The golgin protein Coy1 functions in intra-Golgi retrograde transport and interacts with the COG complex and Golgi SNAREs

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ABSTRACT Extended coiled-coil proteins of the golgin family play prominent roles in maintaining the structure and function of the Golgi complex. Here we further investigate the golgin protein Coy1 and document its function in retrograde transport between early Golgi compartments. Cells that lack Coy1 displayed a reduced half-life of the Och1 mannosyltransferase, an established cargo of intra-Golgi retrograde transport. Combining the coy1Δ mutation with deletions in other putative retrograde golgins (sgm1Δ and rud3Δ) caused strong glycosylation and growth defects and reduced membrane association of the conserved oligomeric Golgi (COG) complex. In contrast, overexpression of COY1 inhibited the growth of mutant strains deficient in fusion activity at the Golgi (sed5-1 and sly1-ts). To map Coy1 protein interactions, coimmunoprecipitation experiments revealed an association with the COG complex and with intra-Golgi SNARE proteins. These physical interactions are direct, as Coy1 was efficiently captured in vitro by Lobe A of the COG complex and the purified SNARE proteins Gos1, Sed5, and Sft1. Thus our genetic, in vivo, and biochemical data indicate a role for Coy1 in regulating COG complex-dependent fusion of retrograde-directed COPI vesicles.

INTRODUCTION Nascent secretory proteins that have been glycosylated and folded in the endoplasmic reticulum (ER) are delivered to the cis-Golgi in a coat protein complex II– (COPII) dependent manner (Brandizzi and Barlowe, 2013; Xu and Ng, 2015). As glycosylated secretory proteins traverse the Golgi, they are sequentially subjected to a series of glycan trimming, extension, and modification reactions. The order of these reactions arises from the compartmentalization of the Golgi, wherein each cisterna is enriched in a distinct set of resident enzymes, transporters, and chaperones (Stanley, 2011). Maintenance of compartmental identity requires that resident Golgi proteins are localized to the correct cisterna, yet many of these proteins are not statically retained to one cisterna (Harris and Waters, 1996; Todorov et al., 2000). Instead, resident proteins flow forward through the Golgi and are recycled back to earlier compartments through a vesicular trafficking pathway referred to as intra-Golgi retrograde transport (Banfield 2011; Fisher and Ungar, 2016).

Early Golgi resident proteins that have flowed to later cisternae are first packaged into coat protein complex I (COPI)-coated vesicles in an Arf1- and Vps74-dependent manner (Lanoix et al., 2001; Martínez-Menárguez et al., 2001; Tu et al., 2008). These vesicles are then directed to specific Golgi compartments by Rab GTPases (Mizuno-Yamasaki et al., 2012) and a subset of coiled-coil proteins known as golgins (Gillingham and Munro, 2016). Once tethered to the appropriate membrane, the vesicular Qc-SNARE Sft1 bundles together with the cisternal Qa-, Qb-, and R-SNAREs Sed5, Gos1, and Ykt6 to drive fusion between the opposing membranes (Bock et al., 2001; Parlati et al., 2002; Xu et al., 2002; Volchuk et al., 2004). Both the tethering and fusion steps depend on the conserved oligomeric Golgi (COG) complex, a member of the multi-subunit CATCHR-family of transport factors (Yu and Hughson, 2010; Climer et al., 2015). Although these components have been well characterized individually, it remains mechanistically unclear how, when, and where they are coordinated with one another to orchestrate vesicle capture and fusion.
Here we report a role for Coy1 in intra-Golgi retrograde transport. Coy1 has been implicated in this transport process based on a genetic interaction between COY1 and gos1Δ. Both COY1 and its human homologue CASP encode a ∼75-kDa Golgi-localized protein with a C-terminal transmembrane domain and a cytoplasmic N-terminus composed of a series of coiled coils (Gillingham et al., 2002). CASP has been proposed to function as a tether for retrograde-directed Golgi vesicles, as recombinant CASP could specifically capture vesicles laden with resident Golgi enzymes (Malsam et al., 2005). However, CASP was unable to nucleate vesicle tethering events in vivo when targeted to an ectopic site (Wong and Murno, 2014). Thus the functions of CASP and Coy1 remain enigmatic.

The identification of Coy1 as a COP II vesicle protein led us to further investigate its role in the early secretory pathway (Margulis et al., 2016). Here we report that Coy1 has no direct effect on ER to Golgi transport in a cell-free transport assay. Instead, our live cell labeling, genetic, and biochemical studies indicate a role for Coy1 in intra-Golgi retrograde transport. Rather than a function in tethering as has been established for most other golgins, our genetic and physical interactions implicate Coy1 in organizing the cisternal Golgi membrane for fusion with incoming retrograde directed vesicles.

RESULTS
Coy1 in ER-Golgi transport
We previously identified Coy1 as an enriched component of COP II vesicles in proteomic analyses (Margulis et al. 2016). To validate this result, a small-scale COP II budding reaction was conducted (Barlowe et al., 1994). Microsomes derived from wild-type (WT) cells were incubated with an energy regeneration system and purified COP II proteins to drive vesicle budding. Vesicles were separated from bulk membranes by centrifugation, and samples of the vesicular fraction were compared with the reaction input by immunoblotting (Figure 1A) (Belden and Barlowe 1996). The ER vesicle protein Env41 was enriched in the COP II vesicle fraction as previously reported (Otte et al. 2001), as was Coy1. A subunit of the translocon, Sec63, was not detectable in the vesicular fraction (Feldheim et al. 1992). The distributions of Env41 and Sec63 were unchanged in the coy1Δ strain. Thus Coy1 is efficiently packaged into COP II vesicles but has no apparent effect on COP II packaging efficiency or stringency.

We wondered whether Coy1 served as a vesicular tether that initiates contact between COP II vesicles and the cis-Golgi. To test this possibility, we monitored anterograde ER-Golgi transport in wild-type and coy1Δ strains using a cell-free assay (Cao et al., 1998). Overall transport (Figure 1B) and tethering (Figure 1C) efficiency in coy1Δ were indistinguishable from the wild-type reaction. As tethering factors at the Golgi are predicted to exhibit some functional redundancy with one another (Munro 2011; Roborti et al. 2015; Gillingham and Munro 2016), we hypothesized that a transport defect would be observed if Coy1 was deleted in addition to other golgins implicated in ER to Golgi transport. We repeated the transport assay using strains lacking Coy1 and either of the putative ER to Golgi tethering factors, Rud3 or Grh1 (VanRheenen et al., 1999; Behnia et al., 2007). No change in transport efficiency was observed in a coy1Δ rud3Δ double mutant (data not shown). Strikingly, a tethering defect apparent in grh1Δ membranes (Behnia et al., 2007) was suppressed by combining with coy1Δ (Figure 1D), suggesting Coy1 can indirectly affect anterograde transport despite being dispensable for this process. Consistent with this interpretation, overexpression of COY1 was
sufficient to induce transport and tethering defects (Supplemental Figure 1).

**Coy1 distribution and trafficking**

To understand how Coy1 could exert such an effect on anterograde transport, we sought to characterize the protein more thoroughly, beginning with investigating its localization and trafficking pattern. Coy1 colocalizes with a cis-Golgi marker via immunofluorescence microscopy (Gillingham et al., 2002), but its precise subcellular distribution is unknown. To address this question, membranes from wild-type cell lysates were centrifuged to generate ER (p13)- and Golgi (p100)-enriched membrane pellets. The membrane pellets were solubilized and separated on 10.5% SDS–PAGE gels. Immunoblotting was conducted with antisera against Coy1, as well as Erv41 and Erv46, both of which actively cycle between the Golgi and ER.

**FIGURE 2:** Coy1 is localized to Golgi membranes. (A) Cell lysate from a WT (CBY740) strain was centrifuged through a 18–60% sucrose density gradient. Fractions were collected from the top to the bottom of the gradient and resolved on 10.5% SDS–PAGE gels. Immunoblotting was conducted with polyclonal antibodies against the early Golgi mannosyltransferase Och1, the ER vesicle protein Erv25, the ER glucosidase Gls1, the vacuolar hydrolase CPY, and Coy1. (B) WT and sec12-4 strains were grown to early log phase at 25°C and treated with 50 μg/ml cycloheximide for 20 min to halt translation. The cultures were then shifted to 38.5°C for 90 min to induce an ER export block in sec12-4 cells. After conversion to spheroplasts, cell lysates were centrifuged to generate ER (p13)- and Golgi (p100)-enriched membrane pellets. The membrane pellets were solubilized and separated on 10.5% SDS–PAGE gels. Immunoblotting was conducted with antisera against Coy1, as well as Erv41 and Erv46, both of which actively cycle between the Golgi and ER.
Coy1 functions in retrograde transport

Coy1 functions in intra-Golgi transport

As Coy1 is dispensable for ER to Golgi anterograde transport in vitro and does not actively cycle back to the ER for a role in Golgi to ER retrograde transport, we wondered if Coy1 instead functioned in intra-Golgi retrograde transport. CASP has been implicated in this transport process, as microinjection of soluble, recombinant CASP inhibits intra-Golgi retrograde transport of the Golgi enzyme GalNAcT2-YFP (Malsam et al., 2005). We tested if Coy1 might contribute to this transport process by examining the retention of an established cargo of intra-Golgi retrograde transport, Och1, by pulse-chase analysis (Harris and Waters, 1996; Bruinsma et al., 2004). Wild-type and coy1Δ cells were pulsed with a mix of radiolabeled [35S]cysteine/[35S]methionine and then chased with an excess of unlabeled amino acids. Lysates from each time point were immunoprecipitated using antiserum against Och1 and a control protein, Kar2. Kar2 levels remained comparable throughout the chase period in both strains (Figure 4, bottom panel). By the end of the chase, Och1 levels were slightly reduced in the wild-type strain, whereas Och1 was rapidly depleted in the coy1Δ strain (Figure 4). The difference in the half-life of Och1 between the two strains was statistically significant (p < 0.05, unpaired Student’s t test). We interpret this to indicate an Och1 retention defect in coy1Δ, resulting in missorting and degradation of Och1.

As Och1 initiates outer-chain mannose extension, one might expect a defect in the retention of this enzyme to result in hypoglycosylation. However, no such glycosylation defects are apparent in coy1Δ cells, and total Och1 protein levels are similar between wild-type and coy1Δ cells (data not shown). We wondered if such a phenotype was being buffered against by other redundant transport components. Three mammalian golgins, GMAP-210, TMF, and golgin-84, are sufficient to initiate capture of vesicles laden with resident Golgi proteins in vivo (Wong and Munro 2014). The former two golgins have orthologues in yeast, called Rud3 and Sgm1, that have also been implicated in retrograde Golgi transport (Kim et al., 1999; VanRheenen et al., 1999; Siniossoglou and Pelham, 2001; Kim, 2003). If Coy1 functions in this process alongside Rud3 and Sgm1, cells could exhibit growth and glycosylation defects as more of these golgins are lost. To test this possibility, serial dilution assays were conducted using strains lacking one, two, or all three of these golgins (Figure 5A). No apparent phenotype was observed in coy1Δ rud3Δ, consistent with prior reports (Gillingham et al., 2002), while a mild growth defect was apparent in coy1Δ sgm1Δ. A more severe phenotype was observed in rud3Δ sgm1Δ. The triple knockout grew similarly to rud3Δ sgm1Δ at lower temperatures but was completely unable to grow at the restrictive temperature (Figure 5A). Expression of Coy1 on a plasmid was sufficient to suppress this phenotype (Supplemental Figure 2C). These results indicate that COY1 is required for viability at high temperatures when RUD3 and SGM1 are both absent.

A similar pattern of growth was observed when the same strains were grown in the presence of calcoflour white, a cell wall stressor used to identify glycosylation mutants (Ram et al., 1994). The rud3Δ strain exhibited mild sensitivity to calcoflour white (Figure 5B), whereas the coy1Δ and sgm1Δ strains were not affected. The coy1Δ sgm1Δ mutant exhibited a growth defect on media containing calcoflour white, as did the rud3Δ sgm1Δ strain. The triple knockout
failed to grow at any temperatures beyond the first dilution. Viability of the triple knockout on media with calcofluor white was restored when COY1 was provided on a plasmid. Notably, a plasmid encoding Coy1ΔTM failed to rescue growth of this strain, indicating that the transmembrane domain of Coy1 is functionally important (Supplemental Figure 2B). These results suggested that glycosylation defects become exacerbated as more of these golgins are lost. To validate this interpretation, the glycoproteins Gas1 and CPY were examined in whole-cell lysates derived from each strain (Figure 5C). Gas1 exits the ER as a 105-kDa protein and is further glycosylated in the Golgi to 69 kDa, and is proteolyzed in the vacuole to its mature 61-kDa form (Hasilik and Tanner, 1978; Stevens et al., 1982).

The molecular weight of Gas1 was unchanged relative to the wild-type strain in each of the single knockouts but was reduced in the coy1Δ sgm1Δ and rud3Δ sgm1Δ double mutants. Notably, the

FIGURE 4: The half-life of the Golgi mannosyltransferase Och1 is reduced in coy1Δ cells. Early log phase cultures of wild-type (CBY740) or coy1Δ (CBY2660) cells were pulsed with a radiolabeled cysteine/methionine mix and then chased with an excess of unlabeled amino acids. Samples were taken at the indicated time points and were subsequently lysed and then proteins were immunoprecipitated with antibodies against Och1 or Kar2. Immunoprecipitates were resolved on 10.5% SDS-PAGE gels and visualized by autoradiography. The half-life of Och1 was quantified from three independent experiments. In the wild-type strain, the half-life of Och1 was 98.5 min with a SD of 10.1, while the half-life of Och1 in coy1Δ was 40.3 (+/– 27.9, n = 3).

FIGURE 5: COY1 genetically interacts with tethering factors involved in intra-Golgi retrograde transport. (A) Tenfold serial dilutions were set up on YPD plates at the indicated temperatures with the following strains: WT (CBY740), coy1Δ (CBY2660), rud3Δ (CBY2678), sgm1Δ (CBY3874), coy1Δ sgm1Δ (CBY4102), coy1Δ rud3Δ (CBY2692), rud3Δ sgm1Δ (CBY4101), and coy1Δ rud3Δ sgm1Δ (CBY4103). (B) Serial dilutions were set up as in A on YPD + 5 μg/ml Calcofluor White. (C) Lysates derived from midlog phase cultures of the indicated strains were separated on a 10% SDS–PAGE gel and immunoblotted for Gas1, CPY, and Yet3. A darker exposure of the CPY blot is also shown to highlight accumulation of the p2 precursor. (D) Semi-intact cells were centrifuged at 100,000 × g to resolve cytosolic proteins in the supernatant fraction (S) from membrane bound proteins in the pellet (P). Equivalent amounts of each sample were resolved on 11% SDS–PAGE gels and immunoblotted for Cog3, Och1, Sec19, and Yet3. (E) Quantification of the percentage of Cog3 and Sec19 present in the supernatant fraction as depicted in D (means ± SD, n = 10 independent experiments, *p < 0.05, **p < 0.005, one-way ANOVA followed by Tukey’s test). (F) Wild-type (CBY740), coy1Δ (CBY2659), cog3-2 (CBY3881), and cog3-2 coy1Δ (CBY4788) strains were diluted onto YPD plates at the indicated temperatures. (G) Cultures from F were grown to early-log phase at 30°C and then shifted to 38.5°C for 1 h. Cells were separated from the medium by centrifugation, and secreted proteins were concentrated by trichloroacetic acid (TCA) precipitation while cell lysates were prepared by bead-beating. Media and cellular proteins were resolved on 10.5% SDS–PAGE gels and immunoblotted for CPY and Gas1, with Kar2 and Yet3 serving as loading controls.
molecular weight of Gas1 was reduced further still in the triple knockout relative to the double mutants. CPY was also underglycosylated in lysates from the coy1Δ sgm1Δ, rud3Δ sgm1Δ, and coy1Δ rud3Δ sgm1Δ, with the latter two strains also exhibiting an accumulation of a precursor form of CPY, indicative of a transport defect. These results solidly a role for Coy1 in maintaining the fidelity of glycosylation, along with Rud3 and Sgm1.

We next sought to identify a mechanistic cause for these additive phenotypes. In a yeast two-hybrid assay, every identified mammalian golgin protein interacts with the COG complex (Miller et al., 2013), an octameric peripheral membrane protein complex that is implicated in vesicle tethering and fusion throughout the Golgi (Fisher and Ungar, 2016). These interactions led us to test whether golgin proteins collectively function to recruit the COG complex to the Golgi membrane. We focused our analysis on coy1Δ and sgm1Δ because a strain carrying both mutations exhibited a glycosylation defect (Figure 5, B and C) and to avoid confounding effects caused by disrupted ER to Golgi trafficking that may arise from rud3Δ. Semi-intact cells derived from wild type, coy1Δ, sgm1Δ, and coy1Δ sgm1Δ cultures were diluted in buffer and centrifuged at 100,000 × g for 20 min to separate cytosol from cellular membranes. The integral membrane proteins Yet3 and Och1 were both solely detected in the membrane pellet fraction, while Sec19, a protein that cycles between membranes and the cytosol (Garrett et al., 1994), was similarly enriched in the cytosol in each strain (Figure 5, D and E). Cog3 appeared equally split between the membrane and cytosolic fractions in the wild-type setting. This distribution significantly shifted toward the cytosol in the coy1Δ and sgm1Δ strains, and this redistribution was more pronounced in the coy1Δ sgm1Δ strain (Figure 5, D and E). Thus the glycosylation defect in the coy1Δ sgm1Δ strain can be partially attributed to disrupted COG complex membrane association.

| A | WT + pRS426 | 22°C | WT + pRS426-COY1 | 30°C |
|---|-------------|------|------------------|------|
| sed5-1 + pRS426 | | | |
| sed5-1 + pRS426-COY1 | | | |

| B | WT + pRS426 | | WT + pRS426-COY1 | |
|---|-------------|------|------------------|------|
| sly1-ts + pRS426 | | | sly1-ts + pRS426-COY1 | |

FIGURE 6: COY1 genetically interacts with sed5-1 and sly1-ts. (A) The sed5-1 (CBY263) and isogenic wild-type strain (CBY267) were transformed with either an empty vector (pRS426) or a 2μ COY1 overexpression vector (pRS426-COY1). Two independent isolates per transformations were serially diluted on selective media and grown at the indicated temperatures. (B) The sly1-ts strain (CBY268) and isogenic wild-type strain (CBY267) were transformed with either an empty vector or the COY1 overexpression vector and subjected to growth assays as in A.

Attenuated membrane association of the COG complex in the coy1Δ strain suggested an interaction between Coy1 and the COG complex. To address the physiological relevance of this potential interaction, a haploid strain lacking COY1 and bearing the cog3-2 point mutation was constructed by mating, and its growth was compared with that of the parent strains (Figure 5D). The cog3-2 strain grows poorly at lower temperatures and is inviable at an elevated temperature (Wuestehube et al., 1996). Strikingly, deletion of COY1 suppressed this growth defect, and the double mutant remained viable at the restrictive temperature, although its growth was reduced relative to the wild-type and coy1Δ strains. Expression of COY1 on a plasmid was sufficient to reintroduce a strong growth defect in the cog3-2 coy1Δ strain (Supplemental Figure 2D). However, the glycosylation and transport defects of cog3-2 were not corrected, as Gas1 and CPY ran at a reduced molecular weight in cell lysates from both the cog3-2 and cog3-2 coy1Δ, stains and an accumulation of p2 CPY was apparent in both strains as well. Furthermore, secretion of p2 CPY was elevated an average of 1.46 ± 0.17 SD, n = 3) and 1.37 ± (0.28-fold relative to the wild-type strain in the coy1Δ and cog3-2 single mutants and was elevated 2.3-fold (± 0.39) in the double mutant (Figure 5E). These results functionally implicate COG complex and Coy1 with one another but also indicate that the latter does not simply act as a negative regulator of the former.

The COG complex and Coy1 both affect SNARE function. The COG complex physically and functionally interacts with the early Golgi syntaxin Sed5 and its cognate SM protein Sly1 (Suvorova et al., 2002; Shestakova et al., 2007; Laufman et al., 2009), while overexpression of COY1 results in a severe growth defect in a strain lacking the retrograde Golgi SNARE Gos1 (Gillingham et al., 2002). We wondered whether these SNARE interactions could explain how coy1Δ suppresses the growth defect of cog3-2. We observed that overexpression of COY1 was toxic in strains bearing either the sed5-1 or sly1-ts mutations (Figure 6, A and B). Milder growth defects were also apparent when COY1 was overexpressed in the sec18-1 and bet1-1 strains, while no phenotype was observed when COY1 was overexpressed in strains bearing mutations in Golgi-associated tethering factors, including grh1Δ, imh1Δ, rud3Δ, sgm1Δ, trs33Δ, and ypt1-3 (Supplemental Table 1). These genetic interactions implicate Coy1 in a functional network with the COG complex, Sed5, and Sly1.

Coy1 interacts with intra-Golgi retrograde tethering and fusion machinery

We next examined whether Coy1 could be detected in complex with any of the components suggested by genetic interactions. To this end, Coy1 was immunoprecipitated from digitonin-solubilized spheroplasts derived from wild-type cells with anti-Coy1 polyclonal antibodies. Coy1 was enriched in the immunoprecipitate, and Cog3, Sly1, and Gos1 were all pulled down with Coy1 antibodies (Figure 7A). These interactions were specific, as none of the immunoprecipitating proteins were detected in control reactions in which the preimmune serum was used.
FIGURE 7: Coy1 interacts with Sly1, Cog3, Gos1, and Sed5. (A) Wild-type (CBY740) semi-intact membranes were solubilized with digitonin and immunopreципitated (IP) with magnetic beads that had been preincubated with either Coy1 antiserum (αCoy1 IP) or preimmune serum from the same animal (Preimmune). Immunoprecipitates were washed, eluted in sample buffer, and resolved on 10.5% SDS–PAGE gels. Input represents a sample of the solubilized membranes prior to incubation with the antibody-bound beads. Immunoblotting was performed with polyclonal antiserum directed against Coy1, Sly1, Cog3, and Gos1, with the vacuolar SNAREs Vam3 and Vam7 and the ER resident protein Yet3 serving as negative controls. Positions of the MW ladder are indicated to the right of the blots. (B) Immunoprecipitation was conducted as described in A with antibodies directed against Cog3 or with preimmune serum. Blotting was conducted for Cog3 and Coy1, with Yet3 and Gos1 as negative controls. (C) Immunoprecipitation was conducted as described in A, using αSed5 preimmune and immune serum. Sed5 could not be detected in the immunoprecipitate, further supporting specificity. (D) Immunoprecipitation was conducted as described above with αSly1 antiserum or preimmune serum. Blotting was conducted against Sly1 and Coy1, with Yet3 and the vacuolar SNARE Nyv1 serving as negative controls. (E) Immunoprecipitates were prepared and processed as described above, using antiserum against Gos1. As Gos1 preimmune serum was not available, magnetic beads alone served as a negative control (−antiserum). Blotting was conducted for Gos1 and Coy1, with the golgin Bug1 and the vacuolar SNARE Vam7 functioning as negative controls.

Moreover, the ER resident protein Yet3 and the vacuolar SNAREs Vam3 and Vam7 were not detected in the immunoprecipitate, further supporting specificity. In reciprocal experiments, Coy1 could be coinmunoprecipitated with antibodies against Cog3 (Figure 7B), Sed5 (Figure 7C), Sly1 (Figure 7D), and Gos1 (Figure 7E). Thus Coy1 appears to exist in complex with Cog3, Sed5, Sly1, and Gos1.

We addressed whether Coy1 could directly interact with proteins that were recovered in coinmunoprecipitates through a series of in vitro binding assays. First, the cytosolic domains of a panel of soluble Golgi SNAREs were expressed and purified as glutathione S-transferase (GST) fusion proteins (Supplemental Figure 3A; Tsui and Banfield, 2000). Then yeast cells overexpressing Coy1ΔTM-HA were lysed and centrifuged at 120,000 × g to generate a Coy1-enriched high-speed supernatant fraction. The fusion proteins were then immobilized on glutathione agarose resin and incubated with this supernatant fraction. The resin was washed, bound proteins were eluted with sample buffer, and Coy1ΔTM-HA was detected by immunoblotting with a monoclonal αHA antibody. Coy1 was not appreciably pulled down by GST alone or by GST fusions to the ER-to-Golgi Qc-SNARE Bet1 or the retrograde Golgi R-SNARE Ykt6. In contrast, nearly 15% of the total Coy1ΔTM-HA added to the reaction could be recovered by GST-Gos1, GST-Sed5, and GST-Sft1 (Figure 8A). These results indicate that Coy1 specifically interacts with intra-Golgi retrograde Q-SNAREs.

To test whether Coy1 physically interacts with the COG complex, Lobe A of the COG complex, which consists of Cog1, Cog2, Cog3, and Cog4, was expressed and purified from bacteria as previously described (Supplemental Figure 3B; Lees et al., 2010). Recombinant, purified COG was then immobilized on NiNTA resin and incubated with the Coy1ΔTM-HA enriched supernatant fraction. This fraction was also incubated with untreated NiNTA resin to measure background binding of Coy1ΔTM-HA. The resins were washed and eluted as described above. Less than 1% of the Coy1ΔTM-HA added to the reaction was recovered on the resin alone, whereas 20% of Coy1ΔTM-HA could be recovered with the COG complex bound to the resin (Figure 8B). Thus Coy1 physically interacts with the COG complex in vitro.

As the COG complex directly interacts with Sed5 (Suvorova et al., 2002), we sought to test whether the interaction between Coy1ΔTM-HA and Sed5 was attributable to any residual COG complex in the cytosolic input used as a source of Coy1ΔTM-HA. We observed that Coy1 was readily resolved from the COG complex by gel filtration chromatography (Figure 8C). Coy1 eluted prior to Cog3 and another control protein, Sec13, when detergent-solubilized semi-intact wild-type cells were injected onto a Superose 6 column. A similar elution pattern was observed when cytosol from the Coy1ΔTM-HA (CBY3484) strain was gel filtered, although the Coy1 peak was shifted one fraction later, perhaps attributable to reduced protein abundance of Coy1ΔTM-HA relative to full length Coy1.

Yet3 and Gos1 as negative controls. (C) Immunoprecipitation was conducted as described in A, using αSed5 preimmune and immune serum. Sed5 could not be detected in the immunoprecipitate, and its partner SM protein Sly1 served as a positive control instead. (D) Immunoprecipitation was conducted as described above with αSly1 antiserum or preimmune serum. Blotting was conducted against Sly1 and Coy1, with Yet3 and the vacuolar SNARE Nyv1 serving as negative controls. (E) Immunoprecipitates were prepared and processed as described above, using antiserum against Gos1. As Gos1 preimmune serum was not available, magnetic beads alone served as a negative control (−antiserum). Blotting was conducted for Gos1 and Coy1, with the golgin Bug1 and the vacuolar SNARE Vam7 functioning as negative controls.
Proper functioning of the Golgi complex requires compartmentalization of this organelle, yet many resident Golgi proteins are not restricted to one cisterna. Instead, these resident proteins dynamically cycle through the Golgi complex and are retrieved to earlier compartments by dedicated vesicular transport pathways. How these vesicles are targeted to and fused with specific cisernae remains an active area of investigation. In this report, we establish a role for the golgin protein Coy1 in intra-Golgi retrograde transport based on genetic and biochemical evidence. We find that the half-life of Och1, a resident Golgi protein known to undergo COPII- and COG-dependent retrograde transport (Harris and Waters, 1996; Spang and Schekman, 1998; Brunsma et al., 2004), is reduced in a coy1Δ strain (Figure 4), indicating a defect in the retrieval of this mannosyltransferase. We expect that this role in intra-Golgi retrograde transport is conserved, as microinjection of CASP, the human homologue of Coy1, inhibits retrograde transport of GalNac-T2 without affecting retrograde transport of ERGIC53 in BSC-1 cells (Malsam et al., 2005). Glycosylation defects were not apparent in coy1Δ or in cells lacking other golgins (Sgm1 and Rud3) putatively involved in tethering retrograde-directed COPI vesicles (Kim et al., 1999; Kim, 2003; VanRheenen et al., 1999; Fridmann-Sirkis et al., 2004; Siniosoglou, 2005; Wong and Munro, 2014). However, cells lacking Sgm1 and either COY1 or RUD3 exhibited glycosylation and growth defects, and these phenotypes were further exacerbated in the triple knockout (Figure 5, A–C). These glycosylation defects are mechanistically attributable in part to disrupted membrane association of the COG complex, as the COG complex redistributes from the membrane to the cytosolic fraction in coy1Δ and sgm1Δ cells, and this shift is intensified in coy1Δ sgm1Δ double mutant cells (Figure 5, D and E). These results establish Coy1, Rud3, and Sgm1 as collectively important for the maintenance of glycosylation fidelity and reveal a shared role for the golgins in recruiting the COG complex to Golgi membranes.

These findings could be interpreted to indicate functional redundancy among Coy1, Rud3, and Sgm1, but several features set Coy1 apart from these other golgins. First, golgins are considered to share the same general topology: an N-terminus that extends into the cytoplasm and a C-terminus that is anchored to the Golgi membrane association of the COG complex, as the COG complex redistributes from the membrane to the cytosolic fraction in coy1Δ and sgm1Δ cells, and this shift is intensified in coy1Δ sgm1Δ double mutant cells (Figure 5, D and E). These results establish Coy1, Rud3, and Sgm1 as collectively important for the maintenance of glycosylation fidelity and reveal a shared role for the golgins in recruiting the COG complex to Golgi membranes.

Coy1ΔTM-HA was also resolved from Coy3 when cystosol from pGAL-Coy1ΔTM-HA was centrifuged on a Superose 6 column (GE Healthcare). Fractions were resolved on 10.5% SDS–PAGE gels and immunoblotted for Coy1, Coy3, and Sec13. Middle, Cytosol from Coy1ΔTM-HA (CYB3484) semi-intact cells diluted in binding buffer was injected onto a Superose 6 column and fractions were immunoblotted as above. Bottom, Cytosol from pGAL-Coy1ΔTM-HA was centrifuged and fractionated as described above. D) Cytosol from pGAL-Coy1ΔTM-HA cells was resolved on a Superose 6 column. Fractions 8 and 9 were pooled and then aliquoted into tubes containing immobirolized GST-Bet1 or GST-Sed5. Binding assays and immunoblotting were conducted as described in A.

**DISCUSSION**

Proper functioning of the Golgi complex requires compartmentalization of this organelle, yet many resident Golgi proteins are not restricted to one cisterna. Instead, these resident proteins dynamically cycle through the Golgi complex and are retrieved to earlier compartments by dedicated vesicular transport pathways. How these vesicles are targeted to and fused with specific cisternae remains an active area of investigation. In this report, we establish a role for the golgin protein Coy1 in intra-Golgi retrograde transport based on genetic and biochemical evidence. We find that the half-life of Och1, a resident Golgi protein known to undergo COPII- and COG-dependent retrograde transport (Harris and Waters, 1996; Spang and Schekman, 1998; Brunsma et al., 2004), is reduced in a coy1Δ strain (Figure 4), indicating a defect in the retrieval of this mannosyltransferase. We expect that this role in intra-Golgi retrograde transport is conserved, as microinjection of CASP, the human homologue of Coy1, inhibits retrograde transport of GalNac-T2 without affecting retrograde transport of ERGIC53 in BSC-1 cells (Malsam et al., 2005). Glycosylation defects were not apparent in coy1Δ or in cells lacking other golgins (Sgm1 and Rud3) putatively involved in tethering retrograde-directed COPI vesicles (Kim et al., 1999; Kim, 2003; VanRheenen et al., 1999; Fridmann-Sirkis et al., 2004; Siniosoglou, 2005; Wong and Munro, 2014). However, cells lacking Sgm1 and either COY1 or RUD3 exhibited glycosylation and growth defects, and these phenotypes were further exacerbated in the triple knockout (Figure 5, A–C). These glycosylation defects are mechanistically attributable in part to disrupted membrane association of the COG complex, as the COG complex redistributes from the membrane to the cytosolic fraction in coy1Δ and sgm1Δ cells, and this shift is intensified in coy1Δ sgm1Δ double mutant cells (Figure 5, D and E). These results establish Coy1, Rud3, and Sgm1 as collectively important for the maintenance of glycosylation fidelity and reveal a shared role for the golgins in recruiting the COG complex to Golgi membranes.

These findings could be interpreted to indicate functional redundancy among Coy1, Rud3, and Sgm1, but several features set Coy1 apart from these other golgins. First, golgins are considered to share the same general topology: an N-terminus that extends into the cytoplasm and a C-terminus that is anchored to the Golgi membrane association of the COG complex, as the COG complex redistributes from the membrane to the cytosolic fraction in coy1Δ and sgm1Δ cells, and this shift is intensified in coy1Δ sgm1Δ double mutant cells (Figure 5, D and E). These results establish Coy1, Rud3, and Sgm1 as collectively important for the maintenance of glycosylation fidelity and reveal a shared role for the golgins in recruiting the COG complex to Golgi membranes.

Coy1ΔTM-HA was also resolved from Coy3 when cystosol from pGAL-Coy1ΔTM-HA lysate was centrifuged on the Superose 6 column. Coy1ΔTM-HA was efficiently captured by GST-Sed5 when the gel-filtered fractions depleted of the COG complex were used as the input for this binding reaction (Figure 8D). We conclude that the interaction between Coy1 and Sed5 is not mediated by the COG complex.

**FIGURE 8**: Coy1 directly interacts with multiple Golgi SNAREs and with lobe A of the COG complex. (A) Lysate from pGAL-Coy1ΔTM-HA (CYB3588) was diluted in binding buffer and centrifuged at 120,000 × g to generate a supernatant enriched in Coy1ΔTM-HA. This supernatant fraction was incubated with GST fusion proteins purified from E. coli and immobilized on glutathione-agarose resin. This resin was then washed, eluted with sample buffer, and resolved by SDS–PAGE. Immunoblotting was conducted using a monoclonal αHA antibody to monitor Coy1 recovery and with antiserum raised against GST-Ykt6, while ∼15% of Coy1ΔTM-HA is recovered by GST-Gos1, GST-Sed5, and GST-Sft1. (B) Supernatant fractions enriched in Coy1ΔTM-HA were prepared as in A and incubated with untreated NNTA resin or purified COG1-4 (purified from E. coli and immobilized on NINTA resin). Samples were incubated, washed, and eluted as in B. Immunoblotting was conducted with monoclonal antibodies against the HA epitope and with polyclonal antiserum against Cog3 and Cog2. Trace amounts of Coy1ΔTM-HA are brought down with the untreated resin (≤1%), while 20% of the input is recovered with the NINTA-bound COG complex. (C) Top, Triton X-100 solubilized semi-intact cells from WT (CYB740) cultures were injected onto a Superose 6 column (GE Healthcare). Fractions were resolved on 10.5% SDS–PAGE gels and immunoblotted for Coy1, Coy3, and Sec13. Middle, Cytosol from Coy1ΔTM-HA (CYB3484) semi-intact cells diluted in binding buffer was injected onto a Superose 6 column and fractions were immunoblotted as above. Bottom, Cytosol from pGAL-Coy1ΔTM-HA cells was centrifuged and fractionated as described above. D) Cytosol from pGAL-Coy1ΔTM-HA cells was resolved on a Superose 6 column. Fractions 8 and 9 were pooled and then aliquoted into tubes containing immobilized GST-Bet1 or GST-Sed5. Binding assays and immunoblotting were conducted as described in A.
Rud3 and Sgm1, because most golgins are considered to primarily function as vesicle tethers (Kim et al., 1999; VanRheenen et al., 1999; Siniossoglou, 2005).

CASP was initially thought to function as a tether for retrograde directed Golgi vesicles, but this conclusion has come into dispute. CASP and golgin-84 were proposed to interact with one another on opposing membranes as a means of tethering retrograde directed vesicles to the Golgi, as these proteins coimmunoprecipitated with one another and golgin-84, but not CASP, was readily incorporated into COPI vesicles in an in vitro budding assay. Further support for this hypothesis was provided in vitro when CASP was demonstrated to capture vesicles laden with resident Golgi proteins, including golgin-84 (Malsam et al., 2005). This result stands in contrast to a more recent report that demonstrated that mitochondrially anchored CASP is unable to redistribute any resident Golgi proteins to this ectopic site, including its proposed tethering partner, golgin-84 (Wong and Munro, 2014). How can these discrepancies be resolved? It is notable that CASP could only capture vesicles that had shed their COPI coat (Malsam et al., 2005). Although it is not clear when the COPI coat is removed during intra-Golgi retrograde transport, there is a precedent for coat proteins serving as substrates for tethers (Cai et al., 2007; Angers and Merz, 2009; Ren et al., 2009; Zink et al., 2009). Under scouring this possibility, the COG complex and the COPI coat copurify with one another, and COPI-coated vesicles accumulate in the cytoplasm of HeLa cells upon knockdown of COG3 (Suvorova et al., 2002; Zolov and Lupashin, 2005). Therefore, we propose that retrograde-directed COPI-coated vesicles are first captured by other golgins, either alone or in combination with the COG complex (Wong and Munro, 2014; Willet et al., 2013). Once tethered, removal of the coat could permit transfer of the vesicle to CASP. What might CASP bind that is only accessible after uncoating? The vesicular SNARE GS15 represents a promising candidate, as our in vitro binding assays establish a direct interaction between Coy1 and Sft1, the yeast homologue of GS15 (Figure 8B). As Coy1 also physically interacts with Sed5 and Gos1 (Figures 7, A, C, and E, and BB), an exciting implication of these findings is that Coy1 could serve to distally connect the v-SNARE to a cognate set of t-SNAREs. Our results offer a potential means of reconciling seemingly contradictory results.

These physical interactions also contextualize the genetic interactions exhibited by COY1. Overexpression of COY1 is toxic in gos1Δ (Gillingham et al., 2002), sed5-1, and sly1-ts cells (Figure 6), while the deletion of COY1 suppresses the growth defect and partially suppresses the temperature sensitivity of cog3-2 without rescuing this strain’s glycosylation and transport defects (Figure 5, F and G). As Coy1 has no direct role in anterograde transport (Figure 1B), we speculate that Coy1 binds Sed5 to reserve this syntaxin for use in intra-Golgi retrograde transport, perhaps by enhancing interactions between Sed5 and either Gos1 or the COG complex (Suvorova et al., 2002; Shestakova et al., 2007). By sequestering Sed5 for use in retrograde transport, Coy1 would also indirectly diminish the incorporation of Sed5 into the anterograde SNAREpin (Tsui et al., 2001; Parlati et al., 2002). Consistent with this assumption, a mild anterograde transport defect is observed in a strain overexpressing COY1 (Supplemental Figure 1), while microinjection of soluble CASP into BSC-1 cells also modestly slows anterograde transport of V5-G (Malsam et al., 2005). If retrograde transport is dysfunctional (as in gos1Δ or cog3-2) or if components required for both anterograde and retrograde transport pathways at the early Golgi have been compromised (sed5-1 or sly1-ts), then this proposed function of Coy1 would prove deleterious, as Sed5 would be restricted to a transport process that is either inefficient or less essential, respectively (Liu et al., 2015; Papanikou et al., 2015). Removing Coy1 from this context would dysregulate transport through the Golgi (consistent with elevated CPY secretion in cog3-2 coy1Δ) while freeing up Sed5 for use in anterograde trafficking. As secretary protein transport and biosynthesis are coordinated with one another (Lipschutz et al., 2003; Toikkanen et al., 2003; Cancino et al., 2014; Luini et al., 2014), we speculate that the loss of COY1 restores viability to cog3-2 cells by enhancing the delivery of nascent proteins to the Golgi, obviating the need for a robust recycling pathway.

The suppression of the growth defects and temperature sensitivity of cog3-2 by coy1Δ appears to parallel the rescue of the grh1Δ tethering defect by coy1Δ. Both the COG complex and Grh1 were initially proposed to function in anterograde transport because anterograde transport efficiency is diminished when the functionality of these proteins is absent (VanRheenen et al., 1998, 1999; Behnia et al., 2007). The detrimental effect on ER-to-Golgi transport for COG mutants has since been explained as an indirect consequence of disrupted retrograde Golgi transport (Ballew et al., 2005). Similarly, a direct role for Grh1 in tethering COPII vesicles has been contested based on the absence of an effect on vesicle accumulation between the TER and Golgi in a grh1Δ strain of Pichia pastoris (Levi et al., 2010). As knockdown of the mammalian homologues of GRH1 or COG3 results in fragmentation of the Golgi, disrupted anterograde transport in grh1Δ, cog3-2, and cog2-1 could arise as an indirect consequence of aberrant Golgi morphology rather than a loss of factors directly required for COPII vesicle capture (Zolov and Lupashin, 2005; Puttenveedu et al., 2006). As our data implicate Coy1 in regulating Golgi SNARE availability and function, we speculate that the loss of Coy1 would suppress Golgi fragmentation by increasing the promiscuity of vesicle consumption at the Golgi or enhancing the rate of homotypic cisternal fusion (Bhave et al., 2014).

Although much remains to be determined on how Coy1 affects the function of COG complex and SNAREs, our present work establishes a role for this golgin at the interface between vesicle tethering and fusion. As our understanding of how tethering proceeds to fusion remains limited at present, Coy1 represents an exciting case study on how these processes may be coordinated with one another. To this end, it will be essential to determine which regions of Coy1 are required to interact with the COG complex and SNARE proteins and if Coy1 can interact with COG subunits and one or more SNAREs simultaneously. Integrating genetic and phenotypic analyses with in vitro reconstitution experiments should further elucidate our understanding of Coy1 and its role in intra-Golgi retrograde transport.

MATERIALS AND METHODS

Yeast strains and media

Yeast strains used in this study are listed in Table 1. Unless otherwise noted, strains were cultured on YPD (1% bacto-yeast extract, 2% bacto-peptone, 2% glucose [Becton, Dickson, and Company, Franklin Lakes, NJ]) or YMD for auxotrophic selection (0.7% yeast nitrogen base without amino acids, 2% glucose, Complete Supplement Mixture [MP Biomedicals, Solon, OH]). Strain growth was monitored by absorbance at 600 nm (OD600) on a GeneSys 105 UV-Vis spectrophotometer (Thermo Fisher, Waltham, MA).

Strain and plasmid construction

The lithium acetate method was used for yeast transformation (Ito et al., 1983). The pRS426-COY1 plasmid was previously described (Gillingham et al., 2002). To generate Coy1-HA (CBY2674), CBY740
| Strain  | Genotype                                      | Reference                          |
|---------|-----------------------------------------------|------------------------------------|
| RSY248  | MATα his4-619                                 | Kaiser and Schekman (1990)         |
| RSY263  | MATα sec12-4 ura3-52                          | Kaiser and Schekman (1990)         |
| RSY944  | MATα ura3-52 lys2-801 bet1-1                  | Cao et al. (1998)                  |
| CBY263  | MATα trp1-1 ade2-1 ura3-1 leu2-3,112 can1-100 sed5-1 | Cao et al. (1998)                  |
| CBY267  | MATα trp1-1 ade2-1 ura3-1 leu2-3,112 can1-100 | Mizuta and Warner (1994)           |
| CBY268  | MATα trp1-1 ade2-1 ura3-1 leu2-3,112 can1-100 sly1-1s | Mizuta and Warner (1994)           |
| CBY474  | MATα trp1-1 ade2-1 ura3-1 leu2-3,112 can1-100 ypt1-3 | Cao et al. (1998)                  |
| CBY740 = BY4742 | MATα his3Δ1 leu2Δ0 lys2Δ0 ura3Δ0 | Brachmann et al. (1998)            |
| CBY742 = BY4741 | MATα his3Δ1 leu2Δ0 met15Δ0 ura3Δ0 | Brachmann et al. (1998)            |
| CBY1702 | CBY740 with sec18-1                           | Heidtman et al. (2003)             |
| CBY2009 | CBY740 with grh1Δ::kanMX6                     | Research Genetics                  |
| CBY2030 | CBY740 with imh1Δ::kanMX6                     | Research Genetics                  |
| CBY2289 | CBY740 with trs33Δ::kanMX6                    | Research Genetics                  |
| CBY2659 | CBY740 with coy1Δ::kanMX6                     | Research Genetics                  |
| CBY2660 | CBY740 with coy1Δ::kanMX6                     | Research Genetics                  |
| CBY2666 | CBY740 with cog6Δ::kanMX6                     | Research Genetics                  |
| CBY2674 | CBY740 with COY1::TM-3xHA-HIS3MX6             | This study                         |
| CBY2678 | CBY740 with rud3Δ::kanMX6                     | Research Genetics                  |
| CBY2692 | CBY2660 with rud3Δ::kanMX6                    | This study                         |
| CBY3178 | CBY2009 except grh1Δ::natMX4                  | Research Genetics                  |
| CBY3258 | CBY3178 with coy1Δ::kanMX6                    | This study                         |
| CBY3481 | CBY740 with pGAL-COY1::kanMX6                 | This study                         |
| CBY3484 | CBY3481 with COY1::3HA-HIS3MX6                | This study                         |
| CBY3588 | CBY3484 with kanMX6-pGAL::COY1ATM-3xHA        | This study                         |
| CBY3874 | CBY740 with sgm1Δ::kanMX6                     | Research Genetics                  |
| CBY3881 | CBY740 with cog3Δ::cog3-2natMX6               | This study                         |
| CBY4101 | CBY2678 with sgm1Δ::hphMX6                    | This study                         |
| CBY4102 | CBY2660 with sgm1Δ::hphMX6                    | This study                         |
| CBY4103 | CBY4101 with coy1Δ::natMX6                    | This study                         |
| CBY4788 | CBY3881 with coy1Δ::kanMX6                    | This study                         |

A strain expressing Coy1ATM-HA under the GAL1 promoter was constructed by transforming CBY3481 with the pFA6α-3HA-His3MX6 cassette to truncate COY1 after the region encoding amino acid residue 613 (Longtine et al., 1998). The cog3-2 mutant allele in the CBY740 background (CBY3881) was constructed using established methods (Tong and Boone, 2006; Wilson and Barlowe, 2010). First, the cog3-2 ORF plus 179 downstream base pairs was amplified using genomic DNA isolated from CBY559 (VanRheenen et al., 1999). The nourseothricin resistance gene was then amplified from pFA6α-kanMX6 using primers designed to fuse this cassette immediately downstream of the amplified cog3-2 ORF (Hentges et al., 2005). These constructs were then transformed into CBY740. Isolates were screened for nourseothricin resistance, temperature sensitivity, and truncation of the 98-kDa Cog3 protein via immunoblotting (Wuestehube et al., 1996; VanRheenen et al., 1998; Kim et al., 1999). The SGM1 gene was disrupted in the rud3Δ (CBY2678) and coy1Δ (CBY2660) backgrounds using the Hygromycin B resistance gene from the plasmid pFA6α-hphMX6.

TABLE 1: Yeast strains used in this study.
yielding rud3Δ sgm1Δ (CBY4101) and coy1Δ sgm1Δ (CBY4102). COY1 was then disrupted in the rud3Δ sgm1Δ (CBY4101) strain with the nourseothricin resistance gene using pFA6a-natMX6 plasmid (Hentges et al., 2005). The cog3-2 coy1Δ strain (CBY4788) was constructed by mating coy1Δ (CBY2659) with cog3-2 (CBY3881). Spores from this cross were plated on YPD + geneticin + nourseothricin. Antibiotic-resistant haploid cells were screened by immunoblotting with anti-Cog3 and anti-Coy1 antibodies to confirm the mutant alleles.

The prS426 Coy1ΔTM plasmid was constructed by amplifying the PHO5 promoter region through the first 1842 base pairs of the COY1 ORF from the prS426 COY1 plasmid (Gillingham et al., 2002) with a Xhol restriction site encoded on the forward primer (GTAATTATAACTCAGGAGTCGAGCTCCTCCCTT) and a pair of stop codons followed by a SacI restriction site on the reverse primer (GAAAGTGTGTAGCCTTATTCACCTGCTATTATTTTTTATTATTTG- TAAAATGACTTTTGCAAAAC). The amplified fragment and the prS426 plasmid were digested with Xhol and SacI-HF restriction enzymes and ligated together. Ligation reactions were transformed into DH5α, and the sequence of the plasmid was verified by Sanger sequencing. Expression of Coy1ΔTM from this plasmid was verified by immunoblotting lysates derived from a coy1Δ (CBY2660) strain transformed with this plasmid with the polyclonal Coy1 antibody.

Antibodies and immunoblotting

Polyclonal antibodies have been previously described for CPY (Rothblatt et al. 1989), Gas1 (Fankhauser and Conzelmann 1991), Sec63 (Feldheim et al., 1992), Kar2 (Brodsky and Schekman 1993), Sec13 (Salama et al., 1993), Env25 (Belden and Barlowe 1996), Sec19 (Barlowe 1997), Sly1 and Sec5 (Cao et al. 1998), Vam3 (Uengernann et al. 1998), Och1 and Env41, Env46 (Otto et al., 2001), Yam7 (Merz and Wickner 2004), Nvy1 (Thomgren et al. 2004), Coy2 and Coy3 (Ballew and Barlowe 2010), Gls1 (Inadome et al. 2005), Yet3 (Wilson and Barlowe 2010), and Gls1 (Shibuya et al. 2015). Monoclonal antibodies against the HA epitope (HA.11) were purchased from BioLegend (San Diego, CA). Primary antibodies were used at a 1:1000 dilution. Horseradish peroxidase–conjugated donkey anti-rabbit and sheep anti-mouse secondary antibodies were utilized for detection of the polyclonal and monoclonal antibodies, respectively, at a dilution of 1:10,000 (GE Healthcare, Chicago, IL). Supersignal West Pico Chemiluminescent substrate (Thermo Fisher, Waltham, MA) and a G:BOX Chemi XR5 (Syngene, Frederick, MD) were used to detect chemiluminescence. Immunoblot signal intensity was quantified by densitometric analysis using the Syngene GeneTools analysis software.

Polyclonal anti-Coy1 antiserum was raised against a glutathione S-transferase (GST) fusion protein, GST-Coy1, containing amino acid residues 1–613 of Coy1. The GST-Coy1 fusion was expressed as a 100-kDa protein in Escherichia coli BL21 cells and was contained in inclusion bodies that fractionated in the 15,000 rpm pellet of cell lysates. Inclusion body pellets were washed with Tris-buffered saline (TBS) containing 1.0% Triton X-100 and then washed and resuspended in TBS. The resuspended pellet was used as antigen in rabbits (Covance, Denver, PA).

To generate polyclonal antiserum against Bug1, full-length Bug1 was purified from E. coli by Intein Mediated Purification using the Affinity Chitin-binding Tag system from New England Biolabs (Ipswich, MA). Bug1 was cleaved from its affinity tag after purification and used as an antigen in rabbits (Covance, Denver, PA).

Membrane preparations and subcellular fractionation

Strains were converted to spheroplasts as previously described (Baker et al., 1988). Sucrose gradient centrifugation experiments were conducted as described by Powers and Barlowe with minor modifications (1998). Spheroplasts were resuspended in 10 mM HEPES, pH 7.4, 1 mM EDTA, 12.5% sucrose, and 1 mM phenylmethylsulfonyl fluoride (PMSF) and lysed by 10 strokes of a chilled Potter-Elvehjem homogenizer. The lysate was centrifuged for 4 min at 3500 rpm at 4°C in an SS-34 rotor, after which the supernatant was recovered and layered atop a t10-step sucrose gradient (from 54–18% on top of a 2-ml 60% sucrose cushion in 10 mM HEPES, pH 7.4, 1 mM MgCl2). Gradients were centrifuged in an SW-40 rotor at 36,000 rpm for 3 h at 4°C, and 785-μl fractions were collected from the top to the bottom of the gradient. Samples from each fraction were diluted with an equal volume of sample buffer, boiled at 75°C for 10 min, resolved on 10.5% SDS–PAGE gels, and immunoblotted using polyclonal antibodies against Coy1, Gls1 (as an ER and vacuolar marker), Env25 (Golgi and ER marker), Och1 (Golgi marker), and CPY (vacuolar marker).

ER- and Golgi-enriched membrane pellets were prepared following an ER export block as previously described with modifications (Heidtmann et al., 2003, Mukherjee and Barlowe, 2016). Wild-type (RSY248) and sec12-4 strains (RSY263) were grown over night at room temperature, back-diluted to an OD600 of 0.1, and grown to mid log phase at 25°C. Cycloheximide was added to 50 μg/ml to minimize protein translation (Todorow et al., 2000). Cell cultures were incubated for 20 min at 25°C and then transferred to a 38.5°C water bath for 90 min. Cells were harvested by centrifugation and converted to spheroplasts. The spheroplasts were resuspended in 2.5 ml JR lysis buffer (20 mM HEPES, pH 7.4, 0.1 M sorbitol, 50 mM potassium acetate (KOAc), 2 mM EDTA, 1 mM PMSF, and 1 mM dithiothreitol [DTT]), homogenized with six strokes of a Potter-Elvehjem homogenizer, and centrifuged at a SS-34 rotor at 4000 rpm at 4°C for 5 min to pellet nuclei and unbroken cells. The supernatant was recovered and centrifuged at 14,000 rpm for 10 min at 4°C (Eppendorf 5417R) to generate the ER-enriched p13 pellet. The supernatant from this spin was recovered and centrifuged at 60,000 rpm for 15 min at 4°C in a TLA 100.3 rotor, yielding a Golgi-enriched p100 pellet. The p13 and p100 pellets were resuspended in 100 μl JR lysis buffer, diluted 1:1 with 5x sample buffer, boiled, and separated on 10.5% SDS–PAGE gels. Immunoblotting was conducted with polyclonal antibodies against Coy1, Env41, and Env46.

For COG membrane association assays, yeast semi-intact cells from wild type (CBY740), coy1Δ (CBY2660), sgm1Δ (CBY3874), and coy1Δ sgm1Δ (CBY4102) cultures were diluted in 20 mM HEPES, pH 6.8, 250 mM sorbitol, 150 mM KOAc, 5 mM Mg(OAc)2 (B88) + 1 mM PMSF + 5 mM EDTA + 1 mM DTT to a final concentration of 6 A280 units/mL. Samples were centrifuged at 54,000 rpm for 20 min in a TLA 100.3 rotor. Samples of the supernatant fractions were diluted in sample buffer and the remaining supernatant was aspirated off. After resuspending the pellet in B88, an equivalent volume of the pellet was diluted into sample buffer, and the fractions were resolved on 10.5% SDS–PAGE gels. Immunoblotting was conducted with antisera against Cog3, Sec19, Och1, and Yet3. Statistical analyses were performed with GraphPad Prism 7 software.

In vitro vesicle budding, tethering, and transport assays

Analytical scale COPII budding assays were performed as previously described (Barlowe et al., 1994) using washed microsomes derived from wild-type cells (Wuestehube and Schekman, 1992). Tethering and fusion assays were performed as described by Cao et al. (1998). The average of duplicate measurements is shown in Figure 1, B–D, with the error bars representing the range.
Indirect immunofluorescence microscopy

The localization of Coy1-HA and Coy1TM-HA was monitored by immunofluorescence microscopy as described by Powers and Barlowe (1996). Cells were fixed with formaldehyde, converted to spheroplasts, and adhered to slides coated with polylysine. Cells were washed, incubated at room temperature in blocking buffer (1% BSA and 0.5% Triton X-100 in phosphate-buffered saline [PBS]), and then incubated with primary antibodies. The anti-HA monoclonal antibody (HA.11) and the anti-Och1 antibody were both used at a dilution of 1:200, while the Kar2 antibody was used at a dilution of 1:300. Secondary Alexa Fluor 488 conjugated anti-mouse antibodies and Alexa Fluor 594 conjugated anti-rabbit antibodies were used at a dilution of 1:1000 (Thermo Fisher, Waltham, MA). Images were obtained at room temperature using a DeltaVision Imaging System (GE Healthcare, Chicago, IL), comprising a customized inverted wide-field microscope (IX-71; Olympus, Central Valley, PA), an UPlanS Apochromat 100x/1.40 NA lens (Olympus) with DeltaVision immersion oil (n = 1.516; GE Healthcare), a camera (CoolSNAP HQ2; Photometrics, Tucson, AZ), and an Insight solid-state illumination unit. Images were deconvoluted in SoftWoRx (GE Healthcare) and prepared with ImageJ (National Institutes of Health, Bethesda, MD) and Photoshop (Adobe, San Jose, CA).

In vivo labeling

The pulse-chase experiments were performed as previously described (Belden and Barlowe 1996; Bue et al., 2006). Wild-type (CBY740) or coy1Δ (CBY2660) cells were cultured at 30°C in YMD with reduced sulfate to an OD of 0.5, harvested, then washed and resuspended in YMD without sulfate to an OD of 5. Cultures were incubated for 5 min in a 30°C water bath and then pulsed with [35S] Promix (GE Healthcare, Chicago, IL) at a concentration of 8.3 μCi/OD of 6.5. The chase was initiated by addition of excess unlabeled methionine and cysteine. After 0, 5, 10, 20, and 40 min, 1-ml samples of cells were taken and diluted into 4 ml ice-cold 20 mM NaN_3. Cells were then harvested by centrifugation, washed, and resuspended in 225 μl cold NaNO_3 and then diluted with 225 μl 2x lysis buffer (100 mM Tris, pH 7.4, 2% SDS, 10 mM EDTA, 2 mM PMSF). Ice-cold glass beads were then added to the resuspended cells. Lysis was performed by vortexing the samples for 2 min, incubating at 95°C for 2 min, allowing the samples to cool to room temperature, vortexing for 5 min, and boiling again at 95°C for 1 min. After clearing the lysed samples by centrifugation at 14,000 rpm (Eppendorf 5415 C) for 1 min, 100-μl samples of lysate were diluted with 1 ml IP-SDS buffer (15 mM Tris, pH 7.5, 150 mM NaCl, 1% Triton X-100) and incubated overnight with 40 μl of a 20% Protein A sepharose slurry (GE Healthcare) and either 0.5 μl anti-Och1 or 1 μl anti-Kar2 antiserum. The resin was then washed twice with 1 ml IP-SDS buffer and eluted by boiling in sample buffer. Immunoprecipitates were resolved on 10.5% polyacrylamide gels and visualized by autoradiography using a phosphor screen on a Typhoon 8600 Variable Mode Imager and, later, on film. Half-lives were quantified using the ImageQuant 5.2 software (GE Healthcare).

Immunoprecipitations

Immunoprecipitations were conducted as described by Bue and Barlowe (2009) with minor modifications. A 5% digitonin stock was prepared by adding 50 mg digitonin (EMD Millipore, Billerica, MA) to 1 ml sterile water, which was then boiled at 95°C for 10 min and allowed to cool at room temperature. Twenty microliters of Pierce Protein A magnetic beads (Thermo Fisher, Waltham, MA) was washed twice in B88-8 (20 mM HEPES, pH 8.0, 250 mM sorbitol, 150 mM KOAc, 5 mM Mg(OAc)_2 + 0.05% digitonin, resuspended to 1 ml, and incubated for 45 min with 1 μl of antibody. Beads were washed twice in B88-8 + 0.05% digitonin and set aside until the soluble extract had been prepared. Aliquots of wild-type spheroplasts (at 7.5 A_260 units per aliquot) were thawed, resuspended in 1 ml B88, and centrifuged once at 10,000 rpm for 2 min at 4°C to wash (Eppendorf 5417R). The supernatant was aspirated and the pellet was resuspended to 220 μl in B88-8 + 1 mM PMSE + 5 mM EDTA. The samples were incubated for 2 min in a 24°C water bath, after which 220 μl solubilization buffer was added (B88-8 + 1 mM PMSE + 5 mM EDTA + 2% digitonin). Samples were returned to the 24°C water bath, incubated for 15 min and centrifuged at 10,000 rpm for 2 min at room temperature (Eppendorf 5424). A sample of this supernatant was diluted in sample buffer as a means of monitoring the reaction input. The remaining supernatant was transferred to a fresh, chilled Eppendorf tube and diluted with 700 μl B88-8 + 1 mM PMSE + 5 mM EDTA + 0.5% digitonin. Washed, antibody-bound magnetic beads were added to the diluted samples and incubated for 2 h for the αCoy1 IP, or for 16 h for the reciprocal IPs. The reactions were then applied to a magnet, washed three times with B88-8 + 0.05% digitonin, and eluted by boiling in 30 μl sample buffer. Samples were separated on 10.5% polyacrylamide gels, and immunoblotting was conducted using polyclonal antibodies.

Protein purification, binding assays, and gel filtration chromatography

GST fusion proteins were purified as described with some modifications (Parlati et al., 2000; Tsui and Banfield, 2000; Peng and Gallwitz, 2002). Each plasmid (including pGEX-2T, pGST-BET1, pGST-GOS1, pGST-SED5, pGST-SFT1Δ11, pGST-YKT6, and pQLinkH COG1-4) was transformed into E. coli strain C43 DE3 for expression (Tsui and Banfield, 2000; Lees et al., 2010). For the GST fusion proteins, starter cultures were back diluted 1:100 into 1 l lysisogeny broth (LB) (1% tryptone, 0.5% yeast extract, 0.5% NaCl) with 50 μg/ml ampicillin. For GST, GST-Bet1, GST-Gos1, GST-Sft1, and GST-Ykt6, expression was induced with 0.5 mM IPTG at an OD_{600} of 0.6 for 4 h. Cells were harvested by centrifugation at 3500 rpm for 15 min at 4°C in a F108-6 x 500 rotor. Pellets were washed with 50 ml lysis buffer (25 mM HEPES, pH 7.5, 150 mM KCl, 0.05% Tween-20) and resuspended in 40 ml lysis buffer + 5 mM DTT + 1 mM EDTA + one Complete protease inhibitor tablet (Roche, Indianapolis, IN). The GST-Sed5 culture was grown to an OD_{600} of 0.6 and then induced with 0.5 mM isopropyl-β-D-thiogalactoside (IPTG) and shifted to 16°C for 18 h. GST-Sed5 was washed and lysed in a buffer composed of 25 mM HEPES, pH 7.5, 400 mM KCl, 10% glycerol, 1% Triton X-100, 5 mM DTT + 1 mM EDTA + one Complete protease inhibitor tablet. Cells were lysed by French press, and lysates were centrifuged at 14,000 rpm for 15 min at 4°C in a Sorval SS34 rotor. Supernatants were incubated for 2 h at 4°C with 0.1 g preswollen glutathione-agarose resin (Sigma-Aldrich, Waldrich, MA). Unbound material was discarded and the resin was washed with 50 ml lysis buffer. Bound proteins were eluted in 10 ml lysis buffer with 10 mM glutathione. Proteins were further purified by anion exchange (Mono Q HR 5/5; GE Healthcare, Chicago, IL) and gel filtration (Superose 12 HR 10/30; GE Healthcare). Aliquots were then prepared, each corresponding to 50 μg of purified protein.

To purify lobe A of COG (Cog1-4), the same methodologies described by Lees et al. (2010) and Ha et al. (2016) were employed, with minor modifications. Cultures of E. coli C43(DE3) bearing pQ-Link COG1-4 were grown overnight at ambient temperature in LB and then back diluted 1:100 into 1 l tertic broth (1.2% tryptone, 2.4% yeast extract, 0.4% glycerol, 17 mM KH_2PO_4, and 72 mM
K₂HPO₄) with 100 μg/ml ampicillin. This culture was grown at 37°C to an OD₆₀₀ of 0.4, at which point it was shifted to a 22°C incubator and grown to an OD of 0.7. Cultures were then induced with 0.5 mM IPTG and incubated for 18 h. Cells were harvested by centrifugation and then washed and lysed by French press in COG lysis buffer (20 mM Tris, pH 8.0, 150 mM NaCl, 20 mM imidazole, 10 mM β-mercaptoethanol). Lysates were centrifuged at 45,000 rpm for 30 min at 4°C in a 60Ti fixed-angle rotor. The clarified supernatant was incubated for 90 min with 3 ml NINTA agarose (Qiagen, Hilden, GE) that had been equilibrated in COG lysis buffer. The lysate and resin was then applied to an Econo-Column (BioRad, Hercules, CA) to separate unbound material from the resin. The resin was then washed with 100 ml COG lysis buffer, and bound material was eluted with 10 ml COG lysis buffer + 500 mM imidazole. The eluate was applied to an anion exchange column (MonoQ HR 5/5) that had been equilibrated in 25 mM NaCl, 20 mM Tris, pH 8.0, and 10 mM β-mercaptoethanol and fractionated on a linear gradient rising to 1 M NaCl. COG-containing fractions were pooled, concentrated on a 30-kDa NMWCO Amicon 15 centrifugal ultrafiltration device (EMD Millipore, Billerica, MA), and injected onto a gel filtration column (Superose 6 HR 10/30, GE Healthcare) equilibrated with 20 mM Tris, pH 8.0, 150 mM NaCl, and 10 mM β-mercaptoethanol. Aliquots corresponding to 15 μg of purified COG were prepared and stored at -70°C.

To generate a Coy1-enriched yeast lysate, the GAL1-regulated Coy1ΔTM-HA strain (CBY3588) was grown overnight in 50 ml of YES + 1% galactose + 1% glucose. 100 OD₆₀₀ units from this culture were recovered, washed once with YES, and added to 1 l YES + 2% galactose + 0.1% glucose. The culture was harvested at late log phase (OD₆₀₀ = 2), washed with BB8, and lysed with liquid nitrogen in a blender. Aliquots corresponding to 50 mg of the culture wet weight were then prepared and stored at -70°C.

For binding assays, aliquots of either the purified GST fusion proteins or of the COG complex were thawed, diluted with binding buffer (150 mM KOAC, 25 mM HEPES, pH 7.5, 250 mM sorbitol, 0.01% Triton X-100), and immobilized on glutathione agarose or NINTA resin. Meanwhile, aliquots of the Coy1ΔTM-HA enriched lysate were thawed, diluted in Coy1 binding buffer + 1 mM PMSF + 1 mM DTT (and 20 mM imidazole, for the COG binding assay), and centrifuged at 55,000 rpm (TLA 100.3). Five hundred microliters of Coy1 binding buffer, and immobilized on glutathione agarose or NINTA resin. Alternatively, Coy1 binding buffer rather than with binding buffer. All samples from the gel filtration and binding assays were analyzed on 10.5% SDS–PAGE gels.

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