Fission Yeast Sec3 Bridges the Exocyst Complex to the Actin Cytoskeleton

Isabelle Jourdain1,*, Hannah C. Dooley1,2 and Takashi Toda1,*

1 Cell Regulation Laboratory, London Research Institute, Cancer Research UK, 44 Lincoln’s Inn Fields, London WC2A 3LY, UK
2 Current address: Secretory Pathways, Cancer Research UK, 44 Lincoln’s Inn Fields, London WC2A 3LY, UK
*Corresponding author: Isabelle Jourdain, isabelle.jourdain@cancer.org.uk or Takashi Toda, takashi.toda@cancer.org.uk

The exocyst complex tethers post-Golgi secretory vesicles to the plasma membrane prior to docking and fusion. In this study, we identify Sec3, the missing component of the Schizosaccharomyces pombe exocyst complex (SpSec3). SpSec3 shares many properties with its orthologs, and its mutants are rescued by human Sec3/EXOC1. Although involved in exocytosis, SpSec3 does not appear to mark the site of exocyst complex assembly at the plasma membrane. It does, however, mark the sites of actin cytoskeleton recruitment and controls the organization of all three yeast actin structures: the actin cables, endocytic actin patches and actomyosin ring. Specifically, SpSec3 physically interacts with For3 and sec3 mutants have no actin cables as a result of a failure to polarize this nucleating formin. SpSec3 also interacts with actin patch components and sec3 mutants have depolarized actin patches of reduced endocytic capacity. Finally, the constriction and disassembly of the cytokinetic actomyosin ring is compromised in these sec3 mutant cells. We propose that a role of SpSec3 is to spatially couple actin machineries and their independently polarized regulators. As a consequence of its dual role in secretion and actin organization, Sec3 appears as a major co-ordinator of cell morphology in fission yeast.

Key words: actin, endocytosis, exocyst, morphology, Schizosaccharomyces pombe

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Two trafficking events, exocytosis and endocytosis, orchestrate the remodeling of the plasma membrane. Exocytosis involves the fusion of intracellular secretory vesicles with the plasma membrane, thereby providing a source of lipid moieties for membrane extension. It is also used to deliver integral membrane proteins at the cell surface and to release material in the extracellular space. Typically, in yeasts, the secretion of hydrolytic enzymes is necessary to dissolve the cross wall (primary septum) between the two daughters cells and complete cytokinesis (1). Most of our current knowledge of polarized secretion comes from studies conducted in the budding yeast Saccharomyces cerevisiae. Cargoes emanating from intracellular organelles are transported by motor proteins along cytoskeletal tracks towards polarized areas of the plasma membrane. Sites of vesicle targeting are defined by factors present on the vesicle and by polarity cues at sites of growth (2). The initial contact between the vesicle and the plasma membrane is mediated by tethering factors that are thought to bridge SNAREs (soluble N-ethylmaleimide-sensitive factor attachment protein receptor) on apposing membranes. This paired trans-SNARE complex docks the vesicle to the receiving membrane and finally induces lipid fusion (3).

The tethering of secretory vesicles before docking and fusion with the plasma membrane is mediated by the exocyst, a conserved octameric protein complex consisting of Sec3, Sec5, Sec6, Sec8, Sec10, Sec15, Exo70 and Exo84 (4–6). The assembly of the exocyst complex is a sequential process. Sec15 is first loaded onto maturing secretory vesicles and may serve to recruit additional exocyst proteins (7,8). Upon delivery of the vesicle at the cell periphery, this pre-complex interacts with membrane-bound Sec3 and Exo70, thus tethering the vesicle to the plasma membrane (9–12).

Endocytosis is responsible for the uptake of extracellular material and the recycling of lipids and surface proteins. Clathrin-mediated endocytosis is a conserved type of endocytosis but contrary to the situation in mammals, membrane invagination in yeasts shows an absolute requirement for actin (13–15). Specialized structures, the actin patches, are dedicated to pulling the forming vesicles inside the cytoplasm (16,17) and are specifically nucleated by the Arp2/3 complex (18,19). In all fungi, actin patches concentrate at sites of active growth and division (20–22). The factors and mechanisms that restrict the localization of actin patches remain largely unknown.

The characteristic rod-like shape of the Schizosaccharomyces pombe constitutes an excellent tool by which to study plasma membrane remodeling and cell morphogenesis (23,24). The long, straight axis of fission yeast cells is defined by factors deposited at the cell ends by the microtubule cytoskeleton. In situations where the function or localization of these factors is hindered, cells become curved or develop an ectopic cell tip. However, they remain cylindrical and still grow from their poles.
**Figure 1: sec3 temperature sensitive mutants have a septation defect.** A) Serial dilutions of wild-type, sec3-913 and sec3-916 cells were spotted on YE5S plates and incubated at the indicated temperatures. sec3-916 is more sensitive to high temperatures than sec3-913. B) Lectin (blue) and calcofluor (red)-staining of wild-type, sec3-913 and sec3-916 cells cultured in parallel at 27°C and 36°C. Mutant strains accumulate septated cells at the restrictive temperature and septa appear thick. Bar = 5μm. C) The percentage of septated cells is shown as a function of incubation time at 36°C. sec3-916 (green) has a stronger phenotype than sec3-913 (red).
Figure 2: Schizosaccharomyces pombe Sec3 is involved in exocytosis. A) Activity of acid phosphatase secreted from wild-type (blue diamonds), sec3-913 (red squares), sec3-916 (green circles) and sec8-1 (black crosses) cells. The cumulative activity of the enzyme was measured at time points after the cells were transferred to fresh medium and switched from 27°C to 36°C (t = 0). Results are shown relative to cell density (OD600nm). Error bars indicate the variation between two independent experiments and are SD of the mean. B) EM images of wild-type and sec3-913 cells. A septated cell and an interphase cell are shown. Note the thick septum in sec3-913. Higher magnifications of the boxed areas emphasize the presence of accumulated secretory vesicles in the mutant. Bars = 2 μm. C) Fluorescence microscopy images of the vesicular SNARE Syb1 tagged with GFP and expressed in wild-type and sec3-913 cells. D) Sec8-GFP and Sec3-tdTomato colocalize at the cell tip(s) and at a medial single or double ring. E) Reciprocal co-immunoprecipitation (IP) between Sec8-GFP and Sec3-3PK or the 3PK-tagged product of sec3-913. At equal amounts of whole cell protein extract (WCE) loaded on a gel, the mutant protein was less detectable than wild-type Sec3. For this reason, the amount of protein used in the IP experiment was increased up to 25 times. F) Myo52-GFP and Sec3-tdTomato partially colocalize. Except in panels (D) and (F) for which cells were observed at 27°C, results shown are for cells grown at 36°C. In panels (C, D, F), bars = 5 μm and enlargement bars = 1 μm.

the time frame of the experiments presented in this study.

Both mutants had a cell separation defect (Figure 1B and C). In sec3-913, the septation index increased gradually over time at 36°C while in sec3-916 the percentage of septated cells remained constantly high (Figure 1C). By contrast, the septation index of the wild-type strain remained at approximately 11% throughout the experiment. Multiple septated cells were observed in the mutants, with each compartment containing a nucleus (data not shown). Calcofluor-labeled primary septa appeared much thicker in the ts mutants than in the wild-type (Figure 1B). Further analysis of sec3-913 septa by electron microscopy (EM) showed that septa thickened soon after forming and continued to accumulate cell wall material throughout closure (Figure 2B). Final septum closure and dissolution were also delayed occasionally, leaving an apparent slim cytoplasmic bridge between the two non-separated daughter cells. No cell wall deposition invagination or malformation was otherwise observed at the cell periphery (Figures 1B and 2B).

SpSec3 is the missing component of the fission yeast exocyst complex

The septum defects described above could reflect a defect in the delivery and release of hydrolytic septum-specific glucanases (41,42). To test if the secretory pathway was compromised in sec3 ts mutants, we monitored the activity of acid phosphatase secreted into the medium...
and reported it relative to cell density (Figure 2A). Similar to the exocyst mutant sec8-1 (36), the sec3-913 cells, and to a lesser extent the sec3-916 cells, secreted less acid phosphatase than the wild-type. A close observation of sec3-913 cells by EM revealed the cytoplasmatic accumulation of vesicles at the cell tips in growing cells and at the cell equator in cells undergoing septation (Figure 2B). Such vesicles were not seen in the wild-type, nor in sec3-913 at the permissive temperature, presumably due to a higher rate of vesicle fusion with the plasma membrane. We are confident that these structures were post-Golgi secretory vesicles because the v-SNARE Syb1, which normally localizes to scattered, punctate, cytoplasmic structures (Figure 2C, left) (43) was enriched at the cell tips and septum area in sec3-913 at 36°C (Figure 2C, right). Thus, SpSec3 is involved in the trafficking of secretory vesicles.

To further characterize Sec3 in fission yeast, we chromosomally tagged it at the C-terminus with fluorescent proteins (GFP or tdTomato) or with 3PK. In a wild-type background at 27°C, SpSec3 localized to cortical dots capping the growing tips(s) in interphase cells (Figures 2D and S2). At cytokinesis, SpSec3 relocated to the middle of the cell where it first was organized as a single ring. This occurred a few minutes after the contractile actomyosin ring (CAR) was assembled but prior to septin recruitment and primary septum formation (Figure S2). The SpSec3 ring was not contractile and instead split into two parallel rings at the end of CAR constriction. This localization overlapped perfectly with that of other members of the exocyst complex (Sec8-GFP, Figure 2D and Sec10-GFP, data not shown). To check if SpSec3 and the exocyst complex physically interacted, we performed reciprocal immunoprecipitations and found that SpSec3-3PK specifically co-immunoprecipitated with Sec8-GFP (Figure 2E). Genetic data also supported a close functional relationship between SpSec3 and the exocyst complex, as sec3-913 was synthetic lethal with exo70Δ and with sec8-1 (Table S2). Hence, SpSec3 colocalizes and physically and genetically interacts with the exocyst complex.

To verify that SpSec3 was present at the receiving end of the secretory pathway, we tested whether it colocalized with Myo52, a type V myosin that delivers cargoes from the Golgi apparatus to the plasma membrane (43,44). Indeed, Myo52-GFP and SpSec3-tdTomato did partially colocalize at the cell cortex (Figure 2F). Overlapping signals presumably illustrate the moment a cargo was delivered by the myosin to the exocyst complex containing SpSec3, whereas non-coincident signals may reflect the trafficking and recycling of myosin motors toward or from the exocyst complex.

The above results suggest that S. pombe Sec3 is a member of the exocyst complex. During the course of this study, Baek et al. (45) published the crystal structure of the N-terminal domain of S. cerevisiae Sec3. Using this new fold as a template, the authors predicted that S. pombe SPAC17G8.12 notably, codes for a Sec3 protein. To unequivocally show that S. pombe Sec3 is a functional ortholog of other Sec3 proteins, we asked whether we could substitute SpSec3 by human Sec3/EXOC1 (Figure 3). Expression of SpSec3 under the control of the thiamine-repressible nmt1 promoter rescued the growth phenotype of sec3-913 at the semi-restrictive temperature 34°C. By contrast, the empty plasmid had no effect. Consolidating our findings with the predictions of Baek’s et al. (45), human EXOC1 expression partially rescued the growth phenotype of the S. pombe sec3 ts mutant.

Together, our results show that SpSec3 is the missing component of the fission yeast exocyst complex, which tethers vesicles carried on actin filaments from the Golgi apparatus by a type V myosin to the plasma membrane.

S. pombe Sec3 shares many properties with its orthologs but does not define the site of exocyst assembly

Sec3 was proposed to be a landmark for secretion at the bud tip in S. cerevisiae (11). In this model, Sec3 localization is independent of the secretory machinery,
including vesicle production and transport. To assess if this is also the case in *S. pombe*, we observed *SpSec3-GFP* in *for3Δ* cells which lack actin cables (33). We also considered the possibility that actin patches rather than cables may control Sec3 localization and observed it in cells with mis-placed or aberrant patches (*sla2Δ*, *wsp1Δ3*) (30,46). Finally, we used the *myo52Δ* strain in which vesicular transport along actin filaments is abolished (44) and the *sec8-1* mutant cells, in which vesicle tethering is hindered (36). In all cases *SpSec3-GFP* localization appeared identical to that observed in the wild-type (Figure 4A(ii)). Hence, in *S. pombe*, as in *S. cerevisiae*, Sec3 localization is not dependent upon either actin or the late secretory machinery.

A crucial aspect of the landmark model is that Sec3 defines the site of exocytosis by targeting other exocyst components to specific sites of the plasma membrane. Challenging this model, *Sec8-GFP* and other subunits of the exocyst complex were still properly targeted to the tips of *sec3-913* cells incubated at the restrictive temperature for up to two generations (*Sec8-GFP*, Figure 4B; *Sec6-GFP*, data not shown). We were unable to detect *Sec3-913-GFP* at cell tips and middle at the restrictive temperature (data not shown). This could be either due to a technical difficulty to observe *Sec3-GFP* species at 36 °C, the fact that *Sec3-913* is delocalized, or the poor stability of the mutant protein. The third was suggested by the fact that *SpSec3-913-3PK* was less immunodetected than...
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SpSec3-3PK in whole cell extracts (WCE, Figures 2E and 7D). Moreover, the re-expression of SpSec3-913 in the sec3-913 background using the strong nmt1 promoter integrated at the endogeneous locus, rescued the ts phenotype of this strain (Figure 4C). Hence, the low levels of Sec3-913 are responsible for the phenotype of sec3-913 cells. Furthermore, Figure 2E (right panel) shows that SpSec3-913-3PK is in any case unable to interact with Sec8-GFP. This finding eliminates the possibility that residual levels of mutant Sec3-913 would still be able to recruit Sec8-GFP. We conclude from these experiments that Sec8-GFP localizes properly independently of Sec3. Thus, S. pombe Sec3 does not define the site of exocyst assembly.

Sec3 proteins are known to bind directly to the phosphatidylinositol 4,5-bisphosphate (PI(4,5)P2) present in the plasma membrane (12,47). PI(4,5)P2 is produced by phosphorylation of PI4P, a modification catalyzed in S. pombe by the phosphatidylinositol-4-phosphate 5-kinase Its3 (48). In the its3-1 ts mutant, which has reduced levels of PI(4,5)P2, SpSec3 was displaced from the cell poles and cell equator (Figure 4A(ii)).

The polarization of Sec3 at the S. cerevisiae bud tip is mediated by upstream GTPases including Cdc42 (12). Cdc42 is also a major orchestrator of morphogenesis in fission yeast (24) we asked whether it could regulate SpSec3 localization. As expected, Sec3-GFP was detached from the cell tips and middle in the cdc42-2-3 mutant (49) (Figure 4A(ii)).

In a search for other factors that may control the polarization of SpSec3 at cell tips, we identified SpYpt3. SpYpt3 is a Rab11-type GTPase involved in endocytic recycling, sorting and transport (50,51). As shown in Figure 4A(ii), SpSec3-GFP was excluded from the apex of ypt3-i5 cell tips and aggregated at the vicinity of the plasma membrane. We did not notice any modification of SpSec3-GFP localization at the equator of these cells.

We conclude from these experiments that SpSec3 localization and polarization at the plasma membrane are dependent on phosholipids and on the GTPases Cdc42 and Ypt3. They are, however, independent of secretory vesicle transport, tethering and the actin cytoskeleton.

sec3 mutants have a shape phenotype and are hypersensitive to actin poisons

Besides a septation phenotype, the sec3 ts mutants also displayed a shape phenotype (Figure 5A; see also Figure 1B). The diameter of sec3-913 cells increased after one generation at 36°C and continued to enlarge upon longer incubation. The more severe sec3-916 mutant displayed the same phenotype even at the permissive temperature. At the restrictive temperature, this phenotype worsened and cells became misshaped. The extent of this enlargement was quantified by measuring the ratio cell length versus cell width (Figure 5B). sec8-1 cells also showed a shape phenotype, albeit modest, but exo70Δ cells did not.

Because this phenotype was reminiscent of an actin defect, we tested the ability of mutant colonies to grow in the presence of the actin antipolymerizing drug Latrunculin-B (Lat-B; (52)). In the growth test in Figure 5C, for3Δ cells were used as a positive control. sec3-913 appears more sensitive to 2.5 μM Lat-B at the semi-restrictive temperature 32°C, than the wild-type. Thus, Sec3 plays a critical role in actin-mediated cell morphogenesis.

In yeasts, F-actin is found in three independent structures, namely the CAR, actin cables and actin patches. We therefore set out to investigate which of these structures could account for the hypersensitivity of sec3 mutants to the actin poison.

SpSec3 orchestrates CAR constriction and disassembly

We first followed actomyosin ring assembly, contraction and disassembly by time-lapse imaging of the myosin light chain Rlc1. In control conditions during mitosis, Rlc1-mCherry organized as a ring from a broad band of nodes, which constricted until the ring was totally closed and looked like a single dot in the middle of the cell. The CAR then disassembled and the Rlc1-mCherry signal vanished. At 36°C, sec3-913 cells initially formed an apparently normal actin ring but its constriction was substantially delayed (Figures 5D and S3). At the end of constriction, the ring failed to close completely, leaving a cytoplasmic bridge between the two daughter cells. Furthermore, Rlc1-mCherry was seen in filamentous extensions that projected from the unclosed ring toward the cytoplasm of each daughter cell (Figure 5E).

SpSec3 mediates actin cable organization

To image actin cables, we used the F-actin marker LifeAct-GFP (Figure 5F). In control conditions, actin filaments run along the long axis of the cells. Interestingly, in the sec3 ts mutants at the restrictive temperature, filaments appeared faint or absent. This result was confirmed by staining of fixed cells with the F-actin probe Bodipy-Phallacidin (Figure S4). Consistent with an absence of actin cables, time-lapse imaging showed that the long linear translocation of Myo52-GFP along actin cables was abrogated in the sec3 ts alleles (Movie S1). Unlike in sec3-913, sec3-916 and for3A cells, and as previously reported (35), we could detect actin cables in sec8-1 cells, but it is possible that those were remnants of the CAR filaments (Figure 5F).

Actin filament nucleation is achieved by the formin For3 (53). Although For3-3GP was localized at the tips and equator of wild-type cells, in sec3-913 cells the signal was detached from these regions (Figure 6A). This was not a general consequence of altered cell
Figure 5: Sec3 controls actin organization and cell morphology in fission yeast. A) Phase contrast pictures of wild-type, sec3-913 and sec3-916 cells grown at the permissive temperature 27°C, or switched to the restrictive temperature 36°C for one generation. Besides a septation phenotype mutant cells also show an increased diameter. See also Figure 1B. B) Shape phenotype of indicated strains, expressed as the ratio between cell length and cell width. Dark and pale colors show results obtained for cells grown at 27°C or 36°C, respectively. Boxes enclose 50% of the data and lines within the box represent median ratio. Whiskers mark minimum and maximum values. Results are representative of at least two independent experiments. N cells >45. C) Serial dilutions of wild-type, sec3-913 and for3Δ spotted on plates containing either DMSO or 2.5 μM Lat-B and incubated at 27°C or 32°C. The sec3 mutant is hypersensitive to the actin poison. D) CAR constriction rate measured in wild-type (left panel) and sec3-913 (right panel) cells grown at 36°C. Results are representative of at least two independent experiments. See also Figure S3. E) Whole cells pictures showing Rlc1-mCherry-containing rings at the end of constriction. Pictures of one wild-type cell and three representative sec3-913 cells were extracted from time-lapse movies. F) LifeAct-GFP localization in live cells. Actin cables are reduced in sec3 mutants and patches are delocalized. Cytoplasmic projections of an unclosed ring described in (E) can be seen in the sec3-913 cytokinetic cell. sec3-913 cells frequently show short, entangled, actin filaments accumulated at cell tips (higher magnifications of boxed areas). See also Figure S4. Bars = 5 μm and enlargement bar = 2 μm.

polarity because the localization of For3-associated and polarisome members Bud6 and Tea1 (54) were not altered in these cells (Figure 6A and data not shown). Reciprocal co-immunoprecipitations further showed that For3-4Myc and Sec3-3PK specifically interact (Figure 6B). This interaction is weak and possibly transient, which may explain why a colocalization between For3-3GFP and Sec3-tdTomato is difficult to visualize in vivo. We were nevertheless able to find occasional overlaps between the two proteins (Figure 6C). These results suggest that

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SpSec3 docks the formin For3 at the plasma membrane and controls the polarized assembly of actin cables.

Finally, using both LifeAct-GFP (Figure 5F, enlargements) and Bodipy-Phallacidin staining (Figure S4), we noticed an accumulation of entangled actin filaments at the tip of sec3-913 cells. The nature of these structures and the reason for their aggregation are at present unknown but may reflect defects to release or crosslink newly formed filaments at the cell tip.

**SpSec3 interacts with the endocytic machinery and controls patch internalization**

LifeAct-GFP and Bodipy-Phallacidin also marked the actin patches, which were restricted to the growing tip(s) and to the cell center in control cells. Strikingly, in sec3-913 at 36°C and in sec3-916 at any temperature, patches were spread throughout the cell surface (Figures 5F and S4).

Patches in yeasts function in endocytosis and their mis-localization in sec3 mutants suggested that endocytosis may be affected in these cells. Endocytosis was monitored by following FM4-64 uptake over time (Figure 7A). As described previously (55), upon addition to a culture of wild-type cells, FM4-64 first associates with the plasma membrane at zones of active growth and is incorporated into endosomes which later fuse to the vacuolar membranes. Typically, in wild-type cells the signal at the cell poles decreases rapidly and within 15 min following the addition of FM4-64 ~15% of the cells still showed a tip signal. By contrast, FM4-64 incorporation in sec3-913 endosomes was delayed and virtually all cells still showed tip localization after up to 1 h. Hence, sec3-913 cells are defective in endocytosis.

To further explore the role of SpSec3 in endocytosis, we analyzed the dynamic behavior of the actin patch component Sla2 (30) in live cells. At 36°C, patches in sec3-913 were twice as long-lived as in the wild-type (wild-type = 21.23 ± 4.04 seconds; sec3-913 = 48.67 ± 6.98 seconds; n > 81; Figure 7B). sec3-913 patches also moved more slowly than wild-type patches (at 36°C, wild-type = 76 ± 5 nm/second; sec3-913 = 49 ± 7 nm/second; n > 81). Similar results were obtained with other actin patch markers (Arc3-mCherry, Chc1-mCherry, Fim1-EGFP, data not shown) (56). To determine which step of endocytosis is defective in the sec3-913 mutant, we generated kymograph representations of single whole cells over time. In agreement with results presented above (Figure 5F), Sla2-GFP patches were scattered along the long axis of the sec3-913 cells compared to polar or equatorial localization in wild-type cells (Figure 7C). The kymographic profile of wild-type patches matched previous descriptions, which included the local recruitment of the patch machinery at the cell cortex, the directional internalization of the vesicle away from the cortex and towards the cell interior, and the dissociation of the endocytic machinery from the internalized vesicle events (56). In sec3-913 cells, the patches were properly assembled at the cell cortex but were not pulled into the cytoplasm nor were they disassembled (Figure 7C). The same conclusions were drawn using Arc3-mCherry (data not shown).

We next asked if the role of SpSec3 in endocytosis is direct and if it can interact with actin patch components. Figure 7D (left panel) shows that SpSec3-3PK co-immunoprecipitated with Sla2-GFP. A weak interaction was also detected with Arc3-3HA (data not shown).
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Figure 7: SpSec3 controls actin patches localization and internalization. A) Wild-type and sec3-913 live cells grown at 36°C were incubated in the presence of FM4-64 and photographed at regular intervals for up to 1 h following addition of the dye. Two selected time points are shown. The accumulation of FM4-64 at the tip of sec3-913 cells and its absence from endosomes and vacuolar membranes are indicative of a defect in endocytosis. Bar = 5 μm. B) Quantification of the average lifetime (top panel) and average velocity (bottom panel) of Sla2-GFP patches from wild-type (gray) and sec3-913 (black) cells cultured at the indicated temperatures. N patches > 81; N cells > 4. Error bars are SEM of individual cells. C) Kymographs of Sla2-GFP signals in representative wild-type (top panel) and sec3-913 (bottom panel) cells. The x and y axis represent cell length and time, respectively. Higher magnifications of the boxed areas show that patches are not internalized in sec3-913 cells. D) Co-immunoprecipitation (IP) of Sec3-3PK with Sla2-GFP. Negative controls include IPs of protein extracts in which either Sec3 or Sla2 was un-tagged (left panel). WCE: whole cell extract. Asterisk: unspecific band. Note that mutant Sec3-913-3PK, although less expressed than wild-type Sec3-3PK, remains capable of interacting with Sla2-GFP (right panel).

Although mutant Sec3-913 remained capable of interacting with Sla2 (Figure 7D, right panel), the reduced level of the mutant protein likely explains the dispersion of patches (Figures 2E and 7D).

Taken together these results indicate that Sec3 is involved not only in actin patches localization but also in internalization.

Discussion

The exocyst is a conserved complex of eight proteins that tethers post-Golgi secretory vesicles to the plasma membrane before docking and fusion. Orthologs of all but one have been characterized or predicted in S. pombe. In this study, we identified SpSec3, the missing component of the fission yeast exocyst complex. We show that SpSec3 has a role in vesicle tethering that does not involve the recruitment of other exocyst components. SpSec3 is also required for the organization of all three yeast actin structures, the actin patches, actin cables and CAR. We discuss below how this dual role may be geographically and functionally coordinated and how we envision the role of Sec3 family proteins in membrane trafficking and cell morphogenesis.

Sec3 is not a landmark for secretion in fission yeast

In the currently accepted mechanism of secretion, Sec3 and Exo70 are present at specialized sites of the plasma membrane where they interact with the six other exocyst subunits associated with the arriving vesicle, thus capturing vesicles and targeting secretion (9–11,47). Our data argue against this simple ‘landmark model’. Indeed, we found that the exocyst subunit Sec8 is polarized independently of its interaction with Sec3. Firstly, Sec8 was properly localized in cells expressing low levels of Sec3-913 mutant protein. Secondly, the ts phenotype of the sec3-913 strain was rescued by over-expressing the mutant protein, which is otherwise unable to interact with Sec8. This strongly indicates that the secretion defects observed in sec3 ts mutants are not due to a failure to polarize the exocyst complex. The idea that Sec3 may not

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target the exocyst to sites of polarized growth is not novel. Similar to our findings, in a budding yeast strain in which the PH domain of Sec3 was deleted, the truncated product was absent from the plasma membrane but the exocyst complex was properly localized (57,58). Furthermore, the polarized recruitment of other exocyst subunits to the S. cerevisiae daughter cell is unaffected in a sec3 strain deleted of its exocyst-binding region (59), and even in a sec3Δ strain (60). The exocyst complex may be recruited by Exo70 in the absence of Sec3 (61). However, the possibility of a redundancy between Sec3 and Exo70 is itself disputed (58). The allosteric model proposes that the polarization of the exocyst machinery is achieved by the local activation of a pre-assembled exocyst by regulatory GTPases present at key cortical sites (58). In fact, most, if not all, members of the exocyst complex appear to depend on one another for their localization and/or assembly (6,59,62,63). It is likely that the highly regulated delivery, recruitment and ordered assembly of subunits by one another are responsible for the formation of the whole complex and its targeting to sites of active growth. The role of each subunit and of Sec3 in particular essentially remains to be determined.

**SpSec3 is required for cell morphology**

The involvement of the exocyst complex in defining a direction of growth and, hence polarity has been established in a range of cell types from budding yeast to growing plant cells and from neurons to cilia (64–66). In fission yeast, cell polarity is not only a simple function of exocytosis but also requires actin cables (35,67,68). In line with this, our sec3 mutants, which are defective in secretion and possess no actin cables also display a shape phenotype. The current model proposes that the exocyst complex and actin cables are two independent and parallel morphogenesis pathways, both downstream of Pob1 and under the lid of Cdc42 (35,67,68). Our results on the contrary show that the two structures are physically and functionally bridged by Sec3 (Figure 8).

Like SpSec3, the Boi-family protein Pob1 interacts with both For3 and the exocyst complex (68,69). Pob1 recruits Sec6, Sec8 and Sec10 to cell tips, and our preliminary data suggest that it also positions Sec3. SpSec3 does not control the recruitment of other exocyst subunits to the plasma membrane. Thus, the function of Pob1 on the exocyst complex is unlikely to be solely through Sec3. Alternatively, Sec3 and Pob1 may play complementary and overlapping roles in polarized growth (68).

The sec8-1 allele has a much stronger phenotype than any of our two sec3 mutants (our personal observation), but we and others (35) were able to visualize actin cables in these cells, whose diameter was only mildly enlarged. Similarly, exo70Δ cells are not misshapen (35,36). This raises an intriguing question as to whether SpSec3 is unique among the exocyst complex components. It is formally possible that the lack of other exocyst mutants has precluded a comprehensive analysis of the role of the exocyst as a whole in cell morphology. On the other hand, although the actin patches looked essentially normal in sec8-1 cells, equatorial but not polar endocytosis was shown to be inhibited in these cells (55). We envisage that SpSec3 performs multiple tasks by forming various subcomplexes with some, but not all, members of the exocyst complex. Multiple combinations of exocyst components could be a means to provide functional diversity besides secretion, and possibly target different functions to different cellular locations (70).

**Specificity of the actin phenotypes observed in the sec3 mutants**

One of the important findings of this study is that all three fission yeast actin structures were affected in sec3 ts mutants. The constriction and disassembly of the actomyosin ring were delayed; actin patches were delocalized and failed to invaginate; and actin cables were faint or absent.

The actin phenotype of the sec3 mutants is not a secondary consequence of their defective secretion. Although not tethered/fused with the plasma membrane, secretory vesicles in the sec3 mutants accumulated at the cell poles. Thus, if the actin patch and cable components were transported by vesicles, they would also amass in these regions, in a manner similar to Syb1. Instead, we found that they were randomly distributed throughout the cell.

We considered the possibility that SpSec3 could be responsible for the localization or regulation of some major polarity cues, like Cdc42 or phosphoinositides. However, PI(4,5)P2 and their generating enzyme Its3, are not polarized in fission yeast and unlikely to be controlled by Sec3 (48). Moreover, the polarized localization of active Cdc42-CRIB is not affected in sec3 and sec8 mutants (data not shown, (35)). Finally, we have shown that, except for actin, a range of other polarity markers (exocyst, v-SNARE, Myosin V, polarisome) were properly localized at zones of growth and division, which would not be the case if Cdc42 was mis-targeted. Hence the role of SpSec3 in cell morphology is probably not through Cdc42.

Another possibility to explain the ubiquitous effect of Sec3 on actin could be that it specifically controls one structure only, which in turn would indirectly affect the two others. The non-disassembled CAR may sequester a pool of G-actin that would therefore not be available for polymerization into actin filaments or patches (71). This could result from different actin-binding proteins competing for the formation and stabilization of preferential types of actin structures (72,73).

Alternatively, Sec3 may control all three actin structures individually. We favor this hypothesis because SpSec3 physically interacts both with actin patches components and with the actin cable nucleation machinery. This is in accord with other reports that have shown an association of some members of the exocyst with the endocytic or
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Figure 8: Model for the dual role of SpSec3 in secretion and actin organization. As part of the exocyst complex (green), Sec3 (red) has a canonical role in tethering secretory vesicles (v) with the plasma membrane, before docking by SNAREs (gray) and fusion. It also physically binds actin-binding proteins (For3, Sla2 and probably others) and clusters them at the vicinity of their activators (see text). Sec3 therefore bridges the secretion machinery to all three actin structures in fission yeast, the actin cables (red), actin patches (blue) and actomyosin ring (orange). Upstream polarity cues orchestrate the presence of the recruiter Sec3 and the activators in the same areas of the plasma membrane.

actin nucleation machineries (74,75). We speculate that Sec3 may also interact with some CAR components, via a mechanism that is common to all three actin structures (see below).

SpSec3 couples the formin For3 with its activators

For3 is a typical formin, which contains a Dia inhibitory domain (DID) and a Dia autoregulatory domain (DAD). In a closed conformation, the DID and DAD regions interact with one another and inhibit For3. Upon binding of the DID domain with Pob1 and Cdc42 and of the DAD domain with Bud6, For3 adopts an open and active conformation (69,76). Once de-inhibited, For3 nucleates short actin filaments that will later be bundled into long actin cables (76).

Our results identify SpSec3 as a new factor which is essential for the localization of For3 at the plasma membrane. The function of For3 is intrinsically linked to its localization and accordingly the sec3 mutants had no actin cables. For3, but not Bud6 or Cdc42, was delocalized in the sec3-913 mutant. We therefore conclude that SpSec3 is at least necessary to tether For3 to the polarized areas of the plasma membrane and to couple it to its activators. It is also possible that Sec3 binds to the DID or the DAD domains and directly relieves For3 autoinhibition in a manner similar to Bud6 and Pob1/Cdc42.

SpSec3 is at the meeting point between endocytosis and exocytosis

A role of Sec3 as a link between the cell surface and the actin machinery is also demonstrated by the dispersion of actin patches in the sec3 mutants.

The delocalization of the patches in the sec3 mutants is not a consequence of an absence of actin cables. Indeed, patches are hardly delocalized in for3Δ cells (33,55,77) and endocytosis is insensitive to concentrations of Latrunculin-A that depolarize cables (55). Moreover, the patches assemble, invaginate and disassemble locally at the plasma membrane (Figure 7C), which supports the idea that patches exist without cables.

Membrane proteins and lipids that have been internalized by endocytosis can be recycled back to the cell surface. A number of studies conducted in various organisms have reported a localization of exocyst subunits in recycling endosomes and their regulation by recycling endosome-specific GTPases such as Arf6 and Rab11 (75,78–81). Although we have not been able to visualize Sec3 in endosomes, its interaction with endocytic patches was detected by immunoprecipitation. Moreover, its localization appeared to be dependent on the Rab11 GTPase Ypt3. It is intuitive that Sec3 drives the tethering of recycling vesicles with the plasma membrane along with the rest of the exocyst complex. How it may control patch internalization is however less clear. It is possible that sec3 mutants simply fail to recycle factors that define the localization and functionality of actin patches.

All early and late endocytic markers (Chc1, Sla2, Arc3, Fim1 and actin, (56)) that we have tested seemed to assemble into (misplaced) patches. Yet it is clear that these structures were not functional. Their increased lifetime and lower speed in the sec3 mutants suggest that the patches were waiting for a signal to progress through internalization. The propelling force behind patch invagination is provided by actin. If Sec3 plays a similar role in endocytic patches as it does in actin cables, one can imagine that compromised Sec3 causes the declustering (e.g. by poor recycling) of the assembly-competent actin machinery away from its activator(s). As human Sec3/EXOC1 is able to rescue Spsec3 mutants, it would be of particular interest to see whether vertebrate Sec3 plays an analogous role in actin organization.

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Materials and Methods

Yeast genetics and culture

Media, growth, genetics and maintenance of strains were as described in (82). Cells were cultured to mid-log phase in YESS at 27°C and subsequently shifted to 36°C for 3.5 h before observation unless otherwise stated. For simplicity and unless specified, only experiments carried out at 36°C are shown.

For spot tests all strains were adjusted to the same cell concentration of 2 x 10^7 cells/mL and 10-fold dilutions were spotted on plates. Plates were incubated at a temperature range of 27–36°C. Lat-B (Calbiochem) was prepared in dimethyl sulphoxide (DMSO) at a concentration of 5 mg/mL (12.84 μM) and diluted into YESS to 0–10 μM prior to pouring into small 60 x 15 mm dishes. For the rescue experiment of sec3-913 by human EXOC1, cells transformed with the individual plasmids were grown in minimum medium (nitrogen-based EMM2 or glutamate-based GMM) containing 15 μM thiamine, washed five times in H_2O and spotted onto EMM or GMM plates with or without 15 μM thiamine. Only selected conditions of temperature, concentration or medium are shown.

Yeast strains and construction of sec3 temperature sensitive mutants

Yeast strains used in this study are described in Table S1.

For isolation of temperature-sensitive (ts) sec3 mutants, sec3 fragments followed by a 3'-HphR cassette were randomly mutagenized by error-prone PCR amplification using LA-Taq DNA polymerase (Takara). Twenty PCR products (50 μL per tube) were individually ethanol-precipitated and transformed into a wild-type strain. Hph-resistant transformants were then replica-plated on YESS + phloxin-B plates and incubated at 27°C and 36°C. Eleven clones that grew at 27°C but not at 36°C were selected. These were back-crossed to ensure proper integration and to remove potential off-site mutations. Two alleles, sec3-913 and sec3-916, were chosen for further characterization. Sequencing of sec3-913 identified two silent mutations (nucleotides A1260G and T1290C), that resulted in amino acid substitution W150R and T541L. sec3-916 had two silent mutations (nucleotides T930C and T1671C) and three missense mutations (nucleotides A1260G, A50C and T1729C) that lead to two amino acid substitutions N17P and C577R.

Carboxy-epitope-tagged proteins were generated via chromosomal integration of PCR-amplified fragments (83,84). Some Sec3-tagged fusion proteins rendered cells sick under a number of conditions and were not fully functional. However, as far as we are aware, Sec3-GFP cells grown in rich medium were healthy at all temperatures.

The strain sec3-913-3pk was generated by transforming a 3pk-kan<sup>R</sup> PCR fragment into a sec3-913-hph<sup>R</sup> strain. Homologous recombination in 3′ of the ORF was checked by selection in the presence of G418, counter-selection on Hph plates and by colony PCR. The presence of the tag did not modify the ts phenotype (not shown) and the mutant protein migrated at the same size as its wild-type counterpart on denaturing gels. Sec3-913-3PK was however less detectable by immunoblotting than wild type Sec3-3PK (see for example Figure 2E).

Similarly, strain mmt1-P3-kan<sup>R</sup>-gfp-sec3<sup>+</sup> and sec3-913, were obtained by transforming an mmt1-P3-kan<sup>R</sup>-gfp PCR fragment into a wild-type and sec3-913-hph<sup>R</sup>, respectively.

Plasmid constructs

DNA coding for S. pombe Sec3 (SPAC17G8.12; NC_003424.3) was amplified by PCR from wild-type genomic DNA. The cDNA of transcript variant 1 of H. sapiens EXOC1 (NP_060731) cloned in pCMV6-XL5 was purchased from Origene and the insert was re-amplified by PCR. In both cases, cloning into pREP1 was achieved by using primers containing NdeI and BamHI restriction sites (Sigma-Proligo). All constructs were verified by sequencing.

Cell imaging and fluorescence microscopy

Live cell imaging was performed in an imaging chamber (CoverWell 20-mm diameter, 0.5-mm deep; Molecular Probes) filled with 800 μL of 20% agarose in YE5S at 27°C and seeded with a 22 x 22 mm glass coverslip. Cells were imaged using an Olympus IX71 wide-field inverted epifluorescence microscope with the DeltaVision-SoftWoRx system (Olympus and Applied Precision Co.), in a temperature controlled environmental chamber. Olympus UPlanSapo 63x or 100x NA 1.4, oil immersion objective were used and images captured with a Coolsnap-HQ digital CCD camera or a Cascade EMCCD 512B camera (Roper Scientific). Pictures of Z-sections were deconvolved and projected. Counts, measurements and image presentations were made using Metamorph (Molecular Devices Corporation) and downloaded to Microsoft Excel or Prism for analysis.

A stock solution of calcofluor white (fluorescent brightener 28 Sigma) was prepared at 5 mg/mL in H_2O vortexed for up to 24 h and centrifuged to eliminate the undissolved powder. One microliter of this suspension was pipetted onto cells pre-applied to an agarose pad, and a coverslip was sealed. Cells were observed immediately.

Where two strains were visualized simultaneously on the same microscope slide, identification of the wild-type was achieved by pre-staining with fluorescent lectin. Soybean lectin AlexaFluor 594 or 488 conjugate (Molecular Probes) was dissolved at 2 μg/mL in H_2O. One microliter of this stock solution was added to 500 μL cells (to a final concentration of 4 μg/μL). After 10–30 min incubation in the dark, cells were washed at least five times with YESS, mixed with the unstained strain and observed immediately.

For live-cell imaging of actin patches speed was favored and three optical sections only, spanning a total of 0.6 μm below the plasma membrane were acquired every 0.5 seconds (Ar3-mCherry) or 1–2 seconds (Sla2-GFP).

Rsc1-mCherry expressing cells were incubated at 27°C or at 36°C for 2 h prior and imaged for another 2 h at the respective temperature. Z-stacks were acquired every 4 min.

Endocytosis was monitored by following FM4-64 uptake over time using a protocol described previously (55,85) with minor modifications. In brief, FM4-64 (Molecular Probes) was dissolved in DMSO at a concentration of 1.64 μM (1 mg/mL). At time 1=0, 2 μL of this stock solution were added to 400 μL cells (to a final concentration of 8.2 μM). The cells were quickly applied to the agarose pad and photographed at various time points. The efficiency of endocytosis was evaluated by scoring the number of cells at each time point that had taken up FM4-64 and delivered it to vacuoles.

For staining of the actin cytoskeleton, cells were fixed for 1 h in 3% paraformaldehyde. After three washes in PEM buffer (200 mM Pipes, 2 mM EGTA, 2 mM MgSO_4, pH 6.9) the cell wall was digested for 20 min at 37°C with 500 μL of 3 mg/mL zymolyase-20T re-suspended in PEMS (PEM + 1.2 μg/mL Zymolyase). Cells were permeabilized with 500 μL of 1% Triton X-100 in PEMS, washed thrice in PEMS and were incubated for 30 min with constant mixing in a solution containing 50 μL PDM and 1 μL BODIPY-FL-Phallacidin 6.6 μM (200 U/μL dissolved in DMSO).

Acid phosphatase secretion assay

Acid phosphatase secretion was assayed as described previously (36,86,87) with minor modifications. In brief, cells were grown to mid-log phase in YE5S at 27°C and all strains were adjusted to the same concentration. Cultures were washed twice in medium and split into two halves for further incubation at 27°C and 36°C. At indicated time points, the absorbance of the culture was measured at 600 nm. In parallel, a
500 μL sample was incubated for 12 min at 30°C in the presence of 500 μL substrate solution (2 mM p-nitrophenyl phosphate, 0.1 M sodium acetate pH 4.0 pre-warmed to 30°C). The reaction was stopped by addition of 500 μL of 1 M NaOH. OD 405nm was measured at each time point, using a sample with no cell as a blank. Values of acid phosphatase activity were expressed as a function of cell density and all curves were adjusted to the same value at t = 0.

Co-immunoprecipitation
Cells were lysed in extraction buffer [50 mM HEPES, 50 mM NaF, 50 mM Na3-glycerophosphate, 5 mM EGTA, 5 mM ethylenediaminetetraacetic acid (EDTA), 0.2% Triton X100, 1 mM Protease Inhibitor Cocktail, 1 mM phenylmethylsulphonyl fluoride (PMSF)] by the mechanical action of acid-washed glass beads (FastPrep FP120 apparatus (Savant. Co.) 2 x 25 seconds, power 5.5). Debris were eliminated by centrifugation for 1 min, then 5 min at 13000 x g at 4°C. Protein concentrations were determined by Bradford assay (Bioread). For co-immunoprecipitations, and unless stated otherwise, 4–10 mg of WCE was incubated for 2 h at 4°C with protein-A Dynabeads coated with antibodies against 3PK (mouse monoclonal Serotec), Myc (rabbit polyclonal, Babco) or GFP (rabbit polyclonal, Invitrogen). Beads were then extensively washed (50 mM Tris–HCl, pH = 7.4, 1 mM EDTA, 150 mM NaCl, 0.05% NP-40, 10% Glycerol, 1 mM DTT, 1.5 mM PNPP, 1 x Protease Inhibitor Cocktail, 0.1 mM PMSF) and boiled for 5 min in Laemmli sample buffer. Thirty micrograms of WCE was similarly treated and used as controls. Proteins were separated on denaturing 4–12% gradient gels (BioRad) transferred onto PVDF membranes which were further blocked and immunoblotted in the presence of 10% nonfat milk. Primary antibodies against 3PK (mouse monoclonal, Serotec), Myc (BE10 mouse monoclonal, Babco or GFP (mouse monoclonal, Roche) were used diluted to 1:1000 in ImmunoShot Solution 1 (2B Scientific). For secondary antibodies, anti-mouse horseradish peroxidase (HRP)-conjugated (GE-Healthcare) or TrueBlot (eBioscience) were used at a dilution of 1:2000–1:4000 in ImmunoShot Solution 2 (2B Scientific). Signals were detected using enhanced chemiluminescence (ECL; GE Healthcare).

Electron microscopy
Samples were cryofixed by high-pressure freezing using a Leica EM PACT2. Freeze substitution was performed in anhydrous acetone containing 0.01% osmium tetroxide, 0.1% glutaraldehyde and 0.25% uranyl acetate for 3 days at −90°C using a Leica EM AF5. Freeze-substituted cells were gradually warmed to −20°C and were finally infiltrated with Epon. Serial thin sections of ~60 nm were cut using a Leica Ultracut UCT microtome and placed on pioloform-coated slot grids. Sections were post-stained with 2% uranyl acetate in 70% methanol for 5 min at room temperature. The reaction was stopped by addition of 5% glycerophosphate, 5 mM EGTA, 5 mM ethylenediaminetetraacetic acid (EDTA) with protein-A Dynabeads (FastPrep FP120 apparatus (Savant. Co.) 2 x 25 seconds, power 5.5). Debris were eliminated by centrifugation for 1 min, then 5 min at 13000 x g at 4°C. Protein concentrations were determined by Bradford assay (Bioread). For co-immunoprecipitations, and unless stated otherwise, 4–10 mg of WCE was incubated for 2 h at 4°C with protein-A Dynabeads coated with antibodies against 3PK (mouse monoclonal Serotec), Myc (rabbit polyclonal, Babco) or GFP (rabbit polyclonal, Invitrogen). Beads were then extensively washed (50 mM Tris–HCl, pH = 7.4, 1 mM EDTA, 150 mM NaCl, 0.05% NP-40, 10% Glycerol, 1 mM DTT, 1.5 mM PNPP, 1 x Protease Inhibitor Cocktail, 0.1 mM PMSF) and boiled for 5 min in Laemmli sample buffer. Thirty micrograms of WCE was similarly treated and used as controls. Proteins were separated on denaturing 4–12% gradient gels (BioRad) transferred onto PVDF membranes which were further blocked and immunoblotted in the presence of 10% nonfat milk. Primary antibodies against 3PK (mouse monoclonal, Serotec), Myc (BE10 mouse monoclonal, Babco or GFP (mouse monoclonal, Roche) were used diluted to 1:1000 in ImmunoShot Solution 1 (2B Scientific). For secondary antibodies, anti-mouse horseradish peroxidase (HRP)-conjugated (GE-Healthcare) or TrueBlot (eBioscience) were used at a dilution of 1:2000–1:4000 in ImmunoShot Solution 2 (2B Scientific). Signals were detected using enhanced chemiluminescence (ECL; GE Healthcare).

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Supporting Information
Additional Supporting Information may be found in the online version of this article:

Table S1: Schizosaccharomyces pombe strains used in this study

Table S2: Genetic interactions between sec3 and other genes

Figure S1: Partial alignment of the primary sequences of S. pombe and S. cerevisiae Sec3. S. pombe Sec3 (Sp, 603 aa) is approximately twice as short as its budding yeast ortholog (Sc, 1136 aa). Only regions of special interest are shown. Arrows indicate the PH domain (aa 71–241 in Sc; (45)). Crystalized (Sc) or predicted (Sp) secondary structures are boxes in pink (α-helices) and blue (β-strands; (45)). Experimentally characterized amino acids responsible for Rho1-binding are shown in red (88). The three clusters of phosphoinositides-interacting residues are shown in red (88). (Sp) and (Sc) in orange (12). Mutations in the fission yeast sec3 mutants used in this study are highlighted in cyan (sec3-913) and in magenta (sec3-916).

Figure S2: Localization of Sec3 during the fission yeast cell cycle. Time-lapse of Sec3-tdTomato imaged together with the CAR marker Myo2-GFP (A) the septin Spn3-GFP (B) or the calcofluor-stained cell wall (C). For each marker three different panels show three different cells at progressive stages of mitosis and cytokinesis. Sec3-tdTomato is localized at growing tips in interphase (arrows) and as a single (open arrowhead), then double (plain arrowheads) medial ring at cytokinesis. Time indicated in minutes. Bars = 5 μm.

Figure S3: SpSec3 controls CAR constriction. Time-lapse imaging of the CAR marker Rcl1-mCherry observed in representative wild-type and sec3-913 cells, at 27°C or 36°C. Four minutes time increments are shown. Arrowheads point at projections of CAR filaments into the cytoplasm, also shown in Figure 5E. Bar = 5 μm.

Figure S4: Unlike sec8-1, sec3 ts mutants have weak actin cables and delocalized actin patches. Bodipy-Phallacidin staining of fixed wild-type, sec3-913, sec3-916 and sec8-1 cells grown at 27°C or at 36°C for 3.5 or 6 h. See also Figure 5F. Bar = 5 μm.

Movie S1: Time-lapse imaging of Myo52-GFP in wild-type (left) and sec3-913 (right) cells grown at 36°C for 3.5 h. Arrowheads point at examples of moving motors along actin cables. Note the near absence of long-range movement in sec3-913 cells. Time indicated in sec:min. Bar = 5 μm.

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