Independent Transport and Sorting of Functionally Distinct Protein Families in *Tetrahymena thermophila* Dense Core Secretory Granules

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Dense core granules (DCGs) in *Tetrahymena thermophila* contain two protein classes. Proteins in the first class, called granule lattice (Grl), coassemble to form a crystalline lattice within the granule lumen. Lattice expansion acts as a propulsive mechanism during DCG release, and Grl proteins are essential for efficient exocytosis. The second protein class, defined by a C-terminal β/γ-crystallin domain, is poorly understood. Here, we have analyzed the function and sorting of Grt1p (granule tip), which was previously identified as an abundant protein in this family. Cells lacking all copies of *GRT1*, together with the closely related *GRT2*, accumulate wild-type levels of docked DCGs. Unlike cells disrupted in any of the major *GRL* genes, ∆*GRT1* ∆*GRT2* cells show no defect in secretion, indicating that neither exocytic fusion nor core expansion depends on *GRT1*. These results suggest that Grl protein sorting to DCGs is independent of Grl proteins. Consistent with this, the granule core lattice in ∆*GRT1* ∆*GRT2* cells appears identical to that in wild-type cells by electron microscopy, and the only biochemical component visibly absent is Grt1p itself. Moreover, gel filtration showed that Grl and Grt proteins in cell homogenates exist in nonoverlapping complexes, and affinity-isolated Grt1p complexes do not contain Grl proteins. These data demonstrate that two major classes of proteins in *Tetrahymena* DCGs are likely to be independently transported during DCG biosynthesis and play distinct roles in granule function. The role of Grt1p may primarily be postexocytic; consistent with this idea, DCG contents from ∆*GRT1* ∆*GRT2* cells appear less adhesive than those from the wild type.

In eukaryotes, the directional transport of lumenal proteins throughout the network of membrane-bound organelles depends on reversible assembly of multisubunit protein complexes in the cytoplasm. For example, the assembly of a localized clathrin coat at a cell’s surface facilitates both the concentration of specific transmembrane receptors together with their bound ligands at that site and the invagination and budding of the plasma membrane, resulting in endocytosis (18). Similarly, other cytosolic coats assemble and direct traffic at the endoplasmic reticulum (ER) and Golgi apparatus (4). For one protein trafficking pathway in eukaryotic cells, however, the determinative protein self-assembly occurs not in the cytoplasm but within the lumen of the secretory pathway itself. Dense core granules (DCGs) are secretory vesicles whose luminal cargo consists of a condensed polypeptide aggregate. This cargo is secreted when the vesicles fuse with the plasma membrane in response to a specific extracellular stimulus, an event called regulated exocytosis. The aggregation of the cargo occurs progressively within the secretory pathway, beginning in the trans-Golgi network (TGN), and may be promoted by multiple factors including compartment-specific proton and calcium levels (23). Aggregation facilitates the vesicular storage of concentrated secretory proteins but also serves as a sorting mechanism to segregate DCG proteins from proteins that are secreted via other pathways. Evidence for this mechanism includes in vitro experiments showing that some proteins released via constitutive exocytosis remain soluble under TGN-like conditions that promote DCG protein aggregation (10). In vivo, sorting would result if aggregated and soluble proteins exit the TGN in different carriers. Importantly, there is no evidence that sorting of DCG proteins at the TGN requires assembly of cytosolic coat complexes.

While aggregative sorting represents an attractively simple mechanism, relatively little is known about the structure or dynamic properties of the aggregates themselves. This is an interesting issue, as illustrated by several phenomena. First, aggregates in some cell types, like those formed by proinsulin in pancreatic β cells, can become reordered as protein crystals during a multistage process called granule maturation (13). Second, *Aplysia* bag cells can sort different subsets of DCG proteins into distinct granules, suggesting that aggregation can be finely regulated and that different aggregates have different properties in vivo (20). Both of these phenomena have also been observed within the DCGs of unicellular ciliates (3, 14). In addition, ciliate DCGs demonstrate another degree of subtlety in DCG formation because the granule cores in many of these organisms are divided into distinct domains (25). The domain organization indicates that DCG proteins in these cells can segregate from one another even as they are sorted to the same vesicular destination. While the structures of DCGs in many ciliates have been captured by electron microscopy, mo-
lecular studies have advanced in two species, Tetrahymena thermophila and Paramecium tetraurelia (30, 33).

In many ciliates, the individual DCGs are organized in at least two distinct domains within the lumen. First, the bulk of the cargo is organized as a core crystal that expands, spring-like, upon exocytosis (28). This expansion can drive rapid extrusion of the DCG contents, which may be essential for hunting or defensive behaviors (17). In addition, many ciliate DCGs possess a single polarized tip structure that is involved in DCG docking to the plasma membrane and exocytic fusion (25). These tip structures are also filled with condensed, highly organized proteins, which appear by both genetic and morphological criteria to be different from proteins making up the expansible core (1, 21). The proteins that form the distinct domains are beginning to be identified and analyzed. Those that constitute the expansible springs are encoded by homologous families of genes named GRL (granule lattice) in Tetrahymena and tmp (trichocyst matrix) in Paramecium (11, 12, 15). Assembly of Grl proteins begins in the ER with formation of heterooligomers. This is an obligatory step, as shown by the fact that deletion of individual Grl proteins by targeted gene disruption resulted in the ER retention of remaining Grl proteins (12). Further assembly of Grl proteins to form a crystal occurs during DCG maturation and is accompanied by sitespecific proprotein processing (34). Upon exocytosis, the expansion of the crystalline core is controlled by calcium binding to the fully processed Grl proteins (34).

In addition to the GRL-family-encoded proteins, 13 other luminal DCG proteins have been putatively or definitively identified in Tetrahymena, and homologous proteins are predicted in the Paramecium genome (6). The entire set belongs to a gene family that is defined by a carboxy-terminal βγ-crystallin domain, which may function as a DCG-targeting motif (16). Studies of two different members of this family in Tetrahymena, IGRL (induced during granule regeneration 1) and GRT1 (granule tip 1), suggested that these proteins are functionally distinct from the spring-forming Grl proteins. First, whereas gene disruption of any of the highly transcribed GRL genes resulted in grossly aberrant spring formation, no such defect was seen upon disruption of IGRL (16). However, this could be explained by the fact that IGRL1 encodes a relatively low-abundance protein in DCGs, and furthermore its function could be redundant with that of the highly related gene, IGR2.

The second protein in the βγ-crystallin domain family that has been investigated is the 80-kDa product of the GRT1 gene. Grt1p was first detected as one of the most abundant DCG proteins and also that it is packaged intact in DCGs rather than undergoing proteolytic processing (31). Since processing is essential for Grl protein assembly and function, this difference from the spring-forming Grl proteins. First, whereas gene disruption of any of the highly transcribed GRL genes resulted in grossly aberrant spring formation, no such defect was seen upon disruption of IGRL (16). However, this could be explained by the fact that IGRL1 encodes a relatively low-abundance protein in DCGs, and furthermore its function could be redundant with that of the highly related gene, IGR2.

The second protein in the βγ-crystallin domain family that has been investigated is the 80-kDa product of the GRT1 gene. Grt1p was first detected as one of the most abundant DCG components released during exocytosis (32). Biochemical analysis showed that Grt1p differs in its solubility from the Grl proteins and also that it is packaged intact in DCGs rather than undergoing proteolytic processing (31). Since processing is essential for Grl protein assembly and function, this difference appears highly significant. Second, Grt1p accumulates at a single pole of each DCG, corresponding to the tip of the organelle that docks and then fuses with the plasma membrane (5). Two Mendelian mutants with defects in DCG maturation show delocalized Grt1p, and these mutant DCGs can dock but do not appear to undergo exocytosis (5). These results suggested that Grt1p might be involved in forming a DCG tip domain that interacted with the plasma membrane.

We have now investigated the trafficking and function of Grt1p. Our data provide both direct biochemical and cell-biological evidence that Grt1p and Grl proteins form distinct complexes during DCG biogenesis in Tetrahymena. Together with earlier results, our experiments provide genetic evidence that Grl and Grt complexes can be independently trafficked to DCGs. Cells lacking GRT1, together with the closely related GRT2, still show rapid and efficient release of DCG contents upon stimulation with secretagogues, but the released DCG contents are subtly different from those of the wild type, suggesting that Grt1p may primarily serve a postexocytic function.

MATERIALS AND METHODS

Tetrahymena strains and culture conditions. Wild-type CU428.1 T. thermophila (from Peter Bruns, Cornell University) and the SB281 mutant (from Eduardo Orias, University of California—Santa Barbara) were grown at 30°C in SPP medium (1% proteose peptone [BD, Sparks, MD] 0.2% dextrose, 0.1% yeast extract, 0.009% ferric EDTA). All reagents were from Sigma Chemical Co. unless otherwise indicated. Cell densities were measured using a Coulter Counter (Coulter Electronics Limited, Luton, United Kingdom).

GRT1 and GRT2 disruption. A fragment containing the entire coding region of GRT1 and part of GRT2 (residues 1 to 2467), together with the intergenic sequence, was replaced in a genomic clone with the neo3 cassette, using EcoRI sites added to primers on both the 5’ and 3’ ends. The neo3 cassette includes a gene encoding paromomycin resistance under the control of the cadmium-inducible MTT1 promoter (27). The resulting construct was linearized and introduced by biolistic bombardment into vegetative Tetrahymena by particle bombardment to replace the corresponding endogenous sequence by homologous recombination (9). The replacement was driven to completion to generate GRT1/GRT2 double-knockout cells (ΔGRT1 ΔGRT2) by growing the initial transformants in the presence of increasing concentrations of paromomycin (up to 1 mg/mL) with 0.5 μg/mL cadmium chloride.

Southern blotting. Genomic DNA samples from wild-type and ΔGRT1 ΔGRT2 cells were digested with NcoI and probed with an oligonucleotide corresponding to the 5’ flank of GRT1 (nucleotide positions –151 to –42, relative to the translation start). Western blotting. Protein samples were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to 0.45-μm pore-size nitrocellulose (Osmotics, Westborough, MA), under previously described conditions (7). Individual blots were incubated with polyclonal antibodies against either Grt1p (1:500), Grl1p (1:1,000) (32), Grl3p(1:400), or Grl4p (1:250) (12); proteins were detected either using horseradish peroxidase-conjugated goat anti-rabbit antibody at 1:2,000 (Jackson Immunoresearch Laboratories, West Grove, PA), followed by development with Pierce SuperSignal (Pierce Chemical, Rockford, IL) and exposure in an α-Imager (Alpha Innotech, San Leandro, CA), or using Alexa Fluor 600 donkey anti-rabbit immunoglobulin G (IgG) at 1:5,000 (MolecularProbes, Carlsbad, CA) with an Odyssey infrared imaging system (Li-Cor Biosciences, Lincoln, NE).

Immunofluorescence. Cells were fixed and immunolabeled as described previously (7). Grt1p and Grl3p were visualized using monoclonal antibody (MAB) 4D11 (20%, vol/vol) (31) and MAB 5E9 (20%, vol/vol) (5), respectively, followed by development with Pierce SuperSignal (Pierce Chemical, Rockford, IL) and exposure in an α-Imager (Alpha Innotech, San Leandro, CA), or using Alexa Fluor 600 donkey anti-rabbit immunoglobulin G (IgG) at 1:5,000 (MolecularProbes, Carlsbad, CA) with an Odyssey infrared imaging system (Li-Cor Biosciences, Lincoln, NE).

Electron microscopy. Cells were fixed for 30 min with 3% glutaraldehyde in 0.1 M Na cacodylate buffer, pH 7.2, and then washed three times for 10 min each time with 10% sucrose in 0.1 M Na cacodylate, pH 7.2, and postfixed with 1% osmium tetroxide in the same buffer. After samples were washed with water, they were dehydrated by serial passage through solutions of increasing ethanol concentration, from 20 to 100%, and embedded in Epon resin. After polymerization at 60°C for 24 to 48 h, the embedded samples were cut to yield 100-nm sections, which were collected on formvar-coated copper grids and poststained in 2% uranyl acetate and Reynolds’s lead citrate. Electron micrographs were collected on an FEI Tecnai F30 electron microscope interfaced with a high-performance charge-coupled-device camera.

Dibucaine stimulation and purification of DCG contents. Wild-type or ΔGRT1 ΔGRT2 cells were grown to 4 × 10^7 to 6 × 10^7 cells/mL and washed once with 10 mM Na HEPES (pH 7.2) after being pelleted for 30 s at 400 × g in a clinical centrifuge. Loose cell pellets (concentrated ~10-fold relative to the initial culture) were stimulated for 30 s by addition of 2.5 mM dibucaine (from a 25 mM stock). The cells were then diluted at least fivefold with buffer and...
centrifuged at 1,200 × g for 1 min. This resulted in a two-layer pellet, with cells overlaid by flocculent extruded DCG contents. For experiments requiring purification of the DCG contents, the flocculent layer was gently removed with a pipette and dispersed thoroughly, and the centrifugation process was then repeated four to six times until the flocculent contained no visible trapped cells. The final pelleting of flocculent to remove trapped cells was at 10,000 × g for 5 min. The final flocculent was resuspended in buffer containing 1% SDS for electrophoretic analysis. Samples were incubated for 3 min at 100°C, separated by SDS-PAGE, and stained with Coomassie blue G250, and images were captured using an α-imager (Alpha Innotech, San Leandro, CA).

Size exclusion chromatography. Cultures of SB281 (200 ml) (24) were grown overnight to stationary phase (7). When SB281 lysates were analyzed by gel filtration chromatography, a crude membrane fraction from SB281 cells was solubilized in Triton X-100, and the proteins were resolved by electrophoretic analysis. Samples were incubated for 3 min at 100°C, separated by SDS-PAGE, and stained with Coomassie blue G250, and images were captured using an α-imager (Alpha Innotech, San Leandro, CA).

Expression and purification of Grt1p-His6. The coding sequence of GRT1 was amplified using a forward primer containing a PmlI restriction site and a reverse primer containing the coding sequence for His6, followed by a stop codon and an Apal restriction enzyme site. The amplified product was cloned into the pNCVB vector predigested with these two enzymes. The pNCVB vector was designed to integrate a gene of interest at the MTT1 locus, and transformants can be selected based on the presence of a blasticidin resistance cassette in the 3’ untranslated region (5). The linearized insert was used to transform the SB281 Tetrahymena strain by biolistic bombardment. We used the SB281 line rather than the wild type because wild-type cell lysates have a large pool of insoluble or partially soluble DCG proteins released from DCGs. SB281 transformants were selected and passaged in the presence of 60 μg/ml blasticidin and 1.0 μg/ml CdCl2. To purify Grt1p-His6, we first grew 25-ml transformant cultures overnight in the presence of blasticidin and CdCl2. These were then used to inoculate 400-ml cultures grown overnight to stationary phase (7 × 106 cells/ml) in the absence of drug or CdCl2. Transgene expression was subsequently induced by treating the 400-ml cultures with 1.0 μg/ml CdCl2 for 6 h at 30°C. The cells were pelleted and disrupted by suspension in 20 ml of ice-cold lysis buffer (50 mM Na-phosphate, pH 7.0, 200 mM NaCl, 1% Triton X-100, 10 mM imidazole), supplemented with protease inhibitors, for 30 min. Lysates were clarified by centrifugation at 150,000 × g for 45 min in a TLA 100.4 rotor (Beckman, Fullerton, CA), and the supernatants were incubated with 200 μl of Talon resin (BD Biosciences, Palo Alto, CA) for 2 h at 4°C with end-to-end shaking. The resin was washed sequentially with 100 volumes of lysis buffer followed by 3 volumes each of lysis buffer containing 40 mm and 60 mM final concentrations of imidazole. Bound protein was eluted with lysis buffer containing 250 mM imidazole. This purification protocol essentially follows that of Cowan et al. (12).

Expression analysis of GRT1 and GRT2. Transcript levels of GRT1, GRT2, and IGR1 in growing, starved, and conjugating Tetrahymena cells were determined by hybridization to whole-genome microarrays as part of a previous genome-wide expression study (22). The database arising from that study, which is searchable at http://tged.ihb.ac.cn/, was queried to determine the expression profiles of GRT1, GRT2, and IGR1.

RESULTS

Grt1p and Gr1lp are found in nonoverlapping transport complexes. Grl proteins appear to be transported through the Tetrahymena secretory pathway as heterooligomeric complexes (12). To ask whether Grt1p was similarly transported as part of a large complex, we used gel filtration chromatography to resolve Gr1lp- and Grt1lp-containing complexes present in detergent cell lysates. In order to focus on DCG proteins in transit rather than on the proteins that accumulate in DCGs, we exploited a mutant cell line called SB281 that lacks DCGs and in which newly synthesized DCG proteins are rapidly secreted instead of accumulating in a storage compartment (7). A bulk membrane fraction of SB281 cells was prepared and solubilized in Triton X-100, and the proteins were resolved by gel filtration followed by SDS-PAGE and Western blotting. In these samples Gr1lp appears in its proprotein form (pro-Grlp) since there is no detectible enzymatic or morphological DCG maturation in this mutant (32).

When SB281 lysates were analyzed by gel filtration chromatography, pro-Grlp was found predominantly in fractions at or near the column void volume (Fig. 1). This is consistent with previous genetic evidence that pro-Grlp is transported as part of a large oligomeric complex. On the same columns, however, Grt1p was predominantly found in fractions corresponding to much smaller proteins. Indeed, Grt1p, with a monomeric molecular mass of ~80 kDa, eluted shortly after the bovine serum albumin (BSA) marker (67 kDa). There was no significant cofractionation of Grt1p and Gr1lp. This suggests that most of the Grt1p is in a monomeric form not associated with other DCG proteins and that the monomer is very compact.
His6 construct was introduced on a multicopy plasmid, under

roborate these results using a different approach, we fused

flowthrough; lanes 3 to 10, elution with increasing concentrations of

seen in SB281 lysates. Lane 1, lysate applied to column. lane 2, column

band corresponds to Grt1p; the lower band is a cross-reactive species

analyzed by Western blotting using anti-Grt1p antibodies. The upper

panel) Nickel column fractions were subjected to SDS-PAGE and

affinity chromatography as described in Materials and Methods. (Top

expressed in SB281 cells and purified from detergent lysates by nickel-

nickel affinity columns (Fig. 2, top panel). In contrast, none of

proteins (12). As shown in Fig. 2, the His-tagged Grt1p from

induced cell lysates. We previously used this approach to show that His-tagged Grl1p was associated with several other Grl proteins (12). As shown in Fig. 2, the His-tagged Grt1p from SB281 lysates could be readily bound and then eluted from nickel affinity columns (Fig. 2, top panel). In contrast, none of three different Grl proteins bound to the column or coeluted with Grt1p-His6. The results for Grt1p-His6 are consistent with the gel filtration profiles in Fig. 1 and suggest that Grt1p does not interact strongly with Grl proteins under these conditions.

Grt1p has a closely related paralog, Grt2. If Grt1p is not associated with Grl proteins, the proteins may be independently sorted to DCGs and play distinct roles within the DCG core. To test the function and sorting properties of Grt1p, we used homologous recombination to disrupt the endogenous gene. However, a BLAST search with the GRT1 coding sequence revealed that a second, very similar gene was present in T. thermophila, which we named GRT2. GRT1 and GRT2 are adjacent to one another on macronuclear chromosome

CH445588, with 2.5 kb of intergenic sequence (Fig. 3A) (Tetrahymena genome database at http://ciliate.org). The predicted translation products of GRT1 and GRT2 are 83% identical and 92% similar (see Fig. S1 in the supplemental material).

Based on whole-genome microarray analysis of mRNA isolated from Tetrahymena under a variety of conditions, GRT1 is highly expressed in growing, starved, and conjugating cells (22) (Fig. 3B). The profile of GRT1 expression is very similar to that of other DCG protein genes, including GRL1 and IGRI for which the profiles are shown in Fig. 3. The microarray data suggest that GRT2 is transcribed at a much lower level than GRT1 under all conditions.

Gene disruption of GRT1 and GRT2. Since GRT1 and GRT2 may under some circumstances have redundant activities, we designed a single replacement construct to disrupt both genes. We transformed vegetative cells to integrate one or a small number of copies of the knockout construct at the macronuclear GRT1/GRT2 locus and passaged the cells in the presence of paromomycin. Under these conditions, the knock-out construct replaced all endogenous alleles in the macronucleus, leaving intact only the two micronuclear alleles of GRT1/GRT2 (Fig. 4, left blot). Because there is no vegetative transcription in the micronucleus, the resulting strains have functional deletions of GRT1 and GRT2 (∆GRT1 ∆GRT2) under the conditions in which the cells were characterized. As expected, these cells accumulated no detectible Grt1p (Fig. 4, right blot). The ability to derive such cells indicates that GRT1 and GRT2 are nonessential, and indeed ∆GRT1 ∆GRT2 cells grew comparably to wild-type cells (data not shown).

∆GRT1 ∆GRT2 cells are distinct from cells with disruptions in GRL genes. The disruption of any of the genes encoding the six abundant Grl proteins results in Tetrahymena cells whose DCG cores are no longer organized as a lattice (12). Such mutant cores are spherical rather than elongated, a difference that can be seen by indirect immunofluorescence using either of two MAb. The MAB 5E9 recognizes Grl3p while MAB 4D11 recognizes Grt1p (5). In wild-type cells, both antibodies reveal large numbers of DCGs, most of which are docked in linearly arranged sites at the plasma membrane (Fig. 5A and B). As expected, immunofluorescence with MAB 4D11 did not reveal any DCGs in ∆GRT1 ∆GRT2 cells but showed only a diffuse background signal since the protein target of the antibody is absent (Fig. 5B). In contrast, MAB 5E9 labeled DCGs in the ∆GRT1 ∆GRT2 cells that were indistinguishable in both abundance and morphology from wild-type cells (Fig. 5A). The similar accumulation of DCGs in wild-type and ∆GRT1 ∆GRT2 cells was also demonstrated by Western blotting of whole-cell lysates with an antibody against the granule marker Grl1p, which showed that equivalent amounts of this protein accumulated in the wild-type and mutant cells (Fig. 5C). The DCGs in wild-type and ∆GRT1 ∆GRT2 cells were also indistinguishable at the level of electron microscopy (Fig. 6).

To test whether the DCGs lacking Grt1p and Grt2p could undergo exocytosis, we stimulated ∆GRT1 ∆GRT2 cells with two compounds that provoke rapid global DCG secretion in wild-type Tetrahymena. These are Alcian blue and dibucaine. When wild-type cells are treated with the polycationic dye Alcian blue, they become entrapped in robust capsules formed by the dye-dependent cross-linking of the exocytosed DCG contents (29). When we treated wild-type and ∆GRT1 ∆GRT2 cells in parallel with

FIG. 2. Grt1p does not copurify with Grl proteins. Grt1p-His6 was expressed in SB281 cells and purified from detergent lysates by nickel-affinity chromatography as described in Materials and Methods. (Top panel) Nickel column fractions were subjected to SDS-PAGE and analyzed by Western blotting using anti-Grt1p antibodies. The upper band corresponds to Grt1p; the lower band is a cross-reactive species seen in SB281 lysates. Lane 1, lysate applied to column. lane 2, column flowthrough; lanes 3 to 10, elution with increasing concentrations of imidazole, with each fraction representing 1 column volume (fractions 3 and 4, 40 mM; 5 to 7, 60 mM; 8 to 10, 250 mM). Grt1p-His6 elutes primarily in fractions 7 to 9. Arrows on the right indicate the band corresponding to Grt1p and the position of the 67-kDa SDS-PAGE standard, BSA. The remaining three panels show the identical column fractions analyzed by Western blotting with antibodies against Grl1p, Grl3p, and Grl8p. Column fractions correspond to those in the top panel. In all cases, the Grl proteins are found in the column flowthrough and do not copurify with Grt1p-His6.

Purified Grt1p is not associated with Grl proteins. To corroborate these results using a different approach, we fused full-length Grt1p at its C terminus to a His6 tag. The GRT1-His6 construct was introduced on a multicopy plasmid, under the control of an inducible promoter, into the Tetrahymena strain SB281, and the His-tagged protein was purified from induced cell lysates. We previously used this approach to show that His-tagged Grl1p was associated with several other Grl proteins (12). As shown in Fig. 2, the His-tagged Grt1p from SB281 lysates could be readily bound and then eluted from nickel affinity columns (Fig. 2, top panel). In contrast, none of three different Grl proteins bound to the column or coeluted with Grt1p-His6. The results for Grt1p-His6 are consistent with the gel filtration profiles in Fig. 1 and suggest that Grt1p does not interact strongly with Grl proteins under these conditions.
Alcian blue, virtually 100% of cells from both cultures were encapsulated (data not shown). The DCGs in ΔGRT1 ΔGRT2 cells are therefore capable of undergoing regulated exocytosis under these conditions. However, the Alcian blue assay does not allow one to assess the extent of exocytosis, that is, the fraction of DCGs that have released their contents.

A second compound that acts as a potent secretagogue in *Tetrahymena* is the local anesthetic dibucaine (26). When treated with dibucaine, wild-type cells release their DCG contents within seconds. The DCG cores expand during release but remain highly insoluble, and they can be readily pelleted to form a flocculent layer. Measuring the volume of this flocculent layer provides a simple and quantitative assay for exocytosis (12). When wild-type and ΔGRT1 ΔGRT2 cultures were treated in parallel with dibucaine, both produced very similar volumes of flocculent (Fig. 7, left panel). Taken together, our results indicate that ΔGRT1 ΔGRT2 cells accumulate wild-type levels of DCGs, that those DCGs undergo rapid exocytosis in response to dibucaine, and that the expansion of the ΔGRT1 ΔGRT2 cores is comparable to wild type.

During these experiments, we noted a subtle but reproducible difference between the exocytosed material in wild-type and ΔGRT1 ΔGRT2 cells. When *Tetrahymena* cultures are pelleted shortly after dibucaaine addition, a large fraction of

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**FIG. 3.** Organization and expression of *GRT1* and *GRT2*. (A) The organization of *GRT1* and *GRT2* on macronuclear chromosome CH445588. (B) The expression profiles of *GRT1* and *GRT2* at successive time points in growing (L-l, L-m, and L-h), starved (S-0, S-3, S-6, S-9, S-12, S-15, and S-24), and conjugating (C-0, C-2, C-4, C-6, C-8, C-10, C-12, C-14, C-16, and C-18) cultures, as determined by hybridization of stage-specific cDNAs to whole-genome microarrays. Details on the sampling times are found in Miao et al. (22). Also shown are the profiles for two known DCG protein genes, *GRL1* and *IGR1*. 
wild-type cells remains trapped in the expanded DCG cores that form the flocculent layer. This can be seen in the greatly reduced size of the cell pellet in poststimulation, compared to prestimulation, samples (Fig. 7, right panel). The reduction in cell pellet size represents trapping of cells in the flocculent layer and not cell mortality or lysis since the original volume of cell pellet size represents trapping of cells in the flocculent layer and not cell mortality or lysis since the original volume of cells could be recovered by repeated dilution, dispersion, and repelleting of the flocculent layer (data not shown). The major conclusion of the data, combined with results from previous studies, is that the Grl and Grt proteins, though targeted to the same DCGs, are likely to act independently with regard to assembly and sorting and to serve distinct functions for Tetrahymena. The assembly of Grl proteins involves obligate heterooligomer formation in the ER, transport through the secretory pathway in the form of large complexes, and extensive proteolytic processing during DCG maturation (12, 34). None of these appears to be relevant for Grt1p. Previous analysis showed that Grt1p did not undergo processing, and data in the current paper show that it also differs from the Grts in the other properties (5, 31). Gel filtration analysis indicated that Grt1p in the secretory pathway is primarily a monomer, and none of the protein fractionates with the much larger complexes that contain pro-Grt1p. These large complexes are known, from genetic experiments, to contain other pro-Grl proteins (12). Consistent with the absence of Grt1p from these complexes, we found that Grt1p, when affinity purified as a His-tagged variant from Tetrahymena cell lysates, did not copurify with Grl proteins. These biochemical experiments were all done using the SB281 strain, a mutant that fails to synthesize DCGs, in order to focus on protein in transit in the secretory pathway. In wild-type cell lysates, only a small fraction of the Grl or Grt proteins are derived from protein in...
transit while the vast majority reflect the pool released from DCGs. We previously used SB281 lysates to identify heterooligomeric Grl protein complexes, and genetic results strongly supported the existence of the same complexes in wild-type cells (12). We therefore expect that the characterization of Grt1p in SB281 lysates is fully relevant to wild-type cells in which DCGs are formed although we cannot rule out the possibility that there are additional complexes that form in the wild type but are absent in SB281. We do not yet know the genetic lesion in SB281 but a cell biological defect is clearly manifested at the level of the late Golgi compartment/TGN, where there is no sorting of constitutive versus regulated secretory cargo in the mutant (5, 7). If there is a difference between the TGN conditions in the wild-type and SB281 strains, these may have an effect on protein oligomerization in that compartment. Our data, in which protein-protein interactions are analyzed at neutral pH, cannot rule out the possibility that Grt1p and Grl proteins associate in the conditions that exist in the lumen of the ciliate TGN, about which nothing is currently known. Interactions could also be promoted in the TGN by compartment-specific posttranslational modification of Grl proteins or Grt1p though previous work suggests there is no detectible difference in the electrophoretic mobilities of any of these proteins when they are isolated from wild-type and SB281 cells (31, 32). This is an important issue because aggregation is believed to underlie DCG protein sorting in mammalian cells (23).

Importantly, even if there are Grt1p-containing complexes that we cannot detect for technical or biological reasons in SB281 lysates, our composite data in this system argue that such complexes cannot be essential for efficient sorting of either Grt1p or Grl proteins to DCGs in wild-type cells. First, in a thorough study of Grl protein function, we found that the absence of Grl1p had no effect on Grt1p accumulation in DCGs (11). In subsequent studies each of the GRL genes was disrupted, and in every case the resulting DCGs could still be visualized using a MAb directed against Grt1p (12). These results strongly suggested that Grt1p does not depend for sort-

FIG. 5. GRT1 and GRT2 are not essential for DCG formation and accumulation. (A) Visualization of docked DCGs by indirect immunofluorescence using a MAb (5E9) directed against the DCG lattice protein Grl3p. Shown are mid-cell sections of one wild-type (WT) and two ΔGRT1 ΔGRT2 (ΔGRT1-2) (cell line A) cells. Similar numbers of elongated DCGs are present, predominantly docked at the plasma membrane. (B) Visualization of docked DCGs by indirect immunofluorescence using a MAb (4D11) directed against Grt1p. Shown are tangential sections of one wild-type and one ΔGRT1 ΔGRT2 cell. The DCGs, seen en face, are docked at linearly arranged docking sites in the wild-type cell. No DCGs are seen in the ΔGRT1 ΔGRT2 cell since the MAb is directed against Grt1p. (C) Western blot analysis of SDS lysates of wild-type and ΔGRT1 ΔGRT2 (line A) cells using a polyclonal antibody against the granule lattice protein Grl1p. A total of 2 × 10^5 cell equivalents were loaded for each sample, and the bound antibody was detected using the Odyssey system (see Materials and Methods). Lane 1, wild type; lane 2, ΔGRT1 ΔGRT2. Wild-type and ΔGRT1 ΔGRT2 cells accumulate similar levels of Grl1p.

FIG. 6. ΔGRT1 ΔGRT2 DCGs have lattice-organized cores. Wild-type and ΔGRT1 ΔGRT2 (line A) cells were analyzed by thin-section electron microscopy, as described in Materials and Methods. Both images show elongated DCG profiles in which the central core is visibly organized.
the amount of the Grl proteins released from DCGs in (12). Moreover, we found by direct biochemical analysis that previously been shown to result in dramatic exocytosis defects lack any of the Grl proteins since such deficiencies have predisposed DCGs with no exocytosis defects. These DCGs cannot functionally and biochemically more complex than those in Tetrahy-
mema DCGs. This is clear because the complementary evidence that cells lacking Grt1p and the function. The data in the current paper provide the important that Grt1p depends on Grl proteins for sorting but that any Grl protein depends on any individual Grl protein. A remaining possibility is that Grt1p depends on Grl proteins for sorting but that any Grl protein is sufficient, i.e., there is extensive redundancy for this function. The data in the current paper provide the important complementary evidence that cells lacking Grt1p and the closely related Grt2p still sort all of the major Grl proteins to DCGs. This is clear because the △GRT1 △GRT2 cells synthesized DCGs with no exocytosis defects. These DCGs cannot lack any of the Grl proteins since such deficiencies have previously been shown to result in dramatic exocytosis defects (12). Moreover, we found by direct biochemical analysis that the amount of the Grl proteins released from DCGs in △GRT1 △GRT2 cells, as judged by Coomassie staining, was indistinguishable from amounts in the wild type.

The analysis of the △GRT1 △GRT2 cells allowed us to test ideas about potential roles of Grt1p. Prior models for Grt1p function were based on the localization of Grt1p at the tip of DCGs, where docking and fusion occur (5). This suggested that Tetrahymena DCGs might have a tip domain that was specialized for these functions. This model was also inspired by analogy with the DCGs in P. tetraurelia, which are both structurally and biochemically more complex than those in Tetrahymena. Paramecium DCGs have a morphologically distinct protrusion at the tips of the DCGs whose formation is essential for docking and fusion (1). The proteins that compose the Paramecium tip have not been identified, but one strong possibility is that they are encoded by the many Paramecium homologs of GRT1, whose roles have not yet been analyzed. Based on this analogy, we hypothesized that Tetrahymena DCGs possess a functionally important though morphologically undetectable tip domain, comprised in part by Grt1p. However, the △GRT1 △GRT2 cells showed no apparent defects in DCG docking or fusion and so do not support a model in which Grt1p is a determinant of a docking or fusion-specialized tip domain. It remains to be tested whether other members of the crystallin domain family in Tetrahymena, distantly related to GRT1 and GRT2, may play such roles.

GRT1 is the second non-GRL gene in Tetrahymena DCGs to be analyzed, and in both cases gene disruption had no effect on DCG core formation or exocytosis (16). In the case of IGR1 the lack of phenotype could be potentially explained by the low abundance of the protein and also by the fact that a second, closely related gene was subsequently found to be encoded in the genome. In contrast, Grt1p is an abundant protein in Tetrahymena DCGs, and we preempted any potential functional redundancy with the related GRT2 gene by generating the △GRT1 △GRT2 double knockout strain. The absence of exocytosis phenotypes for both IGR1 and GRT1 is therefore consistent with the idea that both Igr1p and Grt1p primarily serve postexocytic functions as secreted proteins, for example, by binding proteins or other molecules present in the cell’s environment. Since Grt1p remains associated after exocytosis with the Grl-based expanded core, as judged by its pelleting in the flocculent layer, the dense cores of ciliate DCGs may serve not only as extrusion machines during exocytosis but also, by virtue of associated proteins like Grt1p, as biologically active complexes that are released into the medium. Since Grt1p remains associated after exocytosis with the Grl-based expanded core, as judged by its pelleting in the flocculent layer, the dense cores of ciliate DCGs may serve not only as extrusion machines during exocytosis but also, by virtue of associated proteins like Grt1p, as biologically active complexes that are released into the medium. Since Grt1p is localized at the docking tip of the DCG, it is positioned to be among the first DCG contents to become externalized upon exocytosis. The function of DCGs in Tetrahymena is not yet known, but DCG exocytosis can protect Paramecium from predators wielding paralyzing toxins (17, 19). This could involve the rapid externalization of toxin-binding proteins at the tips of DCGs during a defense response. We have suggestive evidence that
Gr1p is a sticky protein since the released contents of DCGs lacking Gr1/Grt2 appeared to be less adhesive than the contents of the wild-type DCGs. This conclusion is based on the observation that fewer cells were trapped in the extruded ΔGrt1 ΔGrt2 DCG contents than in the wild-type DCG contents in a pelleting assay. This model of DCG function in Tetrahymena can be contrasted with a second possibility, in which the role of expansible Grl-based cores is to rapidly disperse soluble molecules, such as peptides derived from large stretches of the Grl proproteins that are proteolytically degraded during DCG maturation. The models are not mutually exclusive, and the two mechanisms may contribute to multiple distinct functions of DCGs in these cells.

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