Membrane remodeling during embryonic abscission in Caenorhabditis elegans

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Abscission is the final step of cytokinesis and results in the physical separation of two daughter cells. In this study, we conducted a time-resolved series of electron tomographic reconstructions to define the steps required for the first embryonic abscission in Caenorhabditis elegans. Our findings indicate that membrane scission occurs on both sides of the midbody ring with random order and that completion of the scission process requires actomyosin-driven membrane remodeling, but not microtubules. Moreover, continuous membrane removal predominates during the late stages of cytokinesis, mediated by both dynamin and the ESCRT (endosomal sorting complex required for transport) machinery. Surprisingly, in the absence of ESCRT function in C. elegans, cytokinetic abscission is delayed but can be completed, suggesting the existence of parallel membrane-reorganizing pathways that cooperatively enable the efficient severing of cytoplasmic connections between dividing daughter cells.

Introduction

In metazoan cells, cytokinesis is initiated by signals from the anaphase spindle, triggering the assembly of an actomyosin ring that constricts the plasma membrane to generate a narrow intercellular bridge containing two bundles of antiparallel microtubules that overlap in the central zone called the midbody (Glotzer, 2005; Green et al., 2012). The intercellular bridge is subsequently severed on both sides of the midbody at secondary constrictions, and the midbody remnant is released and subsequently engulfed by one of the daughter cells (Ettinger et al., 2011; Chai et al., 2012; Agromayor and Martin-Serrano, 2013; Elia et al., 2013; Lafaurie-Janvore et al., 2013; Crowell et al., 2014). Several models have been developed to describe the mechanism of abscission (Schiel and Prekeris, 2010; Mierzwa and Gerlich, 2014; Gould, 2016; Schönberg et al., 2017). Although previous studies suggest that internal membrane compartments fill the intercellular channel and template membrane fusion reactions that divide the cell (Finger and White, 2002; Xu et al., 2002; Low et al., 2003), more recent evidence suggests a role for Rab11/FIP3-positive recycling endosomes and Rab35-bound vesicles in delivering key factors, including OCRL, SCAMP2/3, and p50RhoGAP, which regulate actin organization and help to establish secondary constrictions at future abscission sites (Kouranti et al., 2006; Dambournet et al., 2011; Schiel et al., 2012). Microtubules play an essential role in this process by serving as tracks along which vesicles are transported toward constriction sites, but they must be depolymerized before membrane scission, which is thought to be mediated by the ESCRT (endosomal sorting complex required for transport) machinery (Caballe and Martin-Serrano, 2011; Guizetti et al., 2011; Schiel et al., 2011; Mierzwa and Gerlich, 2014; Hurley, 2015; Schönberg et al., 2017). Specifically, ESCRT-III filaments exhibit the ability to remodel lipid bilayers, facilitating membrane constriction and fission (Elia et al., 2012; McCullough et al., 2013; Schuh and Audhya, 2014; Hurley, 2015; Schönberg et al., 2017). In general, current models on cytokinesis are based on cytological changes observed in cultured mammalian cells grown on artificial substrates, begging the question of what actually occurs in an in vivo setting.

The Caenorhabditis elegans early embryo offers several advantages to study abscission in a native setting. First, embryo development is highly stereotyped, enabling accurate correlation of live-cell imaging studies with ultrastructural analysis of cryoimmobilized cells (Pelletier et al., 2006). Second, the reproducible timing and duration of cytokinesis in the one-cell C. elegans embryo also permits definition of the acute impacts of disrupting individual cytoskeletal elements at various time points during furrow ingression (Oegema and Hyman, 2006). Third, the C. elegans germline is particularly sensitive to protein depletion after RNAi, allowing identification of the roles of specific factors implicated in cytokinesis during the first time abscission takes place (Oegema and Hyman, 2006). This contrasts with depletion studies in mammalian tissue culture cells, in which protein elimination requires multiple days and cells...
continue to divide, potentially in an aberrant manner, under conditions in which protein levels are partially reduced.

Previous work using the early *C. elegans* embryo has helped to define discrete steps that take place during cytokinesis (Green et al., 2013). However, a high-resolution ultrastructural view of these events has been lacking. To understand how the topology of the intercellular bridge evolves during cell division, we used a correlative light and electron microscopic approach. Our findings highlight conserved mechanisms that contribute to the abscission process but also demonstrate unique features that distinguish cytokinesis in an embryonic system from that observed in mammalian cells in culture.

**Results and discussion**

**Ultrastructural analysis of the first embryonic cytokinesis in *C. elegans***

To characterize the first embryonic cell division in *C. elegans*, we initially used spinning-disk confocal microscopy and conducted time-lapse imaging of transgenic zygotes coexpressing probes for the plasma membrane (GFP::PH [pleckstrin homology]) and microtubules (mCherry::β-tubulin). These studies reproducibly demonstrated the timing of various events relative to the onset of furrow ingression. In particular, we found that completion of furrow ingression occurred within 170 s (± 28 s; n = 11; Fig. S1 A). Thickening of the plasma membrane followed (360 ± 49 s; n = 13) and was accompanied by microtubule disassembly shortly thereafter (389 ± 45 s; n = 15), as indicated by a local increase in the intensity of the GFP::PH probe and a loss in mCherry::β-tubulin fluorescence, respectively (Fig. 1, A and B; and Fig. S1 A). We also consistently observed the asymmetric internalization of GFP::PH-labeled membranes into the cytoplasm of the posterior cell (876 ± 117 s; n = 12; Fig. 1 A). Finally, complete engulfment of the midbody into the posterior cell occurred ~1,088 s after the onset of furrow ingression (± 137 s; n = 13; Fig. 1 A).

Previous work defined abscission as a two-step process involving cytoplasmic isolation between daughter cells followed by midbody release (Green et al., 2013). However, this work relied largely on light microscopy to define stages of abscission, which provides insufficient resolution to determine changes in membrane topology and cytoskeletal organization during each stage of cytokinesis. We therefore performed correlative light and electron microscopy on isolated embryos that were high-pressure frozen at specific time points after furrow ingression began, and then we collected electron tomograms from serial semithick sections (Fig. 1 C and Videos 1, 2, 3, 4, 5, and 6). A representative early intercellular bridge, frozen 240 s after onset of furrow ingression, was 0.75 µm in length, 1.5 µm in diameter, and contained an array of midzone microtubules. Electron tomograms also revealed an accumulation of vesicles, which were on average 55 ± 4 nm in diameter and distributed throughout the intercellular bridge, as well as tubulovesicular organelles in the surrounding area and numerous membrane ridges protruding from the midbody. In many ways, the structure of the *C. elegans* intercellular bridge at this time point resembles that of a HeLa cell in early telophase (Schiel et al., 2011). At a later time point before abscission (375 s after initiation of furrow ingression), microtubules exhibited a more asymmetric distribution (Fig. 1 C). In some cases, the majority of the microtubules originated in the anterior cell and extended into the intercellular bridge, whereas in other cases, the posterior cell exhibited a higher density of microtubules. The positioning of vesicles within the region mirrored the asymmetry in microtubules, suggesting that they arrive using microtubule tracks and contribute to the delivery of vesicle-associated signaling factors during the early stages of cytokinesis before abscission. To test this idea, we depleted animals of PERM-1 (to permeabilize the embryonic eggshell) and treated one cell–stage embryos with the microtubule-depolymerizing drug nocodazole upon initiation of furrow ingression. Under conditions where microtubules were absent within the intercellular bridge based on both light and electron microscopy, we failed to observe an accumulation of membranes near the midbody, and the intercellular bridge exhibited a reduced diameter (350 nm) in comparison with control embryos (~240 s after onset of furrow ingression; Fig. S1, B and C). Nonetheless, abscission was able to complete under these conditions, consistent with prior work both in *C. elegans* and mammalian cells (Guizetti et al., 2011; Schiel et al., 2011; Green et al., 2013). In the absence of nocodazole, PERM-1–depleted embryos exhibited normal timing of all events associated with abscission (Fig. S1 D).

We next used electron tomography to define the ultrastructure of the intercellular bridge during abscission, analyzing serial semithick sections to ensure that small connections between daughter cells were not overlooked. These data demonstrated that membrane sealing around the midbody occurred between 450 and 850 s after initiation of furrow ingression (Fig. 1, A and C), subsequent to cytoplasmic isolation but well before midbody release, which was previously suggested to correlate with abscission (Green et al., 2013). At no point during this time period did we observe microtubules within the intercellular bridge (Fig. 1 C). In addition, we analyzed multiple cases in which one of the daughter cells had been abscised, whereas the other daughter cell remained connected to the intercellular bridge. In some instances, the anterior cell underwent abscission before the posterior cell (four out of eight embryos; Fig. 1 C′, 445 s), but in others, the posterior cell was abscised first (four out of eight embryos; Fig. 1 C′′, 650 s). Our findings suggest that the first abscission event occurs randomly and without preference for one cell over another. Additionally, to our surprise, we did not detect helical filaments or other abscission-specific structures identified previously in mammalian tissue culture cells (Guizetti et al., 2011) in any serial section during the time period of abscission in *C. elegans* embryos (Fig. 1 D).

After completion of plasma membrane sealing, the residual body of the intercellular bridge, termed the midbody remnant, remained between the daughter cells (Fig. 1 C, 765 s). At this stage, we observed small membrane buds emanating from the midbody remnant, which appeared to be undergoing engulfment into the cytoplasm of the posterior cell (Fig. 1 C). The appearance of these nascent membrane budding structures correlated with the time point of GFP::PH-labeled membrane internalization observed using light microscopy. Approximately 900 s after the onset of furrow ingression, 60% of the embryos (n = 15) displayed a midbody remnant that was completely enclosed by the plasma membrane and engulfed or actively undergoing engulfment into the posterior cell (Fig. 1 C).

The ESCRT machinery is required for membrane internalization after abscission. Although initially described for their role in endosomal cargo sorting, components of the ESCRT machinery have been
Figure 1. **Time-resolved analysis of cytokinesis by correlative light and electron microscopy.** (A) Timing of abscission events during the first embryonic division. The start of cleavage furrow ingression is defined as the 0 s time point. Single time points for each analyzed abscission event are indicated by symbols. A red line highlights the mean time point for each. The time point of abscission was obtained by tomographic analysis as shown in C. n = 11–15, depending on the step analyzed. (B) Determination of the time point of microtubule disassembly. Analysis of fluorescence intensity (mCherry::β-tubulin) in the region of the cleavage furrow (position 1) and in the cytoplasm (defined as background; position 2). The background fluorescence was subtracted from the fluorescence intensity at the cleavage furrow. The mean values of 10 single embryos were plotted over time. The time point of microtubule (MT) disassembly is indicated (red line). Error bars show SD. r.u., relative units. (C) 3D reconstruction of intercellular bridges at defined time points after initiation of furrow ingression as indicated in the right panels (top right corner, seconds). A light microscopy (LM) image (first column, GFP::PH in green and mCherry::β-tubulin in red), an electron microscopy image (second column, tomographic slice), the corresponding 3D model (third column), and a cartoon illustration (fourth column) are shown for each time point. Color coding is as follows: gold, plasma membrane of an open side of the intercellular bridge; red, microtubules; white, open microtubule ends; blue, closed microtubule ends; green, vesicles; dark gray background, cytoplasm at the open bridge; light gray background, cytoplasm at the disconnected bridge; nude, anterior cell membrane; and purple, posterior cell membrane. The midbody remnants (765 s and 900 s) are also shown in gold. (D) Additional tomographic z slices of an embryonic intercellular bridge (650 s; a thickening of the membrane is indicted by arrowheads). Bars, 500 nm.
implicated in a wide variety of membrane remodeling events, including those that occur during cytokinesis (Schuh and Audhya, 2014; Hurley, 2015; Christ et al., 2017). However, the precise role for the ESCRTs during abscission remains controversial. To address this issue, we first examined the distribution of the ESCRT-I subunit TSG-101 during embryonic cell division in C. elegans. Initially, TSG-101 localized to the mitotic spindle and was subsequently detected at the intercellular bridge ~300 s after initiation of furrow ingression (Fig. 2 A). Accumulation of TSG-101 plateaued ~300 s later and remained constant at the midbody thereafter (Fig. 2 B). Consistent with these findings, endogenous TSG-101 colocalized with the midbody component ZEN-4 both before and after abscission (Fig. 2 C). In contrast, the ESCRT-III component VPS-32 was distributed largely at the edges of the midbody remnant at these time points in a manner dependent on the presence of TSG-101 (Fig. 2, C and D).

In vitro analysis previously described that recombinant C. elegans VPS-32 assembles into spiral filaments with a thickness of 4.2 nm (Shen et al., 2014). To determine whether the accumulation of VPS-32 near the midbody correlated with the presence of spiral filaments as observed previously in mammalian cells undergoing abscission (Guizetti et al., 2011), we again took advantage of electron tomography. Although we failed to observe filamentous structures throughout the abscission process (Fig. 1, C and D), we consistently identified filamentous structures at the necks of membrane buds emerging from the midbody remnant ~800 s after the onset of furrow ingression (Fig. 2 E). These membrane bud–associated filaments exhibited a diameter of 12 nm and a spacing of 26 nm, slightly thicker than the recombinant VPS-32 spiral polymers observed by cryo-electron microscopy (Shen et al., 2014) but consistent with the idea that these filaments correspond to ESCRT-III heteropolymers. Upon depletion of TSG-101, which inhibits downstream recruitment of the ESCRT-III complex (Fig. 2 D), GFP::PH-labeled membrane internalization into the posterior cell was no longer observed (12/12 control embryos exhibited GFP::PH-labeled membrane internalization into the posterior cell was no longer observed (12/12 control embryos exhibited membrane open- ing 550 s after initiation of furrow ingression, 500 s later (Fig. 3 B). To define the role of actin during this time interval, we permeabilized the embryos after abscission normally completes (Fig. 3 A; Singh and Pohl, 2014), and fluorescence intensity of the probe peaked ~425 s later (Fig. 3 B). To define the role of actin during this time interval, we permeabilized the embryonic eggshell using PERM-1 depletion and treated one cell–stage embryos with latrunculin A upon furrow closure, after intercellular bridge assembly was complete (~170 s after initiation of furrow ingression; Fig. 3 C). The embryos accumulated TSG-101 at the midbody normally, but neither membrane internalization nor engulfment of the midbody remnant was detected (Fig. 3 C, arrowheads). Additionally, we observed a persistent membrane opening adjacent to the midbody ~550 s after onset of furrow ingression in 80% of the embryos (~2.8 µm in length and 1.5 µm in width on average; n = 15; Fig. 3 C, white arrow). These data suggest that actin polymers play a role late during the abscission process.

To further explore this phenotype, we analyzed the ultrastructure of intercellular bridges in latrunculin A–treated embryos, frozen ~1,020 s after onset of furrow ingression and long after abscission normally completes (Fig. 3 D). Under this condition, serial semithick section electron tomography revealed that depletion of actin polymers prevents abscission. Instead, the intercellular bridge appeared fragmented (Fig. 3 D, arrowheads) and exhibited multiple thin cytoplasmic connections bridging the embryonic daughter cells. In addition, the membrane opening as visualized by light microscopy could be detected (Fig. 3 C, C and D, arrows). We observed a similar phenotype by treating

Actin polymers play an active role during the late stages of abscission

Although actin plays a well-described role in directing cortical contractility during cytokinesis (Mishima, 2016), its roles during late stages of abscission remain unclear (Guizetti et al., 2011; Prekeris, 2011; Schiel et al., 2012; Mierzwa and Gerlich, 2014). Although several lines of evidence indicate that actin is required for initial stabilization of the intercellular bridge, actin filaments must ultimately be cleared for the final abscission step to take place (Dambournet et al., 2011; Schiel et al., 2012). Using transgenic embryos expressing LifeAct-GFP to visualize actin filaments during cytokinesis, we observed an initial signal at the intercellular bridge ~375 s after initiation of furrow ingression (Fig. 3 A; Singh and Pohl, 2014), and fluorescence intensity of the probe peaked ~425 s later (Fig. 3 B). To define the role of actin during this time interval, we permeabilized the embryonic eggshell using PERM-1 depletion and treated one cell–stage embryos with latrunculin A upon furrow closure, after intercellular bridge assembly was complete (~170 s after initiation of furrow ingression; Fig. 3 C). The embryos accumulated TSG-101 at the midbody normally, but neither membrane internalization nor engulfment of the midbody remnant was detected (Fig. 3 C, arrowheads). Additionally, we observed a persistent membrane opening adjacent to the midbody ~550 s after onset of furrow ingression in 80% of the embryos (~2.8 µm in length and 1.5 µm in width on average; n = 15; Fig. 3 C, white arrow). These data suggest that actin polymers play a role late during the abscission process.

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Figure 2. The ESCRT machinery promotes membrane internalization after abscission. (A) Representative images of ESCRT\textsuperscript{TSG-101} accumulation at the intercellular bridge by fluorescence light microscopy. Bar, 10 µm. (B) Quantification of ESCRT\textsuperscript{TSG-101} accumulation. Error bars show SD. r.u., relative units. n = 10. (C) Confocal images showing the accumulation of MKLP\textsuperscript{ZEN-4} together with ESCRT\textsuperscript{TSG-101} or ESCRT\textsuperscript{III} before and after abscission. Arrowheads indicate the distinct accumulation of MKLP\textsuperscript{ZEN-4} and ESCRT\textsuperscript{III} after abscission. Line scan analysis within the boxed regions was used to measure the distributions of ESCRT components relative to ZEN-4 in each image. n = 6 embryos for each condition. Bar, 2 µm. (D) STED microscopy indicating the accumulation of ESCRT\textsuperscript{III} in control [arrowhead] and tsg-101(RNAi) embryos. n = 6 embryos for each condition. Bar, 1 µm. (E) Tomographic slice of a midbody remnant showing membrane internalization (left, arrow; bar, 500 nm) and a series of tomographic slices of the same membrane bud (right;
permeabilized embryos with Blebbistatin, an inhibitor of myosin II, after completion of furrow ingression (~170 s after onset of furrow ingression; Fig. S3 A). Specifically, serial semithick section electron tomography of a Blebbistatin-treated embryo frozen ~1,100 s after onset of furrow ingression showed an intercellular bridge with thin cytoplasmic connections (Fig. S3 B). A previous fluorescence-based imaging study using a temperature-sensitive allele suggested that myosin II is dispensable after completion of furrow ingression (Davies et al., 2014), but our studies provide evidence to the contrary, again highlighting the importance of conducting corroborative electron microscopy studies. The specific roles of actin and myosin in executing the final step of cell cleavage are currently unclear. However, this pathway may represent a redundant mechanism to ESCRT-mediated membrane abscission in embryonic systems. Alternatively, because a primary function of actin is to stabilize the cleavage furrow, fragmentation of the intercellular bridge upon actin depolymerization may have indirect consequences on membrane remodeling or positioning required for final membrane scission.

Dynamin-mediated endocytosis is required before abscission

Based on our finding that the ESCRT machinery plays a key role in membrane removal during the late stages of cell division, we decided to explore the impact of eliminating another essential regulator of membrane internalization during embryonic abscission. The large GTPase dynamin, which is encoded by DYN-1 in C. elegans, functions in the severing of nascent endocytic vesicles to generate free transport carriers (Ferguson and De Camilli, 2012). Although implicated in cytokinesis, its precise function during cell division remains unclear. Consistent with previous work (Thompson et al., 2002), we demonstrated that DYN-1 initially accumulates at the intercellular bridge ~525 s after onset of furrow ingression (Fig. 4 A, arrowheads). Fluorescence intensity measurements of DYN-1::GFP showed a linear accumulation of the protein with a peak ~350 s later (Fig. 4 B). To define the function of dynamin at the intercellular bridge, we conducted a series of depletion experiments. Midzone architecture in DYN-1–depleted embryos was indistinguishable from control embryos throughout furrow closure, microtubule disassembly, and membrane thickening. However, ~602 s (~36 s; n = 5) after onset of furrow ingression, the furrow began to regress (n = 5; Fig. 4, C and D, arrowheads). Based on our finding that myosin levels remained normal on the ingressing furrow in DYN-1–depleted embryos (Fig. S3, C and D), this phenotype was unlikely to be a result of a defect in actomyosin ring stability, as was observed previously in Dictyostelium discoideum after dynamin inhibition (Masud Rana et al., 2013).

To analyze membrane topology immediately before furrow regression, we froze DYN-1–depleted embryos ~515 s after onset of cleavage furrow ingression and examined them using electron tomography (Fig. 4 E). These studies revealed broad openings in the daughter cells with diameters of ~1.25 µm in the anterior cell and ~0.48 µm in the posterior cell. Both openings appeared to be larger than any observed in control embryos at this time point. Interestingly, membrane ridges protruding from the midbody could not be identified in DYN-1–depleted embryos. However, we observed an accumulation of nascent vesicles on the plasma membrane, consistent with the function of dynamin in releasing endocytic transport carriers as well as free vesicles within the intercellular bridge (Fig. 4 E). These data suggest that DYN-1 plays a key role before abscission to clear excess membrane, which may otherwise interfere with generating a membrane topology conducive to abscission.

Collectively, our findings support a model in which actomyosin-based contractility not only supports cleavage furrow ingression, but also stabilizes the intercellular bridge to appropriately position the juxtaposed plasma membranes before abscission. Surprisingly, scission of the intercellular bridge after the first mitotic division in C. elegans does not appear to exhibit an absolute requirement for the ESCRT machinery, arguing instead for the existence of a parallel, potentially actin-dependent pathway that supports this process. Our findings also highlight the substantial amount of membrane that is transported to the furrow during early embryonic divisions, consistent with previous work examining the first cytokinesis of sea urchin embryos (König et al., 2017). In contrast, cell division in mammalian tissue culture cells involves relatively little change in total plasma membrane surface area (Figard and Sokac, 2014), which may help to explain some of the differences in cytokinesis observed in vitro as compared with embryonic divisions in vivo. Importantly, our data argue that excess membrane must be removed during multiple stages of cell division for the process to proceed normally both in a dynamin-dependent manner and also via function of the ESCRT machinery. Recent work in mammalian cells also described the formation of small membrane blebs near the intercellular bridge during cytokinesis (Sherman et al., 2016), suggesting that this requirement is not unique to worm embryos. In the absence of ESCRT-dependent membrane removal, the membrane surrounding the midbody remnant expands, potentially altering membrane tension, which was shown previously to play an important role in the timing of abscission (Lafaurie-Janvore et al., 2013). In addition, changes in membrane tension may disrupt normal phagocytosis of the midbody remnant into the posterior cell (Masters et al., 2013; Figard and Sokac, 2014). How membrane removal orchestrated by the ESCRT machinery directly contributes to the timing of abscission and midbody internalization represents an important issue to be addressed in the future.
Materials and methods

C. elegans strains

The following strains were used in this study: TH155 (unc-119(ed3)-III; ltIs38[pAA1; pie-1p::GFP::PH(PLC1Δ1);unc-119+]; mCherry::β-tubulin); MSN707 (unc-119(ed3)-III; HzIs169[pie-1p-mCherry::ZEN-4; unc-119+]; ltIs38[pAA1; pie-1p::GFP::PH(PLC1Δ1);unc-119+]; TMR08 (unc-119(ed3)-III; ddIs186[WRM0631C_D06::S000126_R6K-pCFJ496-000032-mos1(tsg-101::S000138_R6K-2×TY1-wSNAP-eGFP 3×FLAG); unc-119+]; ltIs44[pAA173; pie-1p-mCherry::PH(PLC1Δ1) + unc-119+]; TMR20 (unc-119(ed3)-III; ddIs86[pie-1p::LifeAct::GFP; unc-119+]; ltIs44[pAA173; pie-1p-mCherry::PH(PLC1Δ1) + unc-119+]; TMR22 (dqls3[pie-1p::dyn-1B::GFP + unc-119+]; ltIs44[pAA173; pie-1p-mCherry::PH(PLC1Δ1) + unc-119+]; and MSN608 (unc-119(ed3)-III; HzIs100[pie-1p::GFP::CAY-1; unc-119+]; eri-1(mg366)]. Maintenance of C. elegans strains was performed according to standard procedures (Brenner, 1974). For all experiments, L4 larvae were shifted to 25°C 24 h before imaging. For all ultrastructural studies, either strain TH155 or MSN608 was used for staging of early embryos before cryoimmobilization for electron microscopy.

RNAi

RNAi by feeding was performed as described previously (Timmons and Fire, 1998). In brief, L4 larvae or young adult animals were seeded onto bacteria, which had been induced to express specific double-stranded RNAs using IPTG. The following feeding clones were obtained from the Ahringer laboratory and grown at 25°C on feeding plates: T01H3.4 (perm-1; 20–24 h), C09G12.9 (tsg-101; 48 h), and C2C6.1a (dyn-1; 12 h). Using oligonucleotides containing T3 and T7 promoters, double-stranded RNAs were also prepared to deplete TSG-101 by injection: oligonucleotide 1, 5′-TAA TAC GAC TCA CTA AAG GTG TTA TTT GGC GTT-3′, and oligonucleotide 2, 5′-AAT TAA ACG TCA AGG TAT CCA AAT GGG AGC TTC TGG-3′. Regions of TSG-101 were amplified by PCR using N2 genomic DNA as a template. PCR samples were purified using the PCR cleanup kit (QIAGEN) and eluted with 50 µl of water. The Ambion kit was used for the T3 and T7 transcription reactions. The reaction mix was incubated at 37°C for 5 h. To remove the DNA template, 1.3 µl DNase was added to each reaction and incubated for 15 min at 37°C. The samples were then purified using the RNasy kit (QIAGEN). Each reaction was eluted with 30 µl RNase-free water. To anneal the RNAs, equal amounts of the T3 and T7 reactions were combined and incubated at 68°C for 10 min and then at 37°C for 1 h. To ensure full depletion of TSG-101 for electron microscopic studies, an eri-1(mg366) mutant background was used, which exhibits hypersensitivity to RNAi (Kennedy et al., 2004).

Live-cell imaging

For live-cell imaging, adult hermaphrodites were transferred to a droplet of M9 buffer and placed on a 20 × 64-mm high-precision cover glass (Roth). Hermaphrodites were dissected to release embryos. Imaging at 20°C was performed on a spinning-disk microscope (ZEISS) using
a C-Apochromat 63×/1.2-W objective lens and a CSU-X1 scan head (Yokogawa Electric Corporation). The 405-, 488-, and 561-nm lasers were used for illumination. For time-lapse acquisitions, a double beam splitter 527/54 + 645/60 was used in front of an electron-multiplying charge-coupled device camera (Evolve 515; Roper Technologies). Images were acquired using ZenBlue software (ZEISS) and processed with FIJI (ImageJ; National Institutes of Health; Walter et al., 2010).

For quantification of membrane internalization, a threshold was established to identify GFP::PH-labeled membranes that were internalized in control embryos. After depletion of TSG-101, the same threshold was used, and we were unable to observe membrane internalization.

**Permeabilization of embryos and drug treatment**

To perform drug treatment experiments, eggshells of early embryos were permeabilized by perm-1(RNAi) treatment (Carvalho et al., 2011). Permeabilized embryos were gently aspirated into cellulose capillaries to allow a transfer of embryos into the medium containing the drug (König et al., 2017). Embryos were initially imaged without addition of a drug but in the presence of 2 µg/ml Hoechst to verify the permeability of the eggshell of each selected embryo. At the time point of interest, the capillary with the enclosed embryo was transferred to a new coverslip with medium containing the drug, and imaging was continued. The following drugs were added: 10 µM latrunculin A after completion of furrow ingression to inhibit actin polymerization, 2.5 µM Blebbistatin after completion of furrow ingression to selectively inhibit nonmuscle myosin II, and 10 µg/ml nocodazole (Sigma-Aldrich) at onset of cleavage furrow ingression.

**Specimen preparation for electron microscopy**

Sample preparation for electron microscopy was performed as described previously (Müller-Reichert et al., 2007). In brief, single *C. elegans* embryos were aspirated into cellulose capillary tubes. Capillaries of ~1 mm in length containing early embryos were imaged under the light microscope and then placed in 100-µm-deep membrane carriers containing M9 buffer + 20% BSA (Sigma-Aldrich) as a filler. Cryoimmobilization was...
performed using an EM PACT2 with a rapid transfer system high-pressure freezer (Leica Microsystems). Freeze substitution was performed for ~48–72 h at ~90°C in anhydrous acetone containing 1% OsO4 and 1% uranyl acetate. Freeze substitution cocktails for drug treatment experiments contained 0.5% OsO4 and 0.5% uranyl acetate in anhydrous acetone. Specimens were then infiltrated and thin-layer embedded in epon/Araldite. Thin (90 nm) or semithick (300–350 nm) serial sections were collected on Formvar-coated copper slot grids and poststained with 2% uranyl acetate in 70% methanol followed by Reynolds’s lead citrate.

Electron microscopy

Thin-section electron microscopy was performed on a 120-kV microscope (TECNAI Bio-Twin; FEI) using a 4 × 2.7–K charge-coupled device camera (Orius SC1000W; Gatan). In preparation for electron tomography, 15-nm colloidal gold particles (BBI Solutions) were attached to both surfaces of serial semithick sections to serve as fiducial markers for image alignment, and regions of interest were preselected by using a transmission electron microscope (EM 906; ZEISS) operated at 80 kV and equipped with a charge-coupled device camera (Tröndle Sharp Eye). The specimens were then placed in a tilt-rotate specimen holder (650; Gatan), and tomographic datasets were recorded using an intermediate-voltage transmission electron microscope (TECNAI F30; FEI) operated at 300 kV and equipped with a 2 × 2–K charge-coupled device camera (US1000; Gatan). The SerialEM program (Boulder Laboratory for 3D Electron Microscopy) was used to automatically acquire images every 1° over a range of ±62° at a pixel size of 1.4 nm. For dual-axis tomography, the grids were imaged in one tilt series and then rotated 90°, and a similar tilt series was acquired. Electron tomograms were computed for each tilt axis using the R-weighted back-projection algorithm (Gilbert, 1972). For double-tilt datasets, the two tomograms were aligned with each other and combined (Mastronarde, 1997). Using the IMOD software package (Boulder Laboratory for 3D Electron Microscopy), electron tomograms were displayed to generate 3D models of intercellular bridges of early embryos frozen at distinct time points throughout cytokinesis. In 3D, we modeled membranes, microtubules, microtubule ends, and vesicles. Cellular components were quantified using the program IMODInfo.

Specimen preparation and imaging by stimulated emission depletion (STED) microscopy

For STED imaging of fixed embryos, *C. elegans* control and tsg-101 (RNAi) animals were dissected onto poly-l-lysine–coated slides, rapidly frozen in liquid nitrogen, subjected to a freeze–crack method to permeabilize the eggshell of embryos, and fixed in 100% MeOH at −20°C for 20 min as described previously (Desai et al., 2003). Antibodies directed against VPS-32 (Shen et al., 2014) and ZEN-4 (Audhya et al., 2005) were conjugated to Alexa Fluor 680 and Cy3 dyes, respectively, and incubated on the slides overnight at 4°C. Samples were mounted with ProLong Diamond and allowed to cure for 24 h at room temperature. Images were collected at room temperature using a 100× STED white oil objective (1.4 NA) on a STED 3x microscope (TCS SP8) equipped with HyD SP GaAsP detectors (Leica Microsystems) using two depletion lasers (660 and 775 nm). Image acquisition was performed using LAS X software (Leica Microsystems). Raw image files were deconvolved using Huygens Professional software (Scientific Volume Imaging), and parallel confocal images were also acquired using a white light laser.

Online supplemental material

Fig. S1 shows light and electron microscopic analysis of an embryo after treatment with nocodazole, demonstrating that microtubules are not detectable within the intercellular bridge after such treatment. Fig. S2 shows serial thin sections of tsg-101(RNAi) embryos ~1,110 s after initiation of furrow regression, demonstrating that ESCRT function is dispensable for the first embryonic membrane scission event in *C. elegans*. Fig. S3 shows light and electron microscopic analysis of an intercellular bridge after treatment with Blebbistatin, demonstrating that myosin II function is required for cytokinetic abscission in *C. elegans*. Videos 1, 2, 3, 4, 5, and 6 show the tomographic reconstruction and 3D modeling of the first intercellular bridge in the *C. elegans* embryo and correspond to Fig. 1 C. The time points of freezing after initiation of furrow regression are Video 1, 240 s; Video 2, 375 s; Video 3, 445 s; Video 4, 650 s; Video 5, 765 s; and Video 6, 900 s.

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