

# Septins guide microtubule protrusions induced by actin-depolymerizing toxins like *Clostridium difficile* transferase (CDT)

Thilo Nölke<sup>a,1</sup>, Carsten Schwan<sup>a,1</sup>, Friederike Lehmann<sup>a,b,c</sup>, Kristine Østevold<sup>a,d</sup>, Olivier Pertz<sup>e</sup>, and Klaus Aktories<sup>a,f,g,2</sup>

<sup>a</sup>Institute of Experimental and Clinical Pharmacology and Toxicology, University of Freiburg, 79104 Freiburg, Germany; <sup>b</sup>Spemann Graduate School of Biology and Medicine (SGBM), University of Freiburg, 79104 Freiburg, Germany; <sup>c</sup>Faculty of Chemistry and Pharmacy, University of Freiburg, 79104 Freiburg, Germany; <sup>d</sup>Faculty of Biology, University of Freiburg, 79104 Freiburg, Germany; <sup>e</sup>Institute of Cell Biology, University of Bern, 3012 Bern, Switzerland; <sup>f</sup>Centre for Biological Signalling Studies (BIOS), University of Freiburg, 79104 Freiburg, Germany; and <sup>g</sup>Freiburg Institute of Advanced Studies (FRIAS), University of Freiburg, 79104 Freiburg, Germany

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**Hypervirulent *Clostridium difficile* strains, which are associated with increased morbidity and mortality, produce the actin-ADP ribosylating toxin *Clostridium difficile* transferase (CDT). CDT depolymerizes actin, causes formation of microtubule-based protrusions, and increases pathogen adherence. Here, we show that septins (SEPT) are essential for CDT-induced protrusion formation. SEPT2, -6, -7, and -9 accumulate at predetermined protrusion sites and form collar-like structures at the base of protrusions. The septin inhibitor forchlorfenuron or knockdown of septins inhibits protrusion formation. At protrusion sites, septins colocalize with the GTPase Cdc42 (cell division control protein 42) and its effector Borg (binder of Rho GTPases), which act as up-stream regulators of septin polymerization. Precipitation and surface plasmon resonance studies revealed high-affinity binding of septins to the microtubule plus-end tracking protein EB1, thereby guiding incoming microtubules. The data suggest that CDT usurps conserved regulatory principles involved in microtubule-membrane interaction, depending on septins, Cdc42, Borgs, and restructuring of the actin cytoskeleton.**

microtubules | septins | actin | bacterial toxin | *Clostridium difficile*

The pathogen *Clostridium difficile* is the causative agent of pseudomembranous colitis and antibiotic-associated diarrhea (1). Recently emerging hypervirulent strains (e.g., NAP1/027) have been associated with severe courses of *C. difficile* infections with increased morbidity and mortality (2). These strains have deletions in regulatory genes, resulting in overexpression of the Rho-inactivating glycosylating toxins A and B (3). Additionally, they produce the binary toxin *C. difficile* transferase (CDT) (2, 4). CDT ADP ribosylates actin at arginine-177, thereby inhibiting actin polymerization (5–7).

CDT-induced depolymerization of F-actin induces formation of long microtubule-based protrusions, which form a meshwork on the surface of intestinal host cells (8). Clostridia are enwrapped by the protrusions, resulting in increased pathogen adherence. In addition to microtubules, the protrusions contain membrane tubules of endoplasmic reticulum and allow traffic of Rab5- and Rab11-positive vesicles (9). Moreover, CDT induces rerouting of fibronectin-containing vesicles from the basolateral to the apical side of intestinal epithelial cells. Here, fibronectin, a binding protein for *Clostridium difficile*, is released at microtubule-based protrusions, suggesting that the cell-surface extensions participate in host-pathogen interaction (9).

Protrusion formation occurs in membrane areas where CDT causes partial depolymerization of cortical actin (8). The toxin induces the redistribution of microtubule capture proteins such as Clasp2 and ACF7, which are normally associated with cortical actin, from the membrane to the cytosolic fraction, suggesting that CDT alters the functions of the capture proteins. Also, the cholesterol content of membranes was found to be crucial for protrusion formation (10). So far, however, the molecular mechanisms involved in formation of CDT-induced cell protrusions are largely unknown.

We asked whether septins (SEPT), which interact with both F-actin and microtubules, are involved in toxin-induced restructuring of the cytoskeleton and formation of protrusions. Septins are conserved GTP-binding proteins, which form hetero-oligomers and polymers, and are considered as the fourth component of the cytoskeleton in addition to microfilaments, microtubules, and intermediate filaments (11–13). Septins are involved in numerous functions, including cytokinesis, ciliogenesis, neurite development, and exocytosis (14–17). They have scaffold functions, act as membrane diffusion barriers, and are involved in host-pathogen interactions. Septins locate to cell membranes, where they control the stiffness/rigidity and the shape of membranes.

Here, we report that formation of CDT-induced protrusions depends on septins, which are located at the base of the protrusions. Septins are involved in early steps of protrusion formation and guide microtubules from the cytosol into emerging protrusions at the cell membrane, a process that probably involves the plus-end tracking protein EB1 (end-binding protein 1). Cell division control protein 42 (Cdc42) and its effector Borg (binder of Rho GTPases), which are up-stream regulators of septins, control CDT-induced protrusion formation (18). The data indicate that CDT exploits a highly conserved regulatory pathway of microtubule-membrane interaction.

## Results

**Septin Translocation After CDT Treatment.** In human colon adenocarcinoma Caco-2 cells, CDT induced the formation of

### Significance

The bacterial toxin *Clostridium difficile* transferase (CDT) depolymerizes actin and forms microtubule-based cell protrusions suggested to be involved in cell adherence of bacteria. Here we report that GTP-binding septin proteins initiate protrusion formation. Accumulation of septins at sites of protrusion initiation depends on the small GTPase Cdc42 (cell division control protein 42) and its effector Borg (binder of Rho GTPases). CDT-induced septin accumulations at the cell membrane guide microtubules to form cell protrusions. Septins directly interact with the microtubule plus-end tracking protein EB1. Thus, the toxin exploits septin-dependent regulation of microtubules to form cell protrusions.

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The authors declare no conflict of interest.

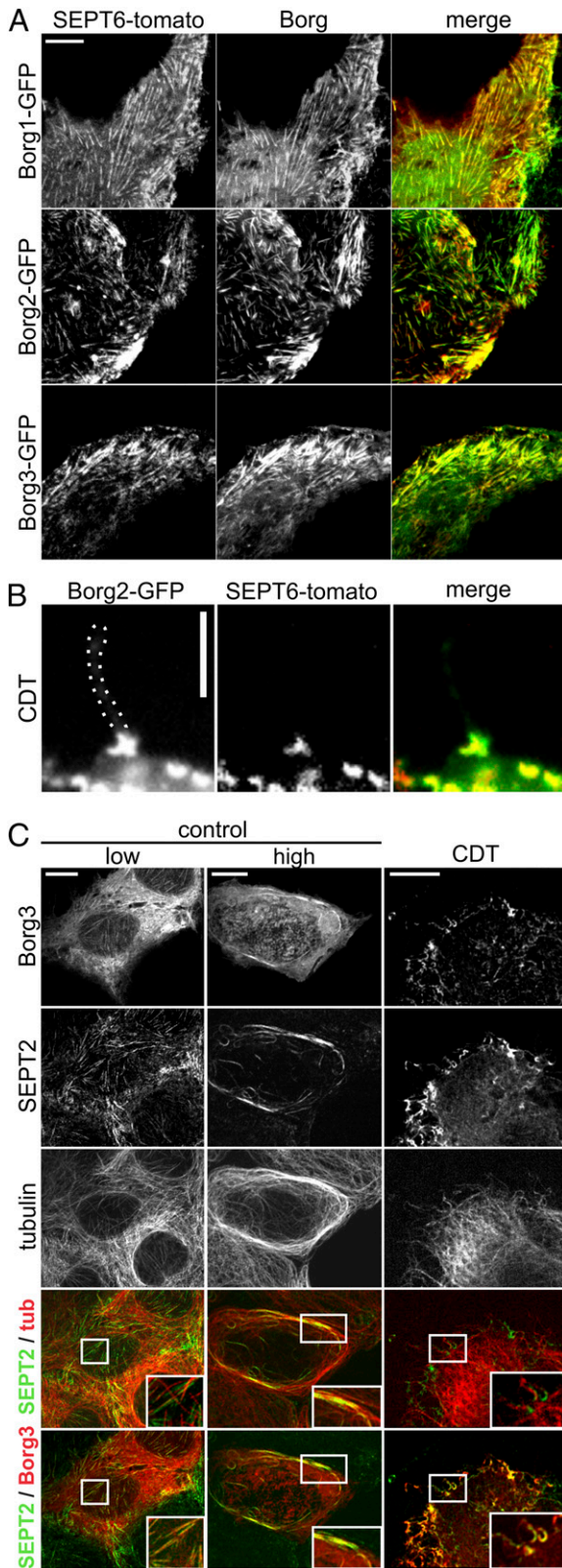
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<sup>1</sup>T.N. and C.S. contributed equally to this work.

<sup>2</sup>To whom correspondence should be addressed. Email: klaus.aktories@pharmakol.uni-freiburg.de.

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**Fig. 2.** Borg proteins colocalize with septins. (A) Caco-2 cells were transfected with SEPT6-tomato and Borg1-, Borg2-, and Borg3-GFP. Cells were treated with CDT (200 ng/mL CDTa and 400 ng/mL CDTb) for 90 min. Borg proteins colocalize with septins. (B) Caco-2 cells were transfected with SEPT6-tomato and Borg2-GFP. Cells were treated as in A. Borg2 colocalizes with SEPT6 at the base of protrusions. (Scale bar, 3  $\mu\text{m}$ .) (C) Caco-2 cells were transfected with

proteins Borg 1, -2, and -3 in filamentous structures (Fig. 2A). After intoxication with CDT, septins formed chevron-like structures at the membrane, and Borgs were localized at the base of toxin-induced protrusions as a part of these structures (Fig. 2B and C). Control cells with a strong Borg expression showed an increased microtubule density and bundling at the cell cortex. These microtubule bundles strongly associated with thick septin filaments (Fig. 2C).

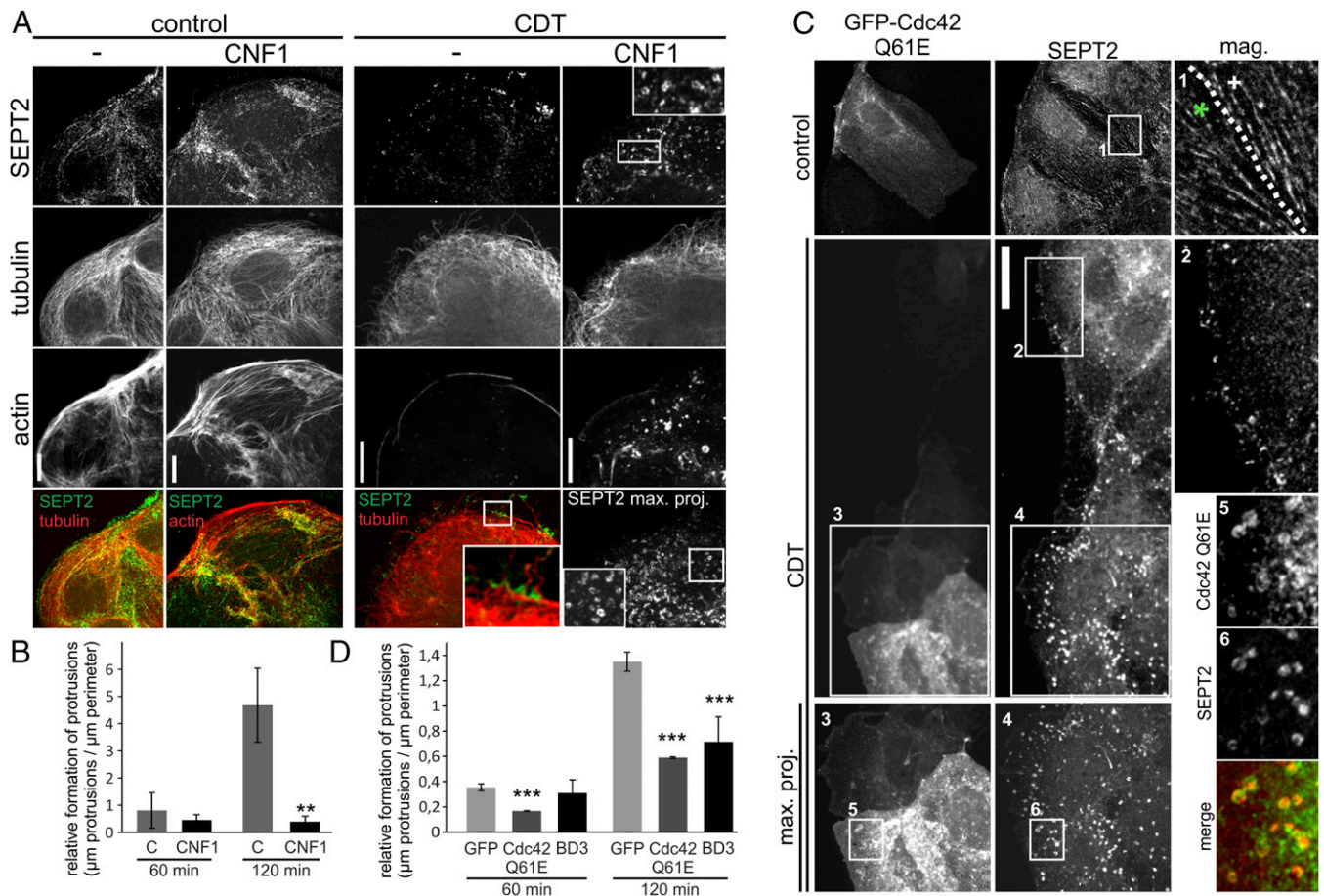
Next, we analyzed the effects of the Cdc42-activating toxin cytotoxic necrotizing factor (CNF1) (22, 23). Whereas CNF1 alone affected the actin cytoskeleton and increased formation of stress fibers, the toxin caused minor or no changes of the septin cytoskeleton. In CNF1-pretreated cells, CDT strongly increased formation of septin rings all over the cell. At the cortex, the chevron-like septin accumulations were changed to an undirected formation of septin rings (Fig. 3A). Concomitantly, CNF1 pretreatment strongly reduced protrusion formation (Fig. 3B), indicating that accumulation and/or functions of septins are downstream of Cdc42 and its effector protein Borg.

Similarly, expression of dominant active Cdc42 Q61E had no effect on septins after 24 h of transfection. However, after CDT-induced partial destruction of the actin cytoskeleton, the overexpression of Cdc42 Q61E had a similar effect as CNF1 with strongly increased formation of septin rings (Fig. 3C). The septin rings colocalized with expressed GFP-Cdc42 Q61E, suggesting that the GTPase Cdc42 is causally linked to ring formation. Moreover, dominant active Cdc42 inhibited the formation of protrusions (Fig. 3D). In line with these findings, we observed that expression of the BD3 domain of Borg2, which was shown to interact with septins, inhibited protrusion formation (Fig. 3D) (21). To analyze the spatiotemporal regulation of Cdc42, cells were transfected with a FRET sensor (24, 25). After 24 h, transfected cells were treated with CDT and the activation of Cdc42 was monitored 1 h after toxin addition. The base of protrusions revealed increased Cdc42 activity in ratio imaging (Fig. 4A and B). In contrast, no increased RhoA activation was observed.

**Septins Guide the Way of Microtubules.** Fig. S8A shows plus-end tracks of polymerizing microtubules, labeled by EB1, which moved along septin filaments. In control cells, microtubules polymerize along septin tracks. Upon CDT-induced actin depolymerization, septins were located at actin-free sites of the membrane, forming unstructured clusters (Fig. 5A and Fig. S8B). These clusters were predetermined sites of protrusion formation. During initial growth and elongation of microtubule-based protrusion, the septin accumulations mature to typical arch- or chevron-like structures (Fig. 1B). After formation of a stable septin base at the protrusion-cell membrane interface, further polymerizing microtubules follow the leading microtubule in the same track (Fig. S8C). Also microtubules that do not polymerize in the same track are redirected at the protrusion base into the protrusion (Fig. 5B and C).

**Septins Interact with the Microtubule Plus-End.** Septins are known to bind microtubules (20). However, the structures involved are largely enigmatic. We asked whether the plus-end tracking protein-1 (EB1), which is a master regulator of protein interactions at the plus-ends of microtubules (21–25), plays a role in septin-microtubule communication. In fact, GST-pull-down experiments with EB1-loaded beads revealed a direct interaction of septins with EB1. Septins were precipitated as purified His-tagged proteins (Fig. S8D and E), as endogenous septins from Caco-2 cell lysates (Fig. 5D–F) or from EB1-GFP-transfected HeLa cells (Fig. 2G and H). To quantify the interaction between EB1 and septin, we used surface plasmon resonance spectroscopy. These studies revealed an EB1-septin interaction affinity of a  $K_d$  of  $80 \pm 12$  nM for SEPT2,

Borg3-GFP and treated with CDT as in A. Cells were stained for SEPT2 and tubulin. (Insets) Septin colocalization with microtubules and Borg in control cells. In CDT-treated cells, insets show chevron-like structures at the protrusion base and colocalization of septins and Borg. (Scale bars, 10  $\mu\text{m}$ .)



**Fig. 3.** Influence of Cdc42 on septin accumulation. (A) Caco-2 cells were pretreated with 150 ng/mL CNF1 for 3 h to activate Rho-GTPases. Cells were subsequently treated with CDT (200 ng/mL CDTa and 400 ng/mL CDTb) for 1.5 h. Cells were stained for SEPT2,  $\alpha$ -tubulin, and actin. To visualize all septin rings in CNF1 plus CDT-treated cells, confocal stacks were projected into one plane. *Insets* show chevron-like structures at the base of protrusions in CDT-treated cells. In CNF1- and CDT-treated cells, *Insets* show septin-ring formation. (Scale bar, 10  $\mu\text{m}$ .) (B) Caco-2 cells were treated as in A. The relative formation of protrusions (" $\mu\text{m protrusions} / \mu\text{m cell perimeter}$ ") was quantified after 60 and 120 min of CDT intoxication. Data are  $\pm$  SEM, fields of view are  $\geq 7$ , and  $n = 3$ . (C) Caco-2 cells were transfected with dominant active Cdc42 (Q61E) for 24 h. Cells were treated with CDT as in A. *Inset* in "control" shows septin filaments in transfected (green asterisk) and nontransfected (white cross) cells. To visualize all septin rings in transfected CDT-treated cells, confocal stacks were projected into one plane. *Insets* show cortical septin structures in CDT-treated cells. In transfected CDT-treated cells, *Insets* show septin ring formation. Septin rings colocalize with dominant active Cdc42. (Scale bar, 10  $\mu\text{m}$ .) (D) Caco-2 cells were treated as in C. Control cells were transfected with GFP only. Cells were also transfected with the BD3 domain of Borg2 fused to GFP. Relative formation of protrusions was quantified after 60 and 120 min. Data are  $\pm$  SEM, transfected cells are  $\geq 13$ , and  $n = 3$ .

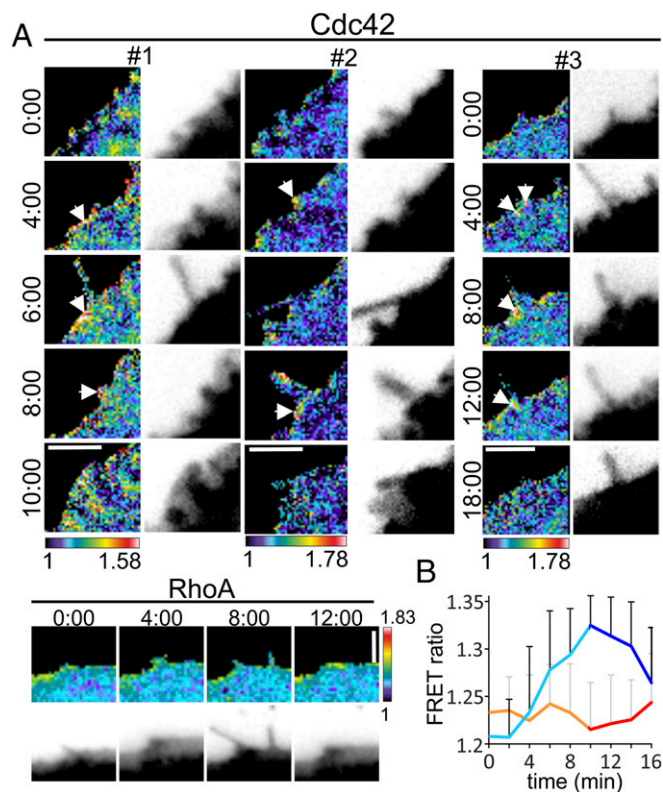
$19 \pm 11$  nM for SEPT6, and  $26 \pm 4$  nM for SEPT7 (Fig. S8 F–H). Thus, septins interact directly and with high affinity with EB1.

## Discussion

Here, we show that septins accumulate at the membrane after cells were intoxicated with the actin-ADP-ribosylating toxin CDT. Septins are well known to interact with anionic phospholipids (e.g., phosphatidylinositol 4,5-bisphosphate and phosphatidylinositol 3,4,5-trisphosphate) (26, 27). In our studies, the sites of septin accumulation are characterized by spots of depolymerized cortical actin, which may allow interaction with membrane phospholipids. During protrusion formation the septin accumulations matured to chevron-like or circular structures at the base of protrusions. Interestingly, a very similar angle-like structure of septins occurs at the base of developing neurite branches (28). However, in neurites, formation of actin filaments are crucial for collateral branching, which is followed by accumulation of microtubules. Although in neurites SEPT6 increases cortactin recruitment, which is essential to trigger formation of actin-based axonal filopodia (28), we did not detect any role of cortactin in CDT-induced protrusion formation (Fig. S9). In our study, CDT causes local depolymerization of actin, which is a prerequisite for septin-dependent, microtubule-based protrusion formation. In fact, several findings indicate that

septins are essential for toxin-induced protrusion formation. While overexpression of septins increased protrusion formation, septin knockdown and treatment with FCF, an inhibitor of septin turnover (20), blocked formation of protrusions. Not only the number but also the length of protrusions was affected by CDT treatment, suggesting that septins are involved not only in initiation but also in growth, dynamics, and stabilization of microtubule-based protrusions. Accordingly, we also observed septins along the protrusion shaft. Although knockdown of SEPT6 and SEPT7 clearly inhibited protrusion formation, knockdown of SEPT2 had no effect, although it was part of the septin structure at the base of protrusions. We assume that functional redundancy of other septins (e.g., SEPT1 and SEPT5) from the SEPT2 group is responsible for protrusion formation after SEPT2 knockdown.

Studies from yeast and mammalian cells indicate that the small GTPase Cdc42 plays an essential role in septin regulation (21, 29). Accordingly, we observed locally increased activity of Cdc42 in FRET experiments before and during protrusion formation. However, persistent activation of Cdc42 was inhibitory. When dominant active Cdc42 or the Rho protein activator CNF1 was used, protrusion formation was strongly inhibited. These findings suggest that the free cycling of Cdc42 from the inactive to the active state is necessary to fulfill septin regulatory functions. CNF1 and also



**Fig. 4.** Cdc42 activation in CDT-induced protrusion formation. (A) Caco-2 cells were transfected with a FRET sensor for Cdc42 or RhoA. Cells were intoxicated with CDT (200 ng/mL CDTa and 400 ng/mL CDTb) for 40–80 min. Sites of protrusion formation have increased Cdc42 activity (arrowheads). Rho activity is not increased at these sites. The nomenclature “#1,” “#2,” and “#3” represents three protrusion formation events with the Cdc42 FRET sensor. Time is indicated on the left (Cdc42) or above (RhoA). Donor images are on the right (Cdc42) or below (RhoA) with inverted contrast. (Scale bars, 5  $\mu$ m.) (B) Quantification of FRET intensity of Cdc42 and RhoA constructs during protrusion formation. Cells were treated as in A. When the protrusions appear, the curve turns from orange to red (RhoA) or from turquoise to blue (Cdc42). Data are  $\pm$  SEM,  $n = 8$  for Cdc42, and  $n = 6$  for RhoA.

dominant active Cdc42 caused increased formation of septin rings in the cytosol, whereas the membrane localization of septins with chevron-like structures were decreased. This is in line with a role of cycling Cdc42 in septin membrane recruitment observed in yeast and mammalian cells (30, 31). Moreover, the persistently active form of Cdc42 may favor uncontrolled ring formation of septins that prevents a spatiotemporal controlled polymerization of septins at the site of protrusion formation.

Cdc42 effectors of the Borg family are suggested to play a pivotal role in septin recruitment and organization (21, 32). Five Borg family members (Borg1–5) are known (18). All Borgs bind directly to septins via their BD3 domain (21). We could colocalize Borg1–3 with septin filaments. After CDT treatment Borgs colocalized with the chevron-like structures at the base of protrusions. The regulation of Borgs by Cdc42 is not well understood. Although Borgs bind to active GTP-bound Cdc42 *in vitro* (18), it was shown that the active Cdc42 can displace Borg from septins. Thus, Borg–septin interaction appears to be negatively regulated by Cdc42 (21). Accordingly, we observed that Borgs were displaced from protrusions after CNF treatment of cells.

Bowen et al. reported that septins track microtubule bundles (33). Similarly, we observed tracking of polymerizing cytosolic microtubules along septins in control cells. In CDT-treated cells, where membrane-associated septin clusters had formed, microtubule tips interacted with the chevron-like structures of septins. Moreover, we observed that septins guided microtubules to form projections and forced new directions of microtubule growth. This was evident when

microtubules contacted septins at distal parts of the chevron-like structures. The septin interaction redirected the tip of microtubules into the center of the septin “funnel,” where microtubules continued to grow beyond the edge of the cell to form protrusions.

We show that septins interact with microtubules via EB1, which is a multifunctional interaction partner at the microtubule tip (21–25). Pull-down experiments and surface plasmon resonance spectroscopy, exhibiting  $K_d$  values in the nanomolar range, revealed that septins bind directly to EB1. Previously, we showed that toxin-induced protrusions contain endoplasmic reticulum, which is connected to microtubules via stromal interaction molecule 1 (Stim-1) (9). Stim-1 is involved in Stim/Orai-dependent store-operated calcium entry, and septins are coordinators of Stim/Orai channels (34). It remains to be studied whether Stim-1 is another functional connection between microtubules and septins.

Taken together, here we show that the actin-ADP-ribosylating toxin CDT of *C. difficile* induces septin accumulations at the cell membrane, which are essential for the formation of toxin-induced microtubule-based protrusions. This effect is regulated by Cdc42 and most likely by Borg proteins. Septins guide growing microtubules and determine the site of protrusion formation at the cell membrane by interaction with EB1 at the tips of microtubules. Thus, the bacterial toxin exploits septin-dependent regulatory mechanism of microtubule organization to eventually form a network of cell protrusions and to increase the adherence of the pathogen during *C. difficile* infection.

## Methods

For detailed methods, see *SI Methods, Cell Culture and Transient Transfections; Preparation of Primary Colon Epithelial Cells; Expression and Purification of Proteins; shRNAs; Antibodies, Fluorescent Dyes, and Fluorescent Proteins; Immunostaining; Live-Cell Imaging; FRET Experiments; Pull-Down Assays and Coimmunoprecipitation; Surface Plasmon Resonance; ADP-Ribosylation Assays; and Statistics.*

**Immunostaining.** Cells were washed with PBS, fixed with 4% (wt/vol) formaldehyde in PBS, washed with PBS, permeabilized with 0.15% Triton X-100 in PBS, and blocked. Incubation with the primary antibody was overnight at 4 °C. Cells were washed and incubated with the suitable secondary antibody for 1 h. Cells were washed, dried, and embedded with Mowiol. Cells were analyzed with the microscope mentioned in *SI Methods*. Images were processed with Metamorph software.

**Live-Cell Imaging.** For live-cell imaging, cells were incubated in a chamber with humidified atmosphere at 37 °C on the microscope (*SI Methods, Immunostaining*). Quantification of microtubule protrusion formation was performed according to ref. 8.

**FRET Experiments.** Procedure for FRET measurements is given in the immunostaining section. At each time point, three images were recorded for CFP, FRET, and YFP. Donor (CFP) and FRET images were acquired sequentially with exposure times  $\leq 300$  ms. Processing of ratiometric images was performed with the Biosensor Processing Software 2.1 (Danuser laboratory: [lccb.hms.harvard.edu/software.html](http://lccb.hms.harvard.edu/software.html)). Subsequent image analysis was performed in MetaMorph. Ratio images were color-coded; warm and cold colors represent high and low biosensor activity, respectively.

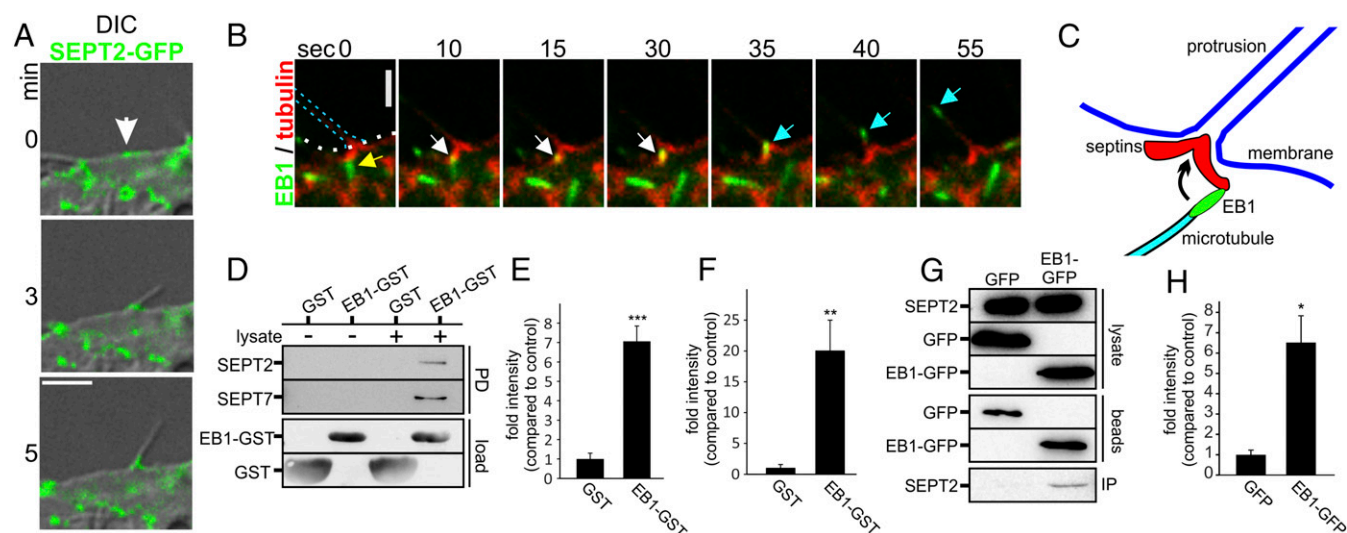
**Pull-Down Assays and Coimmunoprecipitation.** Cells were lysed in buffers supplemented with Complete protease inhibitor (Roche). For pull-down experiments, lysates were incubated with GST or GST-EB1 bound to glutathione Sepharose 4B (GE Healthcare) and subjected to Western blot analysis.

For pull-down of recombinant proteins, GST or GST-EB1 bound to glutathione Sepharose 4B was incubated with purified His-tagged proteins.

For coimmunoprecipitation, lysates were incubated with  $\mu$ Mac5 GFP isolation beads (Miltenyi Biotec). The beads were immobilized to  $\mu$ Columns and after elution, proteins were detected by immunoblot.

Mice were housed and handled in accordance with good animal practice as defined by FELASA ([www.felasa.eu](http://www.felasa.eu)) and the national animal welfare body GV-SOLAS ([www.gv-solas.de](http://www.gv-solas.de)).

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**Fig. 5.** Microtubules are guided along septins. (A) Combined DIC and confocal images of Caco-2 cells. Cells were transfected with SEPT2-GFP and treated with CDT (200 ng/mL CDTa and 400 ng/mL CDTb) for 170 min (white arrow, site of protrusion formation with preexisting septin accumulation). (Scale bar, 5  $\mu$ m.) (B) Caco-2 cells were transfected with SEPT6-tomato and EB3-GFP. The 55-s time-lapse sequence after CDT intoxication (as in A) for 1.5 h. Yellow arrowhead marks an EB3-positive microtubule plus-end, approaching a septin structure at the protrusion base. White arrowhead marks the same plus-end pausing at the base of protrusion. Cyan arrowhead shows redirection and fast entering into the protrusion. (C) Model for redirection of polymerizing microtubules into protrusions. Plus-ends of polymerizing microtubules contact a septin chevron-like structure and are redirected into the protrusion. (D) Representative blot of an EB1-GST pull-down of endogenous SEPT2 and SEPT7 from Caco-2 cell lysates. GST-loaded beads were used as a control. (E) Quantification of blots as in D for SEPT7. Blots were normalized to the amount of SEPT7 in the lysate. Data are  $\pm$  SEM,  $n = 4$ . (F) Quantification of blots as in D for SEPT2. Blots were normalized to the amount of SEPT2 in the lysate. Data are  $\pm$  SEM and  $n = 4$ . (G) Representative blot of a coimmunoprecipitation of SEPT2 along with EB1-GFP. HeLa cells were transfected with EB1-GFP for 24 h. SEPT2 and EB1-GFP from the lysates were coimmunoprecipitated with anti-GFP antibody conjugated to magnetic beads. GFP-transfected cells were used as control. (H) Quantification of blots as in G for SEPT2. Blots were normalized to the amount of EB1-GFP and GFP on beads. Data are  $\pm$  SEM and  $n = 3$ .

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