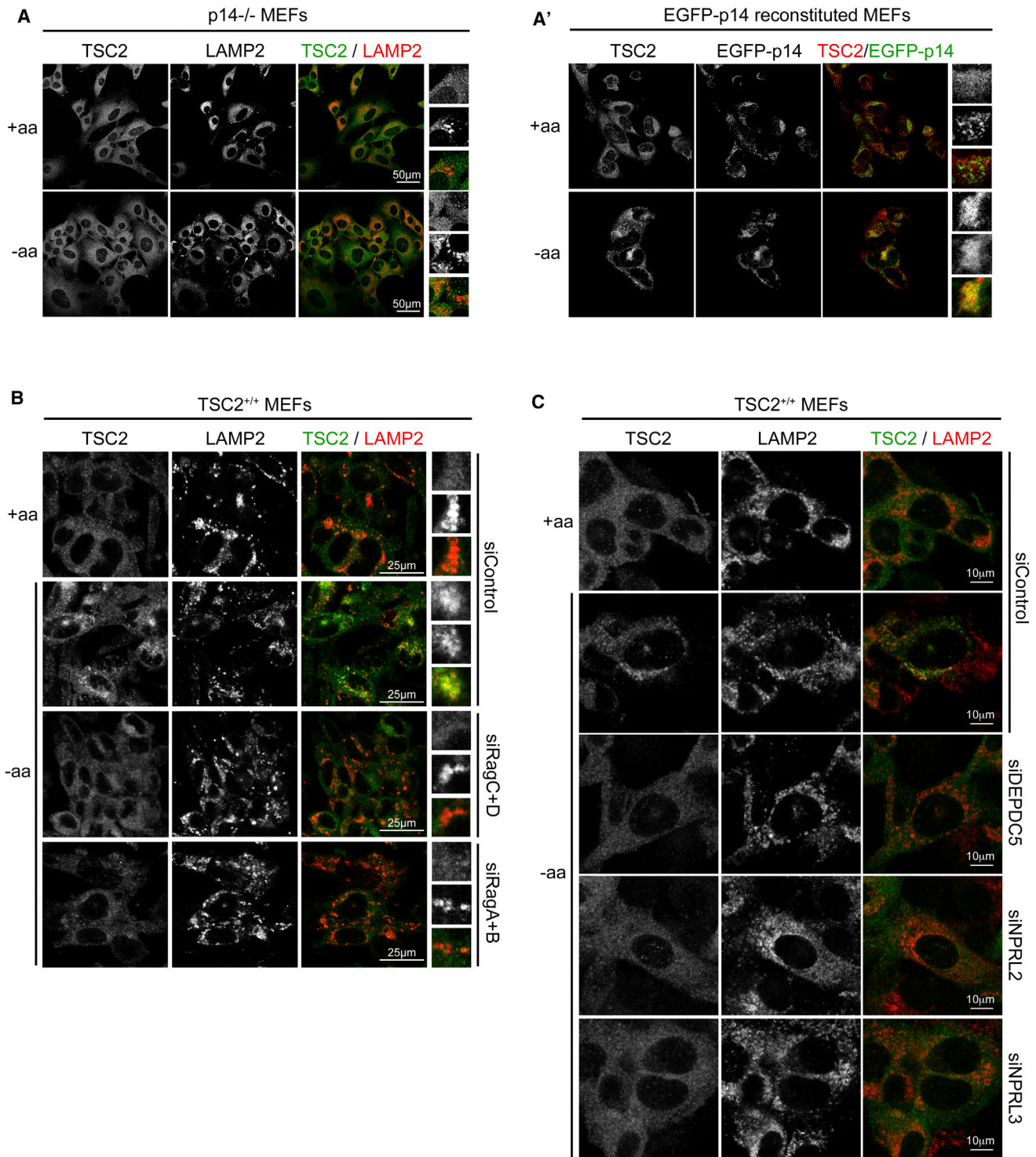


Figure 4. Lysosomal Recruitment of TSC2 Depends on the Rag Proteins

(A and A') Lysosomal localization of TSC2 upon amino acid removal requires the LAMTOR complex. Confocal micrograph of *p14/LAMTOR2* knockout MEFs (A) or control *p14* knockout MEFs reconstituted to express an EGFP-p14 fusion (A'), treated with medium containing or lacking amino acids for 1 hr in the presence of dialyzed FBS. Lysosomes were marked either with anti-LAMP2 antibody or with the lysosomally localized EGFP-p14.

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however, *TSC2*^{-/-} MEFs retained elevated levels of mTORC1 activity in the absence of amino acids for up to 1.5 to 2 days, at which point they started dying, leading to concomitant drops in both phospho-S6K and total S6K signals on the immunoblots (Figure S5B). When phospho-S6K signals are normalized to total S6K levels, one can see that *TSC2*^{-/-} MEFs retain a consistent level of mTORC1 activity up to their death (Figure S5B, bottom). In sum, TSC2 seems to be intimately linked to amino acid sensing by cells, in that (1) TSC2 localization changes upon amino acid removal, and (2) TSC2 is required for mTORC1 to fully turn off in the absence of amino acids.

TSC2 Is Required to Completely Release mTORC1 from the Lysosome upon Amino Acid Starvation

mTORC1 activation requires two regulated steps—one is activation of Rheb, and the other is localization of mTORC1 to the lysosomal surface where it meets Rheb, thereby forming an active complex (Inoki et al., 2003; Tee et al., 2003). In particular, the presence or absence of amino acids is known to regulate mTORC1 localization (Kim et al., 2008; Sancak et al., 2008). Since TSC2 appears to be involved in amino acid signaling to mTORC1, we asked whether TSC2 is required for the proper subcellular localization of mTORC1. As previously described (Kim et al., 2008; Sancak et al., 2008), in control cells, mTOR accumulates on lysosomes in the presence of amino acids and becomes diffusely cytoplasmic in the absence of amino acids (Figure 6A). In contrast, in *TSC2* null MEFs (which, as expected, are larger than control cells), mTOR localization is very strongly lysosomal, both in the presence and in the absence of amino acids (Figures 6A' and 6B, top). This defect can be rescued by transfecting them with a TSC2 expression plasmid, confirming specificity of the phenotype (Figure 6B, bottom, note lysosomally localized mTOR in the starved nontransfected *TSC2*^{-/-} cells, which is rescued in the cells expressing TSC2 + EGFP). As expected from the known effect of TSC1 loss on TSC2 stability, this defect in mTOR localization could also be observed in *TSC1* null MEFs (Figure S6A). Therefore, unexpectedly, the TSC1/2 complex is required for mTORC1 to be released completely from lysosomes upon amino acid withdrawal. Consistent with this phenotype being specific for loss of the TSC1/2 complex, and in agreement with the data presented in Figure 5, mTOR was readily released upon amino acid removal from lysosomes in *PTEN* mutant MEFs, which also start with elevated mTORC1 activation levels (Figure S6B).

Since mTORC1 localization is known to be regulated via the GTP/GDP load of the Rag GTPases (Sancak et al., 2008) and TSC2 has been described as a GAP for other small GTPases (Inoki et al., 2003; Wienecke et al., 1995; Xiao et al., 1997), we hypothesized TSC2 might regulate mTORC1 localization by acting as a Rag GAP. However, we radioactively quantified the relative GTP/GDP load of overexpressed Rag proteins in cells

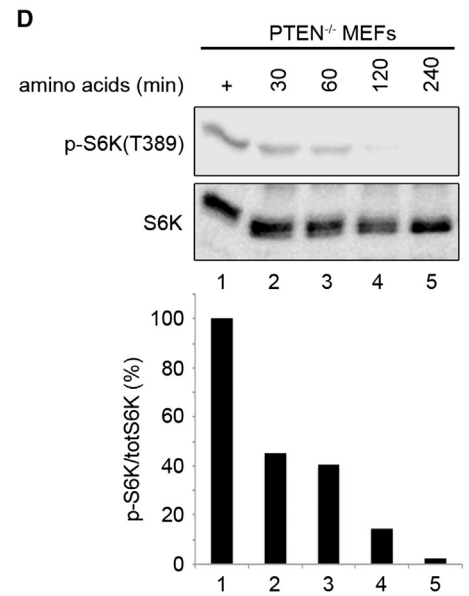
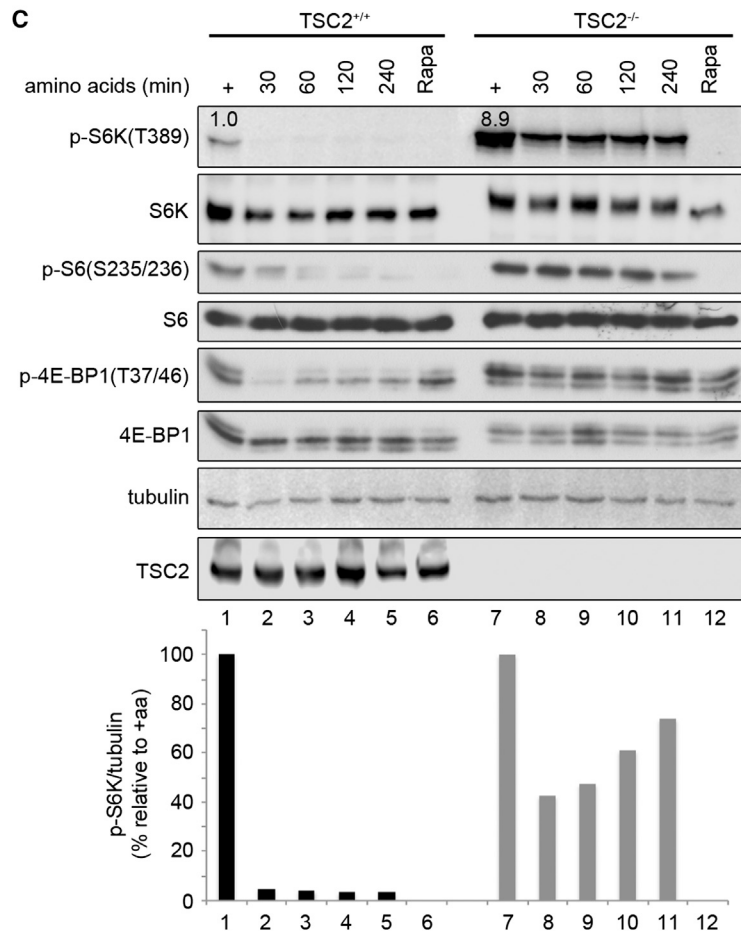
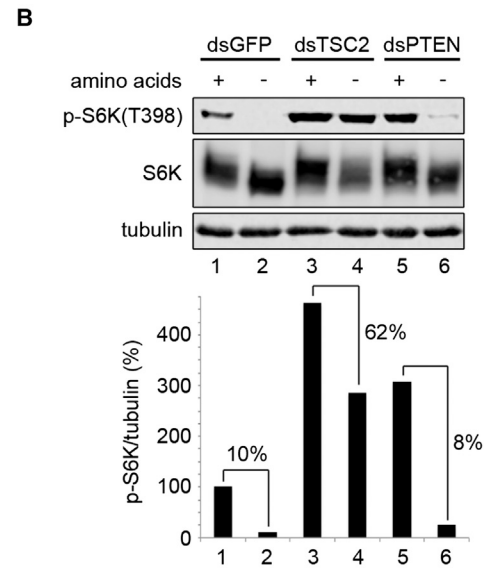
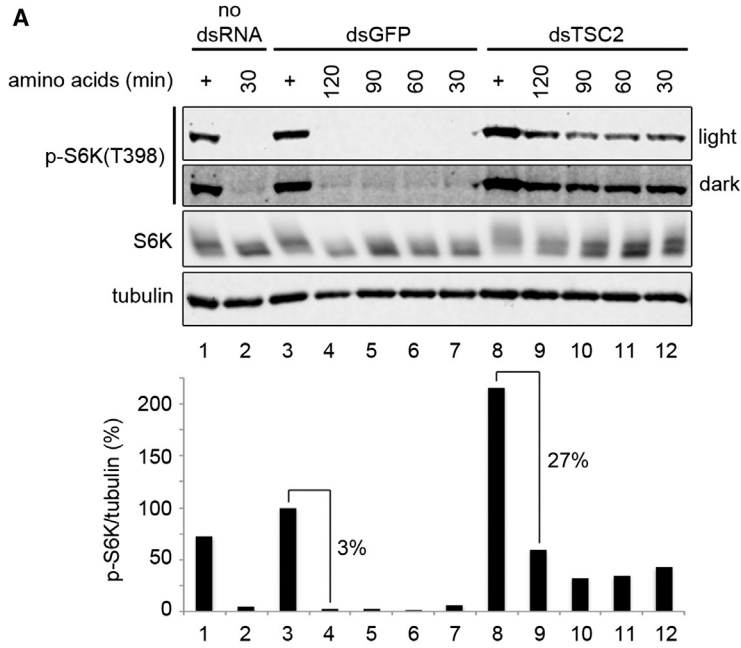
in the presence or absence of coexpressed TSC2 and could find no evidence to support this (data not shown). Another possible explanation could be that TSC2 affects the level of intracellular amino acids, thereby affecting mTORC1 localization. However, this does not seem to be the case: the intracellular levels of ten amino acids drop upon amino acid starvation. For these, the intracellular levels drop equivalently, or even more dramatically, in the *TSC2* null MEFs compared to control cells (Figure S5C). Unexpectedly, the intracellular levels of eight other amino acids either remain constant or increase in control cells upon amino acid removal from the medium (Figures S5D and S5E). For these, the intracellular levels either increase less dramatically or actually decrease in *TSC2* null MEFs compared to controls (Figures S5D and S5E). Because TSC2 acts via Rheb (Inoki et al., 2003) and because Rheb is known to bind the mTORC1 complex (Long et al., 2005), this raised the possibility that Rheb might mediate the effect of TSC2 on mTOR localization by helping to anchor mTOR at the lysosomal membrane in collaboration with the Rag proteins. Indeed, in *TSC2* null MEFs, knockdown of Rheb by siRNA allowed mTOR to be released from lysosomes upon amino acid removal (Figure 6C, controls in Figures S6C and S6D). Equivalent results could be observed in *TSC2* null cells stably transfected with inducible shRNA constructs targeting an independent region of Rheb, confirming specificity of the effect (Figure S6E). Furthermore, overexpression of active, but not inactive, Rheb was sufficient to localize mTOR to the lysosome even in the absence of amino acids (data not shown). In sum, these results identify Rheb as the mTORC1-anchoring activity in the *TSC2* null MEFs.

TSC2 Is Required for Cells to Respond Physiologically to Amino Acid Starvation

The data presented above support a model (detailed below and in Figure S7C) whereby TSC2 is required for cells to respond correctly to the removal of amino acids, allowing mTORC1 to be fully displaced from the lysosomal surface, thereby allowing it to become fully inactivated. We asked whether the inability of *TSC2* null cells to respond correctly to amino acid starvation at the molecular level is paralleled by an equivalent defect at the physiological level. One might expect that, if cells do not fully shut off mTORC1 upon amino acid starvation, this may lead to a metabolic catastrophe, as the demand for nutrients by cellular processes is not correctly reduced to match the reduced supply. To this end, we deprived control and *TSC2* null MEFs of amino acids in the presence of dialyzed serum. Surprisingly, control MEFs were able to survive these conditions for several days; the cells were still visibly present after the treatment, and they recommenced proliferating when given complete medium (Figure 7A), suggesting that they were in a quiescent state. This could also be quantified using a standard viability assay that quantifies ATP levels (blue trace, Figure 7B). In contrast

(B) Lysosomal localization of TSC2 upon amino acid removal requires the Rag proteins. MEFs transfected with control siRNA against *Renilla* luciferase or siRNAs targeting the various Rag proteins, treated with medium containing or lacking amino acids for 1 hr in the presence of dialyzed FBS. Cells were stained and imaged as in (A).

(C) Lysosomal localization of TSC2 upon amino acid removal requires that the Rag proteins change to the inactive conformation. MEFs transfected with control siRNA or siRNAs targeting components of the GATOR1 complex DEPDC5, NPRL2, or NPRL3 required for the Rag proteins to become inactive upon amino acid removal. Cells were stained and imaged as in (A).



(legend on next page)

(and in agreement with data in Figure S5B), *TSC2* null MEFs died within 3 days in response to amino acid removal, indicating a physiologically defective response to amino acid starvation (Figures 7A and 7B). If *TSC2* null MEFs are dying upon amino acid withdrawal due to an incomplete inhibition of mTORC1 activity, then blocking mTORC1 pharmacologically in these cells should improve their survival. Indeed, treating *TSC2* null MEFs with rapamycin rescued their death upon amino acid starvation (Figures 7D and 7E). One catabolic process that is activated in response to low mTORC1 activity to promote cell survival upon amino acid starvation is autophagy. Control MEFs efficiently activated autophagy upon amino acid starvation (seen as accumulation of a faster migrating form of LC3A and degradation of p62, Figure 7C). In contrast, *TSC2* null MEFs were impaired in activating autophagy (Figure 7C), thereby identifying one of probably multiple mTORC1-dependent physiological responses to amino acid starvation that are defective in *TSC2* null cells. In sum, these data support the notion that *TSC2* is required physiologically for cells to respond to amino acid starvation.

DISCUSSION

The data presented here suggest a model whereby, in the presence of amino acids, mTORC1 accumulates on lysosomes due to a dual anchoring activity composed primarily of the Rag proteins but supported by Rheb (Figure S7C). While amino acids are present, binding between the Rag proteins and TSC1/2 is low, causing the TSC1/2 complex to remain cytoplasmic. Upon amino acid removal, the Rag proteins cause mTORC1 to be released from lysosomes via two independent activities, both of which result from the Rag proteins shifting to an inactive conformation. First, the Rag proteins reduce their binding for mTORC1, thereby releasing one of the two activities tethering mTORC1 at the lysosome. Second, the Rag proteins actively recruit TSC2 to the lysosome. This allows TSC2 to act on Rheb, thereby releasing the second tethering activity keeping mTORC1 on the lysosome. In the absence of TSC2, this second activity is unaffected, causing mTORC1 to remain lysosomally localized (Figure S7C, right).

The LAMTOR complex (composed of the p18, p14, and MP1 proteins) has been shown to serve as a docking point for mTORC1 (Sancak et al., 2010) and MEK/ERK complexes (Schaeffer et al., 1998; Teis et al., 2002), regulating their recruitment to late endosomes/lysosomes and their activation status. Our data demonstrate that integrity of the LAMTOR complex is critical for proper *TSC2* subcellular localization upon amino acid withdrawal, therefore highlighting the importance of this scaffold complex for endomembrane-mediated activation/inactivation of signaling pathways.

The data presented here show that the TSC1/2 complex is part of the molecular machinery required for mTORC1 to respond properly to the absence of amino acids. We find that the TSC1/2 complex responds to amino acid starvation by changing its subcellular localization (Figures 3, 4, S3B, and S4E) and that *TSC2* is required for mTORC1 to be fully released from lysosomes and fully inactivated upon amino acid removal (Figures 5 and 6). That said, however, in cells lacking *TSC2*, there is nonetheless a clear initial drop in TORC1 activity upon amino acid removal (Figures 5A and 5C). The remaining activity is then sustained indefinitely (Figure S5B). Hence, mTORC1 appears to consist of two pools or two degrees of activation, one of which requires *TSC2* to become inactive upon amino acid withdrawal and one of which responds independently of *TSC2*. This might explain why previous studies arrived at differing interpretations of their data (Gao et al., 2002; Roccio et al., 2006; Smith et al., 2005) because there is some response of TORC1 to amino acid removal in *TSC2* null cells; however, the response is severely blunted compared to controls. Further work will hopefully shed light on these two pools of activity. Although the impairment in mTORC1 response to amino acids in *TSC2* null cells is partial, it is nonetheless of critical physiological relevance because *TSC2* null MEFs die upon amino acid removal in sharp contrast to control MEFs (Figures 7 and S5B).

Looking at the model in Figure S7C from several perspectives yields various insights. (1) Regulation of mTORC1 activation could previously be rationalized as consisting of two independent, parallel steps: first, regulation of Rheb via TSC1/2 in response to a plethora of signals including stresses and growth factor signaling, and second, regulation of mTORC1 subcellular localization to lysosomal membranes in response to amino acids. Only when mTORC1 is properly localized to meet active Rheb would an active holoenzyme form. The data presented here blur the distinction between these two steps because Rheb also affects mTORC1 localization, and amino acids also signal through TSC1/2. Instead, the two sets of regulatory inputs into mTORC1 appear to be more integrated. (2) Amino acid removal is “dominant” over growth factor signaling, causing mTORC1 to shut off despite the presence of growth factors (Blommaert et al., 1995; Hara et al., 1998). This was previously explained by the fact that, in the absence of amino acids, mTORC1 could not localize near active Rheb to form an active complex. Our dual tethering model is also consistent with this notion but for a slightly modified reason, which is that amino acid starvation acts to sever both the Rag and Rheb lysosomal tethering activities. (3) Seen from the perspective of the Rag proteins, they swap binding partners depending on the state of amino acid signaling, binding preferentially to mTORC1 in the presence of amino acids, and binding preferentially to the TSC1/2 complex in the absence of amino acids.

Figure 5. *TSC2* Loss Causes Insensitivity to Amino Acid Removal in *Drosophila* and Mammalian Cells

(A and B) *Tsc2*, but not *PTEN*, knockdown makes *Drosophila* cells largely insensitive to amino acid removal. (A) Immunoblots of *Drosophila* S2 cells treated with no dsRNA, dsRNA against GFP (control), or dsRNA against *Tsc2*, treated with media containing (+) or lacking amino acids for the indicated times. (B) *PTEN* knockdown S2 cells efficiently inactivate TORC1 upon amino acid removal, despite starting with elevated TORC1 activity levels. (C and D) *TSC2* knockout MEFs do not completely shut off TORC1 upon amino acid withdrawal, whereas *PTEN* knockout MEFs do. Immunoblots from (C) *TSC2*^{-/-} and the respective control *TSC2*^{+/+} MEFs or (D) *PTEN* knock-out MEFs treated with medium containing or lacking amino acids for the indicated times. Lanes 6 and 12 (C), cells treated with 20nM Rapamycin for 1 hr in the presence of amino acids (Rapa). Quantified with a LI-COR imaging system.

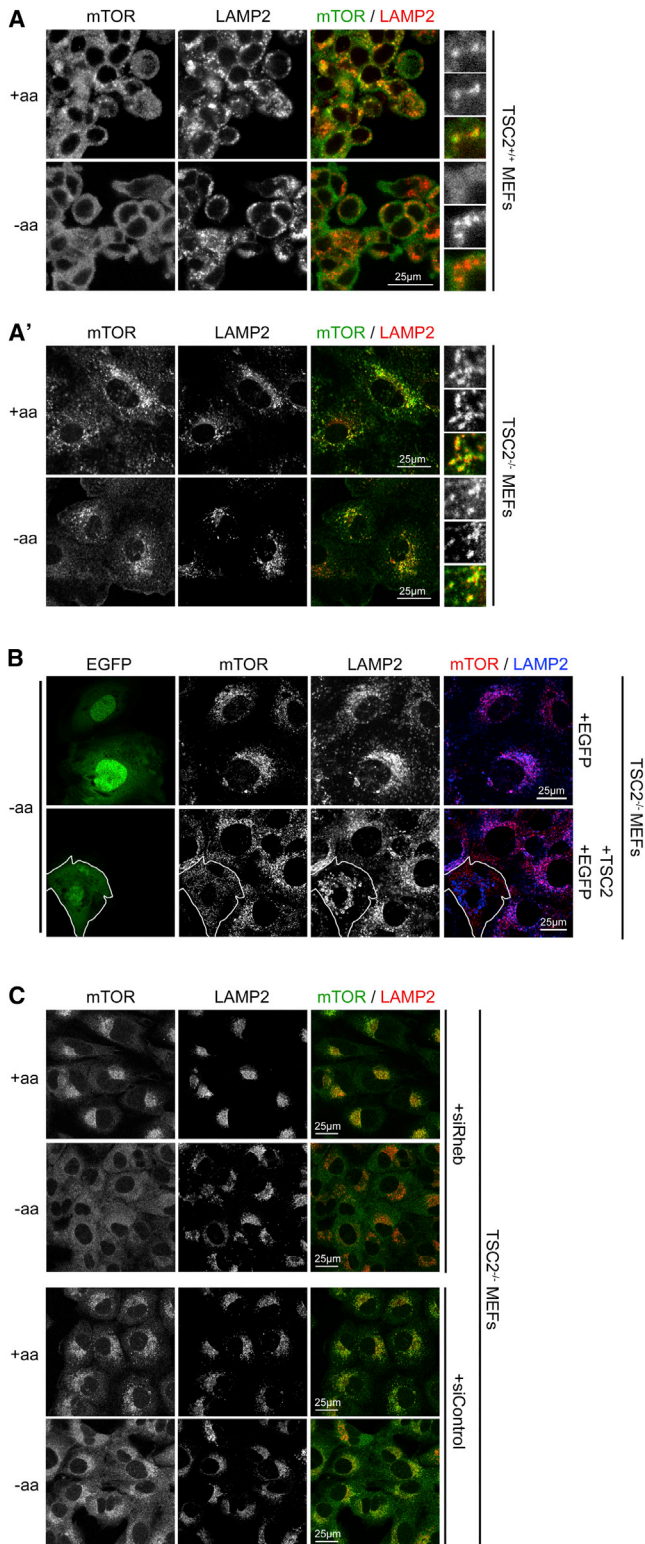


Figure 6. TSC2 Is Required for mTOR to Localize away from Lysosomes upon Amino Acid Removal in a Rheb-Dependent Manner

(A and B) mTOR is released from lysosomes upon amino acid removal in control (A), but not *TSC2* null, MEFs (A'). This is rescued by re-expressing

Consequently, mTORC1 and TSC1/2 also swap subcellular localizations.

We noticed that, in our hands, knockdown of Rag proteins did not result in as strong a reduction in TORC1 activity in the presence of amino acids as was previously reported (Kim et al., 2008; Sancak et al., 2008). The simplest explanation is technical—that our Rag knockdowns are not strong enough to fully abrogate Rag recruitment of mTORC1 in the presence of amino acids but are sufficient to impair Rag recruitment of TSC2 in the absence of amino acids. In that case, optimizing the Rag knockdowns might lead to even stronger effects than the ones we present here. Two alternate biological explanations, however, might be worth investigating in the future. The first is that our data suggest a dual anchoring mechanism of mTORC1 at the lysosomal membrane—one by the Rag proteins and one by Rheb. It is possible that the relative contribution of lysosomal tethering of mTORC1 by the Rag proteins and by Rheb might depend on their relative levels of expression and activation in the system being studied. This balance will likely depend on the cell line and on cell culture conditions. A second possible explanation could be one of biological kinetics, influenced by treatment strategy. The outcome might be quantitatively different if one looks at acute amino acid removal from cells adapted to complete medium (which we do here) or if one looks at amino acid add-back to cells that have equilibrated their signaling to the absence of amino acids. Indeed, we see that, if we knock down the Rag proteins in HEK293FT cells, we do not see a dramatic reduction in mTORC1 activity in the presence of amino acids (Figure S1D, lanes 1–3). However, if we remove amino acids for 1 hr and then readd amino acids for 30 min, the same degree of Rag knockdown causes an obvious reduction in mTORC1 activity (Figure S1D, lanes 7–9). Likewise, in *Drosophila* S2 cells, RagC knockdown only had a mild effect on TORC1 activity in untreated cells (Figures 1A and S1A) but severely blunted the ability of cells to respond to amino acid add-back (Figure S1B). This difference between amino acid removal and amino acid add-back raises the interesting possibility that the Rag proteins are key in recruiting mTORC1 to the lysosome, a process that happens upon amino acid readdition, and that both the Rag proteins and Rheb work together to keep mTORC1 on the lysosome once it is there. Indeed, in agreement with this model, mTOR is able to be recruited to the lysosome upon amino acid readdition in *TSC2* null MEFs in which Rheb is knocked down (Figures S7A and S7B), indicating that, although Rheb tethers mTOR to the lysosome upon amino acid removal (Figure 6C), it is not required for de novo recruitment of mTOR to the lysosome upon amino acid add-back.

TSC2 + EGFP (to mark transfected cells), but not EGFP alone, in the *TSC2* null MEFs (B). Note that the cell expressing *TSC2* and EGFP no longer has mTOR accumulated on lysosomes, whereas the surrounding, nontransfected cells retain lysosomally localized mTOR.

(C) Defective release of mTOR from lysosomes upon amino acid withdrawal in *TSC2* null MEFs is rescued by knocking down Rheb. *TSC2* null MEFs transfected with either Rheb siRNAs (top) or control siRNAs (bottom) treated for 1 hr with medium containing or lacking amino acids in the presence of dialyzed FBS.

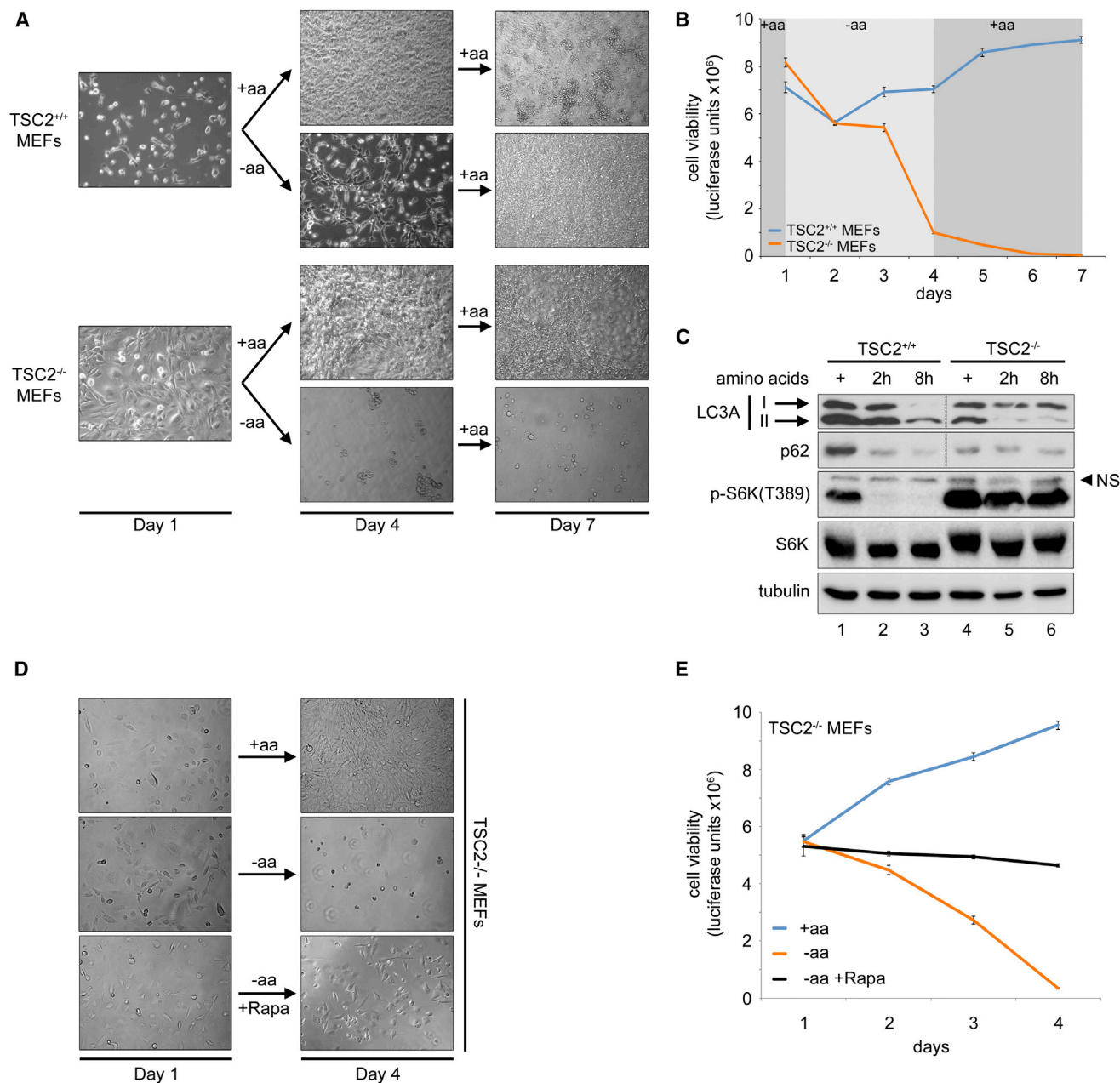


Figure 7. TSC2 Null Cells Are Impaired in Their Response to Amino Acid Starvation, Leading to Their Death

(A and B) Amino acid removal causes cell death in TSC2 null cells. TSC2^{-/-} and the respective TSC2^{+/+} control MEFs were treated on day 1 with medium containing or lacking amino acids in the presence of dialyzed FBS. On day 4, all samples were given fresh medium containing amino acids. (A) DIC images captured on days 1, 4, and 7. (B) Cell viability was determined using the Cell-Titer Glo kit (Promega).

(C) TSC2 null MEFs show defective autophagy induction upon amino acid withdrawal. TSC2^{-/-} and the respective TSC2^{+/+} control MEFs were treated with medium containing (+) or lacking amino acids in the presence of dialyzed FBS for the indicated times. Activation of autophagy visualized on an immunoblot via increased conversion of LC3A to the faster migrating lipidated form II and degradation of p62 protein.

(D and E) Pharmacological inhibition of TORC1 rescues survival of TSC2 null MEFs upon amino acid withdrawal. TSC2^{-/-} mouse embryonic fibroblasts were treated with medium containing (+aa) or lacking (-aa) amino acids with or without 20 nM Rapamycin (-aa/+Rapa) for 3 days in the presence of dialyzed FBS. DIC images (D) and respective cell viability titers were determined using the Cell-Titer Glo kit (Promega) (E). Error bars represent SD.

Previous reports showed that hyperactive mTORC1 signaling or dysregulated translation can lead to a metabolic mismatch in supply and demand, leading to cellular or organismal death

(Choo et al., 2010; Efeyan et al., 2013; Kim et al., 2008; Leprivier et al., 2013; Teleman et al., 2005). We hypothesized that, if TSC2 is required for mTORC1 activity to respond to amino acid

starvation, then TSC2 might also be necessary for cells to respond physiologically to this stress. Indeed, TSC2 knockout MEFs die upon removal of amino acids, whereas control cells do not (Figure 7). The fact that cells with elevated mTORC1 activity due to impaired nutrient sensing die when deprived of amino acids raises the interesting hypothesis that limiting nutrient supply to tumors of certain genotypes might have a beneficial effect on their treatment. Consistent with this effect being due to elevated mTORC1 activity, the death of TSC2 knockout MEFs is rescued by rapamycin treatment (Figure 7). This leads to the unexpected finding that rapamycin can actually promote cell survival under nutrient deprivation conditions, which might have therapeutic implications in mTOR-related malignancies.

We identify here the subcellular localization of TSC1/2 as one mechanism regulating this GAP holoenzyme. In an accompanying manuscript in this issue of *Cell*, Menon et al. (2014) show that TSC2 subcellular localization is also regulated by insulin signaling. They show that, in the absence of FBS, TSC2 is lysosomally localized. We show here that, in the absence of amino acids, TSC2 is lysosomally localized, even in the presence of growth factor signaling (FBS). Hence, the presence of both amino acids and growth factor signaling are required to keep TSC2 in the cytoplasm, and as long as one of the two is missing, TSC2 becomes lysosomally localized. Combined, our findings raise the possibility that TSC2 subcellular localization is a general mechanism for regulating this complex. It would be interesting to study whether the other inputs known to regulate TSC2 also affect its subcellular localization.

EXPERIMENTAL PROCEDURES

Detailed experimental procedures are provided in the [Extended Experimental Procedures](#).

Immunoblot Quantifications

Immunoblots were imaged and quantified with a LI-COR Odyssey FC imaging system, which has a larger dynamic range than what can be shown on the figure images. Data analysis was performed on the raw data and not the image displayed; hence, bands that look saturated on the figures for visualization purposes do not impact the data analysis. Quantifications of colIP efficiency (Figures 2C, 2D, and S2B) were calculated as the amount of Rag in the FLAG-TSC2 IP minus the negative control IP, normalized to the amount of TSC2 in the IP and the amount of Rag in the input.

Cell Imaging

All cell images within one panel were acquired and displayed with the same settings. Details of confocal microscopy can be found in the [Extended Experimental Procedures](#).

Cell Culture

Drosophila S2 cells were cultured in Schneider's medium and supplemented with 10% FBS (PAA). HEK293FT cells (Invitrogen) and MEFs were cultured in high-glucose Dulbecco's modified Eagle's medium (DMEM), supplemented with 10% FBS (Biocrom), except *PTEN*^{-/-} MEFs, which were also supplemented with 2 mM Glutamine. *TSC1*^{-/-}, *TSC2*^{+/+p53}^{-/-}, and *TSC2*^{-/-p53}^{-/-} MEFs were a kind gift by David Kwiatkowski and Michael Hall and were described previously (Kwiatkowski et al., 2002; Zhang et al., 2003). *p14*^{-/-} and *EGFP-p14* reconstituted MEFs were a kind gift of Lukas Huber, described in Teis et al. (2006). *PTEN*^{-/-} MEFs were provided by Hong Wu, described in Lesche et al. (2002).

SUPPLEMENTAL INFORMATION

Supplemental Information includes Extended Experimental Procedures and seven figures and can be found with this article online at <http://dx.doi.org/10.1016/j.cell.2014.01.024>.

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