Segregation of sphingolipids and sterols during formation of secretory vesicles at the trans-Golgi network

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The trans-Golgi network (TGN) is the major sorting station in the secretory pathway of all eukaryotic cells. How the TGN sorts proteins and lipids to generate the enrichment of sphingolipids and sterols at the plasma membrane is poorly understood. To address this fundamental question in membrane trafficking, we devised an immunoisolation procedure for specific recovery of post-Golgi secretory vesicles transporting a transmembrane raft protein from the TGN to the cell surface in the yeast Saccharomyces cerevisiae. Using a novel quantitative shotgun lipidomics approach, we could demonstrate that TGN sorting selectively enriched ergosterol and sphingolipid species in the immunoisolated secretory vesicles. This finding, for the first time, indicates that the TGN exhibits the capacity to sort membrane lipids. Furthermore, the observation that the immunoisolated vesicles exhibited a higher membrane order than the late Golgi membrane, as measured by C-Laurdan spectrophotometry, strongly suggests that lipid rafts play a role in the TGN-sorting machinery.

Introduction

One of the major functions of membrane trafficking in eukaryotic cells is to provide each organelle connected within the system with its functional complement of proteins and lipids (van Meer, 1989; Mellman and Warren, 2000). Biosynthesis of secretory proteins starts in the ER, and the glycosylation of proteins is completed in the Golgi apparatus (Mellman and Warren, 2000). From the TGN, proteins are distributed to the plasma membrane (PM), endosomes, and lysosomes (Griffiths and Simons, 1986; Traub and Kornfeld, 1997; Bard and Malhotra, 2006; Pfeffer, 2007). The lipid composition changes progressively throughout the secretory pathway; the ER displays a relatively low concentration of sterols and sphingolipids, which in turn accumulate toward the PM (Simons and van Meer, 1988; van Meer and Simons, 1988; van Meer, 1989; Zinser and Daum, 1995).

Lipid rafts have been proposed to be involved in the generation of the lipid gradients in the secretory pathway (Simons and Ikonen, 1997). These membrane structures have evolved from controversial detergent-resistant entities to dynamic, nanometer-sized membrane domains formed by sterols, sphingolipids, saturated glycerophospholipids, and proteins (Simons and Vaz, 2004; Hancock, 2006). Lipid raft coalescence can be induced by lipid-protein and protein–protein interactions to form ordered membrane microdomains involved in signal transduction (Simons and Toomre, 2000), virus assembly (Brügger et al., 2006), and membrane trafficking (Schuck and Simons, 2004). Raft involvement in TGN cargo sorting was suggested from work on apical transport in epithelial cells and formed the basis for the raft concept as it was first formulated (Simons and van Meer, 1988; Simons and Ikonen, 1997). However, direct evidence indicating sorting of lipids in membrane trafficking is still lacking, and the function of rafts in the formation of post-Golgi transport carriers remains elusive.

In this work, we aimed to provide answers to the issue of whether the TGN can sort lipids during transport carrier assembly.

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References


Acknowledgments

This work was supported by grants from the Deutsche Forschungsgemeinschaft (Simons, Ku 1552/2-2 and G1/13-1), the European Research Council (Simons, ERC Grant 207947), and the Spanish Ministry of Science and Innovation (Surma and Klemm, BFU2007-68139). We would like to thank the German Research Foundation for funding the Center for Molecular Biophysics (SFB802).

Supplemental Material can be found at: http://jcb.rupress.org/cgi/content/full/jcb.200901145/DC1

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If raft coalescence plays a functional role in the sorting machinery that forms carrier vesicles transporting raft proteins to the cell surface, these secretory vesicles should be selectively enriched in sterols and sphingolipids. However, testing this hypothesis has so far been hampered by numerous shortcomings of biochemical and analytical methodology available for such experiments. In this study, we used the yeast *Saccharomyces cerevisiae* as our experimental system. The molecular lipid composition of *S. cerevisiae* includes ergosterol (in contrast to cholesterol in mammalian cells), three classes of inositol-containing sphingolipids, inositolphosphoceramide (IPC), mannosyl-IPC (MIPC), and mannosyl-di-IPC (M[IP]2C), glycerophospholipids, and glycerolipids (Daum et al., 1998; Holthuis et al., 2001). Similar to epithelial cells, yeast uses at least two distinct cell surface delivery pathways (Simons and Wandinger-Ness, 1990; Wandinger-Ness et al., 1990; Harsay and Bretscher, 1995; Bagnat et al., 2000; Gurunathan et al., 2002; Harsay and Schekman, 2002; Rodriguez-Boulan et al., 2005). The pathway that transports raft proteins from the TGN to the PM is mediated by a population of light density secretory vesicles (LDSVs), which are similar to apical transport carriers in epithelial cells (Harsay and Bretscher, 1995; Bagnat et al., 2000). In a visual genome-wide screen for identifying sorting factors at the TGN, we found that the chimeric raft protein FusMidGFP displayed a trafficking phenotype in mutants with sphingolipid and ergosterol biosynthesis defects (Proszynski et al., 2004, 2005). Thus, we reasoned that the formation of the TGN-derived secretory vesicles could be generated based on a raft clustering mechanism. To purify the vesicles transporting FusMidGFP, we developed an immunoisolation procedure and implemented a shotgun lipidomics approach for quantitative characterization of the lipidome of secretory vesicles and the donor compartment.

**Results**

**Immunoisolation of TGN-derived secretory vesicles**

To isolate TGN-derived secretory vesicles in high purity, we devised an immunoisolation procedure using a transmembrane
raft protein as bait. We engineered the bait from the chimeric protein FusMidGFP, a type I transmembrane O-glycosylated raft protein which is directly delivered from the TGN to the PM in LDSVs (Fig. 1; Proszynski et al., 2004, 2005).

Several experimental factors were important to ensure successful immunoisolation of secretory vesicles. First, we had to localize the bait protein specifically into the TGN-derived transport carriers. Second, the bait protein had to be engineered with a high affinity epitope for immunoisolation. Third, we developed a specific immunoadsorbent with high binding capacity and low unspecific binding. Fourth, we introduced a protease cleavage site into the bait that enabled tobacco etch virus (TEV) protease–specific release of the secretory vesicles from the immunoadsorbent (Aebersold and Mann, 2003).

At the C terminus of the bait protein FusMidGFP, we introduced a high affinity 9× myc (M9) tag. A TEV protease site (T) was inserted between GFP and M9 and flanked by linker regions (L), generating the immunoisolation raft carrier bait FusMidGFPLTLM9, hereafter abbreviated as FusMidp.

To isolate post-Golgi secretory vesicles from living cells, we expressed the bait FusMidp in the temperature-sensitive exocyst mutant sec6-4 (Novick et al., 1980). When expressed at the permissive temperature 24°C, FusMidp reached the cell surface, as observed by fluorescence microscopy (Fig. 1 A). At the restrictive temperature 37°C, FusMidp accumulated intracellularly and did not translocate to the PM, as demonstrated by fluorescence microscopy and biochemistry (Fig. 1 A and Fig. S1 A). Electron microscopy demonstrated that the block of secretion at 37°C led to intracellular accumulation of vesicles (Fig. 2, A and B). By tomography, we observed that the vesicles were spherical with a diameter of ~100 nm (Fig. 2 C and Video 1).

For the specific isolation of TGN-derived FusMidp-vesicles, we combined conventional subcellular fractionation with a novel immunoisolation procedure. Based on previous work, we designed an immunoadsorbent composed of cellulose and sheep anti–mouse antibody (Hales and Woodhead, 1980; Wandinger-Ness et al., 1990). This immunoadsorbent showed high binding specificity, quantitative vesicle pickup, and efficient organelle release after TEV protease cleavage.

Secretory vesicles were isolated by cell lysis, differential fractionation, and organelle prepurification by isopycnic sucrose gradient centrifugation followed by immunoisolation (Figs. S2 and S3). Mouse anti-myc antibody was added to the prepurified organelle fraction enriched in the bait FusMidp and incubated with the sheep anti–mouse immunoadsorbent. Vesicles were specifically eluted by addition of TEV protease, producing the expected truncation of the bait (Fig. 3 A).

Immunodepletion of endosomes reduces organelle cross-contamination of immunoisolated secretory vesicles

To specifically recover FusMidp–secretory vesicles, we implemented the immunodepletion of endosomes by applying a mouse antibody against the endosomal t-SNARE Pep12p (Fig. 3 A).

Figure 2. Accumulation of secretory vesicles in sec6-4 cells at the restrictive temperature 37°C. (A) Transmission electron micrograph of a budding sec6-4 cell cultured at the permissive temperature 24°C. (B) Transmission electron micrograph of a budding sec6-4 cell cultured for 45 min at the restrictive temperature 37°C. White arrowheads indicate accumulated secretory vesicles with a diameter of ~100 nm. (C) 3D reconstruction of a budding cell with accumulated secretory vesicles (red) in a tomogram recorded from a 200-nm thick section prepared from the same culture of cells as shown in B; the PM is reconstructed in blue, the vacuole is orange, and other organelles such as Golgi cisternae or endosomal structures are in yellow. V, vacuole. Bars, 1 μm.
Assessment of the purity of FusMidp-vesicles

To document the relative enrichment of different organelle markers throughout the isolation procedure, we performed Western blot analysis. First, we showed that the bait protein was fully glycosylated (Proszynski et al., 2004) in the immunoisolated vesicles, thus confirming that they were TGN derived (Fig. 3 A). We verified that binding and release of the FusMidp-vesicles were specific because an unspecific mouse antibody did not retain vesicles on the immunoadsorbent (Fig. S3 A, b). Moreover, release of a bait construct lacking the TEV protease site was not possible (Fig. S3 A, c). Importantly, in analyzing whether other potential raft proteins were included in the vesicles, we found that the major glycosyl-phosphatidylinositol (PI [GPI])–anchored protein Gas1p (Mayor and Riezman, 2004) was present in FusMidp-vesicles (Fig. 3 A).

The soluble secreted protein invertase (the RFP fusion protein invertase-RFP [InvRFP]), a marker of an alternative cell surface delivery pathway mediated by heavy density secretory vesicles (HDSVs; Harsay and Bretscher, 1995), was not detectable in the eluate, demonstrating that FusMidp-vesicles are distinct from InvRFP-carriers (Fig. 3 A). We confirmed by Western blotting of the extracellular medium that InvRFP was correctly processed at both temperatures and secreted by sec6-4 cells at 24°C (Fig. S3 B).

Assessing the contamination with ER, we monitored the protein Dpm1p, which was efficiently depleted by subcellular fractionation (in P20; Fig. S1, A and B). Residual amounts of Dpm1p were detected in the sucrose gradient (Fig. S2 A), but Dpm1p was essentially removed from the isolated vesicles (Fig. 3 A). The design of the immunoisolation procedure also depleted Pep12p-containing endosomes (Fig. 3 A). As expected, residual amounts of late Golgi vesicular t-SNARE Tlg1p were present in the isolated FusMidp-vesicles (Fig. 3 A; Holthuis et al., 1998).

In addition, we verified that the sample amounts used for assessment of the vesicle purification gave responses within the dynamic range of the Western blot analysis (unpublished data). We hereby conclude that we have established a robust immunoisolation procedure that, for the first time, allows specific isolation of a pure population of secretory vesicles.

Morphology and integrity of immunoisolated secretory vesicles

To assess whether the immunoisolation procedures preserved the integrity of the secretory vesicles, we first examined the morphology of the FusMidp-vesicles by electron microscopy. We found a homogenous population of spherical vesicles with a diameter of 100 nm (Fig. 4). This result agrees well with the size of the vesicles observed in living sec6-4 cells at 37°C (Fig. 2, B and C). To evaluate the integrity of the isolated vesicles, we performed a floatation experiment as previously described (Harsay and Bretscher, 1995). We bottom loaded the immunoisolated FusMidp-vesicles on a linear Nycodenz gradient and centrifuged to equilibrium. The vesicle marker FusMidp and the GPI-anchored Gas1p floated to the top of the gradient with densities as LDSVs (Fig. S4; Harsay and Bretscher, 1995). In addition, the soluble secretory LDSV marker Bgl2p cofractionated with FusMidp and Gas1p.

The addition of TEV protease released the bait-labeled vesicles specifically because Pep12p was not cleaved and therefore retained endosomes on the immunoadsorbent. This strategy was essential for the successful purification of the FusMidp-vesicles (Fig. 3 A) and their molecular characterization.
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population of 100–300-nm membrane structures and small vesicles of ~40–50 nm (unpublished data). Moreover, the FusMidp-vesicles and TGN/E were morphologically distinct from purified yeast PM (Fig. S1 C).

Because the vesicle isolation was performed at 4°C, which can induce phase separation and vesicle pinch-off in simple model membranes (Lipowsky, 1993), we sought to determine whether the immunoisolation of vesicles from cells could be compromised by an artifactual fragmentation of the donor compartment. To do this, we performed a set of control experiments. First, we examined whether there was a difference in the amount of vesicles produced during subcellular fractionation at 4 and 24°C (Fig. S5 A), and, second, we investigated whether subcellular fractionation at 4 and 24°C affected the abundance of secretory cargo within the TGN/E (Fig. S5 B). These data showed that the immunoisolated secretory vesicles were not contaminated with artifactual membrane entities created by purification at 4°C.

Collectively, these results demonstrated that the immunoisolation procedure recovered a specific population of secretory vesicles from living cells, carrying the raft proteins FusMidp and Gas1p, and that the secretory vesicles had a distinct morphology and composition in comparison with their donor compartment TGN/E and the yeast PM.

FusMidp-vesicles are enriched in ergosterol and sphingolipids

To determine whether the TGN exhibits the capacity to selectively sort lipids, we performed a comparative lipidomics study of FusMidp-vesicles and the TGN/E. To do this, we used a novel quantitative shotgun lipidomics platform for absolute quantification (i.e., picomole or mole percentage) of ergosterol, sphingolipid, glycerophospholipid, and glycerolipid species that yields accurate assessment of the stoichiometric relationship between the membrane lipid constituents (Ejsing et al., 2009). Lipid species were quantified by spiking samples with defined amounts of lipid species.
internal lipid standards for each monitored lipid class, which correct for biased lipid recoveries and differential ionization efficiencies in the subsequent steps of lipid extraction and quantitative mass spectrometric analysis (Ejsing et al., 2006a,b). Comparative lipidome analysis of FusMidp-vesicles, the TGN/E, and yeast cell lysates resulted in the absolute quantification of 83 molecular lipid species constituting 12 lipid classes.

The most abundant lipid species in the FusMidp-vesicles was ergosterol, comprising 22.8 mol% of the lipidome (corresponding to 5.7 pmol/µl), followed by 11.1 mol% M(IP)2C 18:0;3/26:0;1 (Fig. 5). The lipidome of the TGN/E contained 8.5 mol% PI 16:0-16:1 as the most abundant lipid species followed by 9.8 mol% ergosterol. In comparison, ergosterol and M(IP)2C 18:0;3/26:0;1 were 2.3- and 2.6-fold enriched, respectively, in the FusMidp-vesicles as compared with the TGN/E. Furthermore, we observed a similar enrichment of IPC 18:0;3/26:0;1 (2.2-fold) and MIPC 18:0;3/26:0;1 (2.4-fold) in the FusMidp-vesicles. Evaluating the lipid class composition (sum of all lipid species constituting the same lipid class), we observed that PI comprised the most abundant class in both FusMidp-vesicles and the TGN/E (Fig. 6). In addition to the enrichment of ergosterol and sphingolipids, the FusMidp-vesicles also showed a 3.2-fold enrichment of phosphatidic acid (PA). FusMidp-vesicles displayed a concomitant 3.7–1.6-fold reduction in phosphatidylserine (PS), phosphatidylethanolamine (PE), DAG, and phosphatidylcholine (PC) as compared with TGN/E. Comparing the lipid composition of FusMidp-vesicles and the TGN/E with that of total yeast cell extracts (Fig. 6), we concluded that the immunoisolation procedure enabled the specific recovery of subcellular organelles having selective differences in molecular lipid composition in addition to the differences in protein composition and morphology.

The membrane organization of FusMidp-vesicles is more ordered than the TGN/E. Because the lipidome data demonstrated that the FusMidp-vesicles had a composition compatible with raft lipid enrichment (ergosterol and sphingolipids; Simons and Vaz, 2004), we sought to determine whether there was a difference in the membrane structure of the FusMidp-vesicles as compared with the TGN/E. If the enrichment of ergosterol and sphingolipids in FusMidp-vesicles were to reflect a difference in bilayer order caused by raft coalescence, the prediction would be that the bilayer would be more packed than the donor membranes from where the vesicles form (Simons and Vaz, 2004). To measure membrane order, we used a spectrophotometric assay based on fluorescent C-Laurdan (Parasassi et al., 1991; Gaus et al., 2006; Kim et al., 2007). The fluorescence emission spectrum of C-Laurdan is sensitive to membrane order and may be used to calculate a parameter called the general polarization (GP), whose magnitude serves as a relative measure of membrane order. The data demonstrated that the membrane of the FusMidp-vesicles (GP = 0.24 ± 0.01) was more ordered than the TGN/E (GP = 0.09 ± 0.05; Fig. 7). This result indicated that the biogenesis of FusMidp-vesicles involves a modulation of membrane architecture during its formation at the TGN.

Figure 6. Lipid class composition of FusMidp-vesicles, the TGN/E, and total cell extract. The mole percentage of lipid class was calculated as the sum of the mole percentage of lipid species of the respective lipid class. (n = 4 independent experiments; mean estimate ± SD). Cer, ceramide; PG, phosphatidylglycerol.
lipidome analysis demonstrated that FusMidp-vesicles comprised 2.3-fold more ergosterol and sphingolipids as compared with the TGN/E (Fig. 5). These compositional differences between vesicle and donor compartment membranes unequivocally demonstrate that lipid sorting occurs in the TGN. Furthermore, the enrichment of these lipids corroborates the model that the TGN sorting machinery uses a raft-clustering mechanism during the formation of secretory vesicles. In addition to the enrichment of ergosterol and sphingolipids, the lipid analysis also documented differences in other lipid classes between the FusMidp-vesicles and the TGN/E. PA was also elevated in the FusMidp-vesicles (Figs. 5 and 6). This lipid has previously been implicated in post-Golgi secretion through phospholipase D action (Routt et al., 2005). Interestingly, PS, PE, and PC were depleted in FusMidp-vesicles.

The molecular constituents of secretory vesicles suggest a functional role for sphingolipids and sterols in TGN sorting

The results presented in this study corroborate our previous finding that ergosterol and sphingolipids are functionally important for cell surface transport of FusMidp (Proszynski et al., 2005). Several other studies have also implicated ergosterol and sphingolipids in the delivery of other raft-associated proteins, including Gas1p, from the Golgi to the PM in yeast (Bagnat et al., 2000; Eisenkolb et al., 2002; Umebayashi and Nakano, 2003; Gaigg et al., 2005, 2006). Interestingly, sphingolipid mutants identified in the screen produce sphingolipid species with shortened fatty acid moieties (elo3Δ) or species having a dihydrophosphogolosine (sur2Δ) instead of a phytosphingosine moiety (Haak et al., 1997; Oh et al., 1997; Ejsing et al., 2009). These molecular attributes have been implicated in the formation of liquid-ordered domains by augmenting sphingolipid–sterol interactions by hydrogen bonding (sur2Δ cannot hydroxylate the long chain base at C4; Kuikka et al., 2001) and by coupling the two membrane leaflets in raft domains by interdigitation of the very long chain fatty acid moiety from the luminal leaflet into...
the underlying cytoplasmic leaflet (elo3Δ synthesizes C22:0 and C24:0 fatty acids instead of C26:0; Veiga et al., 2001). The effects of these seemingly minor modifications of lipid structure on membrane trafficking only emphasize the importance of lipidomic analysis in studies of membrane trafficking.

We note that the lipid composition of transport vesicles that exit from the Golgi toward endosomes and lysosomes remains to be determined. However, from our finding that the TGN/E contains a lower ergosterol and sphingolipid concentration as compared with the FusMidp-vesicles, we predict that the route to the endosomes is less dependent on these lipids. This prediction is corroborated by the absence of ergosterol and sphingolipid mutants in genetic screens analyzing these pathways (Seaman et al., 1997; Bonangelino et al., 2002). Another important open issue is the lipid composition of the vesicles carrying invertase. Unfortunately, the transmembrane receptor for invertase is unknown, and, therefore, we could not address this aspect.

Mechanistic implications

The functional involvement of raft-based sorting in the assembly of transport carriers at the TGN could imply the formation of stabilized liquid-ordered domains from small rafts in the TGN membrane (Simons and Ikonen, 1997; Hancock, 2006). Raft coalescence would be promoted by an increase in sphingolipid and sterol concentration toward the trans side of the Golgi complex. Work from our laboratory has recently demonstrated that raft coalescence can be induced at a physiologically relevant temperature to form domains selectively enriched in raft proteins and lipids (Lingwood et al., 2008). Thus, induction of raft clustering in some similar way could lead to selective segregation of proteins and lipids in the TGN membrane. Interestingly, the incorporation of the GPI-anchored raft protein Gas1p into FusMidp-vesicles is consistent with a raft-clustering mechanism. So is the observation that FusMidp-vesicles exhibited a higher membrane order than the TGN/E, as measured by C–Laurdan spectrophotometry. This assay corroborates the notion that the vesicle bilayer undergoes a structural change, as predicted by a raft clustering mechanism (Simons and Vaz, 2004).

The lipid-facilitated sorting mechanism underlying the generation of the raft carriers at the TGN would lead to an enrichment of sphingolipids and ergosterol at the PM and enable the establishment of the lipid gradient characteristic of the secretory pathway (Hechtberger et al., 1994; Zinser and Daum, 1995; Schneider et al., 1999; Holthuis et al., 2001). This gradient could be enhanced by retrograde removal of glycerophospholipids by coat protein I (COPI)–mediated transport. Brügger et al. (2000) produced COPI vesicles in vitro from rat liver and COS cell Golgi membranes. Mass spectrometric analysis of COPI vesicles showed a reduction in both cholesterol and sphingomyelin content. If similar segregation of ergosterol and sphingolipids also occurred during yeast COPI vesicle biogenesis, the selective transport of sterols and sphingolipids anterogradely and the recycling of phosphatidylcholine retrogradely could be a basic element in the processes responsible for Golgi sorting functions.

Recently, the cytoplasmic protein complex termed exomer was shown to be important for TGN to PM transport in yeast (Wang et al., 2006). Additionally, peripheral membrane proteins such as BAR (BIN/amphiphysin/Rvs) domain–containing proteins could facilitate the budding process at the TGN (Itoh and De Camilli, 2006). But what are missing are the factors responsible for raft clustering in the TGN. We postulate that luminal or cytosolic proteins are required for induction of raft coalescence. Candidates could be Chs5p, Kes1p, or Rvs161p, which have been previously identified in the screen for sorting factors (Proszynski et al., 2005).

Lipid species–dependent phase segregation in biological membranes contributes to cargo sorting and membrane sculpting

Immiscibility of two liquid phases in membrane bilayers can promote membrane bending (Lipowsky, 1993). Recent work on shiga toxin has demonstrated that when this pentameric protein binds the glycosphingolipid Gb3 on the cell surface, it creates tubular membrane invaginations that internalize the toxin into cells (Romert et al., 2007). This process is energy independent and seems to be driven by a phase segregation process. Similarly, polyoma virus binds to the cell surface ganglioside GM1, and this multimeric binding leads to domain-induced budding into the cell (Lipowsky, 1993; Damm et al., 2005; Ewers et al., 2005). Thus, a new principle is emerging in membrane trafficking that uses lipid–lipid and lipid–protein interactions driving protein and lipid sorting during transport carrier formation. The novel strategy presented in this work will make it possible to isolate and characterize other sorting stations and carriers in the secretory pathway of living cells and thereby contribute to unraveling the molecular mechanisms of this complex distribution machinery.

Materials and methods

Yeast strain and culture media

The yeast strain KSY302 used in this study expressed the temperature-sensitive allele sec6-4 of the exocyst subunit SEC6 (provided by P. Novick, Yale University School of Medicine, New Haven, CT). The genotype of KSY302 is sec6-4; leu2-3, 112; ura3-52. KSY302 was cultured in synthetic complete media containing yeast nitrogen base (Difco), CSM-URA-LEU (BIO101), and 2% raffinose (Sigma-Aldrich). The yeast strain KSY302 used in this study expressed the temperature-sensitive allele sec6-4 of the exocyst subunit SEC6 (provided by P. Novick, Yale University School of Medicine, New Haven, CT). The genotype of KSY302 is sec6-4; leu2-3, 112; ura3-52. KSY302 was cultured in synthetic complete media containing yeast nitrogen base (Difco), CSM-URA-LEU (BIO101), and 2% raffinose (Sigma-Aldrich). Construct expression was induced by adding 2% galactose (Merck).

Plasmids

To produce the vesicle bait FUSMIDGFPITLM9 (plasmid p147), ITLM9 was amplified by PCR and integrated by homologous recombination 3’ of FUSMIDGFP into HindIII-linearized plasmid TPQ55 (based on pRS416; Proszynski et al., 2004) by recombination in yeast. The TGN/E bait, GAP1GFPITLM9 (plasmid p165), was engineered by recombining ITLM9 3’ of GAP1GFP into XhoI-linearized p163 (GAP1GFP, provided by C. Kaiser, Massachusetts Institute of Technology, Cambridge, MA). INVRFP (plasmid p145) was made by restriction digestion of a construct carrying InvRFP using SacI and HindIII and ligated into pRS415. The bait without a TEV site, FUSMIDGFPITLM9 (plasmid p137), was produced as p147 with a PCR fragment of UWF9 not containing the TEV site.

Antibodies

Mouse IgG ChromPure whole molecule (Jackson Immunoresearch Laboratories), mouse anti-human CD3 (clone OKT3; e Bioscience), sheep anti–mouse antibody (Wandinger-Ness et al., 1990), mouse anti-my antibody (c-Myc 9E10; Santa Cruz Biotechnology, Inc.), mouse anti-GFP (Santa Cruz Biotechnology, Inc.), rabbit anti-RFP (provided by M. Zerial, Max Planck Institute of Molecular Cell Biology and Genetics, Dresden, Germany), rabbit anti-Gas1p (provided by H. Pieszman, University of Geneva, Geneva, Switzerland), mouse anti-Pep12p (Invitrogen), mouse anti-Spm1p (Invitrogen), mouse anti-
rabbit anti-Tlg1p (provided by J. Holthuis, Utrecht University, Utrecht, Netherlands; Holthuis et al., 1998), and rabbit anti-Bgl2p (provided by R. Schekman, University of California, Berkeley, Berkeley, CA) were used.

**Microscopy**

Microscopy was performed with a microscope (BX61; Olympus) and a UP-PlanApo 100× NA 1.40 oil immersion objective (Olympus). Pictures were acquired at room temperature in synthetic complete medium with a camera (SPOT; Diagnostic Instruments, Inc.) using MetaMorph software (MDS Analytical Technologies).

**Subcellular fractionation and immunoinosolation**

KSY302 (sec6-4) transformed with the constructs FusMidGFPLTLM9 (p147) and InvRFP (p145) or Gap1GFPLTLM9 (p165) and p145 was cultured at 24°C in 250 ml SC-URA-LEU and 2% raffinose containing 2% galactose and incubated for 16 h (RCF_max 121,262 g, 30 krpm, at 4°C in an SW60 swing-out rotor). The gradient was centrifuged for 16 h (RCF_max 121,262 g, 30 krpm, at 4°C in an SW60 swing-out rotor). The gradient profile was monitored by Western blotting for Tpm1p and Gap1p and against GFP to detect the immunoinosolation baits and against the ER integral membrane protein Dpm1p and the late endosome marker Pep12p.

**Mass spectrometric lipid analysis**

FusMidvesicles, TGN/E, and total cell extracts were subjected to quantitative shotgun lipidomic analysis (Eising et al., 2009). In short, samples were mixed with 20 µl of internal lipid standard mixture, providing a total spike of 15 pmol DAG 17:0-17:0, 29 pmol PA 17:0-17:1, 59 pmol PE 17:0-17:1, 15 pmol phosphatidylglycerol 17:0-14:1, 49 pmol PS 17:0-14:1, 58 pmol PC 17:0-14:1, 62 pmol PI 17:0-17:0 (provided by C. Thiele, Max Planck Institute of Molecular Cell Biology and Genetics, Dresden, Germany), 15 pmol ceramide 18:0/18:0, 23 pmol IPC 18:0/26:0, 20 pmol MIPC 18:0/26:0, 30 pmol MIP[P]C 18:0/26:0, 30 pmol MIP[P]C 18:0/26:0, and 160 pmol stigmasterol,5,7,22-trienol. Samples were subsequently subjected to lipid extraction executed at 4°C (Eising et al., 2009). The lower organic phases were isolated and evaporated in a vacuum evaporator at 4°C. Lipid extracts were dissolved in 100 µl chloroform/methanol (1:2; vol/vol) and subjected to quantitative lipid analysis on both a hybrid i quadropole time of flight mass spectrometer (QSTAR Pulsar; MDS Analytical Technologies) and an LTQ Orbitrap instrument (Thermo Fisher Scientific) equipped with the robotic nanoflow ion source TriVersa NanoMate (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and phosphatidylglycerol species were quantified by negative ion mode multiple precursor ion scanning analysis (Eising et al., 2006a); PC and ceramide species were quantified by consecutive positive ion mode precursor ion scanning m/z 184.1 and multiple reaction monitoring analysis, respectively. IPC, MIPC, and MIP[P]C species were quantified by negative ion mode flow through mass analysis on an LTQ Orbitrap mass spectrometer; ergosterol was quantified after chemical acetylation by multiple reaction monitoring. Automated processing of acquired mass spectra and identification and quantification of detected molecular lipid species were performed by dedicated softwares (MDS Analytical Technologies; Eising et al., 2006a) and ALEX (Analysis of Lipid Experiments) software (Eising et al., 2009).

Lipid species were annotated by their molecular composition. Glycerophospholipid and DAG species are annotated as: lipid class:<number of carbon atoms in the first fatty acid moiety><number of carbon atoms in the second fatty acid moiety><number of double bonds in the second fatty acid moiety>—<number of carbon atoms in the fatty acid moiety><number of double bonds in the fatty acid moiety>—<number of carbon atoms in the first fatty acid moiety>—<number of carbon atoms in the second fatty acid moiety> (e.g., PI 16:0/18:1). Sphingolipid species are annotated as: lipid class:<number of carbon atoms in the long chain base moiety><number of double bonds in the long chain base moiety><number of hydroxyl groups in the long chain base moiety><number of carbon atoms in the fatty acid moiety>—<number of carbon atoms in the fatty acid moiety>—<number of double bonds in the fatty acid moiety>—<number of hydroxyl groups in the fatty acid moiety>—<number of carbon atoms in the fatty acid moiety> (for example, IPC 18:0/3/26:0;1). It is an IPC species containing a C18 phytosphingosine (having three fatty acid moieties) and a C26:0 amide-linked fatty acid moiety with a hydroxyl group.

**C.Laudran fluorescence spectrofluorimetry**

FusMidvesicles and TGN/E were visualized and quantified for total cholesterol content (C.Laudran, Max Planck Institute of Molecular Cell Biology and Genetics, Dresden, Germany), 15 pmol ceramide 18:0/18:0, 23 pmol IPC 18:0/26:0, 20 pmol MIPC 18:0/26:0, 30 pmol MIP[P]C 18:0/26:0, 30 pmol MIP[P]C 18:0/26:0, and 160 pmol stigmasterol,5,7,22-trienol. Samples were subsequently subjected to lipid extraction executed at 4°C (Eising et al., 2009). The lower organic phases were isolated and evaporated in a vacuum evaporator at 4°C. Lipid extracts were dissolved in 100 µl chloroform/methanol (1:2; vol/vol) and subjected to quantitative lipid analysis on both a hybrid i quadropole time of flight mass spectrometer (QSTAR Pulsar; MDS Analytical Technologies) and an LTQ Orbitrap instrument (Thermo Fisher Scientific) equipped with the robotic nanoflow ion source TriVersa NanoMate (Advion Biosciences, Inc.). DAG, PA, PS, PE, PI, and phosphatidylglycerol species were quantified by negative ion mode multiple precursor ion scanning analysis (Eising et al., 2006a); PC and ceramide species were quantified by consecutive positive ion mode precursor ion scanning m/z 184.1 and multiple reaction monitoring analysis, respectively. IPC, MIPC, and MIP[P]C species were quantified by negative ion mode flow through mass analysis on an LTQ Orbitrap mass spectrometer; ergosterol was quantified after chemical acetylation by multiple reaction monitoring. Automated processing of acquired mass spectra and identification and quantification of detected molecular lipid species were performed by dedicated softwares (MDS Analytical Technologies; Eising et al., 2006a) and ALEX (Analysis of Lipid Experiments) software (Eising et al., 2009). Lipid species were annotated by their molecular composition. Glycerophospholipid and DAG species are annotated as: lipid class:<number of carbon atoms in the first fatty acid moiety><number of carbon atoms in the second fatty acid moiety><number of double bonds in the second fatty acid moiety>—<number of carbon atoms in the first fatty acid moiety>—<number of carbon atoms in the second fatty acid moiety> (e.g., PI 16:0/18:1). Sphingolipid species are annotated as: lipid class:<number of carbon atoms in the long chain base moiety><number of double bonds in the long chain base moiety><number of hydroxyl groups in the long chain base moiety><number of carbon atoms in the fatty acid moiety>—<number of carbon atoms in the fatty acid moiety>—<number of double bonds in the fatty acid moiety>—<number of hydroxyl groups in the fatty acid moiety>—<number of carbon atoms in the fatty acid moiety> (for example, IPC 18:0/3/26:0;1). It is an IPC species containing a C18 phytosphingosine (having three fatty acid moieties) and a C26:0 amide-linked fatty acid moiety with a hydroxyl group.
were excited at 385 nm, and emission spectra were recorded from 400 to 530 nm at 23°C on a fluorescence spectrometer (Fluoromax-3; Horiba). Spectra of unstained samples were subtracted from the sample labeled with Laurdan. GP values were calculated according to Parassassi et al. (1991):

$$GP = \frac{I_{400-440} - I_{470-530}}{I_{400-440} + I_{470-530}}$$

where $I_{400-440}$ and $I_{470-530}$ stand for the total fluorescence intensity recorded from 400 to 460 nm and from 470 to 530 nm, respectively.

Electron microscopy

Yeast cells were cryoimmobilized using a high pressure freezer (EM-PACT2 + RTS; Leica; Manninen et al., 2005). Samples were freeze-substituted in an automatic freeze-substitution machine (Leica) at −90°C in 1% osmium tetroxide plus 0.1% uranyl acetate in anhydrous acetone for 24 h. The temperature was increased 4°C/h (27 h) to 20°C. Samples were washed in acetone and embedded at 24°C in epon/Araldite. 70- and 200-nm sections were cut using a microtome (Ultracut UCT; Leica). Sections were collected on Formvar-coated Cu grids and poststained in 5% uranyl acetate in 50% methanol for 3 min and aqueous lead citrate. Thin sections (70 nm) were viewed on a transmission electron microscope (Tecnai 12; FEI) at 100 kV. 200-nm sections for tomographic imaging were prepared identically; however, a solution of 15-nm gold was applied to both sides of the grids as fiducial markers to assist in the tomographic imaging procedure.

Intermediate voltage electron microscopy and tomography

Regions of interest were located on 200-nm sections on a Tecnai F30 (FEI). Identified targets underwent a preirradiation step with a large spread electron beam to minimize the polymer shrinkage during the tilt series acquisition. The collection of the tomographic data was controlled by the program SerialEM (Mastronarde, 2005). Series of 121 micrographs were acquired by rotating the specimen from ±60° around the y axis in 1° tilt increments at a magnification of 9,400x with an accelerate voltage of 300 kV produced by a field emission gun. Electron micrographs were collected on a cooled charge-coupled device camera (US1000; Gatan). 2,048 x 2,048-pixel images were aligned, and the 3D volume was reconstructed by weighted back projection with the IMOD program suite (Mastronarde, 1997). The obtained 3D electron density map is a volume of 1,848 x 1,848 x 206 voxels with anisotropic information. The vesicles and structures were identified in the x–y plane and segmented manually in every 10 planes with 3DMOD (Kremer et al., 1996). The final model was also rendered in 3DMOD.

Conventional chemical fixation of isolated organelles

Membrane pellets of isolated TGN/E vesicles and the TGN/E were fixed in 2.5% glutaraldehyde in PBS for 1 h, rinsed in PBS, and fixed in osmium tetroxide (1% aqueous solution for 1 h), dehydrated in an ascending ethanol series (30–100 with 10% concentration steps) two times for 5 min at each concentration, and embedded in epon/Araldite for microscopy. Sections were cut at 30–70 nm and mounted on Formvar-coated Cu grids, poststained, and examined as described in Electron microscopy.

Online supplemental material

Fig. S1 shows the molecular and morphological characterization of the yeast PM. Fig. S2 shows the organelle marker profile of the sucrose gradient on Formvar-coated Cu grids, poststained, and examined as described in Electron microscopy.

References


