Anillin/Mid1p interacts with the ESCRT-associated protein Vps4p and mitotic kinases to regulate cytokinesis in fission yeast

Imane M. Rezig, Wandiahyel G. Yaduma, Gwyn W. Gould & Christopher J. McInerny

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**ABSTRACT**

Cytokinesis is the final stage of the cell cycle which separates cellular constituents to produce two daughter cells. Using the fission yeast *Schizosaccharomyces pombe* we have investigated the role of various classes of proteins involved in this process. Central to these is anillin/Mid1p which forms a ring-like structure at the cell equator that predicts the site of cell separation through septation in fission yeast. Here we demonstrate a direct physical interaction between Mid1p and the endosomal sorting complex required for transport (ESCRT)-associated protein Vps4p, a genetic interaction of the *mid1* and *vps4* genes essential for cell viability, and a requirement of Vps4p for the correct cellular localization of Mid1p. Furthermore, we show that Mid1p is phosphorylated by aurora kinase, a genetic interaction of the *mid1* and the aurora kinase *ark1* genes is essential for cell viability, and that Ark1p is also required for the correct cellular localization of Mid1p. We mapped the sites of phosphorylation of Mid1p by human aurora A and the polo kinase Plk1 and assessed their importance in fission yeast by mutational analysis. Such analysis revealed serine residues S332, S523 and S531 to be required for Mid1p function and its interaction with Vps4p, Ark1p and Plk1p. Combined these data suggest a physical interaction between Mid1p and Vps4p important for cytokinesis, and identify phosphorylation of Mid1p by aurora and polo kinases as being significant for this process.

**Introduction**

Much is understood about the control and regulation of the cell division cycle, with many critical cell cycle mechanisms identified being evolutionarily conserved, present across the eukaryotic kingdom. One cell cycle stage that has been increasingly studied is cytokinesis where DNA, organelles and cell constituents are partitioned and allocated to two daughter cells during their physical separation. The fission yeast *Schizosaccharomyces pombe* has proven to be an excellent model organism to study the eukaryotic cell cycle [1]. It is especially useful for studying cytokinesis and cell division as, explicit in its name, it divides by medial fission involving a contractile actomyosin ring leading to the process of cell separation, which is similar to these processes in mammalian cells [2].

A number of proteins have been identified that regulate cytokinesis in fission yeast including those in a signal transduction pathway named the Septation Initiation Network (SIN) [3]. The SIN proteins form a pathway that facilitates contractile ring constriction and promotes the formation of a medial cell wall-like structure between the two daughter cells, called the septum. Furthermore, SIN components associate with the spindle pole bodies and link mitotic exit with cytokinesis [4]. Sid2p is one regulator of the SIN-cascade of signaling proteins [5]. It terminates the signaling cascade leading to the transition from the spindle pole bodies to the cell division site promoting the onset of cytokinesis [6]. A recent study identified anillin/Mid1p as a substrate for Sid2p and indicated that the phosphorylation of Mid1p facilitates its removal from the cell cortex during the actomyosin contractile ring constriction [7].

Mid1p forms a ring-like structure at the cell equator that predicts the site of cell separation through the formation of a septum in fission...
yeast [8–10]. In cells containing a chromosomal deletion of the mid1 gene (mid1Δ), a misshaped contractile ring is assembled during anaphase when the SIN pathway becomes active; such observations confirm the important role of Mid1p in directing contractile ring assembly to its correct location [11–13].

Many other groups of proteins have a role in septum formation in S. pombe. Among these the various classes of ESCRT proteins, including the ESCRT-III regulator Vps4p, were found to be required for septation, suggesting that they have a role in cytokinesis in fission yeast [14,15]. Additional experiments suggested that the ESCRT proteins interacted with established cell cycle regulators including the polo kinase Plo1p, the aurora kinase Ark1p and the CDC14 phosphatase, Clp1p to control these processes [14].

In mammalian cells the absence of the ESCRT proteins leads to inhibition of cytokinesis [15–18]. The ESCRT machinery is assembled to catalyze membrane remodeling events allowing for the final separation of the daughter cells [19]. This process is mediated by many cytokinesis proteins including ESCRT-I and III classes of proteins in addition to the ATPase VPS4 [20].

As a way to further understand the regulation of ESCRT proteins during cytokinesis in fission yeast we sought to identify other proteins that interact with anillin/Mid1p. Here we describe a direct physical interaction between Vps4p and the anillin Mid1p which is required for cell viability and for the correct placement of Mid1p. We further show an interaction between Mid1p and the aurora kinase Ark1p which is required for cell viability and identify phospho-acceptor sites in Mid1p and study their role using mutagenesis. Collectively, our observations reveal novel mechanisms by which cytokinesis is regulated by different classes of proteins acting through Mid1p.

Results and discussion

Genetic and physical interactions between Mid1p and Vps4p in fission yeast

Previously, we identified and characterized the requirement and role of ESCRT proteins for cytokinesis in fission yeast [21]. These experiments also revealed the interaction of ESCRT proteins with three cell cycle regulators, the Polo-like kinase Plo1p, the aurora kinase Ark1p and the CDC14 phosphatase Clp1p. These observations offered a framework by which ESCRT proteins are regulated to control cytokinesis. To further explore ways in which the ESCRT proteins might integrate with other cell cycle regulators to control cytokinesis, we examined their interaction with the anillin Mid1p. Mid1p has a central structural role in cytokinesis forming equatorial nodes to create an annular shaped structure that determines the position of the division plane [13,22].

We initiated this by searching for genetic interactions between the mid1 gene and genes encoding ESCRTs and ESCRT-associated proteins. Double mutant fission yeast strains were created containing a chromosomal deletion of mid1 (mid1Δ) and individual chromosomal deletions of ESCRT genes from Classes E-0 to E-III and vps4 and searching for synthetic phenotypes. mid1Δ was combined with sst4Δ (E-0), sst6Δ and vps28Δ (E-I), vps36Δ and vps25Δ (EII), vps20Δ, vps32Δ and vps2Δ (E III), and vps4A [21,23]. In each case double mutants were created which were viable, with no apparent synthetic phenotypes. The exception was mid1Δ vps4Δ which instead failed to form viable colonies, with cells undergoing just a few divisions, and so was synthetically lethal (Figure 1a).

This striking observation demonstrated a genetic interaction between the mid1 and vps4 genes, with one explanation that the two encoded proteins interact to control an essential cellular function. Antibodies to yeast Vps4p are not available to test this hypothesis in intact cells, therefore we expressed and purified a full-length His-tagged version of Vps4p protein from bacteria, using the same method to express and purify three different GST-tagged domains of Mid1p. Full length Mid1p was not purified as it is insoluble under such conditions [22,24].

Full length Mid1p is 920 amino acids in length; the three domains used in pull-down experiments described here encompass the amino acids 1–453 (“N-term”), 452–579 (“Middle”) and 798–920 (“C-term”) [24,25]. Of these three domains, only the C-terminal domain comprised amino acids 798–920 was pulled-down by recombinant Vps4p
This indicates a direct physical interaction between Vps4p and the C-terminus of Mid1p, at least in vitro, and that this interaction involved residues 798–920 of Mid1p.

**Vps4p is required for the correct cellular distribution of Mid1p**

To further understand the interaction between Vps4p and Mid1p we examined the requirement...
for Vps4p to control the cellular distribution of GFP-tagged Mid1p during the cell cycle. Mid1p distribution has been well characterized and shown to move from the nucleus to the equatorial region to form nodes and a medial ring, that predicts the site of cell cleavage during septation [8].

Wild-type cells showed three patterns of GFP-Mid1p: localization to the nucleus, cytoplasmic, and equatorial nodes, as previously reported (Figure 2a) [5,8]). By contrast, GFP-Mid1p in vps4Δ cells showed different patterns. Although cytoplasmic GFP-Mid1p was similar to wild-type, additional plasma membrane localization was observed (Figure 2b). In some cells GFP-Mid1p localized to only one node and the plasma membrane, or localized to three nodes in other cells. The frequencies of these abnormal phenotypes in vps4Δ cells were compared with wild-type and showed significant differences (quantified in Figure 2b for wild-type and Figure 2c for vps4Δ, and compared in Figure 2d). Overall, GFP-Mid1p localization was considerably altered by the absence of vps4+ (Figure 2d).

Figure 2. GFP-Mid1p cellular localization is disrupted in vps4Δ cells. Wild-type (w-t) and vps4Δ (vps4::ura+) S. pombe strains containing GFP-Mid1p grown at 25°C in liquid YE medium to mid-exponential phase and visualized by confocal microscopy. (a) Key to characterized GFP-Mid1p localization phenotypes. Scale bar, 5 µm. Bright-field and fluorescent images of cells of GFP-Mid1p localization phenotypes for wild-type (W-T) (b) and vps4Δ (c) cells. Scale bar, 10 µm. The images presented in (A, B and C) are scanned in a single focal plane using a laser scanning confocal microscope. (d) Two-way ANOVA analysis of frequencies of localization phenotypes in vps4Δ compared to wild-type (W-T). Asterisks (****) denote p values <0.0001 indicating significant differences to wild-type. Error bars, SEM.
These results, in addition to the observation that the mid1 and vps4 genes interact genetically and that the Mid1p and Vps4p physically interact in vitro, suggest that Mid1p and Vps4p coordinate to regulate the S. pombe cell cycle.

**Genetic and physical interactions between Mid1p and Ark1p in fission yeast**

We hypothesized that mid1 and ark1 genes interact to accomplish S. pombe cytokinesis as we and others have shown that aurora kinases modulate ESCRT protein function [17,21,26–28]. To test this hypothesis, we tested whether both genetic and biochemical approaches were taken.

First, a mid1Δ strain was crossed with strains containing temperature sensitive ark1-T11 mutations to generate double mid1Δ ark1-T11 mutant strains. The mid1Δ ark1-TS double mutant failed to form viable colonies (Figure 3a). Such a synthetic lethality indicates a genetic interaction between the mid1 and ark1 genes.

As with mid1 and vps4 such a genetic interaction might be explained by a physical interaction between Mid1p and Ark1p proteins. Since Ark1p is a protein kinase, and any interactions are likely to be low affinity and transient, we tested whether

![Figure 3](image-url)

**Figure 3.** Genetic and physical interactions between mid1 and ark1 in fission yeast. (a) Synthetic lethality in mid1Δ ark1-T11 double mutant cells indicates genetic interaction between mid1" and ark1" genes. Tetrad analysis of h+ ark1-T11 (ark1-T11@Kan) crossed with h- mid1Δ (mid1::ura) to identify mid1Δ ark1-T11 double mutants that show a synthetic lethal growth phenotype. Tetrad created by mating the two strains on solid ME medium, four spores dissected and allowed to grow on solid YE medium until colonies formed. Colonies replicated to solid YE+G418/KanMX4 and EMM-ura medium and incubated to identify growth phenotypes and double mutants. (b) Plk1 and Aurora A kinases phosphorylate Mid1p “N-term” and “Middle” domains. Recombinant GST-tagged Mid1p domain proteins “N-term” (aa 1–453), “Middle” (452–579) and “C-term” (798–920) were purified and analyzed by in vitro phosphorylation assays. In “Exp. A” myelin basic protein (MBP) was used as a positive control for kinase activity. In “Exp. B”, “Exp. C” and “Exp. D” the “N-term”, “Middle” and “C-term” domains of Mid1p were substrates with the kinases Plk1, Aurora A or Aurora B, respectively. Asterisks (*) indicate detected in vitro phosphorylation signals. (c) Coomassie Blue stained gel of the indicated experiments confirms equal protein loading.
Mid1p can be phosphorylated by aurora kinase, using an in vitro phosphorylation approach with recombinant Mid1p domains purified from E. coli. These experiments were performed with the human aurora kinases aurora A and aurora B, and the human polo kinase Plk1 (Figure 3b). Phosphorylation of both the “N-term” and “Middle” Mid1p domains by aurora A and Plk1 was detected, but not by aurora B (Figure 3b, Exp. B and Exp. C). In contrast, no phosphorylation was detected of the “C-term” Mid1p domain by any of the three kinases aurora A, aurora B or Plk1 (Figure 3b, Exp. D).

Ark1p is required for the correct cellular distribution of Mid1p

To further analyze the interaction of Ark1p and Mid1p in S. pombe we examined the effect of two different temperature sensitive ark1 mutants, ark1-T11 and ark1-T8, on Mid1p distribution in cells (Figure 4).

In ark1-T11 cells GFP-Mid1p showed wild-type phenotypes (Figure 4a). But in addition, new localization patterns were apparent, where GFP-Mid1p localization showed nuclear exclusion with the appearance of a ring encircling the nucleus. The frequencies of these phenotypes were quantified and compared to wild-type cells revealing significant differences (Figure 4b). In ark1-T8 cells GFP-Mid1p showed new additional localization patterns (Figure 4d). Cells were observed to be round and shorter than the rod-shaped wild-type cells and GFP-Mid1p localization presented nuclear exclusion with the appearance of a ring that encircled the nucleus. While some cells had one nucleus others showed two, three, or even four nuclei. The frequencies of these phenotypes were quantified and

![Figure 4](https://example.com/figure4.png)

**Figure 4.** GFP-Mid1p cellular localization is disrupted in ark1-T11 and ark1-T8 cells. Wild-type (w-t), ark1-T11 and ark1-T8 S. pombe strains containing GFP-Mid1p were grown at 25°C (the permissive temperature for these temperature sensitive mutants) in liquid YE medium to mid-exponential phase and visualized by confocal microscopy. (a) Key to characterized GFP-Mid1p localization phenotypes. Scale bar, 5 μm. Bright and green fluorescent images of cells (left) and quantitative analysis of GFP-Mid1p localization phenotypes (right) for ark1-T11 (b) and ark1-T8 (d). Scale bar, 10 μm. (c, e) Two-way ANOVA analysis of frequencies of localization phenotypes in ark1-T11 and ark1-T8 compared to wild-type (W-T). Asterisks (****) denote p values <0.0001 indicating a significant difference to wild-type. Error bars = SEM.
compared to wild-type cells revealing significant differences (Figure 4e). These data support the hypothesis that Mid1p and Ark1p coordinate to regulate the *S. pombe* cell cycle.

The growth temperature of the temperature sensitive *ark1* mutants affects the activity of the Ark1p protein but also causes cell cycle arrest; therefore, any phenotypes observed from growing the strains at a restrictive temperature cannot be solely attributed to the absence/inactivation of Ark1p protein as they may instead be a secondary consequence of the cell cycle arrest. To avoid this complication, the yeast cells were grown at the permissive temperature when they complete the cell cycle and divide, so that any phenotypes observed are considered the result of a “partial” absence/inactivation of the Ark1p protein. For this reason, we are unable to determine whether the multinucleated cells phenotype in *ark1*-TS mutants is solely a result of altered GFP-Mid1P localization; however, we note that others have reported the multinucleated cells phenotype in *ark1* mutant strain [29].

**Identification of Mid1p amino acid residues phosphorylated by aurora and polo kinases**

To further explore phosphorylation of Mid1p by aurora and polo kinases a mass spectrophotometry approach was used to identify phospho-acceptor sites. The “N-term” and “Middle” domains of Mid1p gel fragments from *in vitro* kinase assay reactions identical to those shown Figure 3b, but with non-labeled ATP, were excised and subjected to nano-scale liquid chromatographic tandem mass spectrometry (nLC-MS/MS) to generate a Mid1p phospho-acceptor sites map (Figure 5). This analysis identified 35 potential Mid1p residues phosphorylated by either Plk1 or aurora A kinases (S3 Table). In parallel we examined other published databases on fission yeast phosphoproteomes to refine and confirm the number of Mid1p phospho-acceptor sites. Most of these studies used a stable isotope labeling by amino acid in cell culture approach. We selected four studies which studied the *S. pombe* global proteome and identified several additional Mid1p specific phospho-acceptor sites [30–33].

The Mid1p phosphorylation events identified by mass spectrophotometry are summarized in S3 Table, alongside those described by the four published studies. The red highlighted residues represent overlap with the residues identified reported in this work. The phospho-acceptor sites map shows a total of six potential phospho-acceptor

![Figure 5](image_url)
sites distributed along the “N-term” and “Middle” Mid1p sequence, five of which were identified here, and the sixth was added from the published studies (Figure 5b). These are the serine residues S167, S328, S331, S332, S523 and S531.

We used the GPS 5.0 online computational prediction tool to predict phosphorylation sites of Mid1p and it found positions S167, S328 as potential phosho-acceptor sites for Plk1, aurora kinase respectively; S523 and S531, as potential phosho-acceptor sites for aurora and/or Plk1 (http://gps.biocuckoo.cn/download.php).

To confirm the new phosho-acceptor sites identified in Mid1p in vitro phosphorylation experiments involving bacterially expressed phosho-resistant mutant (serine to alanine) forms of the “N-term” and “Middle” domains of Mid1p with aurora A and Plk1 kinases were completed (Figure 6). These experiments reveled markedly reduced phosphorylation of the Mid1p phosho-resistant mutant S167A by both aurora A and Plk1 kinases, and S523A by aurora A alone. These observations support the suggestion that the S167 and S523 residues are phosphorylated by the aurora A and Plk1 kinases, at least in vitro.

**Biological relevance of amino acid residues phosphorylated by aurora and polo kinases in Mid1p**

As another way to examine the role of the aurora and polo kinase phosho-acceptor sites identified in Mid1p we analyzed the effect of their mutation in vivo in fission yeast. Importantly, such an approach allowed confirmation of the relevance in yeast of the phosho-acceptor sites in Mid1p identified using mammalian orthologous protein kinases. *S. pombe* cells exhibit morphology defects in the absence of Mid1p function. Therefore, we tested the effect of mutations of Mid1p phosho-acceptor sites in fission yeast on cell morphology to assay their relevance.

![Figure 6. Reduced in vitro phosphorylation of Mid1p “N-term and “Middle” phosho-resistant mutants by Plk1 and Aurora A kinases. Recombinant GST-tagged Mid1p “N-term” (a) and “Middle” (b) domains phosho-resistant mutants were expressed and purified in E. coli. Proteins subjected to in vitro phosphorylation experiments with Aurora A (Exp. A) or Plk1 (Exp. B) kinases. Asterisks (*) and **) indicate wild-type Mid1p “N-term” and Mid1p “Middle” recombinant proteins used as positive controls. Colored boxes represent predicted proteins: MBP yellow, Mid1p “N-term” species red, Plk1 purple, and all Mid1p “Middle”-related species green. “Multiple” indicates mutations in all four phosho-acceptor sites. GFP-Mid1p is imaged in *ark1* wild-type cells (Fig 2B).](image-url)
Versions of the *mid1* gene containing phospho-acceptor site mutations were generated and integrated in single copy into chromosomal DNA of *S. pombe* mid1Δ cells. Each version of the *mid1* gene had either a phospho-mimetic (S > D) or a phospho-resistant (S > A) point mutation(s) of the serine residues S167, S328, S331, S332, S523 or S531 to create a panel of mutant *S. pombe* strains (S1 Fig). Such *mid1* mutations were made both singly and in combination. As a positive control we integrated the wild-type *mid1*+ gene into mid1Δ cells to create mid1Δ pJK148: *mid1*+. This resulted in cells that behaved and looked identical to wild-type, both on solid medium and in liquid culture (Figure 7 and S1 Fig), confirming that the integration event had not affected Mid1p activity.

**Mid1p phospho-mutants in mid1Δ cells**

Initial experiments examined the effect of Mid1p phospho-mimetic or phospho-resistant mutations in mid1Δ cells. Strains were grown on solid

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**Figure 7.** The S332 phospho-acceptor site is required for Mid1p function. Three phospho-resistant mid1 mutant strains mid1 S332A, mid1 S167A S328A S331A S332A, and mid1 S167A S328A S331A S332A S523A S531A, and the three phospho-mimetic mutants mid1 mutant strains S332D, mid1 S167D S328D S331D S332D, and mid1 S167D S328D S331D S332D S523D S531D streaked to single colonies on solid YE at 25°C to compare growth rates (a) and colony morphology (b). Controls: mid1Δ and wild-type (w-t) cells. Scale bar, 10 μm.
medium, with the formation of colonies observed in all cases. In most strains where serine was changed to either alanine or aspartic acid cell morphology appeared similar to wild-type (Figure 7, S4 Fig, S4 Table and S5 Fig). However, interestingly, mid1 S332A cells (but not mid1 S332D) had defects in morphology similar to those observed in mid1A cells, with slower growth at 25°C (Figure 7). These data indicate a requirement of the phospho-acceptor S332 for Mid1p function.

Furthermore, other phenotypes were revealed when certain Mid1p phospho-mutants were cultured in liquid medium. For example, mid1 S523A and mid1 S531A mutants displayed morphology defects (S2 Fig). In contrast, the equivalent phospho-mimetic mutants mid1 S523D or mid1 S531D did not.

**Mid1p phospho-mutants in ark1-T11 cells**

To examine the role of Mid1p phosphorylation in its interaction with Ark1p we crossed the strains containing phospho-mimetic/resistant versions of mid1 with ark1-T11 mutant cells and searched for synthetic phenotypes in double mutants. Viable colonies were observed in most cases with no synthetic phenotypes detected in many double mutants (S4 Table). However, mid1 S523A ark1-T11 and mid1 S531A ark1-T11 double mutants showed cell morphology defects when cells were grown in liquid culture more severe than the single mutant strains (S3 Fig and S4 Table). These defects were not observed in the equivalent mid1 S523D ark1-T11 or mid1 S531D ark1-T11 double mutants. The cell morphology defects observed in mid1 S523A ark1-T11 and mid1 S531A ark1-T11 double mutants were defined as loss of the rod-like shape of cells. This suggests a role for these two phospho-acceptor sites in Mid1p function during the S. pombe cell cycle to ensure medial division plane placement, and consequently equal sized and rod-shaped daughter cells.

**Mid1p phospho-mutants in plo1-ts35 cells**

To examine the role of Mid1p phosphorylation in its interaction with Plo1p we crossed strains containing phospho-mimetic/resistant versions of mid1 with plo1-ts35 mutant cells and searched for synthetic phenotypes in double mutants. Viable colonies were observed in all cases with no synthetic phenotypes detected for the majority of double mutants (S4 Table). However, colony formation on solid medium was slower for mid1 S332A plo1-ts35 double mutants, and more severe cell morphology phenotypes were observed, compared to the single mutant strains (S5 Fig). These defects were not observed in the equivalent mid1 S332D plo1-ts35 double mutants. Such genetic interactions suggest a link between the regulation of Mid1p and Plo1p.

**Mid1p phospho-mutants in vps4Δ cells**

To examine the role of Mid1p phosphorylation in its interaction with Vps4p we crossed strains containing phospho-mimetic/resistant versions of mid1 with vps4Δ mutant cells and searched for synthetic phenotypes in double mutants, Viable colonies were observed in all cases with no synthetic phenotypes detected (S4 Fig and S4 Table). However, colony formation on solid medium was slower for mid1 S523A S531A vps4Δ double mutants, and more severe cell morphology phenotypes were observed, compared to the single mutant strains. Strikingly, cells had defects in morphology similar to those observed in mid1Δ cells (Figure 8). The same serine residues changed to alanine S523D or S531D had no such effect when combined with vps4Δ (S4 Fig and S4 Table). These genetic interactions suggest a link between the regulation of Mid1p by both Vps4p and Ark1p.

**Conclusions**

In fission yeast the Mid1p protein has important roles during cytokinesis during the cell cycle, with anillin homologues having similar functions in higher eukaryotes. These roles center around its structural function in predicting and controlling the site of cell division in the equatorial region of the cell. Here we have identified new classes of proteins with which Mid1p interacts, and explored the mechanism by which these proteins cooperate and regulate each other and cell growth/morphology.

**Mid1p and Vps4p**

Genetic and biochemical approaches revealed a direct physical interaction of Mid1p with the ESCRT-associated protein Vps4p. Furthermore,
Figure 8. The Mid1p S523 and S531 phospho-acceptor sites regulate its interaction with Vps4p. Strains streaked to single colonies on solid YE medium and grown at 25°C to compare growth rates and colony morphology. Controls: vps4Δ and wild-type (w-t) cells. Scale bar, 10 μm.

Figure 9. Interaction of Mid1p and Vps4p to regulate S. pombe septation. (a) Schematic representation of Mid1p localization during S. pombe cell cycle stages. CR = contractile ring. (b) Mid1p is phosphorylated by Plo1p to promote mitotic entry, during which Rng2p and other proteins interact with the N-terminal domain of Mid1p. (c) In mitosis, Mid1p forms nodes which are attached to the plasma membrane via the lipid binding motifs present within the Mid1p C-terminal domain. Green box represent the overall structure of Mid1p C-terminal region (aa 579–920) containing: the C2 domain (cyan), the connector domain (red) and the PH domain (yellow); dotted lines represent the lipid binding loop. Structure adapted from [13]. The Vps4p interaction with the C-terminal domain of Mid1p potentially regulates this process.
we demonstrated that a chromosomal deletion of the \textit{vps4} gene caused defects in the cellular localization of GFP-Mid1p. These defects included mis-localization of Mid1p-dependent nodes whereby one node is randomly positioned, or three nodes were present. This suggests a role of Vps4p in the Mid1p-dependent node localization pathway, which led us to hypothesize that the function of Mid1p is regulated by Vps4p during nodes attachment to the plasma membrane (Figure 9).

Two types of nodes are involved in \textit{S. pombe} actomyosin ring assembly and contraction (Figure 9, Panel A). Mid1p cortical anchorage first drives the recruitment of cytokinesis proteins, then interactions with myosin filaments causes the condensation of nodes into the actomyosin ring. Mid1p cortical anchorage depends on the PH domain [13] and its potential interaction with Vps4p might stabilize this interaction. Since Vps4p physically interacts with residues within the C-terminal domain of Mid1p, which contains membrane binding motifs, we speculate that Vps4p may facilitate Mid1p cortical anchorage to promote \textit{S. pombe} medial division (Figure 9, Panel C).

We suggest that a physical interaction between Vps4p and Mid1p regulates Mid1p-dependent node attachment to the plasma membrane to determine the division plane in \textit{S. pombe}, and that this interaction directly or indirectly involves Mid1p PH domain cell cortex anchorage (Figure 9, Panel C). It is interesting to note that the domain of Mid1p that interacts with Vps4p \textit{in vitro} (Figure 1) contains the PH domain, suggesting that binding of Vps4p to this region may regulate interaction with the cell cortex (Figure 9).

\textbf{Phosphorylation of Mid1p}

Genetic and biochemical approaches revealed a direct physical interaction of Mid1p with the aurora kinase, Ark1p in fission yeast. Furthermore, human aurora A was shown \textit{in vitro} to phosphorylate the Mid1p the “N-term” and “Middle” domains. Mapping of the amino acids in Mid1p phosphorylated by aurora A and Plk1 revealed 35 potential phospho-acceptor sites with, importantly, some of these sites independently identified in four \textit{S. pombe} global proteomic studies (Figure 5). Such combined analysis suggested six potential serine phospho-acceptor sites in Mid1p: S167, S328, S331, S332, S523 and S531. Subsequent \textit{in vitro} kinase assay experiments confirmed the \textit{in vitro} phosphorylation of the serine S167, S331 and S523 phospho-acceptor sites of Mid1p by aurora A and Plk1 kinases (Figure 6).

As the \textit{in vitro} kinases assay approach used to identify the phospho-acceptor sites in Mid1p utilized human purified aurora kinase and polo kinase it was important to confirm the biological relevance of the sites in fission yeast. This was especially the case for the kinase assay using aurora A, as fission yeast Ark1p more closely resembles aurora B. To do this we examined the effect of mutations of these Mid1p serine phospho-acceptor sites \textit{in vivo} on cell morphology in wild-type, \textit{ark1-T11, plo1-ts35} and \textit{vps4A} \textit{S. pombe} cells (S4 Table and S5 Fig). In wild-type cells defective cell morphology phenotypes were observed for the Mid1p phospho-resistant mutants S332A, S523A and S531A (Figure 7 and S2 Fig), and the phospho-mimetic S523D and S531D when combined with mutations in \textit{vps4}, \textit{ark1}, or \textit{plo1} (Figure 8 and S4 Table). Similarly, defective cells morphologies were detected in \textit{mid1} S523A \textit{ark1-T11} and \textit{mid1} S531A \textit{ark1-T11} double mutants (S3 Fig). Unexpectedly, defective cell growth phenotypes were observed in S332A, S523A and S531A related mutants, but not in S167A and S331A related mutants which instead showed a decreased level of \textit{in vitro} phosphorylation by both aurora and Plk1 or Plk1 only, respectively (Figure 6) when compared to wild-type Mid1p phosphorylation levels by the two kinases (S5 Table). It could be that the effect produced by this mutation is subtle and not detected using this simple assay; alternatively mutation of one phospho-acceptor site might modulate phosphorylation at a distinct site. It will be important in future studies to determine whether Mid1p cellular localization is affected by these mutations, and what the consequence might be for Mid1p function.

Overall, we conclude that the phosphorylation of these amino acid residues is important for Mid1p function in fission yeast and its interaction with these proteins to regulate cell cycle events. It is tempting to speculate that the interaction of Mid1p and Vps4p is regulated by the activity of Ark1p and/or Plo1p, but it is important to note that the regions of Mid1p phosphorylated by these kinases do not include the binding region for Vps4p. Hence, phosphorylation in adjacent
regions may modulate binding via conformational changes in Mid1p, as regions containing the phospho-acceptor sites have been shown to regulate the interaction of Mid1p with other proteins, including Plo1p and Sid1p. Plo1p phosphorylates residues within the first 100 amino acids of Mid1p to trigger Myosin II recruitment during contractile ring assembly [34]. Later at contractile ring constriction Sid1p phosphorylates Mid1p to facilitate its export from the cell cortex [7]. Future experiments will be aimed at unraveling how the Mid1p interactome is modified both by association with Vps4p and by phosphorylation by mitotic kinases.

Materials and Methods

Yeast media and general techniques

General molecular procedures were performed, with standard methodology and media used for the manipulation of S. pombe [35,36]. The yeast strains used in this study are shown in S1 Table. Cells were routinely cultured using liquid or solid complete (YE) or minimal (EMM) medium, at 25°C or 30°C.

DNA constructs

The DNA constructs used in this study are listed in S2 Table. Some were synthesized and cloned by either GenScript or Invitrogen. All constructs were sequenced before use.

Bacterial expression DNA constructs

Four fragments of the mid1+ gene encoding the amino acids 1–453 “N-terminus”, 452–579 “Middle”, 578–799, 798–920 “C-term” [25] were synthesized and cloned into Bam HI/Xho I of pGEX-4 T-1. Full-length vps4+ was synthesized and cloned into Nde I/Bam HI of pET-14b. The C-terminal domain of myo2+ was synthesized and cloned into Nde I/Bam HI of pET-14b.

Mid1 phospho-mimetic/resistant mutants

Eighteen versions of the mid1 gene with different phospho-mimetic/resistant mutations were synthesized, along with a wild-type mid1+ control. All had the wild-type mid1+ promoter in ~1 kbp of DNA upstream of mid1+ open reading frame, and were each cloned into Kpn I/Sac I of pJK148. Integration of the pJK148:mid1 genes (1–19) into S. pombe mid1Δ cells was through homologous recombination after linearization of pJK148:mid1 with Nde I in the leu1+ gene. The resulting panel of phospho-mimetic/resistant mutant strains is listed in S1 Table and S1 Fig.

Recombinant protein purification

GST-Mid1p, 6His-Vps4p or 6His-C-term Myo2p plasmids were grown in BL21 E. coli until an OD of 0.6–0.8, with protein production induced by adding 1 mM IPTG. Mid1p 578–799 was not found to be expressed under these conditions, and so was not used for further experiments.

Bacterial pellets were produced by centrifugation with cells re-suspended in 20 ml re-suspension buffer with EDTA-free protease inhibitors; for GST-Mid1p fusion proteins PBS buffer (137 mM NaCl, 2.7 mM KCl, 10 mM Na2HPO4 and 1.8 mM KH2PO4, pH 7.4) was used, whereas for 6His-Vps4p, HEPES buffer (25 mM HEPES, 400 mM KCl and 10% (v/v) glycerol, pH 7.4) was used. Cells were then lysed by sonication, where a final concentration of 1 mg ml−1 lysozyme was added for cell wall digestion followed by sonication, with 0.1% (v/v) Triton X-100 added prior to lysis. A clear lysate was produced by centrifugation. GST-tagged fusion proteins were purified using 1 ml 1−1 glutathione beads in PBS buffer, while 6His-tagged Vps4p or C-term Myo2p were purified using 500 μl 1−1 Ni-NTA beads in HEPES buffer, either for 2 h or overnight at 4°C. Mid1p was eluted from glutathione beads using Reduced glutathione buffer. Vps4p or C-term Myo2p were washed and eluted from Ni-NTA beads using HEPES buffers. Elution was carried out for 2 h at 4°C. Samples of proteins from each step were subjected to SDS-PAGE to determine elution efficiency.

Pull-down experiments

Pull-down experiments utilized Ni-NTA beads-immobilized bait proteins (6His-Vps4p or 6His-C-term Myo2p) and prey-eluted proteins (GST-Mid1p: “N-terminus,” “Middle”, or “C-term”) to investigate protein–protein interactions. Bait
proteins were loaded onto Ni-NTA beads by incubation in PBS containing 0.01% (v/v) Triton X-100 for 1 h (4°C). After loading, the mixture was washed with PBS containing 0.01% (v/v) Triton X-100, and beads were blocked for nonspecific binding by incubation in PBS containing 0.2% fish-skin gelatin. The beads mixture was incubated with prey protein in PBS with 0.01% (v/v) Triton X-100. Subsequently, beads were washed with PBS containing 0.01% (v/v) Triton X-100 three times, 0.5% (v/v) glycerol and 0.2% (w/v) fish skin gelatin three times, and with PBS alone four times. Proteins were eluted from beads by boiling in Laemmli sample buffer (LSB) (75 mM Tris pH 6.8, 12% (w/v) SDS, 60% (v/v) glycerol, 600 mM DTT and 0.6% (w/v) Bromophenol Blue) and samples were subjected to SDS-PAGE.

In vitro kinase assays

Assays combined either human Plk1 (0.023 µg/µl), aurora A (0.01 µg/µl), aurora B (0.01 µg/µl) and myelin basic protein (MBP; 2.5 µg) (Sigma–Aldrich and Biaffin GmbH) or one of the three Mid1p domains (“N-term”, “Middle” or “C-term” 2.5 µg). Kinase, substrate proteins, 1 µCi [γ-32P] ATP, 10 mM ATP and kinase assay buffer (25 mM MOPS, 25 mM MgCl2, 1 mM EDTA and 0.25 mM DTT, pH 7.2) were mixed in a total volume of 20 µl; the reaction was initiated by adding 5 µl ATP cocktail and incubated at 30°C for 1 h and terminated by the addition of LSB; samples were subjected to SDS-PAGE followed by autoradiography or phaso-imaging. Following detection of in vitro phosphorylation signals, mass spectrometric analysis was carried out by the Dundee FingerPrints Proteomics service (http://proteomics.lifesci.dundee.ac.uk/).

Confocal microscopy

GFP-tagged mid1+ [37] was examined by a He/Ne and Ag laser system of Zeiss LSM microscope using 63X high NA 1.4 objective lens. Cell images were collected using Zeiss Pascal software and processed using ImageJ, Microsoft PowerPoint and Keynote software. Numerical analysis was completed from three independent experiments where 200–250 cells were counted. When GFP-foci were counted, care was taken to scan the entire depth of the cell to ensure that all foci present were quantified. Yeast colonies grown on solid medium were imaged using a Zeiss Axioscope microscope and a Sony DSC-75 camera.

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Disclosure statement

No potential conflict of interest was reported by the author(s).

ORCID

Imane M. Rezig http://orcid.org/0000-0001-6673-8946
Wandialhyl G. Yaduma http://orcid.org/0000-0002-3600-7162
Gwyn W. Gould http://orcid.org/0000-0001-6571-2875
Christopher J. McInerny http://orcid.org/0000-0002-9201-3479

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