Multiple Rab GTPase Binding Sites in GCC185 Suggest a Model for Vesicle Tethering at the Trans-Golgi

Garret L. Hayes,* Frank C. Brown,* Alexander K. Haas,† Ryan M. Nottingham,* Francis A. Barr,† and Suzanne R. Pfeffer*

*Department of Biochemistry, Stanford University School of Medicine, Stanford, CA 94305; and †Cancer Research Centre, University of Liverpool, Liverpool L9 3AT, United Kingdom

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GCC185, a trans-Golgi network-localized protein predicted to assume a long, coiled-coil structure, is required for Rab9-dependent recycling of mannose 6-phosphate receptors (MPRs) to the Golgi and for microtubule nucleation at the Golgi via CLASP proteins. GCC185 localizes to the Golgi by cooperative interaction with Rab6 and Arl1 GTPases at adjacent sites near its C terminus. We show here by yeast two-hybrid and direct biochemical tests that GCC185 contains at least four additional binding sites for as many as 14 different Rab GTPases across its entire length. A central coiled-coil domain contains a specific Rab9 binding site, and functional assays indicate that this domain is important for MPR recycling to the Golgi complex. N-Terminal coiled-coils are also required for GCC185 function as determined by plasmid rescue after GCC185 depletion by using small interfering RNA in cultured cells. Golgi-Rab binding sites may permit GCC185 to contribute to stacking and lateral interactions of Golgi cisternae as well as help it function as a vesicle tether.

INTRODUCTION

The Golgi complex plays a central role in processing and sorting of secreted and membrane proteins. In mammalian cells, it is made up of a set of flattened cisternae that are organized as a stack (Farquhar and Palade, 1998). Within the Golgi, proteins undergo oligosaccharide side chain maturation, and they are sorted to different destinations, including apical or basolateral plasma membranes, prelysosomes, or secretory storage granules, as they exit this compartment.

The Golgi surface is decorated with proteins called Golgins (Short et al., 2005; Sztul and Lupashin, 2006). Golgins are characterized by their predicted, highly extended, coiled-coil structures, and they typically have a high glutamic acid content (Ramirez et al., 2008). Certain Golgins have been implicated as transport vesicle tethers that help capture vesicles and bring them closer to their cognate targets; they are also essential for the maintenance of normal Golgi structure. For example, temperature-sensitive cells lacking GM130 have a disrupted Golgi at the nonpermissive temperature (Vasile et al., 2003); loss of Golgin-84 (Diao et al., 2003), Golgin-245 (Yoshino et al., 2005), TATA modulatory factor (Fridmann-Sirkis et al., 2004), GRASP55 (Feinstein and Linstedt, 2008), GRASP65 (Putheveedu et al., 2006), and the so-called COG complex (Zolov and Lupashin, 2005) all lead to disruption of the Golgi ribbon. Thus, the Golgi complex is highly sensitive to the loss of any one of a large collection of Golgi-associated proteins.

Most Golgins are peripherally associated with this organelle by virtue of interaction with small GTPases of the Rab and/or Arf (Arl) families; a few contain C-terminal membrane anchors (Short et al., 2005). Four human Golgins contain a so-called GRIP domain at their C termini that enables them to bind the Arl1 GTPase at the trans-Golgi network (TGN); Arl1 is essential for the Golgi targeting of multiple GRIP domain-containing proteins (Lu and Hong, 2003; Panic et al., 2003; Setty et al., 2003; Lu et al., 2006). We study GCC185, a GRIP domain-containing protein that uses cooperative, C-terminal interactions with Rab6 and Arl1 to achieve its TGN localization (Burguete et al., 2008). GCC185 came to our attention when we identified it as a Rab9 GTPase binding partner (Reddy et al., 2006). Using small interfering (siRNA) depletion in live cells and in vitro transport assays, we showed that like Rab9, GCC185 is required for the recycling of mannose 6-phosphate receptors (MPRs) from endosomes to the TGN. Cells lacking GCC185 accumulate MPRs in peripheral vesicles that contain Rab9 GTPase. In addition, the elegant and ordered Golgi ribbon fragments into perinuclear mini-stacks (Reddy et al., 2006; Derby et al., 2007).

The accumulation of transport vesicle intermediates in cells depleted of GCC185 is good evidence that this protein functions in the process of transport vesicle tethering (Reddy et al., 2006). We initially proposed that Rab9 present on transport vesicles (Barbero et al., 2002) could recruit cytosolic GCC185 to the vesicle surface and move along microtubule tracks toward the Golgi, in which Rab9 could pass the tether to Golgi-associated Rab6 and Arl1 proteins (Reddy et al., 2006; Burguete et al., 2008). But the Rab9 binding site we identified was located near the C terminus of GCC185, which also interacts with the TGN surface. This left the rest of the 1684-amino acid-long coiled-coil without a clear role. An obvious alternative possibility was that GCC185 contained additional Rab9 binding sites elsewhere along its length to permit vesicle binding at one site and
MATERIALS AND METHODS

Yeast Two-Hybrid Analysis

Yeast two-hybrid analysis was carried out as described previously (Fuchs et al., 2007). Briefly, a library of 57 mutant Rab proteins deficient for guanosine triphosphate (GTPase) hydrolysis was cloned into pCBP9 as a Gal4-binding domain fusion library (Bai and Elledge, 1996). Individual GCC185 trons were cloned into pACT2 as Gal4-activation domain hybrids. Cotransformations of these two constructs into the PJ29-4A yeast strain and plating onto synthetic complete agar plates lacking leucine and tryptophan yielded colonies expressing both fusion proteins. Five individual colonies were picked for each Rab/coiled-coil pairing and streaked onto selective plates (synthetic complete media containing 0.4 mg/ml adenine and lacking leucine, tryptophan, and histidine). A Rab/coiled-coil interaction was indicated by the presence of cell growth on this selective media.

Expression Plasmids, Protein Expression, and Purification

Different cDNAs encoding residues 1-50 and 51-1684 of GCC185 were obtained from Kazusa (Kisarazu, Japan) and American Type Culture Collection (Manassas, VA), respectively. Full-length GCC185 cDNA was constructed by three-piece ligation of polymerase chain reaction (PCR)-amplified products into pCDNA3.1 (+) (Invitrogen, Carlsbad, CA) modified with an N-terminal myc-tag. ΔC1 (358-1684), ΔC2 (752-1684), ΔC22S (890-1684), and ΔC10 (1-155) deletion constructs were assembled via PCR and ligated into pCDNA3.1 (+)-myc. For the full-length GCC185 siRNA rescue construct, GCC185 was cloned into a myc-tagged pCDNA3.1 (+) vector, and eight silent mutations were introduced via QuikChange mutagenesis into the region of GCC185 targeted by the GCC185 siRNA; results were verified by sequencing. This plasmid was used to generate GCC185 (A962-882). CC1 (34-358), CC2 (394-751), and CC3 (805-889) coiled-coil domains were amplified from the full-length sequence and cloned into both the yeast two-hybrid prey vector pACT2 (Clontech, Mountain View, CA) and the glutathione transferase (GST) fusion vector pGEX4T-1 (GE Healthcare, Little Chalfont, Buckinghamshire, United Kingdom); CC3 and CC3 L873A I880A were also inserted into pCDNA3.1 (+)-myc. The CC3 L873 I880A mutant in pGEX4T-1 was generated via QuikChange and confirmed by sequencing. The ΔC123 and C43B (1341-1684) domains were cloned separately into the pACT2 prey vector.

RESULTS

Binding Assays

Binding reactions were carried out similarly to that described previously (Aivazian et al., 2006). Purified Rab proteins were loaded with radioactive [35S]guanosine 5′-O-(3-thio)triphosphate (GTP-γ-S) (PerkinElmer Life and Analytical Sciences, Boston, MA) in loading buffer (50 mM HEPES pH 7.4, 150 mM NaCl, 5 mM MgCl2, 1 mM EDTA, 0.1 mg/ml bovine serum albumin (BSA), 7 μM GTP-γ-S, and 2.7 μM Rab) for 2 h (Rab9A and Rab9B) or 3 h (Rab6A, Rab1A, and Rab33B) at room temperature (Rab9A and Rab9B) or 37°C (Rab6A, Rab1A, and Rab33B). Reactions were then desalted on a PD-10 column (GE Healthcare) into binding buffer (50 mM HEPES pH 7.4, 150 mM KCl, 5 mM MgCl2, and 0.1 mg/ml BSA). Binding reactions were assembled in a 250-μl reaction containing 0.1 μM Rab and 3 μM (for single point binding) or varying (for concentration curve) amounts of GST-coiled-coil binding partners. Glutathione-Sepharose beads (15 μl) equilibrated in binding buffer were added to each reaction and incubated for 30 min at room temperature with rotation. Beads were then washed with 10 column volumes of binding buffer, and the amount of retained Rab protein was detected through liquid scintillation counting of the beads.

Cell Culture

HeLa cells from American Type Culture Collection were cultured at 37°C and 5% CO2 in Dulbecco’s modified Eagle’s medium supplemented with 7.5% fetal bovine serum, penicillin, and streptomycin. For RNA interference (RNAi), HeLa cells were transfected at 50% confluence with fluorescein labeled duplex RNA (Pharmacon RNA Technologies, Lafayette, CO) by using Oligo-Teste (Invitrogen) according to the manufacturer’s instructions. The GCC185 siRNA sequence was located in the first coiled-coil domain and targeted the sequence GGAGTTGGAACAATCACAT. RNAi depletion was carried out for 72 h posttransfection and followed by fixation and staining. For rescue experiments, plasmids containing GCC185 constructs were transfected into HeLa cells after initial siRNA transfection. Glutathione-Sepharose beads (15 μl) were equilibrated in binding buffer and added to each reaction and incubated for 30 min at room temperature with rotation. Beads were then washed with 10 column volumes of binding buffer, and the amount of retained Rab protein was detected through liquid scintillation counting of the beads.

Immunofluorescence Microscopy

Twenty-four hours before fixation, cells were split onto 22-×22-mm cover slips in a six-well plate. Cells were washed twice in phosphate-buffered saline (PBS) and fixed for 20 min in 3% formaldehyde in 200 mM HEPES, pH 7.4 (Warren et al., 1984). After fixation, cells were washed twice and incubated for 15 min in DMEM and 10 mM HEPES, pH 7.4, to quench. Cells were permeabilized for 5 min with 0.2% Triton X-100 in PBS, followed by two washes and a 15-min incubation with 1% BSA in PBS. Cells were incubated with primary antibody, diluted 1:500 (anti GM130), 1:750 (anti myc), or 1:1000 (anti-β-COP or anti-GCC185) in BSA/PBS for 30 min. This was followed by three 5-min washes and a 30-min incubation in secondary antibody diluted 1:1000 in BSA/PBS. The following antibodies were used in this study: rabbit anti-GC185 polyclonal antibodies (301-1684) (Reddy et al., 2006); mouse anti-GM130 (BD Biosciences, San Jose, CA), chicken anti-myc (Bethyl Laboratories, Montgomery, TX), Cy3-goat anti chicken (Jackson ImmunoResearch Laboratories, West Grove, PA), and Cy5-goat anti mouse/rabbit (Invitrogen). A rabbit anti-β-COP polyclonal antibody was generated according to Peupert et al. (1995). Micrographs were acquired using a microscope (Eclipse 80i; Nikon, Tokyo, Japan) fitted with a 60×/numerical aperture (NA) 1.40 plan apochromat objective lens and a charge-coupled device camera (CoolSnapHQ; Photometrics, Tucson, AZ) and controlled with MetaMorph Imaging software (Molecular Devices, Sunnyvale, CA). Pictures were analyzed using MetaMorph and Adobe Photoshop (Adobe Systems, Mountain View, CA) software. Confocal microscopy was performed with a Leica TCS SP2 SE confocal scanner in conjunction with a Leica DM6000 B upright scope (with attached Leica HCX PL, apochromatic 63×/NA 1.4 objective) and a Leica DMR 6000 confocal control box. This setup was controlled by Leica Control software (Leica Microsystems, Heidelberg, Germany). Images taken represent single slices of a confocal Z-stack at 63×.
A library of 57 human Rab proteins (all GTP hydrolysis deficient) representing ~88% of the full complement of human Rabs (~65 excluding nonprenylated Ran related members; Colicelli, 2004) was screened against each of the GCC185 constructs in a yeast two-hybrid assay. As summarized in Figure 1A, as many as 14 different Rab GTPases showed binding to different regions of GCC185 by this method (Supplemental Figure 1). The screen for binding to GCC185 fragment C343 is shown in Figure 1B. As expected, both Rab6A and Rab9A showed robust interaction with the C343 construct, in agreement with previous work that identified Rab6 and Rab9 binding sites within the C-terminal most 110 amino acids (Figure 1B). In addition, C343 interacted with four additional Golgi-associated Rabs: Rab6B, Rab9B (Yoshimura et al., 2007), Rab30, and Rab33B (Table 1). This, coupled with the discrete and distinct Rab interactions for each coiled-coil truncation construct, suggested that the observed interactions were likely significant.

It is of interest to note that yeast two-hybrid screens of full-length GCC185 yielded strong interaction with Rabs1A, 1B, and 35; weak binding was detected for Rabs9B and 6B (data not shown). Most of these Rabs interact with GCC185’s first coiled-coil (Figure 1A). We consider it likely that the GAL4 activation domain at GCC185’s N terminus cannot be brought close enough to the Gal4 DNA binding domain, when Rab partners interact at too great a distance from GCC185’s N terminus. Thus, screens of individual coiled-coils provided additional information regarding possible Rab interactions.

Several of the Rab interactions discovered by yeast two-hybrid (Figure 1) were verified independently using purified proteins and an in vitro binding assay. For this purpose, we purified untagged Rab9A, Rab9B, Rab1A, and His-tagged Rab6A and Rab33B, as well as GST fused to GCC185 CC1, CC2, CC3, and C110 (Figure 2A). Rab proteins were preloaded with [35S]GTPγS, and binding of Rabs to glutathione Sepharose-immobilized GCC185 constructs was tested. As shown in Figure 2B, almost all of the yeast two-hybrid interactions that we tested could be verified using purified components. For example, CC1 showed significant binding to Rab1A but poor binding to Rab9A; CC2 bound Rab6A but not Rab9A; in contrast, CC3 bound Rab9A in

![Figure 1. Yeast two-hybrid analysis of Rab GTPase binding to distinct coiled-coils in GCC185. (A) Top, secondary structure prediction for GCC185 using Paircoils software. Residue number is shown at bottom. Middle, schematic representation of predicted coiled-coils: CC1, CC2, CC3, and fragments representing the remainder of the protein (ΔCC123, C343, and C110). Bottom, summary of Rab binding to each of the domains as determined by yeast two-hybrid analysis. Rab9A is asterisked because it was variably detected by two hybrid; biochemical experiments shown below confirm its binding to CC3. (B) Yeast two-hybrid data for the construct C343. Each box shows five colonies streaked on selective media; numbers represent activated versions of the human Rab proteins.](image-url)

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<td>Allan et al., (2000); Moyer et al., (2001); Haas et al., (2007)</td>
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Table 1. GCC185 binding partners: localization and function
strong preference to Rab6A. Finally, the C-terminal 110 residue fragment bound Rab9A as we have reported previously, in addition to Rab9B, but not Rab33B. Because Rab33B was unable to bind C110, we infer that its binding site lies within the region upstream of C110, but downstream of the C343 N terminus. Unfortunately, it was not possible to obtain soluble C343 protein for in vitro binding studies.

The levels of binding observed in our in vitro binding assays (~1–2 pmol) were comparable with the levels of binding detected for proteins known to interact with C110 at 2–4 μM $K_D$ under similar conditions (Burguete et al., 2008). This is a typical affinity for Rab-effector binding. It should also be noted that at higher concentrations of GST-CC3 (>25 μM), some binding of Rab6A was detected, although it was substantially lower than that of Rab9A (data not shown).

Interaction of Rab9A and Rab6A with the C110 region of GCC185 requires a surface-exposed leucine/isoleucine pair that are found separated by one helix turn, on the same face of the C110 coiled-coil (Burguete et al., 2008). A helical wheel plot of GCC185 CC3 revealed a similar leucine/isoleucine motif near the C terminus of this peptide (Figure 3A). We therefore engineered and purified a GST-CC3 mutant protein containing leucine-to-alanine and isoleucine-to-alanine mutations to test the role of these residues in Rab9A binding. As shown in Figure 3B, the predicted, surface-exposed leucine 873 and isoleucine 880 residues were important for Rab9 interaction with CC3: $K_D$ estimates for the interaction were ~8 μM for the wild-type CC3 and ~29 μM for the mutant CC3. This suggests that Rab9A interacts with two distinct coiled-coils by a similar surface interaction.

Rab9A is localized to late endosomes (Lombardi et al., 1993) and to the transport vesicles that carry MPRs from endosomes to the Golgi complex (Barbero et al., 2002). Because GCC185 seems to be a tether for this transport process (Reddy et al., 2006), it made little sense that GCC185 would both attach to transport vesicles (via Rab9) and to its Golgi target (via Rab6 and Arl1) by using only its C terminus. The presence of a second Rab9A binding site near the middle of...
the protein, in GCC185 CC3, provided a solution to this conundrum and suggested that CC3 might be especially important for MPR transport.

To test this, we used an in vitro transport assay that measures the retrograde transport of cation-dependent (CD) MPRs from late endosomes to the TGN (Itin et al., 1997). Figure 4 shows that purified CC3 inhibited transport in vitro. The ability of CC3 to inhibit transport was concentration dependent (Figure 4B) and with an apparent $K_i$ value similar to the affinity of interaction of Rabs with this fragment. As expected, a control, mutated fragment of GCC185 that contains a coiled-coil but cannot bind Rab9A or Rab6A (C110 I/L $\rightarrow$ A/A) failed to inhibit in vitro transport (Figure 4A). We have shown previously that the wild-type C110 fragment inhibits transport >60% (Reddy et al., 2006). In contrast, the CC3 L/I $\rightarrow$ A/A mutant was still able to inhibit transport (although perhaps slightly less well than wild type), implicating other regions of the CC3 domain in this transport step. Alternatively, Rab9A may bind sufficiently well to the mutant protein (Figure 3B) to still result in inhibited transport under these conditions. Nevertheless, together, these data support the importance of GCC185’s CC3 in MPR recycling.

In an attempt to characterize further the functional importance of CC3, we overexpressed this protein fragment and tested whether it interfered with MPR trafficking as monitored by an increase in lysosomal enzyme secretion. Although the protein was observed to be entirely cytosolic upon expression in HeLa, human embryonic kidney 293, or COS-1 cells, we were not able to obtain high enough expression to detect a change in hexosaminidase secretion (data not shown).

Functional Analysis of GCC185 Coiled-Coil Domains 1 and 2

GCC185’s TGN localization requires Rab6A and Arl1 GTPases and specifically, Rab6A’s precise binding site (I1588, L1595) within the C110 coiled-coil (Burguete et al., 2008). Consistent with these previous observations, a set of GCC185 deletion constructs lacking the first, second, and third coiled-coil domains (ΔCC1, ΔCC12, and ΔCC123) localized normally to the Golgi (Figure 5). As expected, a GCC185 construct lacking the C-terminal 110 amino acids, ΔC110, failed to associate with the Golgi and was completely cytosolic (Figure 5). This confirmed that the first three coiled-coil domains are dispensable for GCC185 Golgi localization. In addition, at least one of the individual coiled-coil domains (CC2) was unable to inhibit transport (although perhaps slightly less well than wild type), implicating other regions of the CC3 domain in this transport step. Alternatively, Rab9A may bind sufficiently well to the mutant protein (Figure 3B) to still result in inhibited transport under these conditions. Nevertheless, together, these data support the importance of GCC185’s CC3 in MPR recycling.

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ability of ΔCC1, ΔCC12, and ΔCC123 to show normal localization indicates that these proteins are not grossly misfolded.

What role do the non–C-terminal, Rab binding domains play for GCC185? To explore whether these domains contribute to Golgi organization, we made use of the striking Golgi fragmentation phenotype observed when GCC185 is depleted from cells by using siRNA (Reddy et al., 2006). GCC185 was depleted from cells by 72-h treatment with fluorescein conjugated siRNA. We then attempted to rescue Golgi fragmentation with plasmids containing different regions of GCC185. These plasmids were constructed so that they would not be targets of the siRNA used (see Materials and Methods) and were expressed at very similar levels as determined by total pixel intensity.

In Figure 6, cells that contained fluorescent, GCC185 siRNA are indicated with an asterisk. As expected, asterisked cells that displayed only green (GM130) fluorescence (and had not received the complementing GCC185 plasmid) showed Golgi fragmentation compared with nonasterisked (nondepleted), green labeled cells (bottom row). Expression of Myc-tagged, full-length GCC185 (bottom row) restored normal Golgi morphology (Figure 6A, top row). In contrast, neither ΔCC1 nor ΔCC12 constructs were capable of reversing the fragmented Golgi phenotype upon siRNA treatment (Figure 6A, middle and bottom rows). Quantitation of these data confirmed the importance of at least CC1 (and possibly also CC2) in contributing to Golgi organization. Similar experiments were carried out using full-length GCC185 that lacked 20 residues in the middle of CC3 as the rescue plasmid. In these cells, Golgi structure seemed restored at the same frequency as seen with wild-type plasmid rescue (data not shown). Therefore, full-length GCC185 is required for proper maintenance of Golgi structure; CC3 residues that contribute to Rab9 binding seem to be dispensable for this aspect of GCC185 function.

Does GCC185 reach across the entire Golgi stack? If GCC185 exists as a fully extended coiled-coil, it would achieve a length of ~240 nm (1.4 Å/residue), a length that is almost sufficient to stretch across the entire Golgi stack in several cell types (200–300 nm; Rambourg et al., 1979; Clermont et al., 1995; Ladinsky et al., 1999) (Figure 7A). In addition, the complement of GCC185-interacting Rabs includes those localized to the trans (Rab6)-, medial (Rab6, Rab33B), and cis/medial (Rab1)-Golgi cisternae. Although these interactions are not required for localization of GCC185 to the TGN (Figure 5; Burguete et al., 2008), they could provide contacts for maintaining Golgi structure with contacts across the height of the stack (Figure 7C). It is intriguing to note that the binding site for the Rab protein that may be located the farthest away from the TGN (Rab1) is located the farthest away from GCC185’s TGN anchoring sites near its C terminus (Figure 1).

To explore this possibility further, we carried out confocal microscopy experiments in which we could distinguish the localizations of GCC185’s N and C termini, independently. Specifically, we compared the distributions of N-terminally myc-tagged, full-length GCC185, and N-terminally myc-tagged, C-terminal C-110 fragment, in relation to the cis-Golgi markers GM130 (Figure 8) and p115 (data not shown). In both cases, very low levels of overlap were observed in terms of the staining of anti-myc antibodies with the cis-Golgi markers.

Overlap was quantified in at least 50 different images for each marker by using a colocalization coefficient according to Manders (R; Manders et al., 1993; Zinchuk and Zinchuk, 2008). Visually apparent colocalization yields Manders R values of 0.6–1.0 (Zinchuk and Zinchuk, 2008). As shown in Figure 8B, in the cells studied, labeling of the endogenous cis-Golgi marker GM130 and the TGN marker GCC185 yielded an R value of 0.55, which represents a baseline for two adjacent, distinct compartments in these experiments. The myc-tagged, C-terminal C110 fragment yielded an R value of 0.53; N-terminally myc-tagged, full-length GCC185 yielded an R value of 0.58. Thus, although the trend was slightly upward, we did not detect a significantly greater degree of colocalization of the N terminus of the

Figure 6. GCC185 coiled-coils one and two are required to rescue Golgi disruption caused by GCC185 depletion. (A) Cells were treated with fluorescein-labeled GCC185 siRNA for 72 h. After 48 h of siRNA treatment, cells were transfected with plasmids encoding the indicated myc-tagged proteins. Golgi localization was determined using mouse anti-GM130 and Cy3-goat anti-mouse antibodies (green); GCC185 rescue construct expression was detected using chicken anti-myc and Cy3-goat anti-chicken antibodies (red). Cells containing siRNA (as determined by detection of fluorescein) are indicated with asterisks. (B) Quantitation of rescue experiments. Golgi fragmentation was scored visually; data represent the average of two experiments in which a total of ≥40 cells were counted for each condition. Error bars represent SE of the mean of two independent experiments. Bar, 10 μm.
full-length protein with cis-Golgi markers than that seen with the much shorter, C-terminal construct.

From these data, we conclude that the majority of GCC185 molecules do not extend fully toward the cis-Golgi; rather, they are likely to reach laterally across the stack to stabilize this dynamic compartment and outward to help in vesicle tethering. This does not preclude a small subset of the proteins from reaching all the way from the TGN to the cis-Golgi. Given the pancake stack-structure of the Golgi (Rambourg et al., 1979; Clermont et al., 1995; Ladinsky et al., 1999), our findings are consistent with most GCC185 molecules facing outward from the bottom of the stack, rather than being localized to the rims of the stack, where they would have the potential to reach upward.

**DISCUSSION**

We have shown here that GCC185 contains at least five and likely six distinct binding sites for multiple Rab proteins that are distributed along its entire length. Yeast two-hybrid screening revealed the ability of 14 different Rabs to bind to these sites; direct, in vitro binding studies confirmed their existence and documented binding for Rabs1A, 6A, 9A, and 9B. In previous work, we showed that Rab6 binds to a C-terminal coiled-coil via a pair of surface exposed, leucine, and isoleucine residues (Burguete et al., 2008). A similar mode of interaction was revealed for Rab9A binding to GCC185’s coiled-coil 3: mutation of this motif resulted in significantly lowered Rab9A affinity. GCC185’s coiled-coil 3 inhibited transport of MPRs from late endosomes to the TGN in vitro, consistent with its importance for GCC185 function. The importance of GCC185’s first and second, N-terminal coiled-coil domains was revealed in experiments that tested the ability of truncation constructs to rescue the Golgi fragmentation phenotype that is seen upon depletion of GCC185. Although full-length GCC185 was able to fully rescue Golgi fragmentation in GCC185-depleted cells, constructs lacking the first or first and second coiled-coil domains failed to promote reestablishment of a normal Golgi ribbon. Thus, length matters—and despite a sequence that at first glance, seems to be just a long coiled-coil, GCC185’s N-terminal domains play important and unexpected functional roles for this protein in live cells.

As summarized in Table 1, the overwhelming majority of Rab proteins identified in this study as GCC185 binding partners are Golgi-localized. Some of the Rabs are cell type specific: for example, Rab27B is present in regulated secretory cells, but localizes to the Golgi when expressed in cultured fibroblasts (Yoshimura et al., 2007). Why might GCC185 interact with so many Golgi-localized Rab proteins? We favor a model in which GCC185 binding to multiple Rab proteins contributes to its role in maintaining normal Golgi
stack morphology. GCC185 is specifically localized to the TGN (Derby et al., 2004). It is hard to imagine why its depletion would lead to the generation of mini-stacks (Reddy et al., 2006; Derby et al., 2007), unless GCC185 could link ministacks together laterally, or contribute to stack stability, vertically. Simple de-polymerization of microtubules is sufficient to generate Golgi mini-stacks, which suggests that the steady state Golgi structure is normally subject to dynamic, lateral fusion and fission, with the fusion process requiring intact microtubules. Under normal conditions, proteins such as GCC185 could reach across laterally, from one mini-stack domain to another, to promote Golgi ribbon stability. Indeed, GCC185’s connection to the microtubule perinuclear endosomal compartments (Strick and Elferink, 2002), and Rab15 and Rab35 that also seem to be present on the Golgi stack. If GCC185 could bind tightly to Rabs located on other Golgi compartments, those interactions might interfere with its ability to mediate vesicle docking at the TGN. Instead, multiple, weak interactions let GCC185 interact reversibly with the Golgi surface, and stabilize this dynamic compartment. Multiple weak interactions could also retain transport vesicles destined for distinct Golgi compartments within the vicinity of the Golgi stack.

A few Rab proteins that bind GCC185 by yeast two hybrid are not Golgi residents. This includes Rab9A, which is present on inbound transport vesicles (Barbero et al., 2002), and Rab15 and Rab35 that also seem to be present on perinuclear endosomal compartments (Strick and Elferink, 2005; Kouranti et al., 2006). It is possible that these Rabs participate in a retrograde transport step that also uses GCC185. Alternatively, Rab35 is most closely related to Golgi-bound Rabs 1A and 1B; binding in the two-hybrid system may reflect interaction with homologous portions of the Rab protein. We are most confident of the interactions that could be recapitulated with purified proteins and include Rabs 1A, 6A, 9A, and 9B, a collection that spans the entire Golgi stack. GCC185 is not the first Golgin shown to bind to multiple Rabs. For example, GM130 binds Rabs 1, 2 and 33B (Short et al., 2005). But, the ability of GCC185 to potentially reach out to incoming transport vesicles and to earlier Golgi compartments seems to be a novel feature of this GRIP domain family member.

A Model for Vesicle Tethering at the TGN

In addition to contributing to Golgi structure, GCC185 is also required for vesicle docking at the Golgi (Reddy et al., 2006; Derby et al., 2007). Figure 7B presents our most current working model for this process. GCC185 exists in the cytosol and at the TGN (Luke et al., 2003). We propose that Rab9, present on late endosome-derived transport vesicles, binds cytosolic GCC185 via CC3 in the middle of GCC185 (and/or C110). Transport vesicles carrying Rab9-GCC185 then move along microtubules toward their targets (Itin et al., 1999; Barbero et al., 2002). Once close to the TGN, GCC185’s C terminus could bind TGN-localized Rab6β, triggering a concomitant Arl1 binding (Figure 7B, step 1). The transport vesicle would then be brought into proximity to the TGN membrane surface, either by dynamic mobility of a stably anchored GCC185, a GCC185 conformational change, or directed transport of the vesicle down the length of the GCC185 molecule (Figure 7B, step 2). Fusion of the Rab9-containing transport vesicle would then follow.

Validation of these steps in our model will require determining the length of TGN-associated GCC185 protein, and characterizing in greater detail each of its Rab binding sites. In the future, single vesicle analyses of vesicle tethering at the TGN should enable us to elucidate the mechanisms by which Rab9-vesicles find and engage their fusion targets.

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