Structure and mechanism of TRAPPIII-mediated Rab1 activation

Aaron M.N. Joiner¹, Ben P. Phillips², Kumar Yugandhar³, Ethan J. Sanford¹, Marcus B. Smolka¹, Haiyuan Yu³, Elizabeth A. Miller², J. Christopher Fromme^{1,4}

¹ Department of Molecular Biology and Genetics/Weill Institute for Cell and Molecular Biology, Cornell University, Ithaca, NY, USA

² MRC Laboratory of Molecular Biology, Cambridge, United Kingdom

³ Department of Computational Biology/Weill Institute for Cell and Molecular Biology, Cornell University, Ithaca, NY, USA

⁴ Corresponding author: jcf14@cornell.edu

ABSTRACT

The GTPase Rab1 is a master regulator of both the early secretory pathway and autophagy. Rab1 activation is controlled by its GEF (guanine nucleotide exchange factor), the multi-subunit TRAPPIII complex. The Trs85 regulatory subunit is critical for robust activation of Rab1 but its mechanistic role within the complex has remained unclear. Here we report the cryo-EM structure of the intact yeast TRAPPIII complex bound to its substrate Rab1/Ypt1. The orientation of the Rab1/Ypt1 hypervariable domain when bound to the complex leads to a model for how TRAPPIII associates with and activates Rab1/Ypt1 at the membrane surface. We identify a conserved amphipathic α -helix motif within Trs85 and demonstrate that this helix is required for stable membrane binding and Rab1/Ypt1 activation by TRAPPIII. Taken together, our results provide a comprehensive analysis of the structure and function of the yeast TRAPPIII complex and reveal that the key function of Trs85 is to serve as a membrane anchor, via its amphipathic helix, for the entire TRAPPIII complex.

INTRODUCTION

The GTPase Rab1 is a master regulator of both the early secretory pathway and autophagy (Jedd et al., 1995; Lynch-Day et al., 2010; Zoppino et al., 2010; Hutagalung and Novick, 2011). In budding yeast the Rab1 protein is encoded by the *YPT1* gene (Segev, 1991; Segev et al., 1988). Once activated by GTP-binding, Rab1/Ypt1 recruits vesicle tethering factors such as Uso1, Atg11, and the COG complex to initiate membrane fusion events (Jedd et al., 1995; Yamakawa et al., 1996; Cao et al., 1998; Suvorova et al., 2002; Lipatova et al., 2012). Proper localization of Rab1/Ypt1 is directly linked to its activation state and due to a slow intrinsic rate of exchange, the key regulatory step of Rab1/Ypt1 activation is catalyzed by its GEF, the multi-subunit TRAPPIII complex.

In the late secretory pathway, TRAPPII (a distinct, but related multisubunit complex) activates the small GTPase Rab11 (Jedd et al., 1995; Jones et al., 2000; Thomas and Fromme, 2016; Thomas et al., 2019). While the TRAPPII and TRAPPIII complexes share a core set of subunits, the distinguishing feature of the TRAPPIII complex is the presence of the Trs85 subunit (TRAPPC8 in metazoans) (Sacher et al., 1998, 2000, 2001, 2019; Lynch-Day et al., 2010). The metazoan TRAPPIII complex possesses three additional subunits that are not present in budding yeast (Bassik et al., 2013; Riedel et al., 2018) (Table S1), but the 7 genes encoding the 8 subunits of the yeast TRAPPIII complex are highly homologous with their metazoan paralogs (Koumandou et al., 2007; Sacher et al., 2019).

Loss of Trs85 results in a major reduction of Rab1/Ypt1 activation *in vivo* and *in vitro*, and causes defects in autophagy and aberrant secretion. TRAPPIII has been

reported to function at many cellular locations important for secretion and autophagy, including COPII vesicles, the Golgi complex, Atg9 vesicles, and the pre-autophagosomal structure in both budding yeast and metazoans (Meiling-Wesse et al., 2005; Nazarko et al., 2005; Lynch-Day et al., 2010; Kakuta et al., 2012; Lipatova et al., 2012; Bassik et al., 2013; Tan et al., 2013; Imai et al., 2016; Lamb et al., 2016; Zhao et al., 2017; Thomas et al., 2018; Stanga et al., 2019; Zhang et al., 2020).

Previous reports have presented crystal structures of the TRAPP core subunits and negative-stain electron microscopy analysis of the TRAPPIII complex, which provided an important foundation for understanding the overall architecture and function of the entire TRAPPIII complex (Jang et al., 2002; Kim et al., 2005, 2006; Tan et al., 2013). The nucleotide exchange mechanism was also elucidated using the core subunits bound to Rab1/Ypt1, and the active site was determined to be comprised of small parts of four subunits (Bet5, Bet3, Trs31, and Trs23) (Kim et al., 2006; Cai et al., 2008). However, the mechanistic role of Trs85 in the TRAPPIII complex has remained unclear, the structure of Trs85 (and the entire TRAPPIII complex) has not been determined, and how the complex interacts with membranes in order to activate Rab1/Ypt1 is unknown.

In order to define the role of the Trs85 subunit within the TRAPPIII complex, we have used a comprehensive approach combining cryo-EM, cross-linking mass-spectrometry, biochemical reconstitution of membrane-binding and nucleotide exchange activity, and *in vivo* functional analysis. Here we report the high-resolution structure of the intact yeast TRAPPIII complex bound to its substrate GTPase Rab1/Ypt1, representing the key intermediate of the nucleotide exchange reaction. The structure

reveals that Trs85 possesses both an N-terminal GTPase-like fold and a C-terminal α -solenoid motif. The α -solenoid connects Trs85 to the TRAPP core, interacting with a core residue mutated in the human disease SEDT. A portion of the flexible “HVD” region of Rab1/Ypt1, which links the GTPase to the membrane, is visible in the structure bound to the Trs31 core subunit. The positioning of the HVD and the electrostatics of the complex suggest a specific orientation of the complex on the membrane surface. We identify a conserved amphipathic α -helix motif on the putative membrane-binding surface of Trs85 and determine that this motif is required for stable membrane binding and Rab1/Ypt1 activation both *in vitro* and *in vivo*. Our results indicate that this region is also required for the function of Trs85 in autophagy, indicating that TRAPPIII uses the same membrane-binding mechanism to activate Rab1/Ypt1 for both secretion and autophagy.

RESULTS

Trs85 anchors the TRAPPIII complex to the membrane surface

We previously reported that the Trs85 subunit is required for robust Rab1/Ypt1 activation at the Golgi complex *in vivo* and on synthetic liposomes *in vitro* (Thomas et al., 2018) (Figure 1A, B). In contrast, Trs85 was dispensable for GEF activity in the absence of membranes (Thomas et al., 2018) (Figure 1C). We hypothesized that a primary function of Trs85 might be to directly interact with the membrane surface. Although we were unable to purify the isolated Trs85 subunit, we compared membrane binding of the purified TRAPPIII complex with and without the Trs85 subunit, using an *in vitro* liposome pelleting assay (Figure 1D). Consistent with our hypothesis, the intact TRAPPIII complex exhibited robust binding to synthetic liposomes, and the Trs85 subunit was required for stable membrane association (Figure 1E).

Cryo-EM structure of the yeast TRAPPIII-Rab1/Ypt1 complex

To establish the mechanism of Trs85 function within the complex, we sought to resolve the high-resolution structure of the intact budding yeast (*S. cerevisiae*) TRAPPIII complex. Furthermore, the published crystal structure of TRAPP core subunits bound to the Rab1/Ypt1 substrate lacks the critical “HVD” (hyper-variable domain) of Rab1/Ypt1 (Cai et al., 2008), which has been shown to play an important role in substrate recognition by the TRAPP complexes (Thomas et al., 2019). We therefore assembled a stable complex in which TRAPPIII was bound to full-length yeast Rab1/Ypt1 in order to also visualize any interactions involving the HVD (Figure S1A). For this purpose, we

used a yeast Rab1/Ypt1 construct in which the two C-terminal cysteine residues that are prenylated *in vivo* were replaced with an affinity tag for purification.

This complex was well-behaved and suitable for high-resolution cryo-EM experiments (Figure S2), but the complex exhibited a strongly “preferred orientation” on frozen grids. Initial datasets indicated that approximately 95% of particles adopted a single orientation, and the other 5% adopted an orientation that was only slightly different from the primary orientation. To capture additional views, we therefore imaged grids tilted by 30°. Although this resulted in a marked improvement, significant orientation bias remained that precluded confident *de novo* building of Trs85 into resulting reconstructions. We therefore collected additional data using grids tilted at 40° and 45°. Although these data were of lower quality due to the higher tilt angle, combination of data from the three tilt angles provided sufficient orientation coverage to generate a 3.7Å reconstruction suitable for *de novo* model building (Figure 2A, B, Table S2, Figure S3). For the core subunits and GTPase substrate, we docked and rebuilt available crystal structures of subunits from yeast and other species (Jang et al., 2002; Rak et al., 2003; Kim et al., 2005, 2006; Pylypenko et al., 2006; Cai et al., 2008). For Trs85, we used trRosetta (Yang et al., 2020) to generate models of Trs85 from budding yeast and several other organisms. The overall topologies of these models were quite similar and matched the 3D-reconstruction reasonably well. The trRosetta model of budding yeast Trs85 was therefore used as a guide for *de novo* building of the Trs85 model.

The TRAPPIII complex adopts a narrow rod-like structure (Figure 2A, B), in agreement with the published negative-stain EM structure (Tan et al., 2013). The structures of each of the core subunits and the GTPase substrate are largely similar to the published crystal structures (Figure S4). Some small differences were noted, for example, we observed covalently-bound palmitate molecules in both Bet3 subunits, whereas only one of the Bet3 subunits appeared to be bound to palmitate in the previous structure of the catalytic core (Cai et al., 2008). Rather than mediate interactions with organelle membranes like the prenyl groups of Rab1/Ypt1, these palmitates are thought to stabilize the folding of the Bet3 subunits (Turnbull et al., 2005; Kümmel et al., 2006). The interaction between the TRAPP active site and the globular domain of Rab1/Ypt1 is essentially the same as that observed in the crystal structure previously determined by the Reinisch group (Cai et al., 2008). However, we observed a novel interaction with the Rab1/Ypt1 HVD that we discuss further below.

To validate the final structural model, we utilized cross-linking mass spectrometry (XL-MS) to identify lysine residues in proximity to each other (Figure 2C, D, Figure S1B, Table S3). From 108 total cross-links, 32 could be mapped to residues modeled in the structure (Figure 2E). 31 of these 32 cross-links were within the maximum distance constraint for disuccinimidyl sulfoxide (DSSO), 30Å, with most occurring between residues less than 20Å apart (Figure 2F). The other cross-link (between Lys78 of Trs31 and Lys85 of Bet3) spanned a distance of 45Å and is likely a false positive because those two cross-linked residues were also far apart in the crystal structure of these subunits bound to Rab1/Ypt1 (Cai et al., 2008). However, we cannot rule out the possibility that this cross-link might have captured some of the dynamic movement of

the complex in solution. The remaining 76 cross-links could not be mapped onto the model because they involved one or more residues located in unmodeled loops (Table S3). For each of these residues, we confirmed that the position of these unstructured loops in the model was consistent with the existence of a cross-link. Overall, the XL-MS results contribute support for the atomic model of the complex.

The C-terminus of Trs85 binds to the core subunits Trs20 and Trs31

The cryo-EM structure reveals that the N-terminal portion of Trs85 adopts a GTPase-like fold and the C-terminal portion folds into a twisting α -solenoid structure comprising five pairs of α -helices (Figure 3A). The α -solenoid connects Trs85 to the TRAPP core through a surface that is well-conserved among Trs85 homologs in related species, and even throughout model organisms and in humans (Figure 3B). This interaction surface is described further below. The GTPase-like fold in Trs85 has multiple insertions relative to a typical GTPase protein, and does not appear capable of binding nucleotide, as key nucleotide-binding residues are not conserved and an α -helix formed by residues 46-55 of Trs85 occupies the region where nucleotide is bound in GTPase proteins.

The structure indicates that the C-terminal α -solenoid region of Trs85 interacts with the TRAPP core. We determined that the C-terminal 198 amino acids of Trs85 (Trs85⁵⁰¹⁻⁶⁹⁸) are sufficient to maintain this interaction during purification of the recombinant complex (Figure 4A, B), and several cross-links were detected between the C-terminal region of Trs85 and either Trs20 or Trs31 in the XL-MS experiments (Figure 2D, E Figure 4D, Table S3).

There are two interfaces between Trs85 and the TRAPP core subunits: an extensive interaction between Trs85 and Trs20 (Figure 4E), involving 31 residues and encompassing a total surface area of roughly 2,250Å², and a much smaller interaction between Trs31 and part of a largely unstructured loop of Trs85. This loop spans residues ~580-612 of Trs85, and appears to be an insertion specific to *S. cerevisiae*.

The interface between Trs85 and Trs20 is quite conserved (Figure 3B), and involves Trs20 Asp46, a residue known to be important for assembly of both the TRAPP II and TRAPP III complexes (Figure 4E) (Zong et al., 2011; Brunet et al., 2013; Taussig et al., 2014) that is mutated in spondyloepiphyseal dysplasia tarda (SED) (Sacher et al., 2019). This and nearby negatively charged residues of Trs20: Glu49, Asp50, and Asp93, interact with multiple positively charged residues of Trs85: Arg618, Arg619, Arg620, and Lys621 (Figure 4E). To test the importance of this interface *in vivo*, we generated alanine substitution and charge reversal mutations for the Trs85 residues and assessed their ability to provide Trs85 function in a sensitized background in which the core Bet3 subunit is tagged at its C-terminus (Sacher et al., 2001). All of the mutants dramatically impaired Trs85 function (Figure 4F). Even mutation of just one of these residues, Trs85 Arg620, to either alanine or glutamate, significantly impaired Trs85 function, with R620E exhibiting a stronger phenotype than R620A (Figure 4F).

To determine whether these phenotypes were due to disruption of the Trs85-Trs20 interaction, we investigated the assembly status of the TRAPP III complex by monitoring the amount of the core subunit Trs23 that co-immunoprecipitated with the Trs85 mutant alleles (Figure 4G). The quadruple mutants and R620E single mutant essentially abolished the interaction between Trs85 and the core subunits, whereas the

R620A substitution mutant exhibited a partial effect. As the severity of complex disruption correlated with the growth phenotypes in the complementation test, these results provide a strong validation of the importance of the Trs85-Trs20 interface observed in the cryo-EM structure (Figure 4F, G).

We previously reported that Trs85 is required for robust activation of Rab1/Ypt1 at the Golgi complex (Thomas et al., 2018). We tested the ability of these complex-disrupting mutants to activate Rab1/Ypt1 at the Golgi in otherwise wild-type cells (Figure 4H). As predicted, the quadruple mutants and R620E single mutant were unable to robustly activate Rab1/Ypt1, as monitored by the loss of punctate RFP-Ypt1 localization. The R620A mutant again exhibited an intermediate phenotype. Interestingly, the complex-disrupting mutations also resulted in mislocalization of Trs85 (Figure 4H), suggesting that Trs85 is not able to stably bind to membranes by itself when not incorporated into the TRAPP III complex.

To examine the importance of the less conserved interface between Trs85 and Trs31 (Figure 2B), we tested a mutant lacking a portion of the partially structured loop designed to disrupt this interaction without disrupting the folding of Trs85 (Trs85^{Δ575-603}). This Trs85 mutant exhibited a minor temperature-sensitive growth phenotype and a partial loss of interaction with the TRAPP core in co-IP experiments (Figure 4F, G). To further examine the effect of this mutant, we attempted purification of recombinant TRAPP III harboring this mutation. The complex remained largely intact during nickel-ion affinity purification but then dissociated during gel filtration chromatography (Figure 4C), suggesting a weakened ability of this Trs85 mutant to stably associate with the core subunits. In contrast, we were unable to purify the quadruple mutant complexes

disrupting the Trs20 interaction, consistent with an inability to purify the isolated Trs85 subunit. Therefore, the loop comprising residues 575-603 plays a subsidiary role in linking Trs85 to the TRAPP core subunits by interacting with Trs31.

A model for the orientation of TRAPPIII on the membrane surface

The TRAPP complexes identify their specific substrates in part through recognition of the GTPase HVD (Thomas et al., 2019). It is likely that other Rab-GEFs also recognize the HVD regions of their GTPase substrates (Chavrier et al., 1991; Dunn et al., 1993; Li et al., 2014), but interactions between GEFs and GTPase HVD regions have not been structurally characterized. In fact, HVD regions are often intentionally omitted from structural studies because of their “unstructured” nature (Dong et al., 2007; Cai et al., 2008). In our cryo-EM reconstruction of the TRAPPIII-GTPase complex, we observed a region of strong but unexpected density on the surface of the complex that we have confidently assigned as corresponding to a portion of the Rab1/Ypt1 HVD (Figure 2B and S4H). There are multiple pieces of evidence supporting this assertion: 1) At lower threshold levels of the reconstruction, there is a nearly continuous tube of density connecting this region of unknown density with the C-terminus of the Rab1/Ypt1 nucleotide-binding domain (NBD) (Figure 5A); 2) In preliminary cryo-EM experiments, we observed 2D classes corresponding to complexes either with or without bound Rab1/Ypt1, and the unknown density was only present when Rab1/Ypt1 was also bound (Figure 5B); 3) XL-MS data identified cross-links between the Rab1/Ypt1 HVD and the Trs31 subunit, occurring in the region between the C-terminus of the Rab1/Ypt1 globular domain and the unknown density (Figure 5C, D, Table S3). Therefore, although

the cryo-EM density is not clear enough to unambiguously assign specific residues of Rab1/Ypt1 to this region, we are confident that it corresponds to some portion of the Rab1/Ypt1 C-terminal HVD.

The Rab1/Ypt1 HVD culminates with two cysteine residues that are prenylated (Pylypenko et al., 2006) and insert into the membrane either prior to or concomitant with activation by TRAPPIII. The observed location of the HVD bound to the surface of Trs31 therefore imposes a significant constraint upon any model for how TRAPPIII binds to the membrane surface. This leads us to propose that the complex binds the membrane in the orientation shown in Figure 5A such that the Rab1/Ypt1 NBD is bound to the surface of TRAPPIII that is most distal to the membrane. Additional evidence in favor of this orientation of the TRAPPIII complex arises from analysis of the electrostatic potential of the surface of the complex, though this analysis has the caveat that some surface residues are unstructured. While there are both positive and negative charges scattered across all surfaces of the TRAPPIII complex, and no single surface contains a concentrated region of positive charge, the proposed membrane-binding surface appears to be the most positively charged of all the surfaces (Figure 5E). In accordance with this idea, we determined that the TRAPPIII complex prefers to bind liposomes with anionic lipid mixtures (Figure 5F, Table S4). A circumstantial piece of evidence supporting the proposed membrane-bound orientation is the extreme preferred orientation adopted by the complex in vitreous ice (Figure S2F). These 2D projections (Figures 5B, S2B) implicate two possibilities for which surface was interacting with the air-water interface within the holes of the EM grid: one is opposite where the Rab1/Ypt1 C-terminus is located, while the other is the same surface we propose TRAPPIII uses to

bind membranes (Figure 5A). As a peripheral membrane protein, we think it is probable that TRAPPIII interacted with the air-water interface via the same surface it uses to bind to organelle membranes within a cell.

A conserved amphipathic helix within Trs85 is required for TRAPPIII membrane binding and Rab1 activation

We were concerned that the proposed membrane binding surface of Trs85 did not appear to possess an overwhelmingly positively charged character, which we expected to be necessary for its interaction with membranes. However, upon closer inspection we identified an unmodeled region on this surface located between the GTPase-like fold and α -solenoid motif (residues 368-409). Though we were unable to model this portion of the protein in our structure because of a lack of clear electron density, the primary sequence is conserved and predicted to fold as an amphipathic α -helix (Figure 6A-C).

In order to determine whether this putative amphipathic helix was important for Trs85 and TRAPPIII function, we made several mutations in this region, disrupting either the charge or both charge and hydrophobicity (Figure 6D). Mutation of 7 residues in this region to alanine (mutant 7A) resulted in severe loss of function (Figure 6E). Removal of the basic charge (mutant 4A) displayed slight growth defects, while charge inversion of the lysine residues by mutation to glutamate residues (mutant 4E) resulted in complete loss of function (Figure 6E).

We considered the possibility that these mutations might prevent Trs85 from associating with the core TRAPP subunits, but both the 7A and 4E mutants were stably

bound to the TRAPP core in co-IP experiments (Figure 6F). We reasoned that if the 7A and 4E Trs85 mutants associate with the core TRAPP subunits but are not functional, then perhaps these residues are indeed part of an amphipathic helix responsible for mediating TRAPPIII membrane-binding, similar to the role of amphipathic helices in many other peripheral membrane proteins (Kahn et al., 1992; Antonny et al., 1997; Huang et al., 2001; Lee et al., 2005; Drin and Antonny, 2010). To test this hypothesis, we purified recombinant mutant TRAPPIII complexes and directly tested their relative membrane-binding affinities using the *in vitro* liposome pelleting assay (Figure 6G). Similar to wild-type, a small amount of each complex pelleted in the absence of membranes. However, in striking contrast to wild-type, both the 7A and 4E mutants did not bind to synthetic liposomes (Figure 6H), indicating that this amphipathic helix is required for TRAPPIII membrane binding *in vitro*.

Consistent with a loss of membrane-binding, both the 7A and 4E Trs85 mutants failed to localize to the Golgi and did not activate Rab1/Ypt1 either *in vivo* (Figure 6I) or in biochemical reconstitution assays in which TRAPPIII GEF activity is membrane-dependent (Figure 6J). Importantly, these mutants retained their ability to activate Rab1/Ypt1 in a membrane-independent GEF assay (Figure 6K), indicating that the active site is unaffected by these mutations. Taken together, these results reveal that the Trs85 amphipathic helix is required for TRAPPIII complex membrane binding and function.

Similar requirements for TRAPPIII function in secretion and autophagy

Prior work has shown that *trs85Δ* cells display a deficiency in autophagy (Meiling-Wesse et al., 2005; Nazarko et al., 2005; Lynch-Day et al., 2010; Kakuta et al., 2012), a defect that can be monitored via the degradation kinetics of GFP-tagged Atg8, the yeast paralog of LC3 (Klionsky et al., 2016). In wild-type cells GFP-Atg8 is degraded as a consequence of rapamycin-induced autophagy, resulting in an accumulation of free GFP. We determined that mutations disrupting the Trs85 amphipathic helix or Trs85 binding to the TRAPP core displayed significant defects in rapamycin-induced GFP-Atg8 degradation (Figure 7A, B). This indicates that the role of the Trs85 amphipathic helix in anchoring the TRAPP core to the membrane surface is critical for both secretion and autophagy (Figure 7C).

DISCUSSION

GEFs control where and when their GTPase substrates are activated by catalyzing nucleotide-exchange. By coupling a localization signal and nucleotide exchange activity within each GEF, the cell can achieve precise spatial and temporal activation of its substrate GTPase at a specific cellular location. The TRAPP complexes activate two of the most important Rab GTPases in the secretory pathway, Rab1 and Rab11. Recent work from our lab and others has clarified the roles of TRAPPIII and TRAPPII as GEFs for Rab1 and Rab11, respectively (Jones et al., 2000; Meiling-Wesse et al., 2005; Morozova et al., 2006; Lynch-Day et al., 2010; Thomas and Fromme, 2016; Riedel et al., 2018; Thomas et al., 2018, 2019). Although the activation of Rab1/Ypt1 by core TRAPP subunits has been extensively studied (Jones et al., 2000; Wang et al., 2000; Cai et al., 2008), the structure and function of the key TRAPPIII regulatory subunit Trs85 has remained elusive. Our results demonstrate that Trs85 serves as a membrane anchor for the TRAPPIII complex by utilizing a conserved amphipathic helix.

The ability of Trs85 to anchor the TRAPPIII complex to a membrane surface is vital to the activation of Rab1/Ypt1 because the stable association of the TRAPPIII complex to the membrane surface establishes the proper spatial geometry and likely increases the frequency of interactions between the active site and the Rab1/Ypt1 substrate (Figure 7C). Simply put, the observed elevated activity of TRAPPIII relative to the TRAPP core is due to the fact that the core does not appear able to adequately bind to membranes, and therefore cannot activate Rab1/Ypt1 at the membrane surface.

A key aspect of our work is the presence of the Rab1/Ypt1 HVD in the cryo-EM structure of the TRAPPIII-Rab1/Ypt1 complex. The HVD terminates with two prenylated

cysteine residues that serve as membrane anchors for the Rab (Pylypenko et al., 2006). At steady-state in cells, inactive (GDP-bound) Rabs are bound to GDI proteins that protect the hydrophobic prenyl groups (Goody et al., 2005; Wu et al., 2010). However, the inactive Rabs dynamically dissociate from GDI and transiently bind to membranes before re-associating with GDI. In this way, inactive Rabs are thought to sample different organelles before encountering their specific GEF to become activated at a particular organelle. The active Rab cannot bind to GDI and is therefore stabilized at its site of activation until inactivated by a Rab-GAP. Accordingly, the structure that we have determined here represents an intermediate of the nucleotide-exchange reaction in which the prenylated cysteines of the HVD are membrane-bound. This structure reveals the HVD is bound by the Trs31 subunit, and its position in the structure provided us with a critical piece of evidence regarding the orientation of the complex on the membrane surface.

Our identification of the TRAPPIII membrane binding surface led to, and was further supported by, the presence of a conserved amphipathic helix within Trs85. We found that mutations disrupting either the hydrophobic face of the helix or inverting the charge of the other face completely abrogated membrane binding of the entire TRAPPIII complex, both *in vitro* and *in vivo*.

The fact that we were not able to model the amphipathic helix provides additional, although perhaps counterintuitive, evidence in favor of its role in membrane binding. Amphipathic helices are very commonly found on the surfaces of proteins, in which they do not perform any specialized function but rather simply are part of the protein fold in which hydrophobic sidechains are on the interior and hydrophilic

sidechains are on the exterior of the folded protein. These α -helices should be readily visible in the structure, as are all of the other predicted α -helices in the TRAPPIII complex structure. However, an amphipathic α -helix that has evolved to insert into the membrane would not be expected to form part of the hydrophilic surface of a folded protein domain (Drin and Antonny, 2010). Rather, it should extend from the globular portion of the protein in order to insert longitudinally into the outer leaflet of the membrane, as is the case with the Trs85 amphipathic α -helix that is required for TRAPPIII membrane binding.

Although there are many examples of amphipathic helices serving as membrane anchors for smaller proteins, such as Arf GTPases and N-BAR domain containing proteins (Drin and Antonny, 2010), TRAPPIII represents a notable example in which a sizeable multi-protein complex is anchored by a single amphipathic helix. Remarkably, the mutation of just four residues in this helix was sufficient to disrupt the binding, and therefore function, of the entire 250kD complex.

The core subunits likely provide additional protein-protein interactions or perhaps supplementary interactions with organelle membranes, as Trs85 localization to the Golgi was abrogated in cells when mutations that diminished its binding to the core were introduced. We determined that the TRAPPIII complex prefers to bind to membranes with a significant anionic charge, but did not identify a specific lipid requirement (Figure 5F, Table S4). Given that TRAPPIII has been implicated to function at many locations within the cell including on COPII vesicles, at the Golgi, on Atg9 vesicles, and at the preautophagosomal structure, it is likely that additional protein binding partners help specify TRAPPIII localization *in vivo* via coincidence detection.

For example, TRAPPIII localization to COPII vesicles likely depends upon an interaction between the TRAPP core and the COPII coat subunit Sec23 (Cai et al., 2007; Lord et al., 2011). Future studies of such functions of TRAPP core subunits are important but made challenging by the need to disentangle observed roles of these subunits in TRAPPIII versus TRAPP II.

Many questions remain regarding the role of both TRAPPIII and TRAPP II in autophagy. Our results indicate that the key function of Trs85, serving as a membrane anchor for the TRAPPIII complex, is important for both normal growth and autophagy (Figure 7C).

MATERIALS AND METHODS

TRAPPIII purification

The full set of TRAPPIII complex genes from *Saccharomyces cerevisiae* were cloned into two plasmids. Two copies of TRS33 were included to compensate for the otherwise substoichiometric expression levels of this subunit. *TRS85* with an N-terminal hexahistidine tag and *TRS33* were inserted into a pETDuet-1 vector (CFB2868). As reported previously, *BET3*, *BET5*, *TRS20*, *TRS23*, *TRS31*, and *TRS33* were inserted into a pCOLADuet-1 vector (pLT21,(Thomas and Fromme, 2016)). These plasmids were co-transformed into the Rosetta2 strain of *E.coli* (Novagen). Multiple liters of culture were grown in terrific broth for 8-12h at 37°C (until the OD₆₀₀ ~2-3). The temperature was then reduced to 16°C and 1 hour later IPTG was added to a final concentration of 300μM to induce protein expression. Following overnight expression (14-18h), cells were collected via centrifugation and resuspended in lysis buffer (40mM Tris pH 8, 300mM NaCl, 10% glycerol, 10mM imidazole, 1mM DTT, 1mM PMSF and 1x Roche PIC). Cells were lysed by sonication and after clearing the lysate, the complex was purified by nickel affinity chromatography (Ni-NTA resin, Qiagen) and eluted with elution buffer (40mM Tris pH 8, 300mM NaCl, 10% glycerol, 250mM imidazole, 1mM DTT). The sample was further purified using anion exchange chromatography with a linear gradient on a MonoQ column (GE Healthcare) (Buffer A: 20mM Tris pH 8, 1mM DTT; Buffer B: 20mM Tris pH 8, 1M NaCl, 1mM DTT). The peak fractions were collected, concentrated, and frozen in liquid nitrogen prior to storage at -80°C. The fractions were thawed (or never frozen) and concentrated for size exclusion chromatography on a Superdex200 10/300 (GE Healthcare) preequilibrated in 10mM

Tris pH 8, 150mM NaCl, 1mM DTT.

Ypt1 Purification

GST-Ypt1-7xHis (gift from T. Bretscher lab) or GST-Ypt1 (pLT50,(Thomas and Fromme, 2016)), with a cleavable N- terminal GST tag in the pGEX-6P vector backbone, were transformed into the Rosetta2 strain of *E.coli* (Novagen). 1-4 liters of culture were grown in terrific broth for 8-12h at 37°C (until the OD₆₀₀ ~2-3). The temperature was then reduced to 16°C and 1 hour later IPTG was added to a final concentration of 300μM to induce protein expression. Following overnight expression (14-18h), cells were collected via centrifugation and resuspended in lysis buffer (1x PBS, 5mM BME, 2mM MgCl₂). Cells were lysed by sonication and the clarified lysate was incubated with Glutathione resin (G-Biosciences) for 2–3h at 4°C to isolate GST-tagged proteins. The resin was washed in lysis buffer, resuspended in PreScission cleavage buffer (50mM Tris pH 7.5, 150mM NaCl, 1mM EDTA, 2mM MgCl₂, and 1mM DTT), and treated overnight with PreScission (3C) protease (~40uL at 1.3mg/mL) at 4°C to remove the GST tag. Cleavage of the GST tag eluted the proteins from the resin. After treatment, the supernatant was collected and analyzed by SDS-PAGE.

TRAPPIII-Rab1/Ypt1 complex formation

Purified TRAPPIII complex (from anionic exchange step) and Ypt1-7xHis were mixed in a 1:3 molar ratio and incubated overnight at 4°C with calf intestinal alkaline phosphatase (Sigma) to hydrolyze nucleotide. After 12-18h, the sample mixture was subjected to size exclusion chromatography on a Superdex 200 Increase 10/300 column (GE Healthcare) in EM buffer (10mM Tris pH 7.5, 350mM NaCl, 1mM DTT). The

trace displayed two dominant peaks corresponding to TRAPPIII-Rab1/Ypt1 complex and free Rab1/Ypt1 (Figure S1). The fractions containing the complex were pooled and concentrated in a 100kDa molecular weight cutoff concentrator (Millipore). The protein was then used immediately to prepare grids for cryo-EM.

Cryo-EM sample preparation, data collection, and data processing

The complex, at 0.1-0.3mg/mL in EM buffer containing either 0.02% Tween-20 or 0.025% amphipol A8-35 (Anatrace) was applied to Quantifoil or UltrAufoil R1.2/1.3 grids (Quantifoil) then blotted for 2s at 4°C with 100% humidity and plunged into liquid nitrogen-cooled liquid ethane using a Vitrobot IV (TFS).

“Flat” data:

Initial imaging data were collected at the MRC-LMB on a Titan Krios operated at 300kV using a Falcon-III detector and 59kX nominal magnification resulting in a 1.34Å/pixel size on the detector. A total of 809 fractionated exposures were obtained over two different 24-hour data collection sessions. RELION versions 2, 3, and 3.1 (Scheres, 2012; Kimanius et al., 2016; Zivanov et al., 2018, 2020) were used for all data processing steps. MotionCor2 (Zheng et al., 2017) was used to perform motion-correction and dose-weighting, and GCTF (Zhang, 2016) was used to estimate micrograph defocus values. After triaging micrographs, particles were autopicked and 2D-classification was used to remove junk particles. The negative-stain structure of the TRAPPIII complex (Tan et al., 2013) was initially used as a reference model for 3D-refinement. After iterative rounds of 3D-classification and 3D-refinement, it was evident

that the resulting 3D-reconstructions were quite anisotropic due to a strong orientation bias of the particles. Alternative particle-picking strategies did not provide additional orientations and therefore we decided to collect data from tilted grids.

30°-tilt data:

A dataset of 430 micrographs was collected on the same microscope with the stage tilted to 30°. An additional 2015 micrographs were collected at the Diamond Light Source / eBIC on a Titan Krios operated at 300kV with a K2 detector and energy filter (Gatan). The image pixel size on the K2 detector was 0.824Å/pixel after binning super-resolution images. Data were processed as described above for the “flat” data, with an additional step of per-particle defocus estimation using GCTF prior to 2D classification.

Each dataset was initially processed independently, producing reconstructions of ~4.5Å resolution using the 0.143 FSC criterion. The datasets were then processed together, subjected to multiple 3D-classification and 3D-refinement procedures, and after CTF-Refinement and Bayesian Polishing ultimately resulted in a 3D-reconstruction with a 0.143 FSC resolution of 3.8Å. However, this 3D-reconstruction was still fairly anisotropic and interpretation of the distal ends of the map was further confounded by some flexibility of the complex that could not be surmounted by focused local refinements or classifications. We therefore decided to collect additional data at even higher tilt angles, as the Lyumkis group demonstrated that 40° tilt was feasible for data collection (Tan et al., 2017).

40°-tilt and 45°-tilt data:

The Cornell CCMR facility Talos Arctica (TFS) with K3 detector and energy filter (Gatan) was used to collect 3062 micrographs on 45°-tilted grids and 1950 micrographs on 40°-tilted grids. The microscope was operated at 200 kV at a nominal magnification of 63kX resulting in an image pixel size of 1.24 Å/pixel. Only a small fraction of these micrographs was suitable for use, either due to extensive beam-induced motion that could not be corrected, or insufficient contrast. CryoSparrc (Punjani et al., 2017) was used to triage micrographs after “Patch-CTF” micrograph estimation. CryoSparrc was also used for 2D classification and initial 3D-classifications to generate a clean particle set that was then imported into RELION with the use of the csparrc2star python script written by D. Asarnow (<https://github.com/asarnow/pyem>). RELION 3.1 was used to combine the particles imaged at 30°, 40°, and 45°, using the scheme shown in Figure S3. Extensive 3D-classification was used to select the best set of particles. In order to deal with apparent flexibility of Trs85 relative to the rest of the complex, we employed Trs85-focused particle subtraction and 3D-classification. CTF-refinement and Bayesian polishing were also used, culminating in a final 3D reconstruction with a 0.143 FSC resolution of 3.7Å.

Density modification using RESOLVE (Terwilliger et al., 2020) in Phenix (Liebschner et al., 2019) was used to facilitate interpretation of the reconstruction. The resulting map was reported by RESOLVE to have a resolution of 3.5Å. In addition, focused local refinement was performed separately on Trs85, the distal region surrounding Trs33, and the central region surrounding Rab1/Ypt1 to generate additional maps to assist with model building.

Particle orientation distributions were assessed and plotted using scripts from A. Leschziner (<https://github.com/leschzinerlab/Relion>) and D. Lyumkis et al. (<https://github.com/nysbc/Anisotropy>).

Model building and refinement

Available crystal structures (Kim et al., 2005; Rak et al., 2003; Cai et al., 2008) for the core subunits and Rab1/Ypt1 were docked into the map and rebuilt using Coot (Emsley et al., 2010). trRosetta (Yang et al., 2020) was used to generate models of Trs85, and portions of these models were docked or used as guides for *de novo* model building. Phenix was used for real-space refinement (Liebschner et al., 2019). Restrained molecular dynamics simulations were performed using ISOLDE (Croll, 2018) within ChimeraX (Goddard et al., 2018) in order to identify and fix model errors arising due to imposition of Ramachandran restraints during refinement. The final refinement was carried out using one half-map and validation was performed using the other half-map. The model-map 0.143 FSC values were: 3.1Å (whole-map), 3.2Å (refinement half-map), and 3.7Å (validation half-map). Model-map FSC curves are shown in Figure S2 and other validation, model, and data statistics are shown in Table 1. PyMol (Schrodinger), Chimera (Pettersen et al., 2004), and ChimeraX were used to produce images for figures.

Surface analyses

Conservation analysis was carried out using the Consurf web server utilizing custom alignments generated with Clustal Omega (Armon et al., 2001; Landau et al.,

2005; Ashkenazy et al., 2010; Celniker et al., 2013; Ashkenazy et al., 2016). The electrostatic surface potential was calculated using the APBS plugin for Pymol (Schrodinger) and the map was colored on a red (-5kT/e) to blue (+5kT/e) scale (Baker et al., 2001; Dolinsky et al., 2007).

Sequence alignment

The multiple sequence alignments were performed using Clustal Omega with default settings (Madeira et al., 2019) and then manually adjusted for fine-tuning of the amphipathic helix region.

Cross-linking mass spectrometry (XL-MS)

~100ug of purified TRAPPIII-Rab1/Ypt1 complex (in 10mM HEPES pH 8, 150mM NaCl, 1mM DTT) was incubated on ice for 1hr with a range of disuccinimidyl sulfoxide (DSSO) concentrations spanning 0.3125mM to 2.5mM. The reaction was then quenched with 10mM Tris pH 8 for 15 minutes at room temperature. 10% of the sample was analyzed by SDS-PAGE (Figure S1B), and the sample cross-linked with 625μM DSSO was chosen for further analysis. To prepare the sample for mass spectrometry, the cross-linked proteins were first dried completely in a vacuum concentrator and then resuspended in 8M Urea, 50mM Tris pH 8, and 5mM DTT. Urea-solubilized proteins were then incubated at 37°C for 30min for denaturation and to reduce disulfide bridges. Reduced cysteines were blocked with 25mM iodoacetamide for 15min. Urea was then diluted to 2M with a solution of 50mM Tris pH 8 and 150mM NaCl. Proteins were digested overnight with gentle nutation at 37°C using 1ug of Promega Trypsin GOLD®.

The next day, the digested sample was acidified with 0.25% formic acid + 0.25% trifluoroacetic acid and desalted using a Waters Sep-Pak column. The desalted, digested sample was dried in a vacuum concentrator and reconstituted in 0.1% TFA + 1 picomole/uL Angiotensin-II peptide (Sigma). 1% of the sample was subjected to LC-MS/MS/MS analysis on an Orbitrap Fusion Lumos mass spectrometer (ThermoFisher Scientific). On-line sample fractionation was performed with linear gradients of 0.1% formic acid (solvent A) and 80% acetonitrile + 0.1% formic acid (solvent B) at a flow rate of 300 nl/min using a home-made 125- μ m x 25-cm capillary column packed with 3- μ m C18 resin. Precursor ions were detected in the Orbitrap mass analyzer (375-1500Th, resolution 60,000 at 375Th). Charge states in the 4+ to 8+ range were selected for MS2 analysis in the Orbitrap (resolution set at 30,000 at 375Th) with 25% CID energy. Peaks with a mass difference of 31.9721Da, indicative of a DSSO cross-link, were subjected to MS3 analysis. For MS3, ions were fragmented in the linear ion trap using higher-energy collisional dissociation (HCD) at 35% HCD energy. The cross-link search was performed using MaXLinker software (Yugandhar et al., 2020). The structure mapping of cross-links was performed using Xlink Analyzer (Kosinski et al., 2015) implemented in UCSF Chimera (Pettersen et al., 2004).

Liposome preparation

Synthetic liposomes were prepared using a mixture of lipids (Table S4). Lipids were combined, dried by rotary evaporation, rehydrated overnight in HK buffer (20mM HEPES, pH 7.4, and 160mM KOAc) at 37°C, then extruded through 100nm or 400nm

filters (Whatman) to generate homogenous liposomes. Synthetic liposomes were stored at 4°C until needed.

Pelleting assays

In 40uL reactions, loaded ~1-2ug of protein (300nM) along with 500μM of 400nm liposomes in HK buffer. Incubated for 15 minutes at RT. Samples were ultracentrifuged for 10 minutes at 4C, 55K rpm (TLA 100.3 rotor). The supernatant and pellet fractions were recovered and evaluated separately via SDS-PAGE. Each assay was performed at least three times. The gels shown are representative of all three experiments. The percent of pelleted protein is calculated by dividing the intensity of the pelleted bands by the total intensity of protein in both the supernatant and pelleted fractions. Then the average amount of protein that pellets in the absence of liposomes is subtracted from each sample. The quantification is based on the averages of all three experiments. All direct comparisons were performed using the same batch of liposomes to account for batch variability.

5-FOA complementation assay

Corresponding plasmids harboring mutant versions of Trs85 were transformed into a sensitized strain background (*trs85Δ::KAN*, Bet3-GFP::HIS (CFY3307)), grown on -LEU plates. Several colonies were resuspended in water then serially diluted and replicated onto -LEU and 5FOA plates. Cells were grown for 48 hours at 26, 30, or 37°C.

Microscopy

Plasmids harboring mutant versions of Trs85 were transformed into a *trs85Δ::KANMX* strain (CFY2692) so that only the plasmid copy of Trs85 would be present. For experiments assessing Rab1/Ypt1 localization, a second plasmid containing mRFPmars-Ypt1 (CFB3511) was co-transformed. Cells were grown at 30°C until log-phase. Cells were collected by centrifugation for 2 minutes at 12,000x g, resuspended in a small amount of media and placed on a microscope slide with coverslip. Cells were imaged at room temperature with a CSU-X spinning-disk confocal microscope system (Intelligent Imaging Innovations) on a DMI6000 microscope (Leica Microsystems) equipped with a CSU-X1 spinning-disk confocal unit (Yokogawa Electric Corporation) and a QuantEM 512SC camera (Photometrics). The objective was a 100× 1.46 NA Plan Apochromat oil immersion lens (Leica Microsystems). Additional components included a laser stack and mSAC (spherical aberration correction; Intelligent Imaging Innovations). SlideBook software was used to control the system (Intelligent Imaging Innovations) and for data analysis. All images within a figure had their minimum or maximum brightness levels adjusted equally for clarity.

Co-immunoprecipitations

Plasmids harboring Trs85 mutants were transformed into a yeast strain containing *trs85Δ::KANMX*, Trs23-6xHis-3xFLAG::TRP (CFY3585). Overnight cultures were grown to saturation and then used to inoculate fresh media. Large cultures grew until log-phase and then cells were collected by centrifugation. Cell pellets were

resuspended in 1mL cold water and then transferred to an Eppendorf tube. Cell pellets were aliquoted to approximately equal weights and then the cells were collected again, flash frozen in liquid nitrogen, and stored at -80°C. Each colP contained roughly 25OD₆₀₀ of cells per tube.

Upon removal from storage, 250uL of lysis buffer [50mM Tris pH 7.5, 0.2% NP40, 150mM NaCl, 1mM EDTA] and 100uL of 0.5mm glass beads were added to each tube. Cells were immediately lysed via harsh vortexing for 2x10min at 4°C, with 2min rest on ice in between. Lysed cell debris was pelleted for 5min at ~21K x g at 4°C and the supernatant was collected. 25uL supernatant was used for SDS-PAGE analysis as the whole cell extract. The remaining lysate was incubated with 12.5uL equilibrated magnetic mNeonGreen-Trap resin (Chromotek) for a minimum of 1 hour rotating at 4°C. After incubation, the resin was collected and washed 3x 1mL with lysis buffer and transferred once to a new tube. Finally, 50uL of 2x SDS sample buffer was added to the resin and incubated at least 10min at 95°C. Samples were subjected to SDS-PAGE and transferred to a PVDF membrane (Millipore) for Western blot analysis probing for HA and FLAG epitopes using 1° antibody = 1:500 mouse αHA (Roche), or 1:500 mouse αFLAG (Roche) and 2° 1:5,000 sheepαmouse (Amersham).

GFP-Atg8 processing assay

Plasmids harboring Trs85 mutants were co-transformed along with GFP-Atg8 (CFB2250) into *trs85Δ::KAN* (CFY2692). Overnight cultures were grown to saturation at 30°C and then used to inoculate fresh media. Once back in log-phase, ~5-10OD₆₀₀ units

of cells were collected cells by centrifugation, 3min at ~3k rpm and processed for TCA precipitation. With the remaining culture, autophagy was induced by addition of rapamycin to a concentration of 0.2 μ M. After 1h of incubation at growth temperature, an additional ~5-10OD₆₀₀ units of cells were collected and processed for TCA precipitation. Samples were resolved by SDS-PAGE and then subjected to Western blot analysis using 1°, 1:1,000 mouse α GFP (Roche) and 2°, 1:5,000 sheep α mouse (Amersham). Autophagy efficiency is quantified measuring the amount of free GFP compared to the total amount of GFP-Atg8 and GFP. The blots shown are representative of all three experiments. The percent of free GFP is calculated by dividing the intensity of the free GFP band by the total intensity of both GFP-Atg8 and free GFP in the rapamycin-induced lane. The quantification is based on the averages of all three experiments.

Prenylated Rab1/Ypt1 preparation

Performed as described previously (Thomas et al., 2018). Briefly, Rab/GDI substrates, the mantGDP-loaded Rab, Gdi1, His₆-Mrs6, and His₆-Bet2/Bet4 were mixed in a 10:10:1:1 molar ratio with a 6-fold excess of geranylgeranyl pyrophosphate (Cayman) in prenylation buffer with 20 μ M mantGDP and incubated at 37°C for 1h. Then, imidazole was added for a final concentration of 10mM. The solution was incubated with a small volume of Ni-NTA resin (Qiagen) at 4°C for 1h to remove enzymes. Stoichiometric Rab/GDI complexes were then isolated using size exclusion chromatography.

Rab1/Ypt1 nucleotide loading for GEF assays

~1mg of purified Rab1/Ypt1 (either prenylated-Ypt1/GDI complex or non-prenylated Ypt1-7xHis) was incubated in the presence of 20 μ M mantGDP (Sigma) and EDTA to facilitate exchange for 30 minutes at 30°C. Afterwards, protein was buffer exchanged in 15x excess buffer to remove leftover nucleotide and EDTA before being aliquoted and stored at -80°C.

In vitro nucleotide exchange (GEF) assays

For all assays, nucleotide exchange was measured by quenched fluorescence of mantGDP (Sigma) (360nm excitation and 440nm emission). Reactions were performed inside of a fluorometer (Photon Technology International) at 30°C with the sequential addition of: 333 μ M synthetic liposomes (100nm), 200 μ M nonfluorescent GTP, and ~250nM prenylated mantGDP:Ypt1 to HKM buffer (20mM HEPES pH 7.4, 160mM KoAC, 2mM MgCl₂). The mixture was then incubated at 30°C for 2 min to allow the Rab/GDI substrate time to equilibrate with the membrane. Nucleotide exchange was monitored after the addition of 20nM GEF complex. For soluble mantGDP:Ypt1-7His, identical conditions were utilized except the liposomes were replaced by HKM buffer. To determine exchange rates, traces of fluorescence versus time were fit to a one phase decay curve with an additional linear drift term using Prism (GraphPad Software). The resulting rate constants were divided by the GEF concentration to obtain the exchange rate.

Software

Structural biology software packages are maintained in our laboratory by SBGrid (Morin et al., 2013).

Data deposition

Data will be deposited in the RCSB PDB, EMDB, and PRIDE databases.

AUTHOR CONTRIBUTIONS

AMNJ – conceptualization and design, data acquisition/investigation (cryo-EM, XL-MS, fluorescence microscopy, cell-based assays, in vitro assays), data analysis/interpretation, writing; BPP – data acquisition (cryo-EM); KY – data acquisition (XL-MS); EJS – data acquisition (XL-MS); MBS – funding acquisition, data interpretation; HY – funding acquisition, data interpretation; EAM – funding acquisition, data interpretation; JCF – funding acquisition, conceptualization and design, data analysis/interpretation, writing.

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FIGURE LEGENDS

Figure 1: The Trs85 subunit anchors TRAPPIII to membranes

- A. Cartoon depicting the TRAPPIII complex subunits and the TRAPP core subunits. Trs85 (purple) is the TRAPPIII-specific subunit.
- B. *In vitro* nucleotide exchange assay of prenylated-Rab1/Ypt1-GDI substrate in the presence of synthetic liposomes. The TRAPPIII complex (purple) is significantly more active than the TRAPP core alone (yellow).
- C. Assays performed as in B, but using a non-prenylated Rab1/Ypt1 substrate and in the absence of synthetic liposomes. Exchange activity with and without the Trs85 subunit are equivalent.
- D. *In vitro* membrane pelleting assay comparing the TRAPPIII complex versus the core subunits. While both complexes exhibit a small amount of pelleting in the absence of synthetic liposomes, in their presence only the TRAPPIII complex is strongly enriched in the membrane-bound fraction. (S=supernatant, P=pellet)
- E. Quantification of D. Average % membrane binding between three assays is plotted, the error bars represent the 95% confidence intervals. The value for each replicate is depicted. The negative values arise from subtraction of the average amount of protein that pellets in the absence of liposomes.

Figure 2: Cryo-EM structure of the TRAPPIII-Rab1/Ypt1 nucleotide exchange complex

- A. Cryo-EM density map colored by each subunit. Bottom panel is rotated 90° relative to the top panel.

- B. Colored as in A, the refined atomic model of the TRAPPIII complex.
- C. Network map depicting all of the 108 cross-links found between the TRAPPIII complex subunits and Rab1/Ypt1. Blue lines are indicative of inter-protein cross-links, red lines denote cross-links detected within a single protein. The relative position of the cross-link represents its location within the primary sequence.
- D. The fraction of total cross-linking sites observed in each polypeptide.
- E. The cross-links mapped onto the refined model. Cross-links are shown as black lines connecting black spheres.
- F. A histogram depicting the length of each cross-link that was mapped to the structural model. The distance was measured between the α -carbons of each pair of cross-linked residues. Only one of the cross-links is beyond the maximum theoretical restraint (30Å) of the cross-linking chemical, DSSO.

Figure 3: Architecture and conservation of the Trs85 subunit

- A. The Trs85 subunit colored from N- (blue) to C-terminus (red). Several features are highlighted and described in the text.
- B. Surface view of Trs85 colored by ConSurf analysis. On the left is the conservation across model organisms and humans, on the right conservation was estimated based on more closely related species. The TRAPP core binding region is highly conserved.

Figure 4: The C-terminus of Trs85 binds to the TRAPP core

- A. Representative Coomassie-stained gel of size exclusion chromatography fractions after purification of recombinant wild-type TRAPPIII complex.
- B. As in A, using a Trs85 mutant containing only the final 198 amino acids. NOTE: Trs85[501-698] is ~25kD in size and now migrates at the same size as some of the core subunits.
- C. As in A/B, using a Trs85 mutant with a truncation of the loop from 575-603. Trs85[Δ575-603] appears to dissociate from the core during chromatography, indicating a reduced affinity for the core subunits.
- D. Network map showing the relative positions of cross-links between the C-terminus of Trs85 and the core subunits Trs31 and Trs20.
- E. View of the cryo-EM structural model showing the interaction surface between Trs85, Trs31 and Trs20. Inset depicts several basic residues from Trs85 that are in close proximity to several acidic residues of Trs20.
- F. Yeast complementation assays performed in a sensitized strain (*BET3-GFP::HIS3, trs85Δ::KANMX*) to assess the functionality of Trs85 mutants within the region shown in E.
- G. Coimmunoprecipitations of Trs23-6xHis-3xFLAG with the same Trs85 mutants.
- H. Live cell fluorescence microscopy of Trs85-mNeonGreen and mRFPmars-Ypt1. Localization of Rab1/Ypt1 to punctate structures is indicative of its activation by TRAPPIII. Scale bar is 2 microns.

Figure 5: The Rab1/Ypt1 HVD points to the membrane binding surface

- A. Cryo-EM density is shown at a low threshold to highlight the almost continuous density connecting the Rab1/Ypt1 nucleotide-binding domain to its HVD. Inset is a closeup of the density corresponding to the Rab1/Ypt1 HVD region. Colored by subunit as in Figure 2, Rab1/Ypt1 is yellow. Density is depicted as it is likely oriented at the surface of organelle membranes.
- B. 2D class averages showing extra density in this region in the presence of Rab1/Ypt1. Circle highlights the Rab1/Ypt1 NBD, and the arrow points to the HVD density.
- C. Network diagram of cross-links observed between Trs31 and Rab1/Ypt1. Many cross-links are observed at the C-terminal region of Rab1/Ypt1. Cross-links involving residues present in the atomic model are colored black, while cross-links between unmodeled residues are colored pink.
- D. As in C, but the cross-links are mapped onto the structure. Cross-links involving residues present in the atomic model are colored black, while cross-links between unmodeled residues are colored pink.
- E. Surface representation of the TRAPPIII-Rab1/Ypt1 complex colored by electrostatic potential. Top panel is oriented as in A, the bottom panel is rotated 90° to visualize the most positive surface which likely lies at the surface of the membrane.
- F. Quantification of *in vitro* TRAPPIII complex membrane pelleting assays performed with various lipid compositions. The strongest binding was observed

with the most negatively charged liposomes. Binding is compared to wild-type on “Golgi” liposomes using t-test with Welch’s correction, * $p < 0.05$, ** $p < 0.01$.

Figure 6: TRAPPIII uses a conserved amphipathic helix to bind the membrane

- A. Refined atomic model of the TRAPPIII complex shown as it likely interacts with membranes. The arrow points to the location of the unmodeled conserved region.
- B. Sequence alignment of the unmodeled region predicted to lie on the membrane-binding surface. Red helix above represents the region predicted to fold as an α -helix in secondary structure analysis. Positively charged residues are highlighted in blue, while bulky aromatic residues are underlined.
- C. Helical wheel representation of the region predicted to be an α -helix. Positively charged residues are colored blue, polar uncharged residues are colored green, and nonpolar residues are colored yellow.
- D. The wild-type Trs85 loop sequence is shown on top, with targets of mutation colored purple and mutants color coded.
- E. Yeast complementation assays performed in a sensitized strain (*BET3-GFP::HIS3*, *trs85Δ::KANMX*) to assess the functionality of Trs85 mutants.
- F. Coimmunoprecipitations of Trs23-6xHis-3xFLAG with the same Trs85 mutants.
- G. *In vitro* membrane pelleting assay examining two Trs85 mutants in the context of the TRAPPIII complex. In the absence of liposomes, all three complexes pellet slightly, however in the presence of membranes, only the wild-type Trs85-

TRAPPIII complex is strongly enriched in the pelleted fraction. (S=supernatant, P=pellet)

- H. Quantification of H. Average % protein pelleted between three assays is plotted, the error bars represent the 95% confidence intervals. The value for each replicate is depicted. The negative values arise from subtraction of the average amount of protein that pellets in the absence of liposomes.
- I. Live cell fluorescence microscopy of Trs85-mNeonGreen mutants and mRFPmars-Ypt1. Scale bar = 2 μ m.
- J. *In vitro* nucleotide exchange assay of prenylated-Ypt1/GDI substrate in the presence of synthetic liposomes. The wild-type Trs85-TRAPPIII complex (purple) is significantly more active than the mutants (red and black) or the core alone (yellow).
- K. Assays performed as in B, but using a non-prenylated Rab1/Ypt1 substrate and in the absence of liposomes. Exchange activity of all complexes are equivalent.

Figure 7: The membrane-binding helix is critical for autophagy

- A. GFP-Atg8 processing is monitored for either the TRAPP core binding mutations or the amphipathic helix (membrane binding) mutations. With functional Trs85, there is an accumulation of free GFP. Samples are from pre-rapamycin treatment (-) and 1hr after addition of rapamycin (+).
- B. Quantification of A. The percent of free GFP for wild-type Trs85 is compared to each mutant using t-test with Welch's correction, ** $p < 0.01$, *** $p < 0.005$. The

average of at least three replicates is plotted with error bars representing the 95% confidence interval.

- C. Model for TRAPPIII-mediated Rab1 activation in cells. Inactive Rab1-GDP/GDI exists in an equilibrium where Rab1-GDP samples various organellar membranes. TRAPPIII is recruited to specific membranes containing negative charges (red circles) and utilizes an amphipathic helix to bind stably to the membrane surface. Rab1 is locally activated by TRAPPIII, then recruits downstream effectors to facilitate trafficking or autophagic events.

Figure S1: Purification of the complex and cross-linking for mass spectrometry

- A. Normalized UV trace from SEC to separate TRAPPIII-Rab1/Ypt1 complex from excess Rab1/Ypt1. Coomassie-stained gel corresponding to the peaks in the trace.
- B. Coomassie-stained gel showing the cross-linked TRAPPIII-Rab1/Ypt1 protein sample over a range of DSSO concentrations. The 625 μ M sample (third lane from the right) was ultimately used for MS3 analysis and database searches.

Figure S2: Cryo-EM data

- A. Example micrograph generated from a flat dataset acquisition, 1.34 $\text{\AA}$ /pixel, -3.0 microns estimated average defocus.
- B. Representative 2D class averages from the flat dataset generated in RELION. The edge of each box corresponds to 482 $\text{\AA}$. ~95% of the particles are found in the 2D classes equivalent to the Ypt1-bound class shown in Figure 5B.

- C. Example micrograph generated from a 40° tilt dataset acquisition, 1.24Å/pixel, - 1.5 microns estimated average defocus.
- D. Representative 2D class averages from the combined tilt datasets generated in RELION. The edge of each box corresponds to 397Å.
- E. FSC curves for each of the final refined particle sets used for 3D reconstructions (see Figure S3).
- F. Orientation distributions of each of the final refined particle sets used for 3D reconstructions (see Figure S3). Generated using a script written by A. Leschziner available on GitHub.

Figure S3: Cryo-EM data processing flowchart

Flow chart showing the steps taken during data processing to generate the high-resolution TRAPPIII-Rab1/Ypt1 3D reconstruction. “Sphericity” is a measure of data isotropy (lower values represent more anisotropy) and was calculated using a 3DFSC script written by D. Lyumkis available on GitHub. The insets show the EM density from each of the final 3D reconstructions, surrounding the portion of Trs85 that contacts Trs20.

Figure S4: Comparisons to published crystal structures

- A. Bet3 subunit from our cryo-EM reconstruction aligned with the Bet3 subunit from pdb:3CUE.
- B. The second Bet3 subunit from our cryo-EM reconstruction aligned to the second Bet3 subunit from pdb:3CUE.

- C. Bet5 subunit from our cryo-EM reconstruction aligned with the Bet5 subunit from pdb:3CUE.
- D. Trs23 subunit from our cryo-EM reconstruction aligned with the Trs23 subunit from pdb:3CUE.
- E. Trs31 subunit from our cryo-EM reconstruction aligned with the Trs31 subunit from pdb:3CUE.
- F. Trs33 subunit from our cryo-EM reconstruction aligned with the TRAPPC6 (Trs33) subunit from pdb:2C0J.
- G. Trs20 subunit from our cryo-EM reconstruction aligned with the Sedlin (TRAPPC2, Trs20) subunit from pdb:1H3Q.
- H. Rab1/Ypt1 subunit from our cryo-EM reconstruction aligned with the Rab1/Ypt1 subunit from pdb:3CUE.

Table S1: TRAPP complex composition

Note: Both TRAPP II and TRAPP III share a core set of subunits (green, middle column) and then contain several unique ones. The yeast names are shown first, with the names of the human proteins in parentheses.

TRAPP III-specific	Shared Core Subunits	TRAPP II-specific
Trs85 (TRAPPC8)	Trs33 (TRAPPC6A, B)	Trs130 (TRAPPC10)
(TRAPPC11)	Trs31 (TRAPPC5)	Trs120 (TRAPPC9)
(TRAPPC12)	Trs23 (TRAPPC4)	Trs65
(TRAPPC13)	Trs20 (TRAPPC2)	Tca17 (TRAPPC2L)
	Bet5 (TRAPPC1)	
	Bet3 (TRAPPC3, C3L)	

Table S2: Cryo-EM model validation statistics

Composition (#)	
Chains	10
Atoms	14935 (Hydrogens: 0)
Residues	1836
Water	0
Ligands	2 palmitate molecules
Bonds (RMSD)	
Length (Å) (# > 4 σ)	0.005 (0)
Angles (°) (# > 4 σ)	0.964 (1)
MolProbity score	2.3
Clash score	17.83
Ramachandran plot (%)	
Outliers	0.06
Allowed	10.09
Favored	89.85
Rama-Z (Ramachandran plot Z-score, RMSD)	
whole (N = 1754)	2.24 (0.19)
helix (N = 842)	0.63 (0.17)
sheet (N = 231)	1.60 (0.33)
loop (N = 681)	2.45 (0.23)
Rotamer outliers (%)	0
C β outliers (%)	0
CaBLAM outliers (%)	1.79
ADP (B-factors)	
Iso/Aniso (#)	14935/0
min/max/mean	
Protein	87.59/179.40/123.81
Ligand	103.94/151.27/127.61
Water	---
Resolution Estimates (Å)	
FSC half-maps (0.143/0.5)	3.7/4.2
FSC model/map (0.143/0.5)	3.1/3.8
Model vs. Data	
CC (mask)	0.82
CC (box)	0.88
CC (peaks)	0.76
CC (volume)	0.82
Mean CC for ligands	0.72

Table S3: All 108 unique cross-links identified via MaxLinker database search.

Peptide A	Protein A	Protein A Position	Peptide B	Protein B	Protein B Position
AMGEEIWkNKTEK	Bet3	19	IYSESLLFkR	Trs31	53
AMGEEIWkNKTEK	Bet3	19	ASAFLSQNESSSkLSNAS NSPGMLANSSTATSASA NER	Trs31	128
AMGEEIWkNK	Bet3	19	GDDFFkEPFTNWNK	Trs23	108
NkTEK	Bet3	21	kVYcLYSDYVMK	Trs23	181
AMGEEIWkNKTEK	Bet3	21	GSVkNDIR	Bet5	64
DFNkVNDHLYSMGYNIGc R	Bet3	51	TckTAGDFETK	Trs31	78
cENLVkTSEVLSK	Bet3	85	GSVkNDIR	Bet5	64
cENLVkTSEVLSK	Bet3	85	SLkAMGEEIWK	Bet3	11
cENLVkTSEVLSK	Bet3	85	QPTIATTDkPDRPK	Trs23	160
GDSQTEIkVK	Bet3	177	SDGYEYTVGPKQAITSEA STTYIPSR	Trs31	29
VkLNR	Bet3	179	LkIEEHHTVDIIR	Trs33	89
VkLNR	Bet3	179	IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18
LSkGSVK	Bet5	60	cENLVkTSEVLSK	Bet3	85
SFYQEVHELyVvKTLMPF YK	Trs20	143	LQEkQTESLSNYITK	Trs31	157
TLMNPFYKITDPIR	Trs20	151	KDNPVYEIEFTNAENPQG FPQDLK	Trs20	11
kHLSK	Trs20	171	LQEkQTESLSNYITK	Trs31	157
kHLSK	Trs20	171	QTESLSNYITkMR	Trs31	168
kHLSK	Trs20	171	IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18
NGGGNNkHTNNEK	Trs23	91	LQEkQTESLSNYITK	Trs31	157
NGGGNNkHTNNEK	Trs23	91	AMGEEIWkNKTEK	Bet3	19
NGGGNNkHTNNEK	Trs23	91	TckTAGDFETK	Trs31	78
NGGGNNkHTNNEK	Trs23	91	IYSESLLFkR	Trs31	53
NGGGNNkHTNNEK	Trs23	91	cENLVkTSEVLSK	Bet3	85
nGGGNNkHTNNEK;NG GGnNNkHTNNEK	Trs23	91	QPTIATTDkPDRPK	Trs23	160
NGGGNnNkHTNNEK;nG GGNNNkHTNNEK	Trs23	91	LGSFKGDDFFkEPFTNW NK	Trs23	108
NGGGNNkHTNNEK	Trs23	91	kMVENLQ	Trs23	213
HTNNEkLGSFKGDDFFK EPFTNWNK	Trs23	97	SNLFDEkVKK	Trs23	210
HTNNEkLGSFKGDDFFK EPFTNWNK	Trs23	97	SNLFDEkVvK;SNLFDEkV KK	Trs23	212
LGSFKGDDFFkEPFTNW NK	Trs23	102	NGGGNNkHTNNEK	Trs23	91
LGSFKGDDFFkEPFTNW NK	Trs23	102	AMGEEIWkNK	Bet3	19

LGSFkGDDFFKEPFTNW NK	Trs23	102	NkTEK	Bet3	21
LGSFkGDDFFKEPFTNW NK	Trs23	102	IYSESLLFkR	Trs31	53
GDDFFKEPFTNWNkSGL R	Trs23	116	cENLVkTSEVLSK	Bet3	85
LGSFkGDDFFKEPFTNW NkSGLR	Trs23	116	ILkDEIPIGED	Bet3	185
QPTIATTDkPDRPK	Trs23	160	GVLSSLGYSSEEVlcLASF IDRPTDRPkTAFPK	Trs33	251
SNLFDEkVKK;SNLFDEK VKK	Trs23	210	nGGGnNNkHTNNEK	Trs23	91
SNLFDEkVK	Trs23	210	kVYcLYSDYVMK	Trs23	181
SNLFDEkVKK;SNLFDEkV KK	Trs23	212	nGGGNNNkHTNNEK;NG GGNNNkHTnNEK	Trs23	91
SNLFDEkVKK;SNLFDEkV KK	Trs23	212	kVYcLYSDYVMK	Trs23	181
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	TSEVLSkcAFK	Bet3	92
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	cENLVkTSEVLSK	Bet3	85
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	LkNQLNLK	Trs85	670
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	GDSQTEIkVK	Bet3	177
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	TLMNPFYkITDPIR	Trs20	151
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	SLkAMGEEIWK	Bet3	11
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	ASAFLSQNESSSkLSNAS NSPGMLANSSTATSASA NER	Trs31	128
IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18	QTESLSNYITkMR	Trs31	168
TckTAGDFETK	Trs31	78	cENLVkTSEVLSK	Bet3	85
TckTAGDFETK	Trs31	78	TVELDGkTVK	Ypt1	55
TckTAGDFETK	Trs31	78	ESMSQQNLNETTQkKED K	Ypt1	184
TckTAGDFETK	Trs31	78	QIkESMSQQNLNETTQK	Ypt1	170
TckTAGDFETK	Trs31	78	EDkGNVNLK	Ypt1	188
ASAFLSQNESSSkLSNAS NSPGMLANSSTATSASA NER	Trs31	128	SLkAMGEEIWK	Bet3	11
ASAFLSQNESSSkLSNAS NSPGMLANSSTATSASA NER	Trs31	128	TSEVLSkcAFK	Bet3	92
ASAFLSQNESSSkLSNAS NSPGMLANSSTATSASA NER	Trs31	128	IYSESLLFkR	Trs31	53
LQEkQTESLSNYITK	Trs31	157	IYSESLLFkR	Trs31	53
LQEkQTESLSNYITK	Trs31	157	ASAFLSQNESSSkLSNAS NSPGmLANSSTATSASA NER	Trs31	128

LQEkQTESLSNYITK	Trs31	157	SDGYEYTVGPKQAITSEA STTYIPSR	Trs31	29
LQEkQTESLSNYITK	Trs31	157	IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18
QTESLSNYITkMR	Trs31	168	ASAFLSQNESSSkLSNAS NSPGmLANSSTATSASA NER	Trs31	128
QTESLSNYITkMR	Trs31	168	SDGYEYTVGPKQAITSEA STTYIPSR	Trs31	29
QISGDVISSDSNVTS ENG NINNMikR	Trs33	86	VkLNR	Bet3	179
LkIEEHHTVDIIR	Trs33	89	ILkDEIPIGED	Bet3	185
FKEMDLLLIMkFicR	Trs33	158	QIDNLkTNHR	Trs33	177
QIDNLkTNHR	Trs33	177	GVLSLGSSEEVIcLASF IDRPTDRPkTAFPK	Trs33	251
FFGGcVSDRDQANEAKV GQHEHEVcDASDSTDSIP K	Trs85	81	NkNLEVPNLSK	Trs85	103
VGQHEHEVcDASDSTDSI PkNK	Trs85	101	NLEVPNLSkK	Trs85	112
cQALAKK	Trs85	278	NkNLEVPNLSK	Trs85	103
cQALAKK	Trs85	278	NLEVPNLSkK	Trs85	112
kFMNK	Trs85	382	KSLFNGAKFFK	Trs85	378
kFMNK	Trs85	382	kSLFNGAK	Trs85	371
GKGkESNSLDVTEPVR	Trs85	502	LIAAKK	Trs85	631
VGAMkK	Trs85	586	NLEVPNLSkK	Trs85	112
DAkDTEdLRGEHK	Trs85	590	VGAMkK	Trs85	586
kWAEQK	Trs85	632	GKGkESNSLDVTEPVR	Trs85	502
WAEQkQWR	Trs85	637	NQLNLkDVDSAPRPSEK NLTR	Trs85	687
WAEQkQWR	Trs85	637	LIAAKK	Trs85	631
DIESTYSEIkFLHGnGLILS K	Trs85	657	LkNQLNLK	Trs85	670
DIESTYSEIkFLHGnGLIL SK	Trs85	657	DAkDTEdLRGEHK	Trs85	590
DIESTYSEIkFLHGnGLIL SK	Trs85	657	TLMNPFYkITDPIR	Trs20	151
DIESTYSEIkFLHGnGLIL SK	Trs85	657	kDNVPVYEIEFTNAENPQG FPQDLK	Trs20	11
DIESTYSEIkFLHGnGLIL SK	Trs85	657	IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18
FLHGnGLILSkLK	Trs85	668	NQLNLkDVDSAPRPSEK	Trs85	676
FLHGnGLILSkLK	Trs85	668	LIAAKK	Trs85	631
FLHGnGLILSkLK	Trs85	668	WAEQkQWR	Trs85	637
LkNQLNLK	Trs85	670	QVSWcLkDIESTYSEIK	Trs85	647
LkNQLNLK	Trs85	670	DVDSAPRPSEkNLTR	Trs85	687
NQLNLkDVDSAPRPSEK	Trs85	676	WAEQkQWR	Trs85	637
NQLNLkDVDSAPRPSEK	Trs85	676	DIESTYSEIkFLHGnGLILS K	Trs85	657
NQLNLkDVDSAPRPSEK	Trs85	676	LIAAKK	Trs85	631
NQLNLkDVDSAPRPSEK	Trs85	676	QVSWcLkDIESTYSEIK	Trs85	647
DVDSAPRPSEkNLTR	Trs85	687	FLHGnGLILSkLK	Trs85	668

DVDSAPRPSEkNLTR	Trs85	687	DIESTYSEIkFLHGnGLILS K	Trs85	657
NQLNLKDVD SAPRPSEk NLTR	Trs85	687	LIAAkK	Trs85	631
DVDSAPRPSEkNLTR	Trs85	687	QVSWcLkDIESTYSEIK	Trs85	647
DVDSAPRPSEkNLTR	Trs85	687	VGQHEHEVcDASDSTDSI PkNK	Trs85	101
DVDSAPRPSEkNLTR	Trs85	687	IIQPSASDQQFPGkSDGY EYTVGPK	Trs31	18
DVDSAPRPSEkNLTR	Trs85	687	kWAEQK	Trs85	632
IkTVELDGK	Ypt1	48	LLLIGNSGVGkScLLLLR	Ypt1	21
TVELDGkTVK	Ypt1	55	QIkESmSQQNLNETTQK	Ypt1	170
LLVGNkcDLK	Ypt1	122	LLLIGNSGVGkScLLLLR	Ypt1	21
cDLkDKR	Ypt1	126	LLLIGNSGVGkScLLLLR	Ypt1	21
cDLKDKR;cDLkDKR	Ypt1	128	LLLIGNSGVGkScLLLLR	Ypt1	21
ESMSQQNLNETTQkK	Ypt1	184	EDkGNVNLK	Ypt1	188
ESMSQQNLNETTQkK	Ypt1	184	TVELDGkTVK	Ypt1	55
kEDKGNVNLK	Ypt1	185	TckTAGDFETK	Trs31	78
kEDKGNVNLK	Ypt1	185	TVELDGkTVK	Ypt1	55
KEDkGNVNLK	Ypt1	188	VkLNR	Bet3	179

Table S4: Composition of synthetic liposomes used in this study

	Golgi	Golgi no cholesterol	Anionic	No PS/PA	No PI	10% PI₄P
PC (mol %)	30	30	51	36	60	58
PE	10	10		10	10	10
PS	3	3	3		3	
PA	3	3	3		3	
PI	29	29	29	29		
PI₄P	1	1	1	1		10
PI_{4,5}P₂			0.8			
CDP-DAG	2	2	2	2	2	
PO-DAG	4	4		4	4	4
DO-DAG	2	2		2	2	2
Ceramide	5	5		5	5	5
Cholesterol	10		10	10	10	10

Note: All lipids are from Avanti Polar Lipids. PC = phosphatidylcholine, PE = phosphatidylethanolamine, PS = phosphatidylserine, PA = phosphatidic acid, PI = phosphatidylinositol, PI₄P = phosphatidylinositol 4-phosphate, PI_{4,5}P₂ = phosphatidylinositol 4,5-bisphosphate, CDP-DAG = cytidine diphosphate diacylglycerol, PO-DAG = palmitoyl-oleoyl diacylglycerol, DO-DAG = dioleoyl diacylglycerol

Table S5: Plasmid list

Name	Description	Vector	Source
pLT21	Trs33, Trs31, Trs23, Trs20, Bet3, Bet5	pColaDuet-1	(Thomas and Fromme, 2016)
pLT35	6xHisTEV-Mrs6	pET28	(Thomas and Fromme, 2016)
pLT40	GST-Gdi1 with cleavable tag	pGEX-6P	(Thomas and Fromme, 2016)
pLT41	6xHisTEV-Bet2, Bet4	pCDF-Duet-1	(Thomas and Fromme, 2016)
pLT50	GST-Ypt1 with cleavable tag	pGEX-6P	(Thomas and Fromme, 2016)
pRS415	Yeast centromeric vector with LEU2 marker	pRS415	(Sikorski and Hieter, 1989)
Ypt1-7His	GST-Ypt1-7xHis cleavable N-terminal tag	pGEX-6P	(McDonold and Fromme, 2014)
CFB2250	GFP-Atg8	pRS416	(Guan et al., 2001)
CFB2730	6xHis-Trs85[501-698]	pETDuet-1	This study
CFB2868	6xHis-Trs85, Trs33	pETDuet-1	This study
CFB2991	Trs85-mNeonGreen-3xHA	pRS415	This study
CFB3125	Trs85[R618E, R619E, R620E, K621E]-mNeonGreen-3xHA	pRS415	This study
CFB3200	Trs85[R618A, R619A, R620A, K621A]-mNeonGreen-3xHA	pRS415	This study
CFB3222	Trs85[Δ507-603]-mNeonGreen-3xHA	pRS415	This study
CFB3223	6xHis-Trs85[Δ507-603], Trs33	pETDuet-1	This study
CFB3511	mRFPmars-Ypt1 in pRS416	pRS416	This study
CFB3710	Trs85[501-698]-mNeonGreen-3xHA	pRS415	This study
CFB4191	Trs85[K378A, K381A, K382A, K386A]-mNeonGreen-3xHA	pRS415	This study
CFB4192	Trs85[K378A, F379A, F380A, K381A, K382A, F383A, K386A]-mNeonGreen-3xHA	pRS415	This study
CFB4193	Trs85[K378E, K381E, K382E, K386E]-mNeonGreen-3xHA	pRS415	This study
CFB4194	Trs85[R620A]-mNeonGreen-3xHA	pRS415	This study
CFB4195	Trs85[R620E]-mNeonGreen-3xHA	pRS415	This study
CFB4227	6xHis-Trs85[K378A, F379A, F380A, K381A, K382A, F383A, K386A], Trs33	pETDuet-1	This study
CFB4229	6xHis-Trs85[K378E, K381E, K382E, K386E], Trs33	pETDuet-1	This study

Table S6: Strain list

Name	Description	Source
SEY 6210	MAT α <i>ura3-52 his3-Δ200 leu2-3,112 lys2-801 trp1-Δ901 suc2-Δ9</i>	(Robinson et al., 1988)
SEY 6210.1	MAT α <i>ura3-52 his3-Δ200 leu2-3,112 lys2-801 trp1-Δ901 suc2-Δ9</i>	(Robinson et al., 1988)
CFY2449	SEY6210.1 Trs85-mNeonGreen:: <i>HIS3</i>	(Thomas et al., 2018)
CFY2692	SEY6210 <i>trs85Δ::KanMX</i>	(Thomas et al., 2018)
CFY3585	SEY 6210 Trs23-6xHis-3xFLAG:: <i>TRP</i> , <i>trs85Δ::KanMX</i>	This study
CFY3307	SEY6210.1 Bet3-GFP:: <i>HIS3</i> , <i>trs85Δ::KanMX</i> , Trs85-mNeonGreen-3xHA:URA	This study

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Figure 1

A

B

C

D

E

Figure 2

A

B

C

D

Percent of total cross-links

E

F

Distribution of mapped cross-links

Figure 3

Figure 4

Figure 5

A

B

C

D

E

F

Figure 7

A

Core binding mutations

Membrane binding mutations

B **C**

Figure S1

A

B

A

Bet3-A cryoEM
Bet3-A pdb:3cue

B

Bet3-B cryoEM
Bet3-B pdb:3cue

C

Bet5 cryoEM
Bet5 pdb:3cue

D

Trs23 cryoEM
Trs23 pdb:3cue

E

Trs31 cryoEM
Trs31 pdb:3cue

F

Trs33 cryoEM
Trs33 pdb:2c0j
(human)

G

Trs20 cryoEM
Trs20 pdb:1h3q
(mouse)

H

Ypt1 cryoEM
Ypt1 pdb:3cue