

REPORT

Vps13-like proteins provide phosphatidylethanolamine for GPI anchor synthesis in the ER

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Glycosylphosphatidylinositol (GPI) is a glycolipid membrane anchor found on surface proteins in all eukaryotes. It is synthesized in the ER membrane. Each GPI anchor requires three molecules of ethanolamine phosphate (P-Etn), which are derived from phosphatidylethanolamine (PE). We found that efficient GPI anchor synthesis in *Saccharomyces cerevisiae* requires Csf1; cells lacking Csf1 accumulate GPI precursors lacking P-Etn. Structure predictions suggest Csf1 is a tube-forming lipid transport protein like Vps13. Csf1 is found at contact sites between the ER and other organelles. It interacts with the ER protein Mcd4, an enzyme that adds P-Etn to nascent GPI anchors, suggesting Csf1 channels PE to Mcd4 in the ER at contact sites to support GPI anchor biosynthesis. CSF1 has orthologues in *Caenorhabditis elegans* (*lpd-3*) and humans (*KIAA1109/TWEEK*); mutations in *KIAA1109* cause the autosomal recessive neurodevelopmental disorder Alkuraya-Kučinskas syndrome. Knockout of *lpd-3* and knockdown of *KIAA1109* reduced GPI-anchored proteins on the surface of cells, suggesting Csf1 orthologues in human cells support GPI anchor biosynthesis.

Introduction

Lipids are exchanged between cellular membranes by both vesicular and nonvesicular mechanisms. Nonvesicular trafficking requires proteins known as lipid transport proteins (LTPs). Until recently, most LTPs in eukaryotic cells were thought to transport lipids by a shuttling mechanism; the LTP extracts a lipid from one membrane, diffuses through the cytoplasm, and delivers the lipid to a second membrane (Chiapparino et al., 2016; Wong et al., 2019). In the past few years, it has been suggested that some eukaryotic LTPs use a different mechanism: they bind two membranes simultaneously and allow lipids to flow between the membranes in a hydrophobic channel formed by the LTP. This mechanism of lipid transport could be used for high-volume lipid movement (Leonzino et al., 2021; Reinisch and Prinz, 2021). Two proteins thought to use this mechanism are vacuolar protein sorting 13 (Vps13) and autophagy-related protein 2 (Atg2; Chowdhury et al., 2018; De et al., 2017; Kumar et al., 2018; Li et al., 2020; Maeda et al., 2019). Structural studies reveal that Vps13 and Atg2 have long rodlike structures with hydrophobic grooves that run from one end of the proteins to the other and could allow lipids to move between

membranes by sliding along the groove (Li et al., 2020; Valverde et al., 2019).

Both Vps13 and Atg2 are conserved from yeast to humans. Atg2 is required for the autophagosome formation (Suzuki et al., 2013). *Saccharomyces cerevisiae* has a single Vps13, while humans have four orthologues; all have been proposed to facilitate lipid transport at membrane contact sites (MCSs), regions where two organelles are closely apposed (Lang et al., 2015; Leonzino et al., 2021; Park et al., 2016). The four Vps13 proteins in humans are implicated in neurodegenerative disorders (Gauthier et al., 2018; Kolehmainen et al., 2003; Lesage et al., 2016; Rampoldi et al., 2001; Seong et al., 2018).

In this study, we show that *S. cerevisiae* has three proteins predicted to be structurally like Vps13 and Atg2: Csf1, Fmp27, and Hob2 (Ypr117w). The functions of Csf1, Fmp27, and Hob2 are not well understood. Fmp27 was recently shown to localize to MCSs between the ER and plasma membrane (PM) and bind phosphoinositides (Neuman et al., 2022). Cells lacking Csf1 have also been shown to have defects in vesicular trafficking of proteins with glycosylphosphatidylinositol (GPI) anchors (Copic

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et al., 2009). GPI anchors are synthesized in the ER by a conserved multistep pathway (Fujita and Kinoshita, 2012). The anchor contains a phosphatidylinositol to which is added glycans and three ethanolamine phosphate (P-Etn) molecules. Anchor assembly and protein attachment occur on the luminal surface of the ER membrane. The P-Etn in GPI anchors is derived from phosphatidylethanolamine (PE; Menon and Stevens, 1992); thus, biosynthesis of each GPI anchor requires three PE molecules. Here, we show that Csf1 is necessary for efficient P-Etn incorporation into GPI anchors.

Results and discussion

Csf1, Fmp27, and Hob2 are structural homologues of Vps13

To identify proteins that are structurally like Vps13 and Atg2 in *S. cerevisiae*, we used the remote homology detection program named HHpred, which identified Csf1, Fmp27, and Hob2. To gain more insight into the predicted structures of Csf1, Fmp27, Hob2, Vps13, and Atg2, we used the protein structure prediction algorithm RoseTTAFold (Yang et al., 2020) to generate structures of each 1,000 residues of the proteins, which we then stitched together into full-length predicted structures using the molecular visualization program PyMol. This analysis revealed the proteins form similar extended β -sheets with long hydrophobic grooves (Fig. 1 A, Video 1, Video 2, Video 3, Video 4, and Video 5). We used the structure comparison program FATCAT (Flexible structure Alignment by Chaining Aligned fragment pairs allowing Twists) to show the predicted structures are similar (Fig. 1, B and C). We compared the predicted structure of the first 1,000 residues of Vps13 and the crystal structure of Vps13 (Protein Data Bank accession no. 6CBC), which is of the first 335 residues (Kumar et al., 2018). The P value is 4.95E-11, indicating the predicted structure aligns well with the solved structure. We also compared the full-length structures of Fmp27, Hob2, and Atg2 we generated with RoseTTAFold with the recently published predicted structures of these proteins generated by the protein structure prediction program AlphaFold (Tunyasuvunakool et al., 2021). Comparisons of the structures using FATCAT indicated they are highly similar (Fig. S1 A). Notably, FATCAT indicated that the most similar portions of the predicted structure are the extended β -sheets found in all the predicted structures, which form long hydrophobic grooves running the length of the proteins (not shown). We found that the central groove that runs the length of these proteins is lined with hydrophobic residues (Fig. S1 B), consistent with the hypothesis that lipids could slide along the grooves.

Csf1, Fmp27, and Hob2 localize to multiple contact sites

If Csf1, Fmp27, and Hob2 are LTPs that function similarly to other bridge-forming LTPs, they should localize to MCSs. A recent study indeed showed that endogenous FMP27-GFP localizes to contact sites between the ER and PM (Neuman et al., 2022). To visualize Csf1 and Hob2, we initially tagged them endogenously with GFP on their C-termini, but the signal was low (not shown). Instead of using a single GFP, we endogenously tagged the proteins with seven copies of GFP11, a fragment of GFP that will self-assemble with the remainder of the protein, GFP1-10

(Makhija et al., 2021). The tagged proteins are functional (Fig. S2 A). We found that GFP-tagged Csf1, Fmp27, and Hob2 form puncta that are on or next to the ER (Fig. 2 A). Most of the puncta are at the cell cortex. The cortical puncta associate with Tcb3 (Fig. 2 B), a protein known to be at ER-PM contact sites (Toulmay and Prinz, 2012), suggesting that Csf1, Fmp27, and Hob2 localize to these contact sites. Not all Csf1, Fmp27, and Hob2 puncta are at the cell cortex; some are adjacent to mitochondria (Fig. 2 C). To determine whether some Csf1, Fmp27, and Hob2 puncta are at ER-mitochondria contact sites, we assessed whether they colocalize with the ER-mitochondria tether ERMES (ER mitochondria encounter structure). GFP-tagged Csf1, Fmp27, or Hob2 was expressed together with mCherry fused to Mmm1, one of the components of ERMES. All three Vps13-like proteins were found to be adjacent to or colocalize with \sim 50% of Mmm1-mCherry puncta (Fig. 2, D and E), suggesting they localize to some ER-mitochondria contact sites. It is not clear why Csf1, Fmp27, or Hob2 localizes to only a subset of ERMES puncta. Together, these findings indicate that Csf1, Fmp27, or Hob2 localizes to ER-PM and some ER-mitochondria contact sites.

Vps13-like proteins are dispensable for bulk lipid transport to mitochondria

It has been suggested that tube-forming LTPs are well suited to facilitate high-volume bulk lipid transport (Leonzino et al., 2021; Reinisch and Prinz, 2021). We investigated their role in non-vesicular lipid transport to mitochondria, which is required for mitochondrial biogenesis (Petrunaro and Kornmann, 2019). We first generated a strain lacking all five Vps13-like LTPs in *S. cerevisiae*: Vps13, Atg2, Csf1, Fmp27, and Hob2. Remarkably, the strain (called 5 Δ here) grew about as well as WT cells at 30°C on the glucose-containing media (YPD; Yeast Extract, Peptone, Dextrose and synthetic complete [SC] medium) and YPG (Yeast Extract, Peptone, Glycerol), which only allows the growth of cells with functional mitochondria (Fig. 3 A). To more directly determine whether there is a defect in bulk lipid transport to mitochondria in 5 Δ cells, we assessed the rate of cardiolipin (CL) synthesis. In *S. cerevisiae*, CL is produced in the mitochondrial inner membrane from phospholipids synthesized in the ER, primarily phosphatidic acid. Therefore, CL production in mitochondria requires phospholipid transport from the ER to mitochondria (Fig. 3 B; Tatsuta and Langer, 2017). To measure CL synthesis, we determined the amount of CL formed in 1 h by labeling cells with [3 H]acetate. The amount of CL formed in 5 Δ cells (expressed as a percentage of the total phospholipid synthesized) decreased \sim 40% (Fig. 3 C). This suggests the five Vps13-like proteins could play a role in transporting CL precursors from the ER to mitochondria but are not required.

It is possible that the ERMES complex compensates for the lack of Vps13-like proteins in 5 Δ cells. ERMES tethers mitochondria to the ER and facilitates phospholipid transport to mitochondria (Kawano et al., 2018; Kornmann et al., 2009). We generated a 5 Δ strain that also lacks the ERMES complex. Because cells lacking the ERMES complex and Vps13 are not viable (Lang et al., 2015), we added an auxin-inducible degron to one of the components of ERMES, Mdm12, to allow conditional

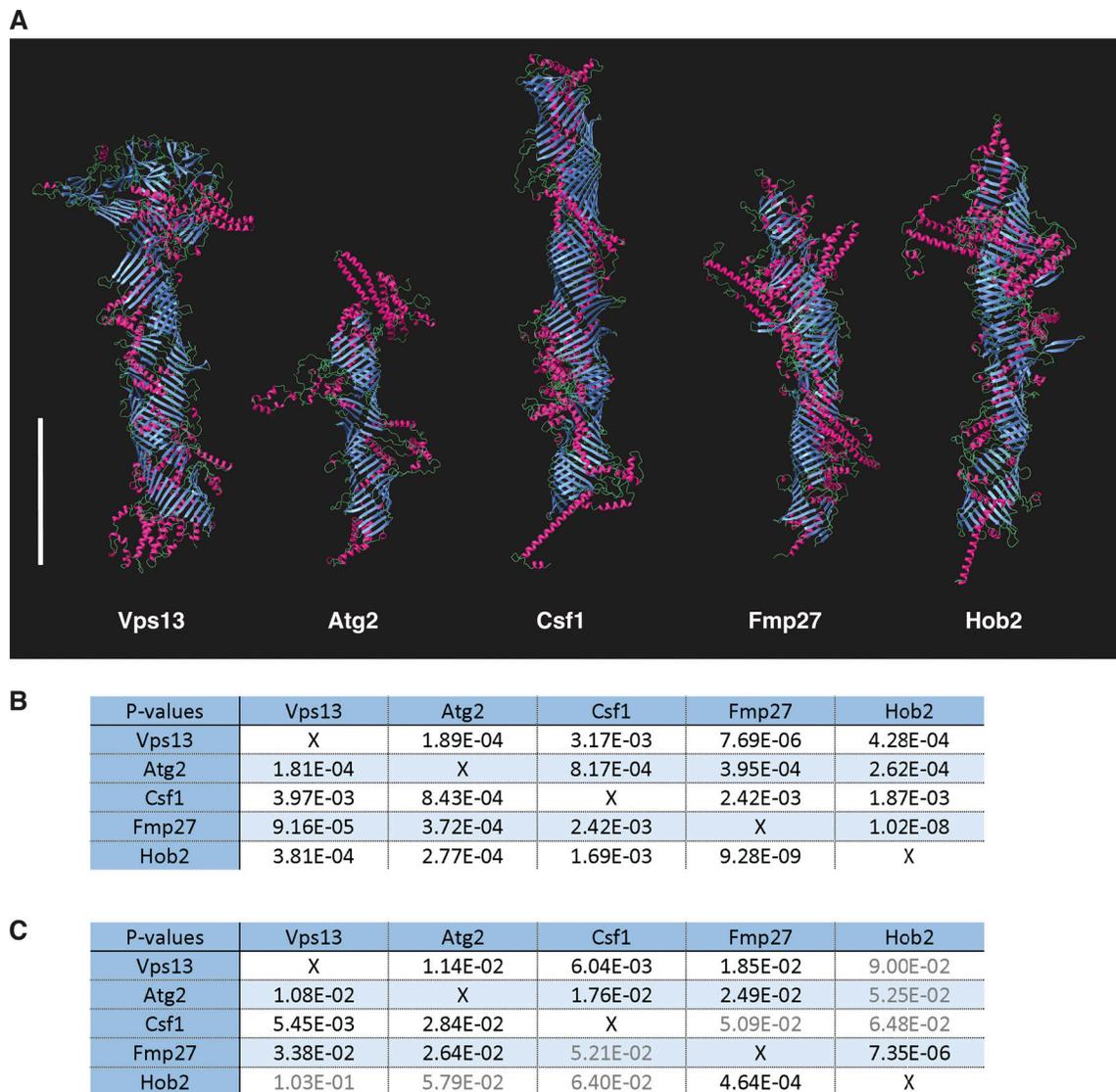


Figure 1. **Structural similarity of Vps13, Atg2, Csf1, Fmp27, and Hob2.** (A) Structure predictions generated with RoseTTAFold: β -sheets (blue), α -helices (red), and coil (green). Bar = 100 Å. (B and C) Significance of the structural similarity between pairs of proteins using FATCAT. P values are from pairwise comparisons of the predicted structures of the first 1,000 amino acids (B) or full-length versions (C) of the proteins. The first structure for each pairwise comparison is listed in the first column. P values >0.05 in gray.

degradation of ERMES (Fig. S2, B–D). Elimination of any of the proteins in the ERMES complex causes it to disassemble (Kornmann et al., 2009). We found the degron causes Mdm12 to degrade within 30 min of adding the auxin to the medium (Fig. S2 B). Mdm12 degradation also caused mitochondria to change shape (Fig. S2 D), indicating that ERMES is not functional after Mdm12 depletion. Interestingly, we found *VPS13* is the only one of the five genes encoding Vps13-like proteins that is required for viability of cells lacking ERMES (Fig. S2 C; compare strains indicated with boxes).

We found that depletion of Mdm12 did not significantly reduce CL synthesis (Fig. 3 C), suggesting ERMES is not required to bring CL precursors to mitochondria. Depletion of Mdm12 in Δ cells did not significantly reduce CL production more than in Δ cells expressing Mdm12, suggesting that the five Vps13-like proteins and ERMES do not play redundant roles in transporting

CL precursors from the ER to mitochondria. Together, these findings suggest that the Vps13-like proteins in yeast do not have an essential role in bulk phospholipid transport to mitochondria.

Csf1 plays a role in PE metabolism

We wondered whether Csf1, Fmp27, and Hob2 might play a role in PE metabolism since phospholipid transport is necessary for some PE metabolism (Fig. 3 D). PE is produced by two pathways in *S. cerevisiae*. One makes PE in the ER and requires ethanolamine (Etn; the Kennedy pathway). The second, called the cytidyl-diphosphate-diacylglycerol (CDP-DAG) pathway, generates PE primarily outside the ER using enzymes that decarboxylate phosphatidylserine (PS) to form PE (Fig. 3 D). The enzymes are in the mitochondrial inner membrane (Psd1) and endocytic compartments (Psd2; Acoba et al., 2020). It should be noted that

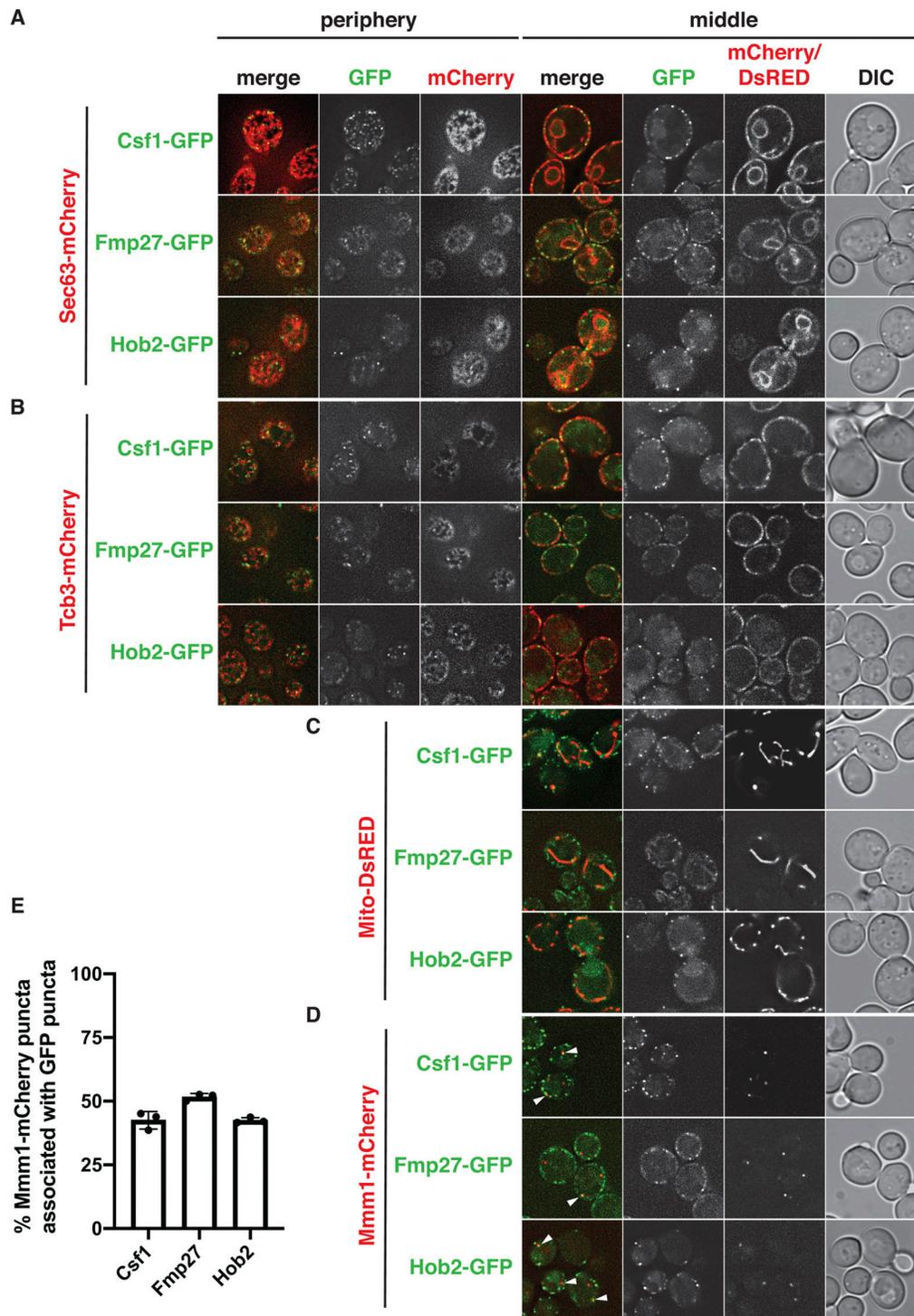


Figure 2. **Csf1, Fmp27, and Hob2 localize to ER-PM and ER-mitochondria contact sites. (A–D)** Cells were visualized live. Cells expressed GFP fusions to the indicated proteins and the ER marker Sec63-mCherry (A), the ER-PM contact site marker Tcb3-mCherry (B), the mitochondrial marker Mito-DsRed (C), or the ERMES marker Mmm1-mCherry (D). Csf1 and Hob2 are endogenously tagged with seven copies of GFP11, and the strains also express GFP1–10; Fmp27 is endogenously tagged with a single copy of GFP. Cells shown in A and B were visualized by focusing on the middle or periphery of the cells. Bar = 5 μ m. Arrows in D indicate colocalization of ERMES and Csf1, Fmp27, or Hob2. (E) Percentage of Mmm1-mCherry puncta associated with GFP puncta (mean \pm SD of three independent experiments; at least 100 Mmm1-mCherry puncta counted per experiment).

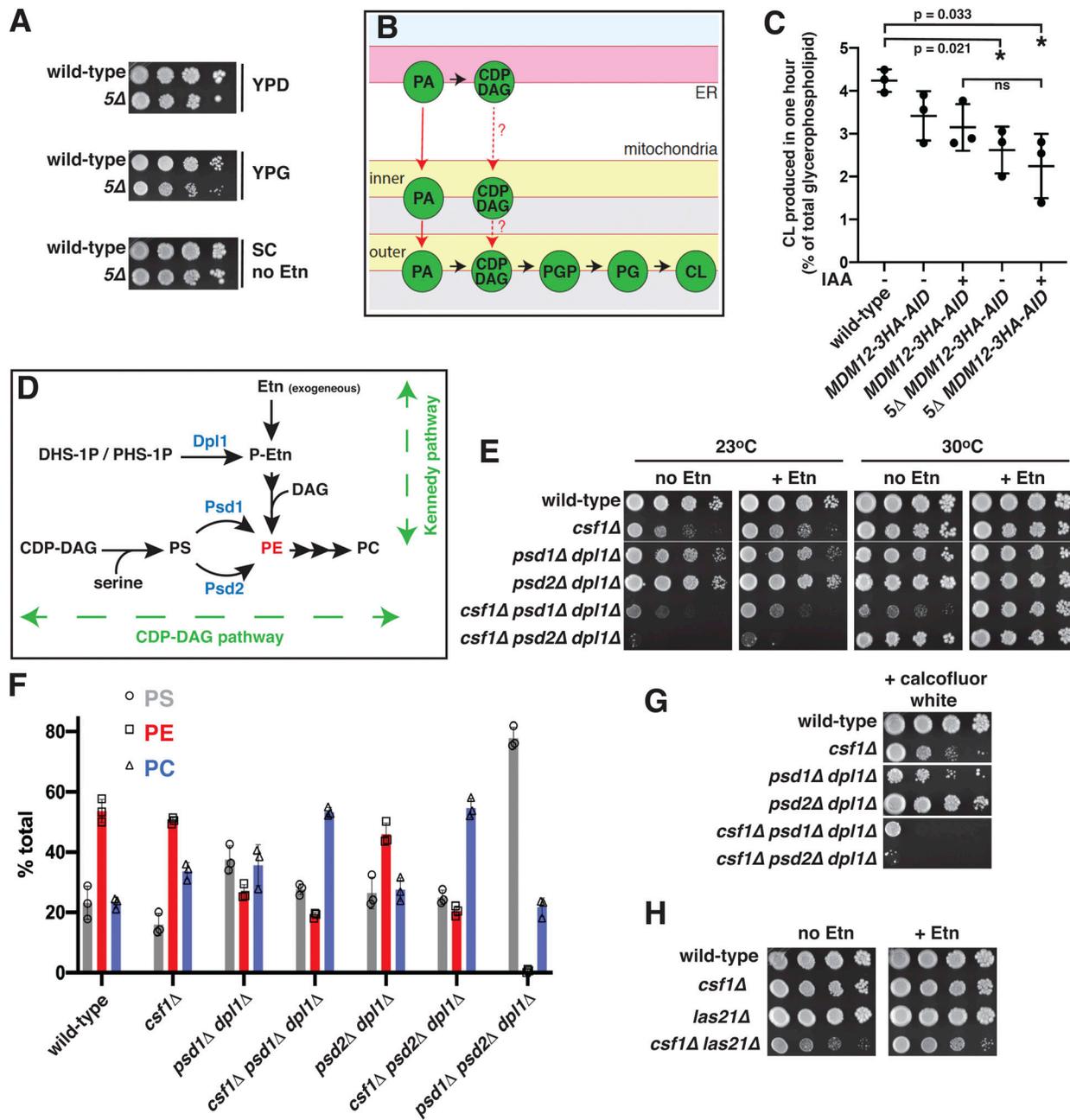


Figure 3. **Vps13-like proteins and ERMES are not required to transport CL precursors to mitochondria.** (A) 10-fold serial dilutions of the indicated strains grown on plates with YPD, YPG, or SC without Etn media. The plates were grown at 30°C for 3 d. (B) Cartoon showing the biosynthetic pathway of CL in *S. cerevisiae*. Phosphatidic acid (PA) and CDP-DAG are produced in the ER membrane (pink). They are transported to the inner membrane of mitochondria (gray), where they are converted to phosphatidylglycerol-phosphate (PGP), phosphatidylglycerol (PG), and CL. (C) The indicated strains were labeled with [³H] acetate, and the amount of CL formed in 1 h was determined. The amount of [³H]CL produced is given as the percentage of the total radiolabeled glycerophospholipids formed. Values from three independent experiments are shown (mean ± SD). Statistical significance was determined by a Student's *t* test (two-tailed, unpaired); *, *P* < 0.05. Strains with *MDM12-3HA-AID* express Mdm12 with a C-terminal fusion to 3HA antigen tags and the auxin-inducible degron (AID). Where indicated, the auxin IAA was added to the media 30 min before [³H]acetate addition to induce degradation of Mdm12-3HA-AID. (D) Cartoon of the Kennedy and CDP-DAG pathways of PE metabolism. Proteins in blue. Abbreviations not listed in text: DHS-1P, dihydrosphingosine-1 phosphate; PHS-1P, phytosphingosine-1 phosphate. (E) 10-fold serial dilutions of the indicated strains on plates with SC with or without Etn. The plates were grown for 3 d. (F) The indicated strains were labeled with [³H]serine for 30 min, and the amount of radiolabeled PS, PE, and PC was determined. The results are the percentage of the three lipids indicated (mean ± SD from three independent experiments). (G and H) 10-fold serial dilutions of the indicated strains grown on SC plates containing 10 μg/ml calcofluor white (G) or SC with or without Etn (H). The plates were grown in 30°C for 3 d.

a fraction of Psd1 is in the ER; therefore, the CDP-DAG pathway may produce some PE in the ER (Friedman et al., 2018). PE produced by the Kennedy pathway requires exogenous Etn or P-Etn produced by dihydrosphingosine phosphate lyase (Dpl1; Fig. 3 D). Cells lacking Dpl1 and grown in media that do not contain Etn can only make PE using the CDP-DAG pathway.

We found Csf1 plays a role in the CDP-DAG pathway of PE metabolism. Cells lacking Csf1, Dpl1, and either Psd1 or Psd2 grow poorly in a medium without Etn (Fig. 4 E). The strains grew normally when Etn was added to the medium, suggesting Csf1 is particularly important for the CDP-DAG pathway of PE metabolism. A role for Csf1 in PE metabolism has also been suggested by a recent preprint from the Kornmann group (John Peter et al., 2021 Preprint). The growth of cells lacking Fmp27 or Hob2 was not reduced in a medium lacking Etn (not shown); therefore, we focused on Csf1.

To determine whether PE metabolism by the CDP-DAG pathway is impaired in cells lacking Csf1, we labeled cells with [³H]serine for 1 h. Cells use [³H]serine to produce [³H]PS, which can be converted into PE and, in turn, into phosphatidylcholine (PC) by the CDP-DAG pathway (Fig. 3 D). We previously showed that following [³H]serine addition to the medium of growing cells, radiolabeled PS, PE, and PC are produced at linear rates (Raychaudhuri and Prinz, 2008). The labelings were done in cells lacking Dpl1 since this prevents the production of radiolabeled P-Etn that could be used to make PE. Surprisingly, we found cells lacking Csf1 have no defect in producing PS (Fig. S2 E) or converting it into PE and PC (Fig. 3 F). This suggests they do not have significant defects in transporting PS to Psd1 and Psd2 or returning the PE produced by these enzymes back to the ER, where it is converted to PC. Together, these findings indicate Csf1 has a role in PE metabolism but does not affect the production of PS, PE, or PC.

Cells lacking Csf1 do not have defects in autophagy

Since Csf1 plays a role in PE metabolism but not in the production of PE or the conversion of PE to PC, we wondered whether Csf1 is necessary for a PE-dependent process. One is autophagy, which requires the conjugation of PE to the protein Atg8, a protein that plays a role in autophagosome formation (Ichimura et al., 2000). During autophagy, Atg8 is conjugated to PE (forming Atg8-PE) and delivered to the vacuole, where it is degraded. When GFP is fused to the N-terminus of Atg8, the Atg8 portion of the fusion is degraded in the vacuole during autophagy while the GFP portion remains relatively stable; thus, the presence of free GFP in cells expressing GFP-Atg8 indicates it was trafficked to the vacuole during autophagy (Shintani and Klionsky, 2004). If Csf1 provides the PE necessary for GFP-Atg8-PE production, cells lacking Csf1 would have reduced levels of free GFP following autophagy induction.

We induced autophagy in cells expressing GFP-Atg8 by shifting them to nitrogen starvation medium for 3 h. This caused the appearance of free GFP in WT cells but not in cells lacking Atg2 or Atg1, a protein kinase required for autophagy (Fig. S2 F). The production of free GFP was not affected in cells lacking Csf1 (Fig. S2 F), indicating Csf1 probably does not play a role in providing PE to produce Atg8-PE.

GPI anchor precursors lacking P-Etn accumulate in cells lacking Csf1

We wondered whether the role of Csf1 in PE metabolism could be to support GPI anchor biosynthesis, since the biosynthesis of each anchor requires three PE molecules (Fig. 4 A). Consistent with this possibility, cells lacking Csf1 have a defect in trafficking a GPI-anchored protein (Copic et al., 2009), and the genetic profile of *CSF1* is like that of many genes required for GPI anchor synthesis (<https://thecellmap.org/>). Cells with defects in GPI anchor synthesis are hypersensitive to the antifungal calcofluor white (Benghezal et al., 1995), and we found cells lacking Csf1 are also hypersensitive (Fig. 3 G). In addition, there is a negative genetic interaction between *CSF1* and *LAS21* (Fig. 3 H), which encodes one of the three Etn phosphotransferases involved in GPI anchor biosynthesis (Fig. 4 A). Together, these findings implicate a potential function of Csf1 in GPI anchor production.

We speculated that if Csf1 plays a role in providing PE for GPI anchor biogenesis, nascent GPI anchors lacking P-Etn would accumulate in cells lacking Csf1. To test this, we labeled various strains with [³H]inositol, extracted phospholipids, and separated them by TLC as previously described (Flury et al., 2000). This study showed that GPI anchor intermediates lacking P-Etn accumulate in cells with mutations in any of the three genes encoding the Etn phosphotransferases were required to produce GPI anchors: *MCD4*, *GPII3*, and *LAS21* (Fig. 4 A). Cells with a conditional *gpiI3-5* allele accumulate the lipid O31a, a GPI anchor precursor that lacks P-Etn (Fig. 4 B; Flury et al., 2000). We found lipid O31a also accumulates in cells lacking Csf1 and proteins in the CDP-DAG pathway of PE biosynthesis, Psd1 or Psd2 (Fig. 3 D; and Fig. 4, C and D). We noted, however, that cells lacking only Csf1 do not accumulate lipid O31a. Since the labeling experiments were done at 30°C, we wondered whether cells lacking Csf1 would accumulate lipid O31a when grown at 23°C, which is known to reduce growth of these cells (Tokai et al., 2000), a finding we confirmed (Fig. 3 E). Indeed, at 23°C, the cells lacking Csf1 accumulated significantly more lipid O31a than WT cells (Fig. 4 E), suggesting Csf1 supports GPI anchor biosynthesis.

To further investigate which are the Csf1-dependent steps in GPI anchor biosynthesis, we characterized strains lacking *Las21*, which accumulate the GPI anchor precursor M4 that has two P-Etns of the three P-Etns normally added to GPI anchors (Fig. 4 B; Flury et al., 2000). We confirmed this finding and discovered that cells lacking Csf1 also accumulate lipid M4, though not as much as cells lacking *Las21* (Fig. 4, F and G). Interestingly, cells lacking both *Las21* and Csf1 have reduced amounts of lipid M4 but more lipid O31a (Fig. 4 G). This finding suggests Csf1 is necessary for a step in GPI synthesis before the one catalyzed by *Las21*, probably the addition of one or both of the first two P-Etns added to GPI anchors. Together, these findings suggest Csf1 is necessary to support the addition of P-Etn to GPI anchor precursors.

Csf1 may channel PE to the Etn phosphotransferase Mcd4

Since Csf1 is likely to be a tube-forming LTP and localizes to MCSs between the ER and other organelles, we hypothesized that it transports PE to the ER to support GPI anchor biogenesis

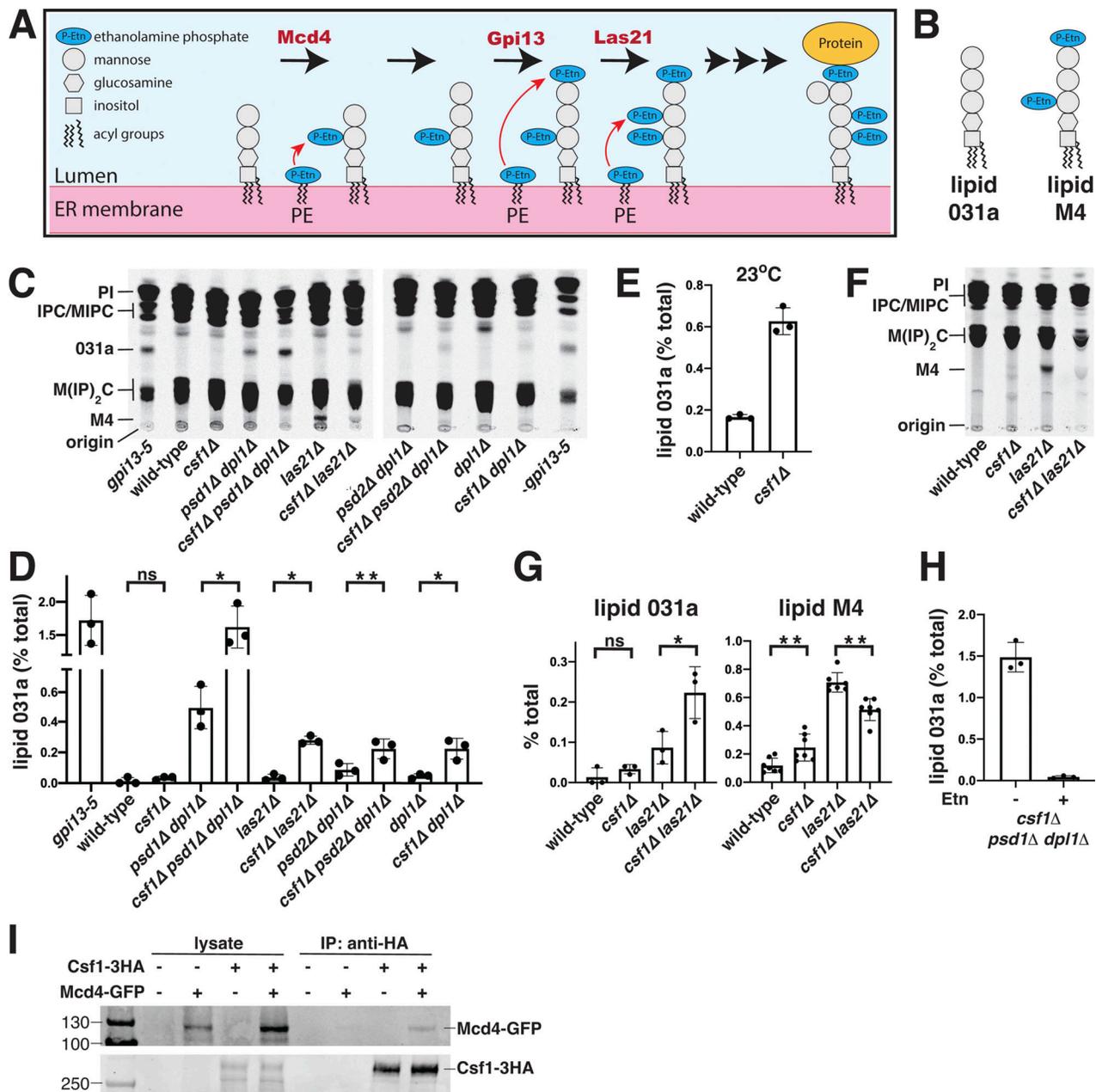


Figure 4. Cells lacking *Csf1* accumulate nascent GPI anchors lacking P-Etn. (A) Cartoon of the three steps in GPI anchor biogenesis where P-Etn is added to nascent GPI anchors. The three Etn phosphotransferases are in red. (B) GPI anchor intermediates (symbols as in A). (C) The indicated strains were labeled with [³H]inositol for 2 h, and lipids were extracted and separated by TLC. All strains were grown in SC at 30°C except *gpi13-5* cells, which were grown at 23°C and shifted to 37°C for 30 min before labeling. Abbreviations not listed in text: IPC, inositol-phosphoryl-ceramide; MIPC, mannosyl-inositol-phosphoryl-ceramide; M(IP)₂C, mannosyl diinositol-phosphoryl-ceramide; PI, phosphatidylinositol. (D) Quantification of the amount of lipid 031a formed in the experiments shown in C. Mean ± SD of three independent experiments. Statistical significance determined by Student's *t* test (two-tailed, paired). *, *P* < 0.05; **, *P* < 0.005. (E) Quantification of the amount of lipid 031a formed in the indicated strains as described in C and D, except the strains were grown at 23°C and labeled for 24 h (mean ± SD). (F) The indicated strains were labeled with [³H]inositol as described in C, except that lipids were separated by TLC using a solvent system that resolves lipid M4. (G) Quantification of the amount of lipid M4 formed as in the experiments shown in F is shown in the right panel. The left panel shows the same data given in D. Mean ± SD of three independent experiments. Statistical significance determined with a Student's *t* test (two-tailed, paired). *, *P* < 0.05; **, *P* < 0.005. (H) Quantification of the amount of lipid 031a formed in the indicated strains as described in C and D, except grown in SC media either with or without Etn (mean ± SD). (I) Lysates from cell expressing *Csf1*-3HA and *Mcd4*-GFP as indicated were incubated with magnetic beads conjugated to an anti-HA antibody. Whole-cell lysates and immunoprecipitated (IP) proteins were separated by SDS-PAGE and immunoblotted with anti-HA and anti-GFP antibodies. Numbers are size of molecular weight markers (kDa). Representative blots from three independent experiments are shown. Source data are available for this figure: SourceData F4.

and that this is particularly important in cells where most PE is made outside the ER. If this is correct, production of PE in the ER by the Kennedy pathway might bypass the role of Csf1 in GPI anchor biosynthesis. We found that addition of Etn to cells lacking Csf1, Psd1, and Dpl1 eliminated accumulation of lipid O31a found in this strain when it is grown without exogenous Etn (Fig. 4 H). Thus, PE production in the ER supports efficient GPI anchor synthesis in cells lacking Csf1.

These findings suggest that in growth conditions where most PE is synthesized outside the ER, there is insufficient PE in the ER to support robust GPI anchor production, perhaps because most PE in the ER is converted to PC. If this is correct, Csf1 may prevent this problem by channeling PE outside the ER to the GPI synthesis machinery. Consistent with this, we used coimmunoprecipitation to show that Csf1 interacts with Mcd4, the Etn phosphotransferase that adds the first P-Etn to nascent GPI anchors (Fig. 4 A). Endogenous Csf1 was tagged with three copies of the HA epitope (Csf1-3HA), and we confirmed Csf1-3HA is functional (Fig. S2 B). We immunoprecipitated Csf1-3HA and found it coimmunoprecipitates with Mcd4 fused to GFP (Mcd4-GFP) expressed from a plasmid under the native *MCD4* promoter (Fig. 4 I). Together, these findings suggest Csf1 channels PE to Mcd4 to support GPI anchor synthesis. This channeling may not be necessary when significant amounts of PE are produced in the ER, which probably increases the amount of PE in the ER available for GPI anchor production.

Csf1 homologues in higher eukaryotes participate in GPI-anchored protein biogenesis

Csf1 has homologues in higher eukaryotes. We wondered whether they are necessary for efficient GPI anchor biogenesis. To test this, we determined whether knocking down or knocking out the homologues in human cells and *Caenorhabditis elegans* decreases the amount of GPI-anchored proteins on the surface of cells.

The human homologue of Csf1 is KIAA1109, or TWEAK. Mutations in TWEAK were recently shown to cause a disorder of brain development and arthrogryposis named Alkuraya-Kučinskas syndrome (Gueneau et al., 2018). We used shRNA to knock down expression of TWEAK and the GPI biogenesis gene GPI Anchor Attachment 1 (GPAA1) in human embryonic kidney 293 (HEK-293) cells. GPAA1 is required to add proteins to GPI anchors (Hiroi et al., 1998). We used FACS to show that knockdown of GPAA1 reduces the amount of the GPI-anchored protein CD55 on the surface of cells (Fig. 5 A). Knockdown of TWEAK with either of two shRNAs that target different sequences similarly reduced the amount of CD55 on the surface of the HEK-293 cells (Fig. 5, B and C). We used quantitative real-time RT-PCR (qRT-PCR) to confirm the knockdown of TWEAK and GPAA1 (Fig. S3 A).

The *C. elegans* genome encodes a single KIAA1109/TWEAK orthologue, *lpd-3*. Four *lpd-3* transcripts have been identified that vary by only a few amino acids. We generated a null allele of *lpd-3* that lacks exons 1–6 (and intervening introns) and introduces three tandem stop codons in each reading frame (Fig. S3, B–D). Homozygous *lpd-3 N-Δ* animals were validated by PCR and Sanger sequencing (Fig. S3, C and D). The *lpd-3 N-Δ* animals

had shortened gonads and a deformed germline (Fig. S3 E). Similar defects are found in GPI anchor-synthesis mutants (Murata et al., 2012). We used fluorescein-labeled proaerolysin (FLAER), which binds GPI anchors, to detect GPI-anchored proteins in the PM of WT and *lpd-3 N-Δ* embryos. Before the first mitotic cell division, embryos have little FLAER signal at the PM (not shown). At the two-cell stage, WT embryos had significant FLAER signal in the PM at the cleavage furrow, while *lpd-3 N-Δ* had significantly less (Fig. 5, D and E). This was also true of four-cell-stage embryos (Fig. S3 F). Taken together, these findings indicate that KIAA1109/TWEAK and LPD-3 play roles in the biogenesis of GPI-anchored proteins and suggest they facilitate GPI anchor biogenesis like Csf1.

Conclusions

We propose that Csf1 channels PE to Mcd4 in the ER to support GPI anchor production (Fig. 5 G). This is consistent with work from the Menon group that suggested PE used for GPI synthesis is preferentially derived from PE produced outside the ER in mitochondria (Vidugiriene et al., 1999). Our results suggest the Csf1 homologues KIAA1109/TWEAK and Lpd-3 support GPI anchor production and probably have the same function as Csf1. Since humans (and other mammals) cannot synthesize Etn and derive it mostly from dietary lipids (Calzada et al., 2016; Patel and Witt, 2017), KIAA1109/TWEAK may be particularly important for GPI biosynthesis when dietary sources of Etn are low. The role of KIAA1109/TWEAK in GPI anchor biogenesis in Alkuraya-Kučinskas syndrome should be investigated.

Our findings strongly suggest that Csf1 transports PE to the ER. However, we cannot rule out that Csf1 indirectly affects GPI anchor synthesis by transporting another lipid or regulating GPI anchor synthesis proteins such as Mcd4. Assuming Csf1 does transport PE to the ER, it surprisingly is not necessary to bring PE to the ER to support PC synthesis but instead seems to direct PE to GPI anchor biosynthesis. It is likely that most PE in the ER is used to produce PC, and Csf1 may ensure there is sufficient PE to support GPI anchor production when PE levels are low in the ER (Fig. 5 G). Understanding the mechanism and regulation of lipid transport by tube-forming LTPs such as Csf1 is an important challenge for the future.

Materials and methods

Strains and cell line

The yeast strains used in this study are listed in Table S1. Strains were grown in SC media containing 2% glucose (Sigma-Aldrich), 0.67% nitrogen base (BD), and an appropriate amino acid dropout mix (US Biological); or YPD media containing 1% yeast extract (BD), 2% peptone (BD), and 2% glucose; or YPG media containing 1% yeast extract (BD), 2% peptone (BD), and 3% glycerol; or nitrogen starvation medium (SD-N) media containing 2% glucose (Sigma-Aldrich) and 0.67% nitrogen base (BD). Where indicated, Etn (Sigma-Aldrich) was added to media to a final concentration of 1 mM from a 1M stock.

The *C. elegans* strains used in this study were N2 (Bristol, WT) and AG645 (*lpd-3[av254]V*. CRISPR/Cas9 Edit, deletion of exons 1–6 and intervening introns).

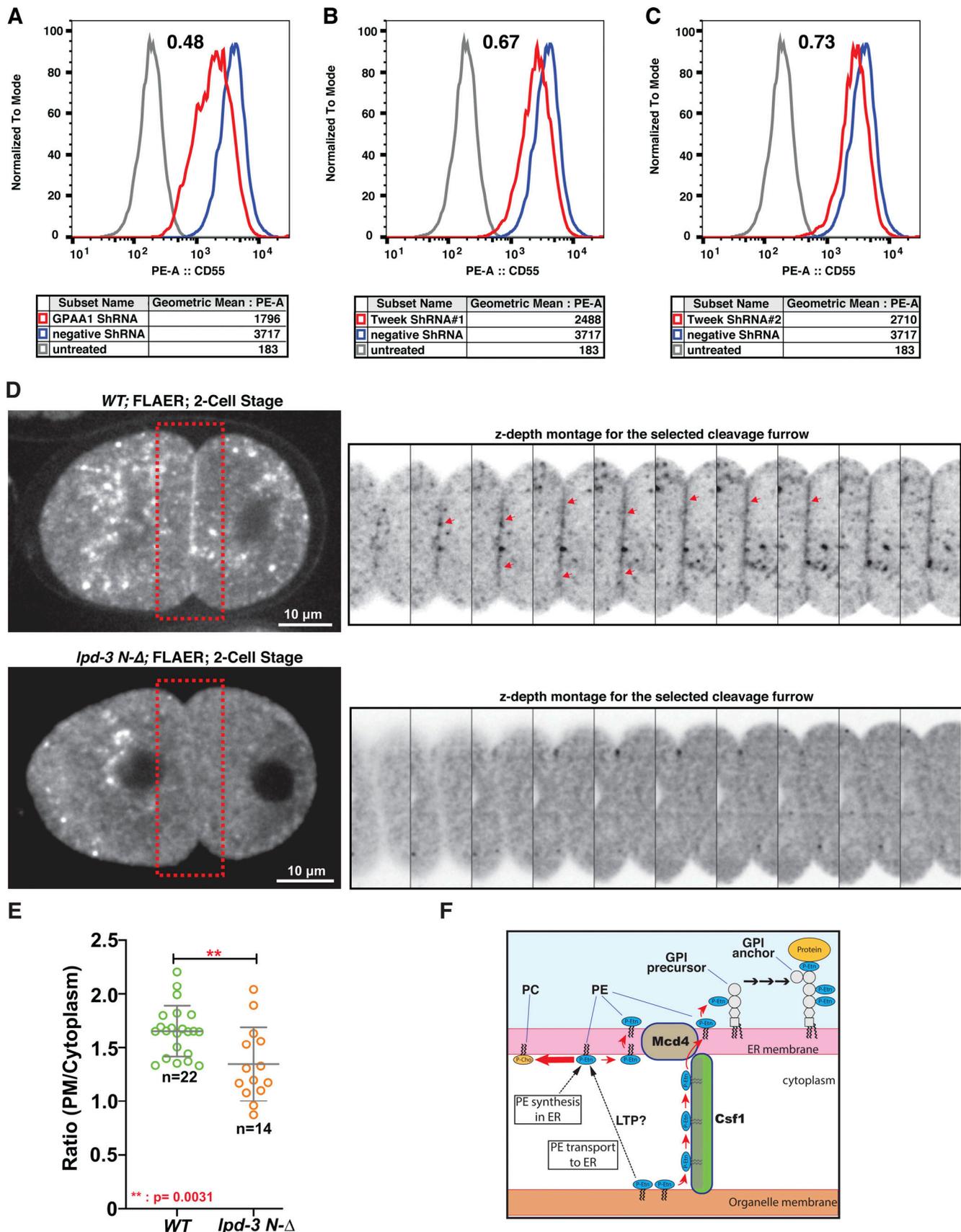


Figure 5. **Csf1** homologues play a role in biogenesis of GPI-anchored proteins. (A–C) Flow-cytometry analysis of HEK-293 cells expressing shRNA targeting GPAA1 (A) or two different shRNAs targeting *KIAA1109/TWEEK* (B and C) using anti-CD55 antibody conjugated to R-phycoerythrin (PE-A). 10,000 cells

were counted, and the geometric mean for negative shRNA (blue), gene targeted by shRNA (red), and untreated (no antibody, gray) shown. Ratio of the geometric mean of fluorescence intensity of the cells treated with shRNA to the targeted genes to negative control are shown in each panel. **(D)** FLAER staining of two-cell-stage *C. elegans* embryos from WT and *lpd-3 N-Δ* animals. Right panels show z-depth montages of the cleave furrows indicated by the red boxes. Bars = 10 μm. **(E)** Quantification of FLAER signals in the cleave furrow of two-cell embryos as shown in D (n = number of tested embryos). P value determined by Student's *t* test (two-tailed, unpaired). **(F)** Model of PE channeling to Mcd4 at ER contact sites. Red arrows indicate movement or metabolism of PE.

HEK-293 cells were cultured with DMEM (4.5 g/liter D-Glucose + L Glutamine; Gibco), 10% FBS heat-inactivated (Quality Biological), and penicillin and streptomycin (Gibco) in incubator at 37°C 5% CO₂.

Serial dilutions to assess growth

Cells were grown in the indicated medium to logarithmic growth phase, washed twice with medium, and adjusted to an OD₆₀₀ of 0.5–1.0. 10-fold serial dilutions were prepared in the medium and spotted onto plates. The plates were incubated for 3 d at the indicated temperature.

Microscopy of yeast

Cells were imaged live at 30°C using DeltaVision Spectris system (Applied Precision), with a wide-field inverted epifluorescence microscope (IX70; Olympus), 100× objective, and a charge-coupled device CoolSNAP HQ camera (Photometrics). Images were deconvolved using the conserved ratio method, 10 cycles, with softWoRx (Applied Precision).

Labeling cells with [³H]serine and analysis of glycerophospholipids

Cells were grown to saturation at 30°C in SC lacking serine (Complete Supplementation Mix; Sunrise Science Products) with 1 mM Etn. 2 ml of a saturated culture was added to 50 ml of fresh medium. The cells were grown for 4.5 h, washed twice with water, and resuspended in 50 ml of fresh medium lacking Etn at an OD₆₀₀ of 0.2–0.4. After 3 h of growth, 0.1 mCi of [³-³H]serine (American Radiolabeled Chemicals) was added, and the cells were grown for 1 h. After labeling, the cells were cooled to 4°C, washed once with water, and lysed in a Precellys24 homogenizer. Lipids were extracted and separated by TLC as described (Kannan et al., 2017). TLC plates were scanned on a RITA Star thin-layer analyzer (Raytest) to the relative radioactive signal from each phospholipid class.

Labeling cells with [³H]acetate and analysis of glycerophospholipids

Cells were grown to saturation at 30°C in YPD. 250–400 μl of the saturated cultures was added to 10 ml of fresh YPD. The cultures were grown at 30°C for 4 h; 5 μl of indole-3-acetic acid (IAA; Sigma-Aldrich; stock = 1 M in ethanol) or ethanol was added; after 30 min, 50 μCi [³H]acetate (American Radiolabeled Chemicals) was added; and the cultures were grown for 1 h. The cells were added to ice-cold water and washed once with water, and the lipids were extracted and analyzed as described in the previous section.

Measurement of autophagy induction

Cells were grown to mid-logarithmic phase in SC at 30°C, washed three times with SD-N medium, and incubated for 3 h in

SD-N at 30°C. At the indicated times, 3 OD₆₀₀ units of cells were withdrawn. The cells were pelleted, resuspended in 50 μl of 1.85 M NaOH, incubated on ice for 10 min, mixed with 125 μl of 20% trichloroacetic acid, and incubated on ice for 30 min. The precipitated proteins were separated by SDS-PAGE and immunoblotted with anti-GFP antibodies (Roche).

Measurement of GPI anchor intermediates

GPI anchor intermediates were measured as previously described (Flury et al., 2000; Reggiori et al., 1997). Briefly, 2.5 OD₆₀₀ units of logarithmic growth phase cells were suspended in 250 μl SC containing 25 μCi myo-[1,2-³H(N)]inositol (American Radiolabeled Chemicals). The cells were incubated for 40 min at 30°C, 750 μl SC was added, and the cells were incubated for 1 h and 20 min. The cells were pelleted and frozen at –80°C. The *gpi3-5* cells were grown at 23°C to logarithmic growth phase in SC, shifted to 37°C for 30 min, and then labeled as described except at 37°C. Cells labeled at 23°C were incubated with 50 μCi myo-[1,2-³H(N)]inositol for 24 h.

The labeled cells were thawed and resuspended in 30 μl ice-cold breaking buffer (60 mM NaPi, pH 8, 44 mM NaCl, and 55 mM glucose). 300 μl glass beads (425–600 mm; Sigma-Aldrich) and 300 μl CHCl₃:methanol(MeOH):H₂O (10:10:3) were added, and the cells were lysed in a Precellys24 homogenizer at 4°C (3 × 6,000 rpm for 30 s). 900 μl CHCl₃/MeOH/H₂O (10:10:3) was added, and the samples were vortexed and centrifuged for 5 min at 15,000 ×g at room temperature. The supernatants were dried in a SpeedVac. The samples were resuspended in 100 μl of water-saturated butanol and desalted by adding 50 μl of water, vortexed, and centrifuged for 3 min at 15,000 ×g at room temperature. The upper phase was removed and saved. The bottom phase was reextracted with 100 μl of water-saturated butanol, and the pooled upper phases were dried under N₂.

The samples were resuspended with 40 μl CHCl₃/MeOH/H₂O (10:10:3) and spotted onto Silica gel 60 TLC plates (Merck). The plates were developed for 1 h 20 min in either CHCl₃:MeOH:0.25%KCl (55:50:10) to quantify lipid 031a or CHCl₃:MeOH:H₂O (10:10:3) to quantify the GPI precursor lipid M4. The TLC plates were scanned on a RITA Star thin-layer analyzer (Raytest) to quantify radioactivity or visualized with a Typhoon FLA 9500 laser scanner (GE Healthcare).

Coimmunoprecipitation

Cells were grown to logarithmic growth phase, and 150 OD₆₀₀ units of cells were washed once with ice-cold water, pelleted, and resuspended in 600 μl of lysis buffer (50 mM Tris, pH 7.4, 150 mM NaCl, 1 mM PMSF, and a proteinase inhibitor cocktail from Roche). The cells were lysed with a Precellys24 homogenizer at 4°C, mixed with an equal volume (600 μl) of 0.5% NP-40 in lysis buffer, and incubated at 4°C for 15 min. 50 μl of each

sample was saved (whole-cell lysate), and the remainder was incubated with 100 μ l of magnetic anti-HA beads (Pierce; 88837) for 1 h at 4°C. The beads were washed five times with lysis buffer and incubated at 4°C for 5 min between washes, and the bound proteins were eluted with 40 μ l SDS-PAGE loading buffer for 15 min at 50°C. 10 μ l of whole-cell lysate and 20 μ l of immunoprecipitated proteins were separated by SDS-PAGE and immunoblotted using anti-HA antibody (C29F4; Cell Signaling Technology; #3724) and anti-GFP (D5.1; Cell Signaling Technology; #2956).

Assessment of CD55 on the surface of HEK-293 cells by FACS

Lentiviruses were produced from pLKO.1 expressing shRNA against GPAA1 (Mission Sigma-Aldrich; TRCN0000242570), KIAA1109 (Mission Sigma-Aldrich; #1 TRCN0000282583 and #2 TRCN0000263340), or negative control (Mission Sigma-Aldrich; nontargeted shRNA control) using standard methods.

HEK-293 cells were grown in DMEM in six-well dishes containing 20 μ l of lentivirus shRNA and 1.6 μ l polybrene (Sigma-Aldrich). The medium was removed after 24 h, fresh medium containing 2 μ g/ml puromycin (Sigma-Aldrich) was added, and the cultures were grown for 10 d.

The abundance of CD55 on the cell surface was determined as described (Knaus et al., 2018; Tanigawa et al., 2017). Briefly, 100,000 cells were washed two times with the DMEM with 1% FBS and incubated with 4 μ l mouse anti-human CD55 conjugated to R-Phycoerythrin (BD Bioscience; 555694) and 16 μ l DMEM with 1% FBS at 25°C. The cells were washed three times with growth medium, and the fluorescence intensity of 10,000 live cells was measured by FACS (ACSymphony Cell Analyzer; BD Bioscience). Results were analyzed with FlowJo 10.6.0 (BD Bioscience).

qRT-PCR

Total RNA was extracted from cells (Pure Link RNA kit; Thermo Fisher Scientific), 1 μ g of total RNA was used to produce cDNA (qScript cDNA SuperMix; Quantabio), and qRT-PCR was performed with the SYBR Green PCR Master Mix (Thermo Fisher Scientific) run on a CFX96 real-time thermal cycler CFX-96 (Bio-Rad). The following primers were used: for GAPDH p368 5'-GTC TCCTCTGACTTCAACAGCG-3' and p369 5'-ACCACCCTGTTG CTGTAGCCAA-3'; for GPAA1 p366 5'-TCAATAGCTTCCGCCAGT AC-3' and p367 5'-CCAAGGACCAGGAGCAAGAA-3'; and for KIAA1109/TWEEK p356 5'-CACTCTGCTATTAAGGGAC-3' and p357 5'-CTCTAATTCACACTGCCATG-3'. All reactions were done in quadruplicate. Relative expression was calculated from the Cq obtained with CFX manager 3.0 software (Bio-Rad) and with the comparative cycle threshold method, using values from the house keeping gene GAPDH and negative control shRNA.

CRISPR used to generate *C. elegans* strain AG645

The Bristol N2 strain was used as the WT for CRISPR/Cas9 genome editing. 20 nucleotide sequences of the *lpd-3*-specific CRISPR RNAs (crRNAs) were selected using a crRNA design tool (https://www.idtdna.com/site/order/designtool/index/CRISPR_CUSTOM). All crRNAs and trans-activating crRNA were synthesized by Horizon Discovery. The single-stranded donor

oligonucleotides of the *lpd-3* repair template were crRNA N-terminus: 5'-GCGAAAACGAAAAAAATGT-3', PAM site: CGG, crRNA C-terminus: 5'-TTATCCATAATATTGACATC-3', and PAM site: TGG and repair template: 5'-ACGGGTTCTAGG CGGCGGCGAAAACGAAAAAAATGTGACTAGATGATGGATAA CGACGGAAAAGGCTTCTATTGGAATA-3'. The oligonucleotides used for genotyping are N-del F1: 5'-AGTCGGCTACTACGG GTTCT-3', N-del R1: 5'-GATCCAGAAATACCGTACCTCA-3', and inner R1: 5'-CGTTTCTCAGTGGCCGAAA-3'. Day 1 gravid hermaphrodites (~20) were injected with mixed CRISPR/Cas9 reagents as described (Iyer et al., 2019). *lpd-3(av254)* was generated by CRISPR/Cas9 mixes that contained two crRNAs at flanking regions of the deleted *lpd-3* coding region. Heterozygous *lpd-3 N-Δ* animals were screened by PCR (Primer N-del gt F1 and Primer N-del gt R1). The PCR amplicon for the deletion is 403 bp. Homozygous mutants were confirmed in subsequent generations.

FLAER staining and microscopy of *C. elegans*

Staining with FLAER (Proaerolysin conjugated Alexa Fluor 488; Cedarlane Corporation; FL1S-C) was adapted from previous studies (Budirahardja et al., 2015; Murata et al., 2012). 23-gauge needles were used to dissect the embryos from the gravid hermaphrodites in M9 buffer, and the embryos were covered with an 18 × 18-mm coverslip. The embryos were then freeze-cracked and fixed in methanol at -20°C for 10 min. The slides were washed twice in PBS for 5 min and blocked with 2% bovine serum albumin solution for 30 min at room temperature. The slides were incubated in 50 nM FLAER staining solution for 2 h. After incubation, the slides were washed twice with PBS for 5 min and mounted in VECTASHIELD (Vector Labs) mounting medium containing 5 μ g/ml DAPI overnight before imaging.

FLAER stained slides were imaged by a spinning disk confocal system that includes a Photometrics Prime 95B EMCCD camera and a Yokogawa CSU-X1 confocal scanner unit. Images were acquired by Nikon's NIS imaging software using a Nikon 60 × 1.2 NA water objective with 0.5- μ m z-step size; 30–50 planes were captured. FLAER signals were quantified using ImageJ/FIJI Bio-format plugin (Linkert et al., 2010; Schindelin et al., 2012). Both the Shapiro-Wilk and Kolmogorov-Smirnov normality tests indicated that all data follow normal distributions.

For differential interference contrast (DIC) imaging of *lpd-3 N-Δ* mutants, animals were immobilized on 7% agar pads with anesthetic (0.01% levamisole in M9 buffer). DIC image acquisition was performed using a Nikon 60 × 1.2 NA water objective with a 1- μ m z-step size; 15–20 planes were captured.

Online supplemental material

Fig. S1 presents material that supports the findings shown in Fig. 1. Fig. S2 presents material that supports the findings shown in Fig. 2, Fig. 3, and Fig. 4. Fig. S3 presents material that supports the findings shown in Fig. 5. Video 1 shows the structure of *S. cerevisiae* Vps13 generated with RoseTTAFold. Video 2 shows the structure of *S. cerevisiae* Atg2 generated with RoseTTAFold. Video 3 shows the structure of *S. cerevisiae* Csf1 generated with RoseTTAFold. Video 4 shows the structure of *S. cerevisiae* Fmp27 generated with RoseTTAFold. Video 5 shows the structure of

S. cerevisiae Hob2 generated with RoseTTAFold. Table S1 lists the *S. cerevisiae* strains used in this study.

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Author contributions: A. Toulmay conceptualized the project, supervised members of the Prinz laboratory, and prepared figures. F.B. Whittle, J. Yang, X. Bai, J. Diarra, and S. Banerjee conducted the investigation. T.P. Levine and A. Golden conceptualized the project. W.A. Prinz wrote the manuscript.

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Supplemental material

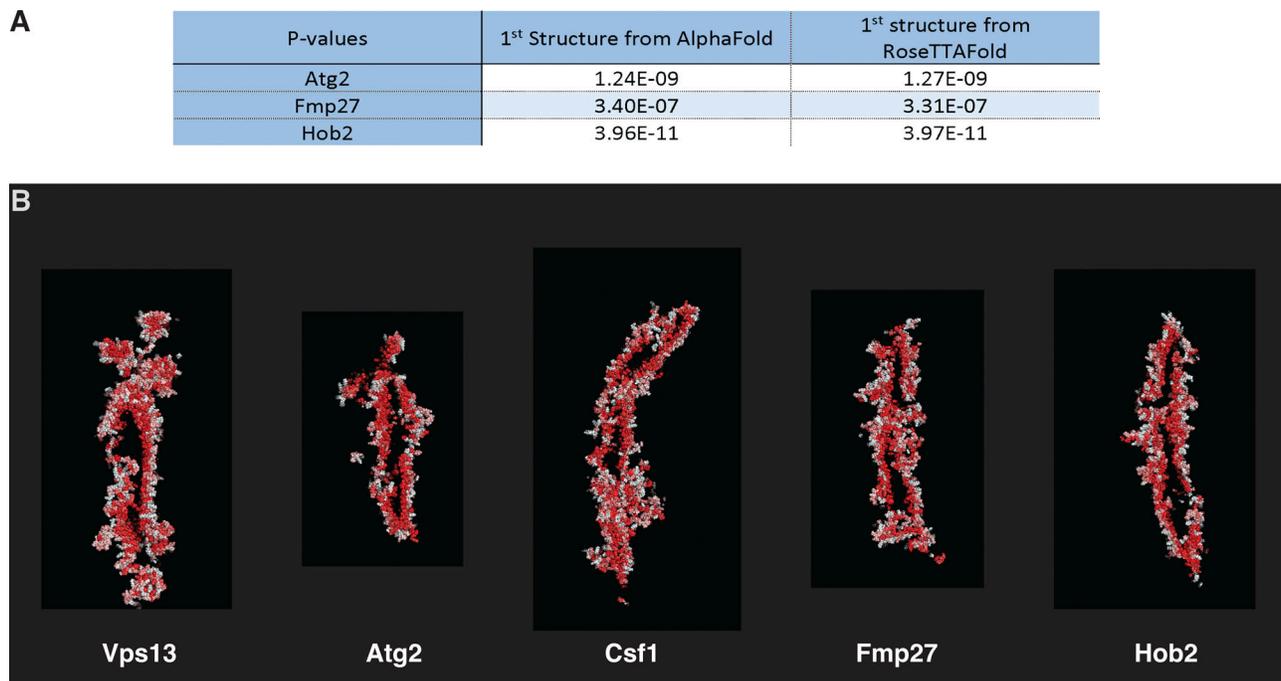


Figure S1. **Similarity of Vps13 homologues.** **(A)** Significance of the structural similarity between pairs of proteins as determined by FATCAT. P values are from pairwise comparisons of the structures of the indicated proteins as determined by RoseTTAFold and AlphaFold. **(B)** Space-filling models of the five Vps13-like proteins in *S. cerevisiae* were generated using PyMol from the predicted structures obtained from RoseTTAFold. Longitudinal cross sections are shown with hydrophobic residues colored red. All five proteins have a central hydrophobic groove.

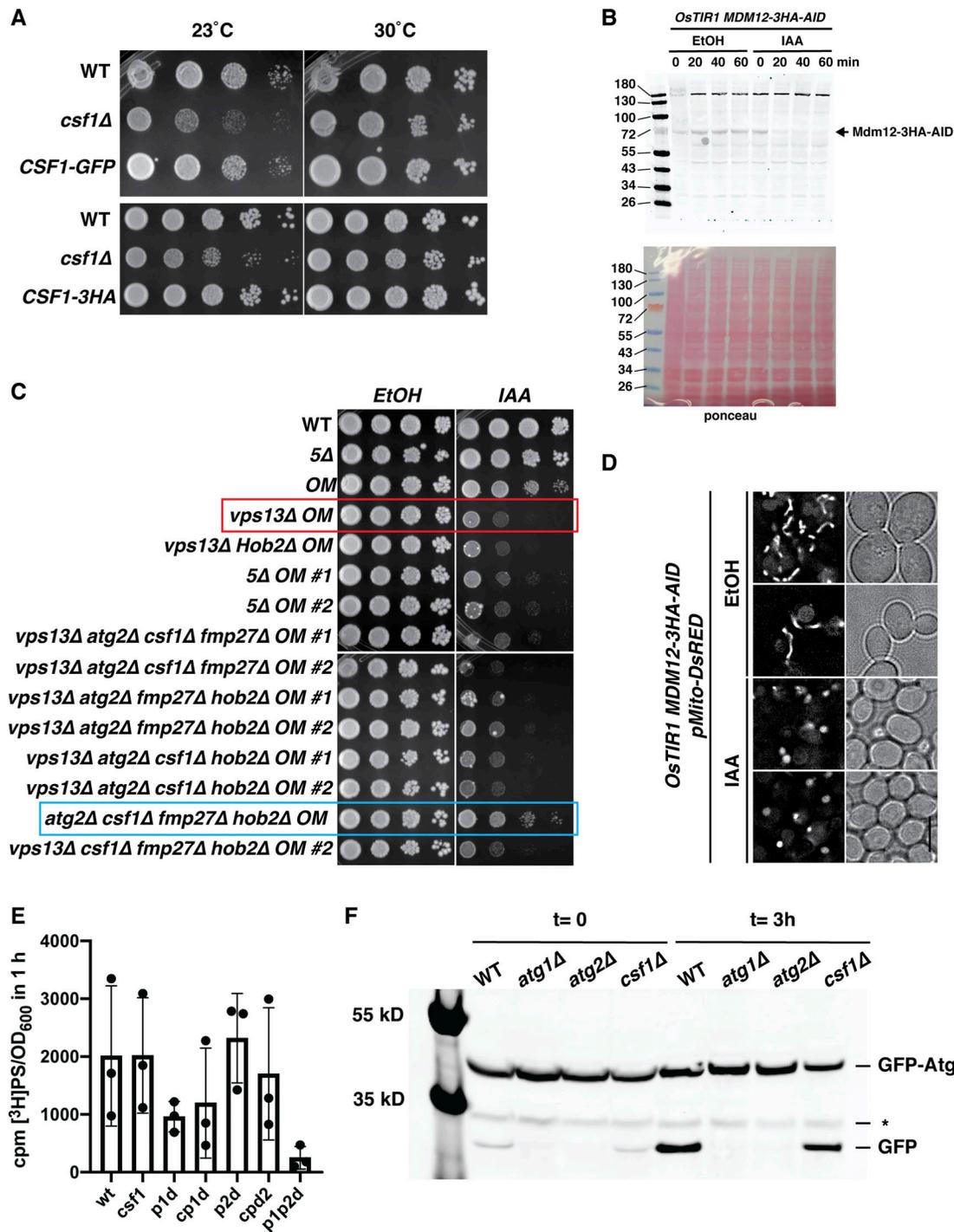


Figure S2. **Csf1 does not play a role in autophagy and other controls for findings in Fig. 2, Fig. 3, and Fig. 4.** (A) 10-fold serial dilutions of the indicated strains grown on SC plates. The plates were grown for 3 d. (B) WT cells expressing *Oryza sativa* TIR1 (OsTIR1) and Mdm12 with a C-terminal fusion to 3HA antigen tags and the auxin-inducible degron (Mdm12-3HA-AID) were grown in SC medium and the auxin IAA or ethanol (EtOH) were added. Samples were taken at the indicated times, and proteins were extracted. The proteins were separated by SDS-PAGE and transferred to a nitrocellulose membrane. The membrane was stained with Ponceau (bottom) and immunoblotted with anti-HA antibodies (top). (C) 10-fold serial dilutions of the indicated strains grown for 3 d on YPD plates containing IAA or ethanol (EtOH). OM, strains expressing OsTIR1 and Mdm12-3HA-AID (described in B). Numbers indicate size of molecular weight markers (kDa). Note that IAA prevents the growth of *vps13Δ OM* cells (red box) but not *atg2Δ csf1Δ fmp27Δ hob2Δ OM* cells (blue box), indicating that Vps13 is the only one of the five Vps13-like proteins required for viability of cells lacking ERMES. (D) Cells expressing OsTIR1 and Mdm12-3HA-AID (described in B) and the mitochondrial marker Mito-DsRed (expressed from the plasmid pMito-DsRed) were grown in SC medium and visualized live 1 h after addition of IAA or ethanol (EtOH) to the medium. Bar = 5 μ m. (E) Total [3 H]PS formed ([3 H]PS + [3 H]PE + [3 H]PC) during the 1-h labeling with [3 H]serine shown in Fig. 3 F (mean \pm SD from three independent experiments). (F) Strains with the indicated genotypes expressing GFP-Atg8 were grown in SC medium to logarithmic growth phase and transferred to SD-N medium (synthetic dextrose, lacking amino acid supplementation). Samples were taken at the indicated times and immunoblotted with an anti-GFP antibody. *, a nonspecific protein bound by the anti-GFP antibody. Source data are available for this figure: SourceData FS2.

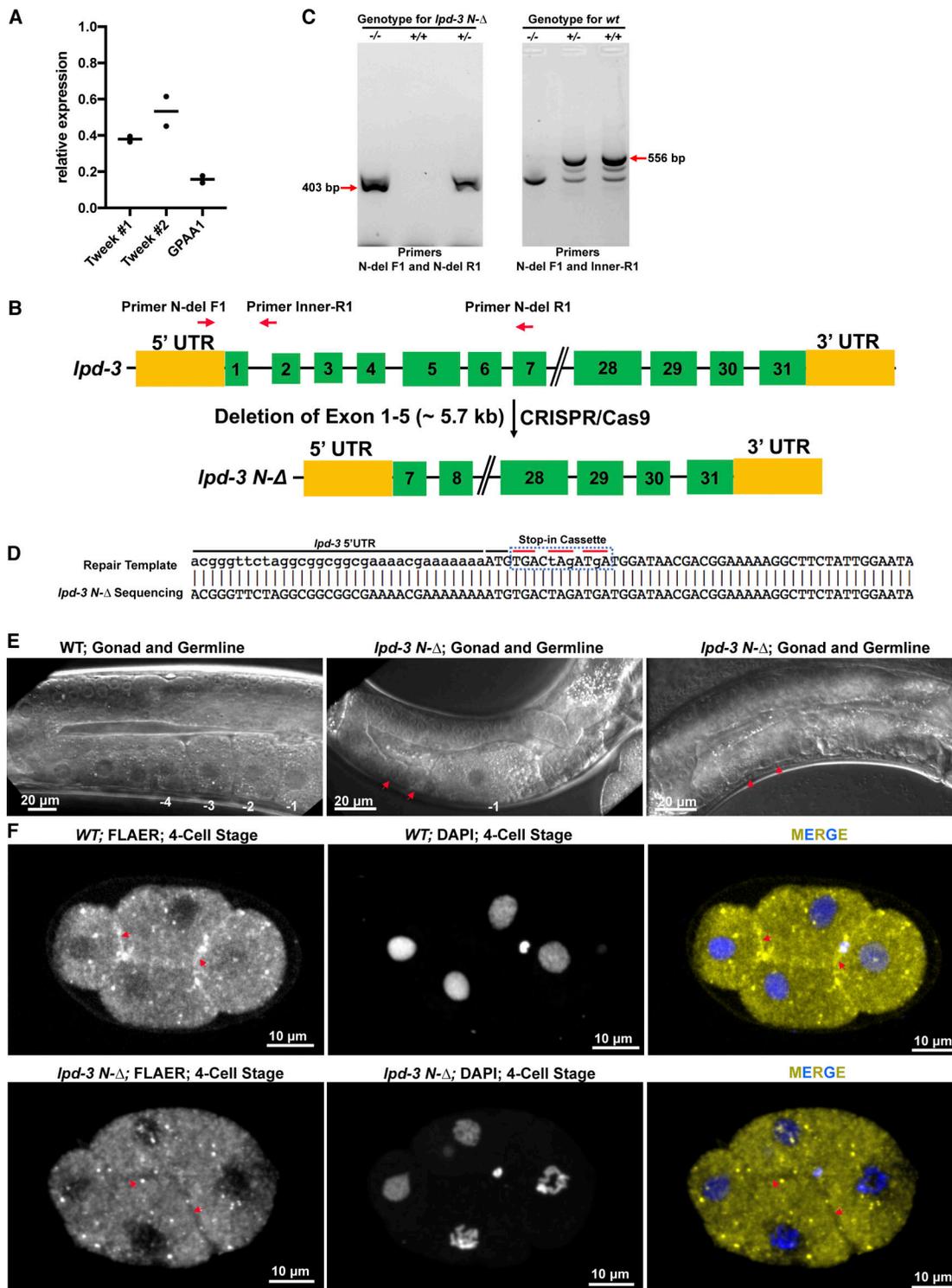


Figure S3. **Supporting information for the results shown in Fig. 5.** (A) Relative expression of the *GPAA1* and *TWEEK* genes for the experiments shown in Fig. 5 A as determined by qRT-PCR. The mean of two independent experiments ($n = 4$ per experiment). (B) Position of the *C. elegans* genotyping primers (N-del F1, N-del R1, and Inner R1) for verifying the knockout allele (red arrows); exons in green and UTRs in yellow. (C) Representative PCR results from genotyping single animals for *lpd-3 N-Δ* candidates. Two primers (two flanking primers N-del F1 and N-del R1 located outside the deleted region) were used to genotype for candidate *lpd-3(av254)* deletion animals. A homozygous deletion (-/-) was indicated by a 403-bp amplicon, which was amplified by N-del F1 and N-del R1. In WT (+/+), the PCR product amplified by N-del F1 and the inner primer Inner-R1 is 556 bp. In heterogenous animals (+/-), both amplicons will be amplified. (D) Homozygous *lpd-3 N-Δ* animals were confirmed by Sanger sequencing, which has a "STOP-IN cassette" (red lines) after start codon. (E) DIC microscopy revealed that deformed gonads and germlines were frequently observed in *lpd-3 N-Δ* animals. Bars = 20 μm. (F) FLAER staining of four-cell-stage *C. elegans* embryos from WT and *lpd-3 N-Δ* animals. The nuclei were stained with DAPI. Red arrows indicate cleavage furrows. Bars = 10 μm. Source data are available for this figure: SourceData FS3.

Video 1. **Structure of *S. cerevisiae* Vps13 generated with RoseTTAFold and ChimeraX: β -sheets (blue), α -helices (red), and coil (green).** The movie shows rotation around the longitudinal axis. Frame rate = 5 frames per second.

Video 2. **Structure of *S. cerevisiae* Atg2 generated with RoseTTAFold and ChimeraX: β -sheets (blue), α -helices (red), and coil (green).** The movie shows rotation around the longitudinal axis. Frame rate = 5 frames per second.

Video 3. **Structure of *S. cerevisiae* Csf1 generated with RoseTTAFold and ChimeraX: β -sheets (blue), α -helices (red), and coil (green).** The movie shows rotation around the longitudinal axis. Frame rate = 5 frames per second.

Video 4. **Structure of *S. cerevisiae* Fmp27 generated with RoseTTAFold and ChimeraX: β -sheets (blue), α -helices (red), and coil (green).** The movie shows rotation around the longitudinal axis. Frame rate = 5 frames per second.

Video 5. **Structure of *S. cerevisiae* Hob2 generated with RoseTTAFold and ChimeraX: β -sheets (blue), α -helices (red), and coil (green).** The movie shows rotation around the longitudinal axis. Frame rate = 5 frames per second.

Provided online is one table. Table S1 lists the *S. cerevisiae* strains used in this study.