Overexpression of a Mutant Form of EhRabA, a Unique Rab GTPase of *Entamoeba histolytica*, Alters Endoplasmic Reticulum Morphology and Localization of the Gal/GalNAc Adherence Lectin\(^*\)

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Received 21 January 2009/ Accepted 7 April 2009

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\(^*\) Published ahead of print on 17 April 2009.

*Entamoeba histolytica* is a protozoan parasite that causes amoebic dysentery and liver abscess. Vesicle trafficking events, such as phagocytosis and delivery of plasma membrane proteins, have been implicated in pathogenicity. Rab GTPases are proteins whose primary function is to regulate vesicle trafficking; therefore, understanding the function of Rab in this organism may provide insight into virulence. *E. histolytica* possesses a number of unique Rabs that exhibit limited homology to host Rabs. In this study we examined the function of one such Rab, EhRabA, by characterizing a mutant overexpressing a constitutively GTP-bound version of the protein. Overexpression of mutant EhRabA resulted in decreased adhesion to and phagocytosis of human red blood cells and in the appearance of large tubular organelles that could be stained with endoplasmic reticulum (ER)-specific but not Golgi complex-specific antibodies. Consistent with the adhesion defect, two subunits of a cell surface adhesin, the galactose/N-acetylgalactosamine lectin, were mislocalized to the novel organelle. A cysteine protease, EhCP2, was also localized to the ER-like compartment in the mutant; however, the localization of two additional cell surface proteins, Igl and SREHP, remained unchanged in the mutant. The phenotype of the mutant could be recapitulated by treatment with brefeldin A, a cellular toxin that disrupts ER-to-Golgi apparatus vesicle traffic. This suggests that EhRabA influences vesicle trafficking pathways that are also sensitive to brefeldin A. Together, the data indicate that EhRabA directly or indirectly influences the morphology of secretory organelles and regulates trafficking of a subset of secretory proteins in *E. histolytica.*
Rabs and thus are considered unique to *E. histolytica* (34). We previously reported that one of these unique Rabs, EhRabA, may be involved in the regulation of polarization, motility, and actin cytoskeletal dynamics (47, 48). In the current study, we demonstrate that overexpression of the putatively GTP-bound form of EhRabA results in the alteration of ER morphology, mislocalization of two subunits of the Gal/GalNAc lectin, and reduced phagocytosis, suggesting that this Rab plays a direct or indirect role in cellular functions that contribute to virulence.

**MATERIALS AND METHODS**

**Strains, culture, and assay conditions.** *E. histolytica* trophozoites (strain HM-1:1MSS) were cultured axenically in TYI-S-33 medium (7) in 15-ml glass screw cap tubes at 37°C. Several assays were performed in serum-free *E. histolytica* medium, TYI-33. Chinese hamster ovary (CHO) cells were grown at 37°C in Dulbecco’s modified Eagle’s medium supplemented with fetal bovine serum (10%, vol/vol), 100 U/ml penicillin-streptomycin, and 1 mM HEPES.

**Antibodies.** Specific antibodies recognizing the subunits of the Gal/GalNAc lectin were gifts of W. A. Petri, Jr. (University of Virginia, Charlottesville). The generation of specific antibodies recognizing EhRabA, calreticulin (gift of N. Guillem, Unité de Biologie Cellulaire du Parasitisme, Institut Pasteur, Paris, France), EhCP2, and SREHP (gifts of S. L. Stanley, Jr., Washington University School of Medicine, St. Louis, MO) has been described elsewhere (9, 38, 48, 50). Monoclonal and polyclonal antibodies recognizing the hemagglutinin (HA) epitope were obtained from Roche Pharmaceuticals (Nutley, NJ) and Zymed (South San Francisco, CA), respectively. The 10C3 mouse monoclonal antibody recognizing the KDEL ER retention signal was obtained from Abcam (Cambridge, MA).

**Western blot analysis.** Western blot analysis of *E. histolytica* cell lysates was performed as described previously (47). Dilutions for anti-EhRabA, anti-Hgl, anti-Igl, anti-KDEL, or anti-SREHP primary antibodies were 1:1500, 1:500, 1:250, 1:2500, and 1:800, respectively. Commercial polyclonal anti-HA antibody was utilized at 2 μg/ml. In all cases, Western blots were normalized by loading equal cell number equivalents to each of the lanes.

**Mutagenesis of the EhRabA cDNA and transfection of *E. histolytica* trophozoites.** Mutagenesis was carried out by a PCR-based method to generate a constitutively GTP-bound version of EhRabA (EhRabAQ84L). In particular, the codon for glutamine (Q) at amino acid position 84 was changed to the codon (CTA) for leucine (L) by using the QuikChange kit (Stratagene, La Jolla, CA) according to the manufacturer’s protocol. Nucleotides encoding an N-terminal HA tag, a 5'-BamHI site, and a 3'-SalI site were also incorporated into the cDNA by PCR as described previously (47). The resulting PCR product was digested with BamHI and SalI and ligated into the *E. histolytica* expression vector pGIR209 (30) (gift of W. A. Petri, Jr., University of Virginia, Charlottesville), which had been digested with BgII and SalI. This vector allows for the inducible expression of exogenous genes when introduced into trophozoites (30). A constitutively GTP-bound (40). Second, to distinguish mutant EhRabA from endogenous EhRabA, the 5’ end of the EhRabA cDNA was modified to include nucleotides encoding the HA epitope (4). The N-terminal fusion was chosen to avoid interference with C-terminal structural elements responsible for membrane association of proteins belonging to the Ras superfamily (36).

The modified cDNA was inserted into the *E. histolytica* expression vector, pGIR209, which confers G418 (neomycin) resistance and allows for tetracycline-inducible expression of exogenous genes when introduced into trophozoites (30). A standard electroporation protocol (43) was utilized to introduce the expression vector into trophozoites which had been previously transfected with an additional plasmid, pGIR308. This companion plasmid encodes the tetracycline repressor protein, which is necessary for tetracycline inducibility. Transfection was confirmed by purification of the expression plasmid from stably transfected cell lines as described previously (43) and sequencing of the cDNA insert (data not shown).

**Results**

**Generation of an *E. histolytica* mutant expressing a GTP-bound form of EhRabA.** To gain insight into the function of EhRabA we generated an *E. histolytica* cell line that conditionally overexpressed a constitutively GTP-bound version of this protein. The cDNA encoding EhRabA was modified in two ways. First, the cDNA was mutagenized to change the codon for glutamine (Q) at amino acid position 84 to that for leucine (L), to generate a mutant version termed EhRabAQ84L. This mutation was chosen based on an identical mutation made in numerous other small-molecular-weight GTPases (5, 28, 35), including a Rab5-like GTPase of *E. histolytica* (35). In other systems, this mutation has been shown to dramatically slow GTPase activity resulting in Rabs that are constitutively GTP bound (40). Second, to distinguish mutant EhRabA from endogenous EhRabA, the 5’ end of the EhRabA cDNA was modified to include nucleotides encoding the HA epitope (4). The N-terminal fusion was chosen to avoid interference with C-terminal structural elements responsible for membrane association of proteins belonging to the Ras superfamily (36).

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Maximum expression of modified EhRabA was achieved by...
the addition of 5 μg/ml tetracycline to the culture medium for 24 h. Western blot analysis using anti-EhRabA antibody revealed two forms of the EhRabA protein in transformed cells after induction with tetracycline: a lower-molecular-mass species (23 kDa) representing endogenous EhRabA, and a higher-molecular-mass species (29 kDa) representing the HA-tagged mutant version (Fig. 1A). The latter exhibited a molecular mass that was slightly higher than expected; however, we cannot rule out the possibility that the mutation alters posttranslational modifications. Decreased prenylation and/or increased methylation, two modifications normally seen on Rabs (13, 37), would have resulted in a slower electrophoretic mobility.

An intermediate-molecular-mass protein was also detected in the mutants by Western blotting with EhRabA-specific antibody. This protein species may represent a degradation product of the higher-molecular-mass species, as it was not recognized by HA-specific antibody (Fig. 1A). The HA-tagged protein was not visible in lysates of wild-type cells or in lysates of mutant cells prior to tetracycline induction (Fig. 1A), confirming the inducibility of the expression system. Longer inductions with tetracycline (48 to 72 h) did not result in higher levels of the mutant protein as determined by Western blot analysis (data not shown). Therefore, for all subsequent studies, expression of mutant EhRabA was induced by the addition of tetracycline 24 h prior to experimentation.

In a previous study we demonstrated that EhRabA localized to small vesicles in the cytoplasm or in extending membranes of motile cells (48). IF staining using antibodies specific for EhRabA or for the HA epitope demonstrated that EhRabAQ84L also localized to both small and large cytoplasmic puncta (a), which do not colocalize with nonspecific HA-staining (b and c). In mutant cells, EhRabA localizes to small and large cytoplasmic puncta. A fraction of EhRabA-positive puncta colocalize with the HA epitope (f and g). Bars, 10 μm.

Expression of GTP-bound EhRabA reduces phagocytosis. Previously, EhRabA was shown to localize to small vesicles in the cytoplasm or in extending membranes of motile cells (48). Since extension of pseudopodia is required for some forms of phagocytosis, and since phagocytosis represents an important virulence function of *E. histolytica*, we measured uptake of hRBCs in mutants expressing EhRabAQ84L. Wild-type and mutant trophozoites were exposed to hRBCs for 10 min, after which uptake of heme was quantified spectrophotometrically. Expression of EhRabAQ84L resulted in a statistically significant reduction in erythropagocytosis (Fig. 2A).

To gain further insight into phagosomal trafficking in the engineered cell line, we exposed wild-type and mutant tropho-

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**FIG. 1.** Expression of endogenous and mutant EhRabA in cell lysates from wild-type cells or transformed cells before (uninduced mutant) and after (induced mutant) induction with tetracycline for 24 h. (A) Cell lysates from wild-type, uninduced, and induced mutant cell lines were resolved by SDS-PAGE and protein was detected by Western blotting using antibodies specific for EhRabA or to the HA epitope. (B) IF microscopy of wild-type (a to c) and mutant (e to g) cells using anti-EhRabA and anti-HA antibodies. Colocalized antigens are shown in yellow (c and g), and corresponding differential interference contrast microscopic images are provided (d and h). In wild-type cells, EhRabA localizes to small cytoplasmic puncta (a), which do not colocalize with nonspecific HA-staining (b and c). In mutant cells, EhRabA localizes to small and large cytoplasmic puncta. A fraction of EhRabA-positive puncta colocalize with the HA epitope (f and g). Bars, 10 μm.
zoites to hRBCs for time periods of up to 40 min. At regular intervals, samples of trophozoites were taken and the level of internalized heme was quantified. Although the cell line expressing EhRabAQ84L internalized less heme overall, both wild-type and mutant cells exhibited an increase in heme accumulation up to 30 min, after which steady-state kinetics were apparent (Fig. 2B). That both cell lines displayed similar kinetics suggests that intracellular trafficking or maturation of phagosomes may not be altered in the mutant. Rather, the reduction in hRBC uptake by the engineered cell line may be the result of a defect in an early stage(s) of erythrophagocytosis, such as adhesion to the target.

Expression of GTP-bound EhRabA alters galactose-sensitive adhesion to hRBCs. To specifically examine early erythrophagosomal events in the mutants, we measured their ability to adhere to erythrocytes by using a standard rosette assay (31). Mutant cells were less efficient at binding hRBCs than wild-type cells (Fig. 3A), suggesting that this early erythrophagosomal event was inhibited by expression of EhRabAQ84L. Since E. histolytica trophozoites rely, in part, on the Gal/GalNAc lectin for adhesion to hRBCs (31), we also determined whether galactose or mannose (negative control) could inhibit the interaction between mutant cells and hRBCs. In wild-type cells, galactose was capable of reducing adhesion to hRBCs by approximately 47%, whereas in mutant cells, the same concentration of galactose inhibited adhesion to hRBCs by 32% (Fig. 3B). Reduced sensitivity of the mutant to galactose inhibition suggests decreased functionality for the Gal/GalNAc lectin in these cells. Mannose was incapable of inhibiting parasite-hRBC interactions for either cell line (Fig. 3B), indicating that the observed inhibition of adhesion by galactose was authentic.
Expression of GTP-bound EhRabA alters the subcellular localization but not levels of subunits of the Gal/GalNAc lectin. The finding that cells expressing EhRabAQ84L were less sensitive to galactose inhibition of adhesion provided the impetus to examine the subcellular localization and the level of Gal/GalNAc lectin subunits in these cells. IF microscopy demonstrated that, in mutant cells, Hgl was minimally localized to the plasma membrane but was found to be enriched in large tubular intracellular compartments (Fig. 4A, panel c). This differed from the pattern observed in wild-type cells, where Hgl was localized mainly to the cell periphery and to puncta throughout the cytosol (Fig. 4A, panel a). Since Hgl possesses the ligand binding site, the altered localization of this subunit in the mutants may lead to their reduced ability to bind hRBCs.

We also examined the subcellular localizations of the other subunits of the Gal/GalNAc lectin. In the mutant cells, Lgl exhibited a similar localization to that of Hgl, concentrating in an intracellular tubular network (Fig. 4A, panel k). Lgl is covalently attached to Hgl through disulfide bridges (14); therefore, it was not surprising to observe an altered localization of Lgl. Interestingly, the distribution of Igl in mutant cells (Fig. 4A, panel g) did not differ from that in wild-type cells (Fig. 4A, panel e). In both cell types, Igl was localized to a significant number of cytoplasmic puncta. Together, these data suggest that EhRabA may directly or indirectly regulate the

FIG. 3. Mutant cells exhibit reduced adhesion to hRBCs. The ability of wild-type and mutant cells to adhere to hRBCs was determined by a standard rosette assay in the absence (control) and presence of galactose or mannose. (A) Mutant cells were less efficient at binding hRBCs compared to wild-type cells. **, P < 0.01. (B) While incubation in the presence of galactose significantly reduced adhesion of both cell lines to hRBC targets, galactose was less efficient at inhibiting adhesion of mutant cells compared to wild-type cells. ***, P < 0.001; **, P < 0.01. Mannose did not significantly (i.e., nonsignificantly [NS]) reduce adhesion of either cell line to host cell targets, indicating that galactose-mediated inhibition of adhesion was authentic. For both panels A and B, the data represent the means ± SD of five trials and are reported as a percentage of adhesion of control cells to hRBCs, which was arbitrarily set to 100%.
trafficking of a subset of vesicles that are also responsible for the localization of Hgl and Lgl, but not of Igl. The data also demonstrate that Igl may be transported by vesicle trafficking routes that differ from that of the Hgl-Lgl heterodimer.

Western blot analysis demonstrated that the levels of at least two Gal/GalNAc subunits, Hgl and Igl, were not altered in the mutant compared to wild-type parental cells (Fig. 4B). Therefore, the differences observed between the mutant and wild-type cell lines were not likely the result of changes in protein levels. Given that antibodies recognize multiple isoforms of Lgl in trophozoites (25), the level of this subunit was not assessed by Western blot analysis.

**Hgl-positive compartments colocalize with markers of the ER but not of the Golgi apparatus.** The novel Hgl-containing organelle observed in the mutants was reminiscent of recently described tubular compartments of the ER of *E. histolytica* (41). Therefore, we counterstained these organelles with an antibody specific for *E. histolytica* calreticulin (9), an ER-resident protein. Since the ER often assumes a perinuclear organization, we simultaneously stained cells with propidium iodide, a nuclear stain. Minimal colocalization of Hgl (Fig. 5A panels a and b) or Lgl (Fig. 5B, panels a and b) and calreticulin was observed in wild-type cells, whereas significant colocalization of the these antigens was observed in mutants (Fig. 5A, panels e and f and B, panels e and f). Significant colocalization of Hgl and Lgl with calreticulin suggests that both subunits are mislocalized to the ER or the same novel compartment derived from the ER. We also observed that the novel organelle was perinuclear in its localization (Fig. 5A, panel g), further supporting the ER origin of this organelle.

To confirm the ER origin of the novel compartment we also used an antibody specific for the ER retention signal, KDEL. This antibody recognizes an array of proteins on Western blots of mammalian cell lysates; however, it prominently stains two mammalian ER-resident proteins, GRP78 (78 kDa) and GRP94 (94 kDa) (27). Likewise, Western blot analysis of trophozoite lysates from untransformed and mutant *E. histolytica* revealed an array of proteins (data not shown) with two prominent bands of 78 and 94 kDa (Fig. 6A). This suggested that this antibody might be used to authentically label the ER in *E. histolytica* trophozoites. While the Western blot assay was performed only to assess the utility of this antibody, it revealed
that the 94-kDa species was more abundant in the mutant than in the wild-type cell line. This may indicate an expansion of the ER as a result of EhRabAQ84L expression. IF microscopy with the anti-KDEL antibody revealed minimal colocalization of Hgl with the KDEL marker in wild-type cells (Fig. 6B, panels a to c) but near-complete colocalization of Hgl with the ER marker in the mutant (Fig. 6B, panels e to g). This provides support for the ER origin of this organelle.

To determine if the novel compartments arose by fusion events between the ER and Golgi complex, we counterstained wild-type and mutant cells with an antibody specific for β-COP. This antibody was previously shown to label Golgi complex-like elements in this organism (46). We observed minimal colocalization of Hgl and the Golgi antigen in the mutant cell line, suggesting that little mixing of these organelles occurred as a result of EhRabAQ84L expression (Fig. 6C).

Expression of GTP-bound EhRabA alters the subcellular localization of a secreted cysteine protease, EhCP2. We have shown that expression of EhRabAQ84L results in aberrant trafficking of at least one transmembrane protein, Hgl. To test the effect of mutant EhRabA expression on another class of secretory proteins, namely, soluble proteins, we looked at subcellular localization of a secreted cysteine protease, EhCP2, by IF microscopy. Like Hgl and Lgl, EhCP2 was mislocalized to large tubular compartments in the mutant cell line (Fig. 7). This suggests that the molecular machinery governing the trafficking of Hgl and Lgl may be shared by this cysteine protease and that EhRabA may directly or indirectly regulate this pathway.

Expression of GTP-bound EhRabA does not alter the localization of SREHP, a serine-rich protein of *E. histolytica*. It has been proposed that a putatively lipid-anchored cell surface serine-rich *E. histolytica* protein, SREHP, also participates in the uptake of hRBCs by phagocytosis (42). Therefore, we examined the cellular levels and localization of SREHP in mutant and wild-type cells. Western blot (Fig. 8A) and IF (Fig. 8B) analyses indicated that both the levels and localization of SREHP remained unchanged in the mutants. In both wild-type and mutant cells SREHP localized to the plasma membrane and to an intracellular network (Fig. 8B). Furthermore, like

FIG. 5. The heavy (A) and light (B) subunits of the Gal/GalNac lectin colocalize with calreticulin (ER marker). IF microscopy results are shown for wild-type (A, panels a to c, and B, panels a to c) and mutant cells (A, panels e to g, and B, panels e to g) using anti-Hgl (A) or anti-Lgl (B) and anti-calreticulin antibodies. The tubular compartment is perinuclear as determined by propidium iodide staining (A, panel g). Colocalized antigens are shown in light blue (A, panels c and g) or yellow (B, panels c and g), and corresponding merged differential interference contrast microscopic images (A, panels d and h, and B, panels d and h) are provided. Bars, 10 μm.
Igl, SREHP did not colocalize with Hgl in the novel compartment (Fig. 8B).

It has been reported that *E. histolytica* trophozoites preferentially phagocytose apoptotic host cells (10) and that SREHP may serve as a receptor for such targets (42). Since the localization of SREHP was unchanged in the EhRabAQ84L-expressing cells, we predicted that these mutants would take up apoptotic targets as efficiently as untransformed cells. Therefore, we tested the ability of the mutants to internalize erythrocytes that were pretreated with calcium. Such treatment induces the exposure of phosphatidylserine on the outer membrane of the hRBCs, which is a hallmark of apoptosis (3). As predicted, phagocytosis of calcium-treated cells was not inhibited by expression of EhRabAQ84L (Fig. 9), suggesting that the mutant cells could efficiently adhere to apoptotic cells. Fixation of mam-
malian cells with paraformaldehyde can also induce exposure of phosphatidylserine on outer membrane leaflets (49), recapitulating the apoptotic phenotype. Therefore, we also tested the ability of mutants and wild-type cells to adhere to fixed epithelial monolayers. Adhesion of mutant cells to paraformaldehyde-fixed monolayers of CHO cells was comparable to that of wild-type parental cells (data not shown), further supporting the idea that expression of EhRabAQ84L does

FIG. 7. Subcellular localization of the Hgl and EhCP2 subunits in wild-type cells and cells expressing EhRabAQ84L. IF microscopy of wild-type (a to c) and mutant (e to g) cells by using anti-Hgl or anti-EhCP2 antibodies. Colocalized antigens are shown in yellow (c and g), and corresponding differential interference contrast (DIC) (d and h) microscopic images are provided. In wild-type cells, Hgl and EhCP2 are localized to the plasma membrane and cytoplasmic puncta. Mutant cells exhibit near-complete colocalization in intracellular compartments, some of which are perinuclear. Bars, 10 μm.

FIG. 8. Subcellular localization of SREHP is unchanged in cells expressing EhRabAQ84L. (A) Western blot analysis of trophozoite lysates from wild-type (WT) and mutant (AQ84L) cells using anti-SREHP antibody. Consistent with a previous report (50), the antibody recognizes a series of proteins ranging in size from 47 to 52 kDa. There were no apparent changes in the level of SREHP in the mutant cell line. (B) IF microscopy of wild-type (a to c) and mutant (e to g) cells using anti-Hgl (a and e) or anti-SREHP (b and f) antibodies. Colocalized antigens are shown in yellow (c and g), and corresponding differential interference contrast microscopic images (d and h) are provided. In wild-type cells, both Hgl and SREHP are enriched at the plasma membrane and localize to intracellular puncta. In mutant cells, SREHP appears to retain its plasma membrane enrichment, whereas Hgl localizes to intracellular and, in some cases, perinuclear compartments. Bars, 10 μm.
not inhibit trafficking of cell surface receptors specific for apoptotic host cells.

Treatment with BFA phenocopies expression of EhRabAQ84L. BFA is a fungal metabolite that can disrupt secretory pathway vesicle trafficking (11). For example, in mammalian cells, this reagent can block egress of vesicles from the ER (21) and can alter the morphology of a population of Golgi-specific vesicles and inhibit the trafficking of a subset of secretory proteins (15). Interestingly, in *E. histolytica*, BFA also induces the formation of tubular structures that colocalize with an ER marker, protein disulfide isomerase (15). These structures are similar to those observed in the EhRabAQ84L-expressing cell line; therefore, we explored whether EhRabA might regulate vesicle trafficking pathways that are also sensitive to BFA treatment. We examined the localization of several cell surface and secreted proteins after BFA treatment of wild-type cells. Interestingly, exposure to BFA recapitulated the EhRabAQ84L-induced phenotype in that Hgl, Lgl, and EhCP2 were abnormally localized to a large intracellular tubular network (Fig. 10A), whereas the distributions of Igl and SREHP were unchanged (Fig. 10B). This suggests that EhRabA participates in vesicle trafficking pathways that are also sensitive to BFA.

**DISCUSSION**

We have demonstrated that overexpression of a GTP-bound mutant of EhRabA results in decreased phagocytosis of nonsequestered red blood cells but not of sequestered red blood cells. Overexpression of the mutant protein also induced the appearance of large tubular and perinuclear organelles that could be stained with ER-specific marker antibodies, suggesting that this novel compartment was of ER origin. The appearance of the tubular organelles in the mutant cells supports the recent finding that *E. histolytica* trophozoites possess a traditional continuous ER (41).

The nature of the phagocytic defect in the mutants was likely related to their inability to adhere efficiently to the targets. This may have been caused by mislocalization of two subunits of the Gal/GalNAc lectin, Hgl and Lgl, to the novel ER-like compartment. In these mutants, at least one other secretory protein, EhCP2, was also mislocalized to the ER-like organelles; however, the cellular localization of two other cell surface proteins, Igl and SREHP, was unchanged. Interestingly, the EhRabAQ84L phenotype was similar to that induced by treatment with BFA. Together, the data suggest that EhRabA directly or indirectly influences the morphological integrity of the ER and the trafficking of a subset of secretory proteins in *E. histolytica*. This report is the first to demonstrate that distinct secretory pathways may exist in *E. histolytica*.

Point mutations in the GTPase domain of Rabs, which result in reduced GTPase activity, can lead to either gain-of-function (18) or loss-of-function phenotypes (35). Constitutively GTP-bound Rabs are generally considered to be hyperactivated and, therefore, have acquired a gain-of-function phenotype. However, in cases where a terminal vesicle fusion event relies on GTP hydrolysis, overexpression of a hydrolase-dead mutant may, in fact, result in a loss-of-function phenotype. It is not known if the engineered glutamine-to-leucine point mutation described here results in gain of function or loss of function for EhRabA.

The localization of EhRabA to membrane protrusions (48) suggests a role in late secretion. If overexpression of the hydrolase-dead version of EhRabA blocks this process (loss of function), it may lead to accumulation of cargo in secretory organelles or an imbalance between anterograde and retrograde trafficking in favor of retrograde movement of secretory proteins toward the ER. Localization of EhRabA near the plasma membrane (48) may also suggest a role in regulating retrograde vesicle trafficking. Expression of the GTP-bound form of this Rab might accelerate retrograde vesicle movement (gain of function) toward the ER. In all of these cases, secretory proteins would accumulate in the ER, which may lead to an expansion of the ER (32).

It is not surprising that alterations in Rab function can lead to changes in organelar morphology. Rabs influence the morphology of organelles by regulating intraorganellar or interorganellar fusion events and/or by regulating the entry in or exit of transport vesicles. Since EhRabA does not localize to the ER (48) it is possible that it is acting in trans through effector proteins to promote intraorganellar or interorganellar fusion of ER-derived compartments. In support of this, overexpression of GTP-bound Rab5 in *Caenorhabditis elegans* results in the appearance of large fused ER-derived compartments reminiscent of those observed in this study (1). Like EhRabA, C. *elegans* Rab5 does not localize to the ER but is thought to promote intraorganellar fusion of the ER through effector proteins.

A striking result of this study was that BFA treatment could recapitulate the novel organellar phenotype associated with overexpression of EhRabAQ84L. BFA is known to block ER-to-Golgi trafficking (21), which could alter the balance between anterograde and retrograde vesicle traffic. The similar phenotype in the EhRabAQ84L-expressing cells may support a model in which trafficking to and from the ER is perturbed. However, there is not complete phenotypic recapitulation by BFA treatment, since others have shown that BFA-treated *E. histolytica* cells do not exhibit reduced phagocytosis of at least one phagocytic target, green fluorescent protein-expressing *E.
coli (15). Therefore, it is likely that the resultant phenotype of the mutant occurred through a different process than that operating in BFA-treated cells. Importantly, BFA induces alterations in the secretory pathway by impairing the association and dissociation of coat proteins on cellular membranes (11), whereas Rabs are not thought to participate in the assembly or disassembly of such protein coats.

It is interesting that the localizations of Igl and SREHP were...
that such GPI-specific secretory pathways similarly exist in unaffected by EhRabAQ84L expression. While Igl is proposed
between these two possibilities will provide significant insight into and other secretory proteins in mutant cell lines is a direct
this study) forms of EhRabA leads to mislocalization Hgl. It
expansion and dysfunction.
Expression of both the GDP-bound (47) and GTP-bound (this study) forms of EhRabA leads to mislocalization Hgl. It
did not exhibit aberrant localization, we do not believe that all secretory trafficking was perturbed in the mutant or that the cells were
under general metabolic stress, which might lead to organelle expansion and dysfunction.

ACKNOWLEDGMENTS
We thank W. A. Petri, Jr. (University of Virginia, Charlottesville), N. Guillaín (Unité de Biologie Cellulaire du Parasitisme, Institut Pasteur, Paris, France), and S. L. Stanley, Jr. (Washington University School of Medicine, St. Louis, MO) for antibody reagents. We thank the members of the Temesvari laboratory for critical reading of the manuscript and helpful discussions.
The project described was supported by grant no. R01AI046414 from the National Institute of Allergy and Infectious Diseases. This material is based upon work supported by CSREES/USDA, under project number SC-1700312 (Technical Contribution No. 5570 of the Clemson University Experiment Station).
The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institute of Allergy and Infectious Diseases, the National Institutes of Health, or the U.S. Department of Agriculture.

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