Compartmentalized Synthesis of Triacylglycerol at the Inner Nuclear Membrane Regulates Nuclear Organization

Highlights
- Nutrients regulate the phospholipid:diacylglycerol acyltransferase Lro1
- Lro1 targets the INM subdomain bordering the nucleolus
- Lro1 is active at the INM and can sustain cell survival during starvation
- Compartmentalized synthesis of nuclear TG is important for nuclear envelope integrity

In Brief
Membrane phospholipids can be used as fatty acyl donors for the synthesis of triacylglycerols by enzymes called PDATs. Barbosa et al. present evidence that, in response to nutrient signals, the yeast PDAT Lro1 is active at the inner nuclear membrane, providing a link between organelle membrane remodeling and lipid storage.
Compartmentalized Synthesis of Triacylglycerol at the Inner Nuclear Membrane Regulates Nuclear Organization

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SUMMARY

Cells dynamically adjust organelle organization in response to growth and environmental cues. This requires regulation of synthesis of phospholipids, the building blocks of organelle membranes, or remodeling of their fatty-acyl (FA) composition. FAs are also the main components of triacylglycerols (TGs), which enable energy storage in lipid droplets. How cells coordinate FA metabolism with organelle biogenesis during cell growth remains unclear. Here, we show that Lro1, an acyltransferase that generates TGs from phospholipid-derived FAs in yeast, relocates from the endoplasmic reticulum to a subdomain of the inner nuclear membrane. Lro1 nuclear targeting is regulated by cell cycle and nutrient starvation signals and is inhibited when the nucleus expands. Lro1 is active at this nuclear subdomain, and its compartmentalization is critical for nuclear integrity. These data suggest that Lro1 nuclear targeting provides a site of TG synthesis, which is coupled with nuclear membrane remodeling.

INTRODUCTION

The internal membrane systems of the eukaryotic cell are highly dynamic, and their regulated remodeling is essential for proper organelle function, during both normal cell growth and stress. For example, the nuclear membrane undergoes dynamic remodeling during the two modes of nuclear division operating in eukaryotes, “open” and “closed” mitosis (Zhang and Oliferenko, 2013; Ungricht and Kutay, 2017), and the accumulation of unfolded proteins in the endoplasmic reticulum (ER) drives significant ER membrane expansion in order to support increased protein folding capacity (Walter and Ron, 2011). Therefore, cells must possess mechanisms to selectively add, remove, or remodel membrane phospholipids at different organelles in response to cell cycle and stress signals, but those remain poorly understood.

Lipid precursors, which are normally directed toward membrane synthesis to promote cell growth, are diverted toward storage during nutrient limitation. The main metabolic energy storage molecules in eukaryotes are triacylglycerols (TGs) which, together with other neutral lipids (e.g., steryl esters), are deposited in lipid droplets (LDs) (Wang et al., 2017). LDs emerge from, and remain associated with, the ER membrane in many cell types and interact with other organelles (Barbosa and Siniossoglou, 2017). Some of these interactions involve LDs with their “client” organelles, such as mitochondria and peroxisomes, which catabolize fatty acids stored in TGs and provide an essential source of energy during starvation (Herms et al., 2015; Rambold et al., 2015). Other interactions suggest a link of LDs to organelle membrane biogenesis. For example, LDs have been proposed to provide lipid precursors for autophagosome membrane biogenesis in yeast and mammals (Dupont et al., 2014; Shplika et al., 2015). Similarly, a specific pool of LDs associate with the expanding prospore membrane that sequesters the meiotic nuclei during sporulation of yeast cells (Ren et al., 2014; Hsu et al., 2017), and LD-mobilized fatty acids are required for bud growth and cell cycle progression in yeast (Kurat et al., 2009).

TG is synthesized by acyl-CoA:diacylglycerol acyltransferases (DGATs) or phospholipid-diacylglycerol acyltransferases (PDATs) (Ruggles et al., 2013). Most eukaryotes express DGATs while PDATs have been described so far in fungi, microalgae, and plants. Whereas DGATs use a fatty acid activated with coenzyme A (FA-CoA) to acylate diacylglycerol (DG), PDATs transfer a fatty acid from a phospholipid directly to DG (Figure 1A). Accordingly, PDATs couple TG synthesis to membrane phospholipid decylation (Dahlqvist et al., 2000; Oelkers et al., 2000) (Figure 1B).

Here, we uncover a phospholipid remodeling pathway that targets a specific subdomain of the inner nuclear membrane (INM). We find that the PDAT Lro1 of Saccharomyces cerevisiae (from hereon called yeast) is imported from the ER to the INM.
abutting the nucleolus. Lro1 is active at this specific nuclear subdomain resulting in the utilization of phospholipid-derived fatty acids to generate TGs and lysophospholipids. Interestingly, targeting of Lro1 is regulated by cell cycle and nutrient signals and is inhibited when the nucleus expands. Notably, we find that synthesis of TG at the INM sustains survival during starvation, suggesting the presence of a pathway that exports TG to the cytoplasmic side of the ER.

RESULTS

Cell Cycle and Nutrient Signals Cause Dynamic Targeting of Lro1 to a Nuclear Membrane Subdomain Associated with the Nucleolus

To determine if PDATs have a role in specific membrane remodeling events during nutrient depletion, we examined the subcellular localization of a C-terminally GFP-tagged Lro1 fusion protein when nutrients start to become scarce. All Lro1 fusions used for localization studies were catalytically active (Figure S1A). Lro1-GFP localizes to the ER during the exponential growth phase (EXP), when lipid intermediates are used to drive phospholipid synthesis to sustain rapid growth, but it relocates to a subdomain of the nuclear envelope as cells face nutrient depletion during diauxic shift (post-diauxic shift [PDS]; Figure 1C; Wang and Lee, 2012). This was observed when plasmid-borne Lro1-GFP was expressed from its own chromosomally integrated nucleolar reporter (NOP10-mCherry) were subjected to the indicated stresses and the percentage of Lro1-GFP targeting to the nucleolar-associated membrane was quantified (n = 3 experiments, at least 600 cells counted per stress condition); comparisons are between 1 or 2 h and PDS.

(Figure 1). Lro1 Targets a Nuclear Membrane Subdomain that Associates with the Nucleolus

(A) Schematic of the major lipid metabolic pathways in yeast; PA, phosphatidate; DG, diacylglycerol; TG, triacylglycerol; FA, fatty acid; LPL, lysophospholipid. (B) Schematic of the PDAT activity; PL, phospholipid. (C) Localization of Lro1-GFP expressed under the control of its own promoter in cells co-expressing an ER (Sec63-mCherry) reporter at the indicated growth phases. (D) Schematic of the organization of the yeast nucleus. (E) Co-localization of Lro1-GFP as in C but with a nucleolar reporter. (F) Left panels: examples of nucleolar enrichment of Lro1-GFP during the exponential phase; right panel, quantification from three experiments, n = 343 cells. (G) Quantification of Lro1 targeting to the nucleolar-associated membrane in response to various stresses. Exponentially growing cells expressing a chromosomally integrated nucleolar reporter (NOP10-mCherry) were subjected to the indicated stresses and the percentage of Lro1-GFP targeting to the nucleolar-associated membrane was quantified (n = 3 experiments, at least 600 cells counted per stress condition); comparisons are between 1 or 2 h and PDS. (H) Immunolabeling of chemically fixed yeast cells expressing Lro1-6xHA. Arrowheads point to gold particles clustering on one side of the nuclear envelope. Stars indicate LDs. MVB, multivesicular bodies; M, mitochondria. Scale bars in (C), (E), and (F), 5 μm; in H, 500 nm. *p < 0.05; **p < 0.01; ***p < 0.001; ns, not significant. See also Figure S1.
Nop1-RFP, we demonstrated that Lro1-GFP indeed accumulates at the membrane bordering the nucleolus (Figure 1E). Interestingly, careful analysis of Lro1 localization during exponential phase also revealed, in addition to its ER localization, an enrichment of Lro1 at the subdomain bordering the nucleolus in 34.0% ± 5.6% unbled and 34.5% ± 2.7% small budded cells, but only in 3.8% ± 5.0% of large budded cells (Figure 1F). This is consistent with Lro1-GFP accumulation at the nucleolus in PDS phase since yeast cells arrest at the G1 phase of the cell cycle at the diauxic shift (Miles et al., 2013). We also observed a similar accumulation of Lro1-GFP at this subdomain during acute glucose starvation, during growth in non-fermentable carbon sources, or when transferring the cells in water but not upon nitrogen deprivation (Figure 1G) or inhibition of rDNA transcription (Figure S1D). Immunoelectron microscopy revealed that an Lro1-6xHA fusion preferentially associated with the perinuclear ER during exponential phase (i.e., 62.6% ± 0.36%), and only part of it was at the cortical and/or peripheral ER (37.3% ± 0.21%), respectively. In the PDS phase, a significant decrease in Lro1 protein levels (see later) reduced the labeling efficiency, precluding statistical quantifications. Nevertheless, in the few cell sections where Lro1-6xHA was detected, this fusion protein was mostly found on one side of the nuclear envelope and always adjacent to LDs (Figure 1H). Taken together, these results show that glucose limitation and cell cycle signals target Lro1 to a subdomain of the nuclear membrane, which is in contact with the nucleolus.

Lro1 is Targeted to the INM

Lro1 is a type II integral membrane protein with a short basic cytosolic N-terminal domain and a larger luminal catalytic domain (Figures 2A and S2A; Choudhary et al., 2011). Expression of its N-domain fused to GFP shows a clear intranuclear localization with enrichment at the nucleolus (Figures 2B, panel 2 and S2B). Notably, the N-domain with the transmembrane segment also accumulates at the membrane in contact with the nucleolus (Figure 2B, panel 4 versus 6). Mutating the K/R residues within two basic clusters into alanines abrogates the nucleolar enrichment of both fusions in PDS phase (Figure 2B, panel 2 versus 3; panel 4 versus 5). Unexpectedly, the same mutations only partially compromise the targeting of Lro1 within the context of the full-length protein, indicating the presence of additional targeting determinants (Figures 2C and S2C). To examine whether these also map in the N-domain, we replaced it with 4 IgG binding domains of Protein A (4xIgGb) and found that this prevented detection of the resulting GFP fusion at the PDS phase, both at the nucleolus and at the ER. Because the stability of Lro1 is controlled by the ubiquitin-protein ligase Hrd1 (Iwasa et al., 2016), we imaged 4xIgGb-Lro1-GFP in an hrd1Δ strain and found out that it could be detected at the ER but not at the nucleolar-associated membrane, at the PDS phase (Figures 2C and S2C). Together, these results show that the N-domain of Lro1 is necessary and sufficient for its efficient targeting to the nuclear membrane subdomain in contact with the nucleolus; when this targeting fails in PDS, Lro1 is unstable in the ER.

Next, we compared the dynamics of the two pools of Lro1 using fluorescence recovery after photobleaching (FRAP). We found that these exhibit different properties: the nucleolar-associated membrane pool of Lro1-GFP was more immobile and its fluorescence recovery was significantly slower than that of the cortical ER, suggesting the presence of a limiting step in its targeting (Figures 2D, S2D, and S2E). Given that its soluble N-domain associates to the nucleolus, we asked whether Lro1 accesses the inner side of the nuclear membrane. We applied two assays to address this question: the first approach was based on the fact that import of integral membrane proteins from the ER to the INM through the nuclear pore is limited by the size of their cytosolic domains, with the cutoff in yeast being 90 kDa (Popken et al., 2015). In support of Lro1 residing in the INM, Lro1-GFP nucleolar targeting was significantly compromised when its extraluminal domain was made larger by appending 1, 2, or 3 copies of the maltose-binding protein (MBP) (Figures 2E and S2F). Moreover, we found that increasing the size of the N-terminal domain of Lro1 and/or preventing its nuclear import results in a significant decrease in its protein levels (Figure S2G), explaining why many cells have low 3xMBP-Lro1-GFP signal at the ER in the PDS phase. To independently determine whether Lro1 can associate with the INM, we used a second assay based on the anchor-away technique (Haruki et al., 2008). This approach requires the co-expression of two chimeric proteins: firstly, the INM protein Heh1 was fused to the FK506 binding protein (FKBP12); secondly, Lro1-GFP, or GFP as control, was fused to the FKBP12-rapamycin-binding (FRB) domain (Figure S2H). FRB and FKBP12 form a high-affinity ternary complex in the presence of rapamycin if they are in close proximity. Following the addition of rapamycin, FRB-GFP changed rapidly (30 min) from a diffuse to a ring-like localization, confirming that the INM anchor is indeed accessible to FRB-GFP (Figure 2F). Addition of rapamycin in the strain expressing FRB-Lro1-GFP resulted in the loss of its cortical ER localization and its accumulation at a perinuclear ring, which is typical of INM proteins. In contrast, FRB-3xMBP-Lro1-GFP retained its cortical ER localization after rapamycin treatment (Figure 2F), consistent with an impairment in nucleolar targeting when the mass of the N-terminal domain increases. Taken together, these data show that Lro1 targets the INM by virtue of its N-domain.

Lro1 is Catalytically Active at the INM in Contact with the Nucleolus

The nuclear membrane associated with the nucleolus has the property of being particularly susceptible to expansion in response to excess phospholipid synthesis (Campbell et al., 2006; Karamanios et al., 2010; Witkin et al., 2012). We therefore examined whether Lro1 is active by following TG synthesis at this membrane subdomain. To do this, we first sought to express Lro1 in a background where it would be the sole source of neutral lipid; hence, we used a mutant with deletions in the two DG acyltransferases (LRO1 and DAG1) and the two steryl acyltransferases (ARE1 and ARE2), henceforth called 4Δ, and which lacks neutral lipids and LDs (Oelkers et al., 2002; Petschnigg et al., 2009). Given that synthesis of neutral lipids is essential for cell survival in stationary phase, 4Δ cells display accelerated cell death in PDS that can be rescued by the expression of Lro1 as the only source of TG. This rescue requires the catalytic activity of Lro1 because mutation of Ser324 within its conserved GHSXG lipase motif abolishes the appearance of LDs (data not shown) and survival in PDS (Figure 3A). Importantly, TG levels rise concomitantly with increases in the levels of Lro1-GFP at the
nucleolar-associated membrane in 4D cells in time course experiments during exit from exponential growth (Figures 3B and 3C). Three-dimensional reconstruction of the LD distribution relative to Lro1-mCherry punctum in the PDS phase (Figure 3D; Video S1). LDs in the vicinity of the nucleolus were also observed during live imaging of 4D cells expressing Lro1 (Video S2). In both experiments, LDs that are not in proximity to the nucleolus can still be detected, these may be derived from Lro1 activity at the ER during the exponential phase or their mobility within the perinuclear ER. Taken together, these results support the notion of Lro1 being active at the INM.

Lro1 Activity Regulates Phospholipid Homeostasis

Next, we investigated the effect of Lro1 activity on membrane phospholipid homeostasis. We hypothesized two scenarios: in the first, the lysophospholipid generated by Lro1 could be reacylated with a different fatty acid, changing the physical properties of the membrane; in the second, the lysophospholipid could be further broken down by a phospholipase, effectively degrading the original phospholipid substrate of Lro1 (Figure 3E).

To discriminate between these possibilities, we first determined the subcellular distribution of the known lysophospholipid acyltransferases (Ale1 and Slc1) or phospholipases B (Nte1 and Lpl1; Plb’s 1 to 3 could not be visualized) when Lro1 localizes to the nucleolar-associated membrane. GFP-fusions of Ale1 and Nte1, and to a lesser degree Slc1, localized to the ER with no apparent co-enrichment with Lro1-mCherry during the PDS phase (Figure 3F). Next, we determined the consequences of Lro1 activity in lipid homeostasis: consistent with its PDAT activity, we found that transient overexpression of Lro1 in wild-type cells caused an increase in TG levels (Figure 3G). Cells lacking Ale1, which has general lysophospholipid acyltransferase activity in yeast (Jain et al., 2007; Riekhof et al., 2007; Tamaki et al., 2007), showed an increase in both lyso-PC and lyso-PE levels.

Figure 2. Translocation of Lro1 to the INM that Associates with the Nucleolus

(A) Schematic of the topology of Lro1. The K/R-rich nucleolar targeting sequences are shown in red. The Ser324 within the GHSXG lipase motif is shown.

(B) lro1Δ cells expressing a nucleolar reporter and the Lro1-GFP mutants shown were imaged at the indicated growth phases. Red stars denote the K/R to A mutations. Arrowheads denote the nucleolus and/or the nucleolar-associated membrane.

(C) Quantification of the subcellular localization of the indicated Lro1-GFP mutants in the specified strains. Red stars denote the K/R to A mutations within the extraluminal domain. Three colonies of each strain were analyzed; at least 200 cells were counted for each strain.

(D) Lro1-GFP was photobleached, either at the nucleolar-associated membrane or the cortical ER (cER), and fluorescence recovery was measured. Data are means ± SD from three independent experiments (seven cells each); arrow indicates the bleaching event.

(E) Nucleolar-associated membrane targeting of 1x-, 2x-, or 3x-MBP-Lro1-GFP fusions during the PDS phase. Right panel: Quantification of the data shown from three experiments, counting only cells with signal in ER or nucleolus; at least 250 cells were quantified for each strain.

(F) Localization of the FRB-GFP control (the outlines of cells are shown; vac, vacuole), and the FRB-Lro1-GFP (middle) or FRB-3xMBP-Lro1-GFP (bottom) fusions, before or after the addition of rapamycin. Arrowheads point to the cortical ER membrane. Scale bars in all micrographs, 5 µm. See also Figure S2.
compared with that seen in the wild-type strain under the same conditions; on the other hand, a mutant strain lacking four known phospholipases B showed no change in lyso-PC and a more modest increase in lyso-PE compared to that of the ale1Δ mutant (Figure 3G). This result is consistent with the Lro1-derived lysophospholipids being directed primarily to re-acylation. However, our data cannot exclude a role for additional enzymes in the processing of Lro1-derived lysophospholipids.

Lro1 Localization Correlates with Changes in Nuclear Morphology

Next, we sought to investigate the nuclear function of Lro1 by mutating proteins required for its INM targeting. To do this, we performed an unbiased screen for factors involved in Lro1 nuclear targeting. We used high content automated microscopy, followed by manual inspection, to determine the localization of Lro1-GFP in 6,000 strains carrying the loss of function mutations in all yeast genes. We focused on mutants with defects in Lro1 targeting during the PDS phase (and hence displaying increased ER localization compared to the wild-type) or decreased overall Lro1-GFP signal due to the degradation of its ER pool in the PDS phase. We identified 137 such mutants, which affect diverse cellular functions (Table S1). Nearly half of the genes of the ontology term “establishment of sister chromatid cohesion” affected Lro1-GFP targeting (Table S1). These included components of the Ctf19 complex of the kinetochore, which display a G2/M delay, consistent with the finding that Lro1 localization is regulated during the cell cycle. In those mutants, Lro1-GFP showed increased ER localization mostly in large budded cells (Figure 4A).

During G2/M mitotic delay, phospholipid synthesis is not halted resulting in expansion of the nuclear membrane that contains the nucleolus (Witkin et al., 2012). We asked whether other conditions that result in the expansion of this membrane domain correlate with loss of Lro1 targeting. During exposure to α-factor mating pheromone, a MAP kinase cascade induces...

Figure 3. Lro1 Is Catalytically Active at the Nucleolar-Associated INM

(A) Wild-type or dga1Δ lro1Δ are1Δ are2Δ (4Δ) cells expressing the indicated plasmids were grown to exponential (EXP) or PDS phases in minimal synthetic medium and spotted on YEPD plates.

(B) 4Δ cells expressing Lro1-GFP and Sec63-mCherry were grown from exponential phase to the indicated densities and imaged.

(C) 4Δ cells expressing Lro1, or an empty vector, were grown to the indicated densities, labeled with BODIPY 493/503 and their fluorescence was quantified by FACS. Data are representative of two independent experiments.

(D) 4Δ cells expressing Lro1-mCherry were grown to the PDS phase and labeled with BODIPY 493/503. Deconvolved through-focus image series were processed to generate 3D image. The full reconstructed field is shown in Video S1.

(E) Model for the Lro1-mediated regulation of phospholipid homeostasis; see text for details.

(F) Co-localization of the indicated GFP fusions with Lro1-mCherry at the PDS phase.

(G) Lipidomic quantifications of TG, LPE, LPC, PE, and PC in wild-type (BY4741), ale1Δ, and plb1Δ plb2Δ plb3Δ nte1Δ lro1Δ (5Δ) cells expressing the denoted plasmids. Cells were grown in galactose for 5 h. Lipid levels were normalized to the corresponding levels of the wild-type strain expressing the empty vector. Data shown are means of at least 5 experiments ± SD. *p < 0.05; **p < 0.01; ***p < 0.001. Scale bars in all micrographs, 5 μm. See also Videos S1 and S2.
the formation of an extended nuclear membrane “pocket” embracing the nucleolus (Stone et al., 2000) (Figure 4B). Lro1-GFP is excluded from this subdomain in α-factor treated cells, while it was still detected in contact with the nucleolus in unperturbed cycling G1/S cells. Thus, under multiple conditions where the nucleus expands, cycling anaphase (Figure 1F), G2/M delayed (Figure 4A) or α-factor-treated cells (Figure 4B), Lro1-GFP is displaced from the nucleolar-associated membrane.

The exclusion of Lro1 from the expanding INM suggests that its presence could modulate nuclear morphology. To test this, we transiently overexpressed Lro1 under two conditions that promote nucleolar membrane expansion (Witkin et al., 2012). Firstly, galactose-driven expression of Lro1 led to a 2.6-fold decrease in the percentage of wild-type cells with expanded nuclei following G2/M arrest with nocodazole (from 79% ± 4% to 30% ± 5%, p < 0.001) (Figure 4C). Secondly, we measured nuclear circularity in large budded rad52Δ cells following galactose-induced Lro1 expression. Circularity ranges from 0 to 1, the latter corresponding to a perfect circle. rad52Δ nuclear circularity increased following Lro1 overexpression (from 0.78 ± 0.14 to 0.86 ± 0.12, p < 0.001). Although Lro1-S324A overexpression also led to an increase in nuclear circularity in both conditions, the catalytic active enzyme was more efficient in restoring nuclear shape (Figures 4C and 4D). We next examined the role of Lro1 at the INM using the anchor-away technique in cells expressing the INM anchor Heh1-FKBP12 in combination with Lro1, FRB-Lro1, or FRB-Lro1-S324A. We first confirmed that FRB-Lro1 is catalytically active when anchored at the INM (Figure S3 A). Next, we incubated the cells with rapamycin to anchor the FRB-Lro1 fusions to the INM and then induced nuclear expansion with nocodazole. Since our data show that Lro1 is controlled through both targeting and DG availability (see later), we deleted DGK1 in this system to increase the DG levels at the INM. We find that FRB-Lro1 activity at the INM is required to increase nuclear circularity (Figure 4E). Collectively, these data show that Lro1 localization correlates with, and impacts, the membrane expansion of this nuclear subdomain.

**Nuclear TG Synthesis Is Sufficient to Sustain Growth during Starvation**

Our data are consistent with a role for Lro1 in TG synthesis at the INM. However, since Lro1 partitions dynamically between cortical and perinuclear ER, it is possible that a pool of Lro1...
remains active at the cortical ER at the PDS phase. Therefore, we asked whether redirecting Lro1 constitutively to the INM would be sufficient to support TG synthesis. We found that the INM-targeting sequence of Heh1 (Meinema et al., 2011) is sufficient to localize an Lro1 fusion (H1-Lro1-GFP) exclusively to a perinuclear ring both in wild-type or 4D cells, which is indicative of INM targeting (Figures 5A and 5B, panels 1 and 2). Consistently, disruption of the Asi ubiquitin ligase complex, which mediates INM protein-specific degradation (Foresti et al., 2014; Khmelinskii et al., 2014), led to an increase in both the protein and the nuclear membrane fluorescence levels of H1-Lro1-GFP (Figure S3B).

Two lines of evidence indicate that H1-Lro1 is active at generating TG at the INM in PDS phase: firstly, expression of H1-Lro1 in 4D cells led to a significant increase of TG levels (Figure 5C) and secondly, H1-Lro1 rescued the lethality of 4D cells in the PDS phase (Figure 5D). Strikingly, TG synthesis at the INM did not compromise long-term survival in stationary phase (Figure 5D).

We noticed, however, that 4D H1-Lro1 cells lack detectable LDs during exponential growth in standard glucose media (Figure 5E, panels 1 and 2). Cellular levels of DG, which is required for the formation of LDs, do not decrease significantly in the exponential phase (Figure S3C). Consistently, disruption of the Asi ubiquitin ligase complex, which mediates INM protein-specific degradation (Foresti et al., 2014; Khmelinskii et al., 2014), led to an increase in both the protein and the nuclear membrane fluorescence levels of H1-Lro1-GFP (Figure S3B). Two lines of evidence indicate that H1-Lro1 is active at generating TG at the INM in PDS phase: firstly, expression of H1-Lro1 in 4D led to a significant increase of TG levels (Figure 5C) and secondly, H1-Lro1 rescued the lethality of 4D in the PDS phase (Figure 5D). Strikingly, TG synthesis at the INM did not compromise long-term survival in stationary phase (Figure 5D).
Nuclear Lro1-Derived TG Is Packed into LDs Associated Mostly with the Outer Nuclear Membrane

If TG, which is generated by Lro1 at the INM, can sustain cell viability during starvation, then this storage lipid has to become available to cytoplasmic organelles where its fatty acids are metabolized. To investigate this, we determined the spatial positioning of LDs with respect to the nuclear membrane in 4Δ H1-Lro1 cells during the PDS phase by using enhanced resolution Airyscan microscopy. As shown in Figure 5F, LDs were detected in the nucleoplasm in close proximity to the nuclear envelope in sequential z-slices that encompass the diameter of the LD. INM-associated LDs were also detected in 4Δ cells expressing wild-type Lro1 at the PDS phase (Figure S3E). In both strains, however, LDs associated with the INM were rare. We found that the majority of LDs in the H1-Lro1 strain associate with the outer nuclear membrane. Electron microscopy using a cryosectioning procedure of chemically fixed cells for the morphological examination of 4Δ H1-Lro1 cells showed that LDs were associated with the outer side of the nuclear envelope but failed to resolve the nature of the LD-nuclear membrane association (Figure 5G). High-pressure freezing electron microscopy confirmed that LDs in this strain associate with the outer nuclear membrane (Figure 5H). Taken together, our data are consistent with a model where TG is exported from the INM and accumulates in the ONM, where it is packed into mature LDs.

Compartmentalization of INM TG Synthesis Is Important for Maintenance of Nuclear Integrity

Our data show that Lro1 can support TG synthesis when it localizes throughout the entire INM (H1-Lro1). In wild-type cells, however, Lro1 activity is restricted to a specific subdomain of the INM. We therefore wondered why yeast cells maintained this localized targeting of Lro1. We hypothesized that, when not confined, LD formation at the nuclear envelope may disrupt nuclear morphology and other nuclear functions, in particular under conditions of enhanced TG synthesis. To test this hypothesis, we first confirmed that the different localization of the Lro1 mutants correlates with a distinct subcellular distribution of LDs in 4Δ cells. Indeed, consistent with its constitutive perinuclear localization, H1-Lro1-derived LDs appeared nearly exclusively associated with the nuclear envelope in the PDS phase or after supplementation of oleate. In contrast, under the same conditions, cells expressing 3xMBP-Lro1 displayed an increased number of cortical LDs (Figures 6A and 6B). Next, we evaluated nuclear morphology in these cells by measuring the size and shape of their nuclei. The average cross-sectional surface of nuclei from the 4Δ H1-Lro1 cells (1.93 ± 0.78 μm²) showed a modest but significant decrease when compared with cells expressing the wild-type enzyme (2.08 ± 0.71 μm²), while the opposite result was observed for 4Δ 3xMBP-Lro1 cells (2.19 ± 0.84 μm²) (Figure 6C). To evaluate the nuclear shape, we measured nuclear circularity during the PDS phase. Circularity was significantly lower in 4Δ H1-Lro1 than in 4Δ Lro1 cells (Figure 6D). The nuclei of 4Δ 3xMBP-Lro1 cells displayed higher circularity than those of 4Δ H1-Lro1 cells although they displayed a modest decrease when compared to 4Δ Lro1 nuclei (Figure 6D). Thus, shifting Lro1 from the cortical ER to the INM decreases nuclear surface, consistent with a role of PDAT activity in organelle remodeling.

DISCUSSION

Eukaryotic cells possess efficient mechanisms for TG synthesis and packing at the cytoplasmic ER. Depending on the cellular metabolic requirements, fatty acids stored in TGs can be used for energy production and/or membrane biogenesis. In budding yeast, the tight association of the nucleolus with the nuclear periphery defines a membrane subdomain, which has been implicated in nuclear shape, nucleophagy, and rDNA anchoring to the nuclear envelope. We find that the yeast acyltransferase Lro1 is active at this subdomain, generating TGs from phospholipid-derived fatty acids, in response to cell cycle and nutrient signals.

Several reports have documented the presence of intranuclear LDs, but the origin of the TG composing them, as well as their roles, remain elusive (Layerenza et al., 2013; Uzbekov and Rinegard, 2013; Cartwright et al., 2015; Grippa et al., 2015; Wolinski et al., 2015; Ohsaki et al., 2016; Romanuska and Köhler, 2018). The DGAT Dga1 makes the majority of TG in yeast cells at the stationary phase (Oelkers et al., 2002; Sandager et al., 2002). Under these conditions, we showed that Lro1 associates with, and concentrates at, the INM subdomain. Thus, our data support the notion of a restricted activity of Lro1 at the nuclear envelope, which may explain its limited contribution in bulk TG synthesis in wild-type cells during starvation (Oelkers et al., 2002). Given its topology, however, we cannot exclude that Dga1 is also contributing to nuclear TG synthesis. In fact, DGAT2 association with intranuclear LDs was proposed to mediate their expansion in hepatocyte-derived cell lines (Ohsaki et al., 2016).

Although it is generally accepted that TG storage in LDs takes place at the ER, our data reveal that cells can survive during
When engineered to depend exclusively on TG generated at the INM. Given that the final destination of the TG-stored fatty acids during starvation are peroxisomes, mitochondria, and the vacuole, this result is consistent with the presence of a pathway that delivers nuclear TG to its cytoplasmic destinations. How such a process could operate will be the subject of future studies, but some scenarios can be hypothesized. Low levels of TG can be accommodated between the leaflets of a phospholipid bilayer and could diffuse through the nuclear pore membrane to the outer membrane, where they would be packed into mature LDs. Alternatively, budding of LDs toward the luminal side could channel them toward the outer nuclear membrane (Figure 6H). Our data on the requirement of nuclear membrane and pore integrity under conditions of enhanced TG synthesis at the INM is consistent with both scenarios. However, LDs can bud toward the nucleoplasm in mutants that affect the phospholipid composition of the ER and that of the LD monolayer, (Cartwright et al., 2015; Grippa et al., 2015; Romanauska and Köhler, 2018). Therefore, membrane phospholipid composition may be critical in determining the directionality of LD budding.

In yeast cells, the nuclear membrane expands to allow anaphase to take place within an intact nucleus. The membrane associated with the nucleolus can also expand in response to excess phospholipid synthesis, resulting in alterations of nuclear shape (Campbell et al., 2006; Witkin et al., 2012). Although the precise mechanisms of this process remain to be determined, one possibility could be that Lro1-mediated generation of either lysophospholipids or re-esterified phospholipids with a distinct fatty-acyl composition modifies the biophysical properties of this membrane subdomain and its dynamics. Because Lro1 targeting to this subdomain is prevented under conditions that require nuclear expansion, it is tempting to speculate that Lro1 plays a role in the regulation of this process. The pool of DG that is required for TG synthesis by Lro1 (Figures 1B and S3D) could be provided by Pah1, which is known to target the nuclear envelope at the diauxic shift (Barbosa et al., 2015), but other
enzymes could act in concert with Lro1 as well. We cannot also exclude the possibility that Lro1 controls nuclear organization through additional mechanisms that are independent of its catalytic activity. The highly basic N-domain of Lro1 faces the nucleoplasm and its interaction with nucleic acid or protein components of the nucleolus could restrict the expansion of its membrane subdomain.

It is conceivable that Lro1-mediated membrane remodeling could also play additional roles at the INM. Changes in nuclear membrane dynamics may be required for the removal of nuclear material during starvation conditions via piecemeal microautophagy (Roberts et al., 2003) and receptor-mediated microautophagy (Mochida et al., 2015; Mostofa et al., 2018). Both processes take place in proximity to the nucleolus. However, we found that at least one of these two processes, receptor-mediated microautophagy, proceeds during glucose starvation in the absence of Lro1 (data not shown). Lro1-mediated membrane remodeling could be also linked to nucleolar functions. For example, rDNA transcription and ribosome biogenesis are energy-consuming processes acutely inhibited during starvation, and they lead to a decrease in nucleolar size (Neumuller et al., 2013). Since the rDNA is physically tethered to the INM in yeast (Mekhail et al., 2008) Lro1 could remodel this membrane subdomain to facilitate nucleolar reorganization during starvation. Future studies will be needed to fully elucidate the role of INM lipid composition in nucleolar functions.

Previous studies reported that PDATs are present in fungi, green algae and plants, and they have a topology similar to that of Lro1 (Stähl et al., 2004; Pan et al., 2015). Our analysis identified PDATs from two additional taxonomic groups, the flagellates Euglenozoa and the fungal-like Oomycetes (Table S2). However, the nucleolar-associated membrane has been described so far only in fungi, raising the question of PDAT function in other taxonomic groups. Notably, the green algal C. reinhardtii PDAT was proposed to use chloroplast membrane lipids to synthesize TG during starvation (Yoon et al., 2012). Similarly, the Arabidopsis thaliana Lro1 ortholog PDAT1, which also generates TG using phospholipids as acyl donors (Stähl et al., 2004), re-localizes from the ER to chloroplasts following starvation induced by light deprivation (data not shown). Therefore, we speculate that PDATs respond to the need of remodeling or turnover of membranes. Given that the requirements of different cell types during stress are often distinct, PDATs may have evolved to target diverse organelles. It will be interesting to define the signals that govern PDAT dynamics in different cell types and examine whether animal cells, which lack apparent PDAT orthologs, maintain the ability to remodel membranes by a combination of phospholipase and acyltransferase activities.

**STAR METHODS**

Detailed methods are provided in the online version of this paper and include the following:

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  - Yeast Lipid Profiling
  - Immunoblotting
  - Flow Cytometry
  - Bioinformatics
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**SUPPLEMENTAL INFORMATION**

Supplemental Information can be found online at https://doi.org/10.1016/j.devcel.2019.07.009.

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**AUTHOR CONTRIBUTIONS**


**DECLARATION OF INTERESTS**

The authors declare no competing interests.

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**SUPPORTING CITATIONS**

The following reference appear in the Supplemental Information: Albert et al., 2016.

**REFERENCES**


Miles, S., Li, L., Davison, J., and Breeden, L.L. (2013). Xbp1 directs global repression of budding yeast transcription during the transition to quiescence and is important for the longevity and reversibility of the quiescent state. PLoS Genet. 9, e1003854.


# STAR METHODS

## KEY RESOURCES TABLE

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**Experimental Models: Organisms/Strains**

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- *S. cerevisiae*: BY4742 MATαhis3Δ1 leu2Δ0 lys2Δ0 ura3Δ0
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- *S. cerevisiae*: BY4741 plb1::hphNT1 plb2::KanMX plb3::NatMX6 nte1::URA3 lro1::HIS3
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- *S. cerevisiae*: BY4742 vps4::KanMX
- *S. cerevisiae*: BY4741 ALE1-GFP::hphNT1

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**LEAD CONTACT AND MATERIALS AVAILABILITY**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Symeon Siniossoglou (ss560@cam.ac.uk).

**EXPERIMENTAL MODELS AND SUBJECT DETAILS**

**Yeast Strains, Plasmids, Media and Growth Conditions**

Unless otherwise specified, reagents were obtained from Sigma (St. Louis, MO). Yeast strains and plasmids are described in the Key Resources Table. Yeast plasmids were generated using standard PCR and cloning techniques. Cells were transformed using the lithium acetate method. Gene deletions and epitope tagging by chromosomal integration were generated by one-step polymerase chain reaction (PCR)-based method (Longtine et al., 1998; Janke et al., 2004; Haruki et al., 2008) and confirmed by PCR. In most experiments, cells were grown overnight at 30 °C in synthetic complete (SC) medium containing 2% glucose, 0.2% yeast nitrogen base (YNB, Difco, BD, Franklin Lakes, NJ), 0.6% ammonium sulfate and amino acids drop-out (60 mg/L leucine, 55 mg/L adenine, 55 mg/L uracil, 55 mg/L tyrosine, 20 mg/L of arginine, 10 mg/L histidine, 60 mg/L isoleucine, 40 mg/L lysine, 60 mg/L phenylalanine, 50 mg/L threonine, 10 mg/L methionine, 40 mg/L tryptophan) to exponential phase (to OD600nm 0.4-0.6), PDS phase (inoculated at...
OD₆₀₀nm 0.05 and grown for approximately 15h to OD₆₀₀nm 4-6), or the indicated OD₆₀₀nm, from fresh pre-cultures, according to the previously determined growth rate in this medium (Barbosa et al., 2015).

For stationary phase survival assays, cells were grown in minimal medium (MM) with 0.17% yeast nitrogen base (Difco, BD, Franklin Lakes, NJ), 0.5% ammonium sulfate and amino acids drop-out (20 mg/L uracil, 20 mg/L histidine, 20 mg/L methionine, 30 mg/L leucine, and 30 mg/L lysine) to PDS phase (OD₆₀₀nm ~ 4), and viability was measured every 24 hours by standard dilutions on YEP plates [2% glucose, 2% bactopeptone (BD, Franklin Lakes, NJ), 1% yeast extract (BD, Franklin Lakes, NJ)] with 2% glucose (YEPD medium) and 2% galactose for the indicated times.

For GAL1/10 promoter-mediated overexpression, cells were first pre-grown in selective SC medium with 2% glucose, grown overnight in selective SC medium with 2% raffinose, and transferred to selective SC medium with 2% galactose for the indicated times.

Plasmid selection. Enrichment of Lro1-GFP at the nucleolus in exponential phase was determined in asynchronous cultures. Rapamycin-induced heterodimerization of FKBP12 with the FRB domain of human mTOR was performed as previously described in the anchor-away technique (Haruki et al., 2008). Briefly, cells were grown in selective SC medium to exponential phase and treated with 1 µg/ml rapamycin for 30 min. In nocodazole-mediated cell cycle arrest experiments, GAL1/10 promoter-mediated overexpression was induced as described above but growing the cells in YEPD media with 2% galactose for 3 h before adding 15 µg/mL nocodazole for 2 h; in the anchor-away strains, cells were first transferred to YEPD media and treated with 1 µg/ml rapamycin for 30 min before adding 15 µg/mL nocodazole for 2 h. For alpha-factor cell cycle arrest, cells were treated with 10 µM alpha-factor for 2 h in YEPD. Lipid droplets were stained with 1.25 µg/mL BODIPY 493/503 for 10 min at room temperature.

Cells grown to the indicated growth phases were pelleted and immediately imaged live at room temperature in a Zeiss AxioImager.Z2 epi-fluorescence upright microscope with a 100× Plan-Apochromatic 1.4 numerical aperture (NA) objective lens (Carl Zeiss Ltd, Jena, Germany). Images were recorded using a large chip sCMOS mono camera for sensitive fluorescence imaging (ORCA Flash 4.0v2, Hamamatsu, Hamamatsu, Japan), saved by Zeiss ZEN2.3 software (Blue edition, Carl Zeiss Ltd, Jena, Germany) and exported to Adobe Photoshop (Adobe, San Jose, CA). Where indicated, cells were mounted on a 1% agarose pad and imaged using a Zeiss LSM880 confocal microscope with Airyscan and the ZEN2 software (Carl Zeiss Ltd, Jena, Germany). Cells were visualized from the periphery by taking 0.2 µm step slices and 50% overlap, and 0.187 µm axial resolution. All microscopy images were captured blindly and quantifications were performed on fields obtained from independent experiments.

For three-dimensional (3D) analysis, through-focus image series were deconvolved with Velocity 6.3 (PerkinElmer, Waltham, MA) using calculated point-spread functions and 3D iterative restoration processing to form 3D image stacks. Stacks were then visualized using Velocity 6.3 (PerkinElmer, Waltham, MA) to generate the 3D images or movies showing the organelles at different positions. For circularity and surface of the nucleus measurements, through-focus image series were deconvolved with Velocity 6.3 (PerkinElmer, Waltham, MA) and merged in 2D images showing the brightest intensity through z, which were exported and analysed in ImageJ 1.47v (NIH, Bethesda, MD).

Time-lapse imaging was performed in a Leica TCS SP8 confocal microscope. Yeast cells (4Δ) expressing Faa4-GFP, Nop1-RFP, and Lro1 under its endogenous promoter in a CEN vector, were grown in selective media to PDS phase and imaged on 2% agar pads. Images were acquired using a 63x oil immersion objective lens with an 8x zoom factor at 30 seconds intervals using white light laser at 488 nm and 578 nm for the excitation of GFP and RFP, respectively. Videos were generated using Imaris at a rate of three frames/second.

Fluorescence Recovery after Photobleaching

Photobleaching experiments were performed on a Leica TCS SP8 confocal microscope with the optional FRAP Booster enabled. Yeast lro1Δ cells expressing LR01-GFP from NOP1 promoter were pelleted and imaged at room temperature at early PDS phase. Images were acquired using a 63x oil immersion objective lens. After acquiring two images at 5-sec intervals, selected regions of interest were photobleached with 3 iterations of 100% laser power (white light laser) at 488 nm. The fluorescence intensity of GFP at the regions of interest was recorded for another 13 frames at 20 sec intervals. For data analyses, the fluorescence intensity of GFP was corrected by the percentage loss of GFP fluorescence intensity obtained from cells under identical conditions but without a photobleaching event. To calculate halftime of recovery (T₁/2), exponential FRAP curve fitting (non-linear regression, one way
Automated Yeast Library Manipulations and High-Throughput Microscopy Screen

Lro1-GFP, expressed from the NOP1 promoter in a CEN/ARS URA3 vector, was introduced in the KanMX deletion and DAmP collections (Giaever et al., 2002; Breslow et al., 2008) by Synthetic Genetic Array, by standard procedures previously described (Cohen and Schuldiner, 2011), using the RoToR bench-top colony arrayer (Singer Instruments, Roadwater, Watchet, UK). Cells were imaged, at room temperature at PDS phase in SC medium lacking uracil, using the automated inverted fluorescent microscopic Scanning system (Olympus, Waltham, Massachusetts, USA), with a 60 x air lens, for GFP (excitation, 490/20 nm; emission, 535/50 nm) and brightfield channels. After acquisition, images were manually reviewed using the ImageJ software (NIH, Bethesda, MD). Cells showing increased Lro1-GFP signal in the ER, or overall decreased targeting to the nucleus without enrichment at other subcellular localizations were selected for further analysis. The data obtained were analysed with GO (gene ontology) term finder of Saccharomyces Genome Database to determine GO term enrichment (https://www.yeastgenome.org/goTermFinder), and selecting the “Process” ontology aspect. Based on this analysis, Lro1-GFP localization was then manually inspected in mutants of interest (see Table S1).

Electron Microscopy

Cells were grown to the indicated growth phases, chemically fixed, embedded in 12% gelatin and cryo-sectioned as described previously (Griffith et al., 2008). Ultrathin cryo-sections were collected with a 1:1 mixture of 2% methylcellulose and 2.3 M, sucrose 120 mM PIPES, 50 mM Hepes, pH 6.9, 4 mM MgCl₂, 20 mM EGTA, and layered on Formvar/carbon coated 100 mesh copper grids. Immunological reactions were performed using a polyclonal anti-HA and a protein A-gold 10 nm conjugate (Cell Microscopy Center, Utrecht, The Netherlands). To determine protein localization, the distribution of gold particles was counted in 100 cells. The relative distribution of Lro1-6xHA to the cortical and the nuclear ER, was performed by counting the number of gold particles associated to these two cellular sub compartiments, on cell profiles randomly screened from immuno-labeled cell sections derived from at least three different grids. Gold particles were assigned to a compartment when no further than 10 nm away from its limiting membrane.

For high-pressure freezing electron microscopy, 4 D H1-Lro1 cells were grown to PDS phase, pelleted and resuspended in a minimum volume of 20% BSA + 5% FBS/PBS to form a paste, and this paste was pipetted into flat specimen carriers and high-pressure frozen with a Leica EM PACT2. Freeze-substitution was performed with a Leica EM AFS2, and pellets were immersed in 0.1% tannic acid acetone at -90°C for 24 hours, before being replaced by 2% osmium/ 0.1% UA (methanolic)/acetone for 48 hours at -90°C. Samples were then gradually warmed as follows; warmed to -56°C at a rate of 5°C/hour, held at -56°C for 12 hours, warmed to -20°C at a rate of 5°C/hour, held at -20°C for 12 hours, warmed to +4°C at a rate of 5°C/hour, held at +4°C for 4 hours. Pellets were washed three times with acetone before being gradually infiltrated with Spurr’s resin over a period of 4 days. Ultrathin sections were cut using a diamond knife mounted to a Reichert Ultracut S ultramicrotome, and floating sections were collected onto copper grids. Grids were poststained first with both 2% uranyl acetate/70% methanol for 4 minutes and then with lead citrate for 4 minutes. Sections were viewed on an FEI Tecnai transmission electron microscope at a working voltage of 80 kV.

Yeast Lipid Profiling

Yeast cells were prepared for mass spectrometry analysis as previously described (Folch et al., 1957) with minor modifications. Briefly, 50 mg of yeast cells were re-suspended in 1 mL of 2:1 chloroform-methanol mixture (v/v), with the addition of 150 μL of the following internal standard solution (approximately 10 to 50 μM in methanol): CE_(18:0), Ceramide_C16:0, FA_C15:0, FA_C17:0, FA_C20:0, LGP_(14:0), PA, PA(C16:0, C18:1), PC, PE, PG, PI, PS, SM, TG, and TAG. The cells were homogenized with 100 mg 0.5 mm diameter glass beads (BioSpec Products, Bartlesville Oklahoma, USA) in a FastPrep-24 instrument (MP Biomedicals, Santa Ana California, USA), using five short pulses at 5 m/s for 1 min; with one minute on ice between each pulse to prevent over heating. Then 400 μL of sterile water was added to the homogenates, vortexed for 1 min, and then centrifuged at 13,200 rpm for 10 minutes. The organic layer was collected in a 2 mL amber glass vial (Agilent Technologies, Santa Clara California, USA) and the remaining mixture was then treated with a second lipid extraction following the same procedure. The two organic layers were combined and air-dried overnight in a fume hood.

For lipid analysis, we used a method previously described with minor modification (Koulman et al., 2009; Lu et al., 2016). Full chromatographic separation of intact lipids was achieved using Shimadzu HPLC System (Shimadzu UK Limited, Milton Keynes, United Kingdom) with the injection of 10 μL onto an Acquity UPLC® CSH C18 column; 1.7 μm, I.D. 2.1 mm × 50 mm, maintained at 55°C. Mobile phase A was 64: acetonitrile and water with 10 mm ammonium formate. Mobile phase B was 9:1, 2-propanol and acetonitrile with 10 mM ammonium formate. The flow was maintained at 500 μL per minute through the following gradient: 0.00 minutes to 40% mobile phase B; 0.40 minutes, 43% mobile phase B; 0.45 minutes, 50% mobile phase B; 2.40 minutes, 54% mobile phase B; 2.45 minutes, 70% mobile phase B; 7.00 minutes, 99% mobile phase B; 8.00 minutes, 99% mobile phase B; 8.3 minutes, 40% mobile phase B; 10 minutes, 40% mobile phase B. The sample injection needle was washed using 9:1, 2-propanol and acetonitrile with 0.1 % formic acid.

The mass spectrometer used was the Thermo Scientific Exactive Orbitrap with a heated electrospray ionisation source (Thermo Fisher Scientific, Hemel Hempstead, UK). The mass spectrometer was calibrated immediately before sample analysis using positive ionization mode.
and negative ionisation calibration solution (recommended by Thermo Scientific). Additionally, the heated electrospray ionisation source was optimised at 50:50 mobile phase A to mobile phase B for spray stability (capillary temperature; 380°C, source heater temperature; 420°C, sheath gas flow; 60 (arbitrary), auxiliary gas flow; 20 (arbitrary), sweep gas; 5 (arbitrary), source voltage; 3.5 kV. The mass spectrometer resolution was set to 25,000 with a full-scan range of m/z 150 to 1200 Da, with continuous switching between positive and negative mode.

Lipid quantification was achieved using the area under the curve (AUC) of the corresponding high resolution extracted ion chromatogram (with a window of ± 8 ppm) at the indicative retention time. The lipid analyte AUC relative to the internal standard AUC for that lipid class was used to semi-quantify and correct for any extraction/instrument variation. The normalized analyte intensities (analyte to internal standard ratios) were then expressed as a percentage of the total lipids extracted from that sample (mol %).

Immunoblotting
Yeast cells (approximately 12 OD600) were pelleted, washed with sterile water, and lysed in 100 μl SDS-sample buffer with 0.5 mm diameter glass beads (BioSpec Products, Bartlesville, OK) by two rounds of boiling for 2 min and vortexing for 30 sec. Protein extracts were centrifuged at 13,000 rpm for 15 min, and the supernatants analysed by Western blot. Western blot signals were developed using ECL (GE Healthcare, Little Chalfont UK).

Flow Cytometry
Quantification of BODIPY by flow cytometry was performed as previously described (Barbosa et al., 2015). Briefly, cells were fixed for 30 min at room temperature with 3.7% formaldehyde, washed once with phosphate-buffered saline, and incubated with 10 μM BODIPY 493/503 for 10 min at room temperature. Labelling was immediately measured using the FL-1 detector of a FACSCalibur flow cytometer (BD Biosciences, San Jose, CA), and the results analysed with FlowJo software, version 9 (Tree Star, Ashland, OR).

Bioinformatics
For the identification of PDATs, the UniProt Knowledgebase (UniProtKB) was searched with the search term phospholipid:diacylglycerol acyltransferase at http://www.uniprot.org/. Partial sequences, sequences with zero or multiple transmembrane domains were removed. In order to extend the number of PDATs, we performed a phmmer search on the HMMER web server at https://www.ebi.ac.uk/Tools/hmmer/ (Finn et al., 2015) using the S. cerevisiae Lro1 (UniProtKB accession P40345) sequence to query UniProtKB. Additional plant PDATs, identified in Pan et al. 2015 were obtained from the Phytozome resource (Goodstein et al., 2012).

QUANTIFICATION AND STATISTICAL ANALYSIS
Unless otherwise stated, data were obtained from at least three independent repeats. The micrographs presented are representative of the results obtained. Signal intensity, surface and circularity were measured using ImageJ 1.47v (NIH, Bethesda, MD). Results were expressed as mean ± standard deviation. Data were analysed with unpaired, two-tailed t tests or one-way ANOVA with Tukey’s multiple comparison test when more than two groups were compared using GraphPad Prism 5 software (GraphPad, La Jolla, CA). Statistical significance was defined as: *, p < 0.05; **, p < 0.01; and ***, p < 0.001.

DATA AND CODE AVAILABILITY
Datasets associated with this study are provided in Tables S1 and S2.