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► **To cite this version:**

Federica Arbizzani, Manos Mavrakakis, Marta Hoya, Juan Carlos Ribas, Sophie Brasselet, et al.. Septin filament compaction into rings requires the anillin Mid2 and contractile ring constriction. *Cell Reports*, 2022, 39 (3), pp.110722. 10.1016/j.celrep.2022.110722 . hal-03835966

**HAL Id: hal-03835966**

**<https://cnrs.hal.science/hal-03835966v1>**

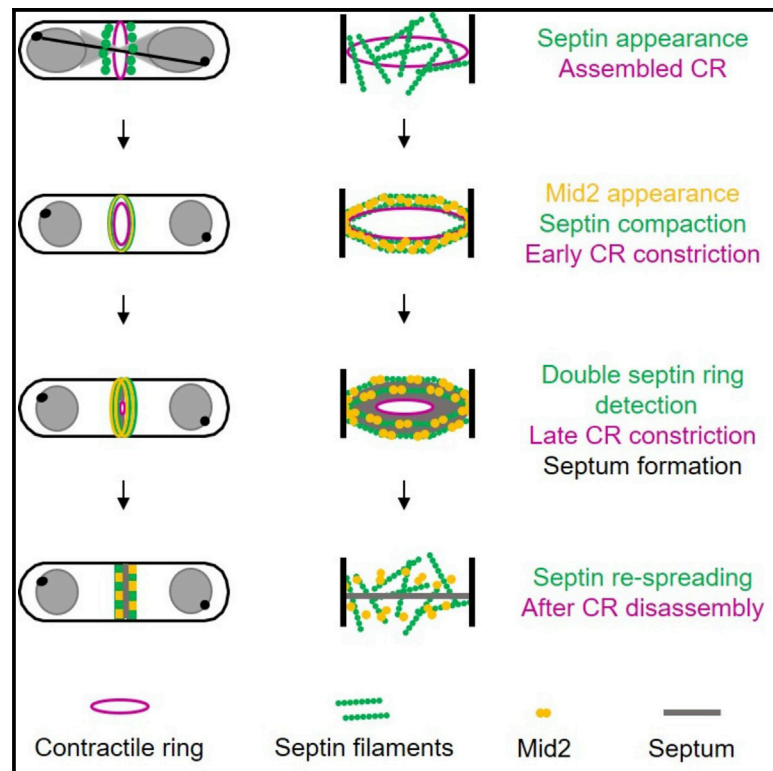
Submitted on 1 Nov 2022

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# Septin filament compaction into rings requires the anillin Mid2 and contractile ring constriction

## Graphical abstract



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## In brief

Arbizzani et al. show that the fission yeast anillin-like protein Mid2 self-interacts to crosslink septin filaments, allowing septin filament orientation parallel to the division plane and compaction into two septin rings delineating the cleavage furrow. This event is triggered by contractile ring constriction.

## Highlights

- Mid2 self-interacts to crosslink septin filaments for septin ring assembly
- Mid2 reorients septin filaments parallel to the division plane
- Mid2-driven septin filament reorientation is triggered by CR constriction



## Article

# Septin filament compaction into rings requires the anillin Mid2 and contractile ring constriction

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<https://doi.org/10.1016/j.celrep.2022.110722>

## SUMMARY

Septin filaments assemble into high-order molecular structures that associate with membranes, acting as diffusion barriers and scaffold proteins crucial for many cellular processes. How septin filaments organize in such structures is still not understood. Here, we used fission yeast to explore septin filament organization during cell division and its cell cycle regulation. Live-imaging and polarization microscopy analysis uncovered that septin filaments are initially recruited as a diffuse meshwork surrounding the acto-myosin contractile ring (CR) in anaphase, which undergoes compaction into two rings when CR constriction is initiated. We found that the anillin-like protein Mid2 is necessary to promote this compaction step, possibly acting as a bundler for septin filaments. Moreover, Mid2-driven septin compaction requires inputs from the septation initiation network as well as CR constriction and the  $\beta(1,3)$ -glucan synthase Bgs1. This work highlights that anillin-mediated septin ring assembly is under strict cell cycle control.

## INTRODUCTION

Cytokinesis is an irreversible process that must precisely partition an equal set of the replicated chromosomes and organelles into each daughter cell. Defects in this process can lead to chromosome mis-segregation or aneuploidy, a hallmark of cancer (Lacroix and Maddox, 2012). Cytokinesis completion requires the participation of different components of the cytoskeleton, namely, the actin cytoskeleton, which assembles into a contractile ring (CR) that pinches in the plasma membrane to create the cleavage furrow, microtubules involved in division plane signaling in animal cells, and septins, present at the division site from yeast to mammals.

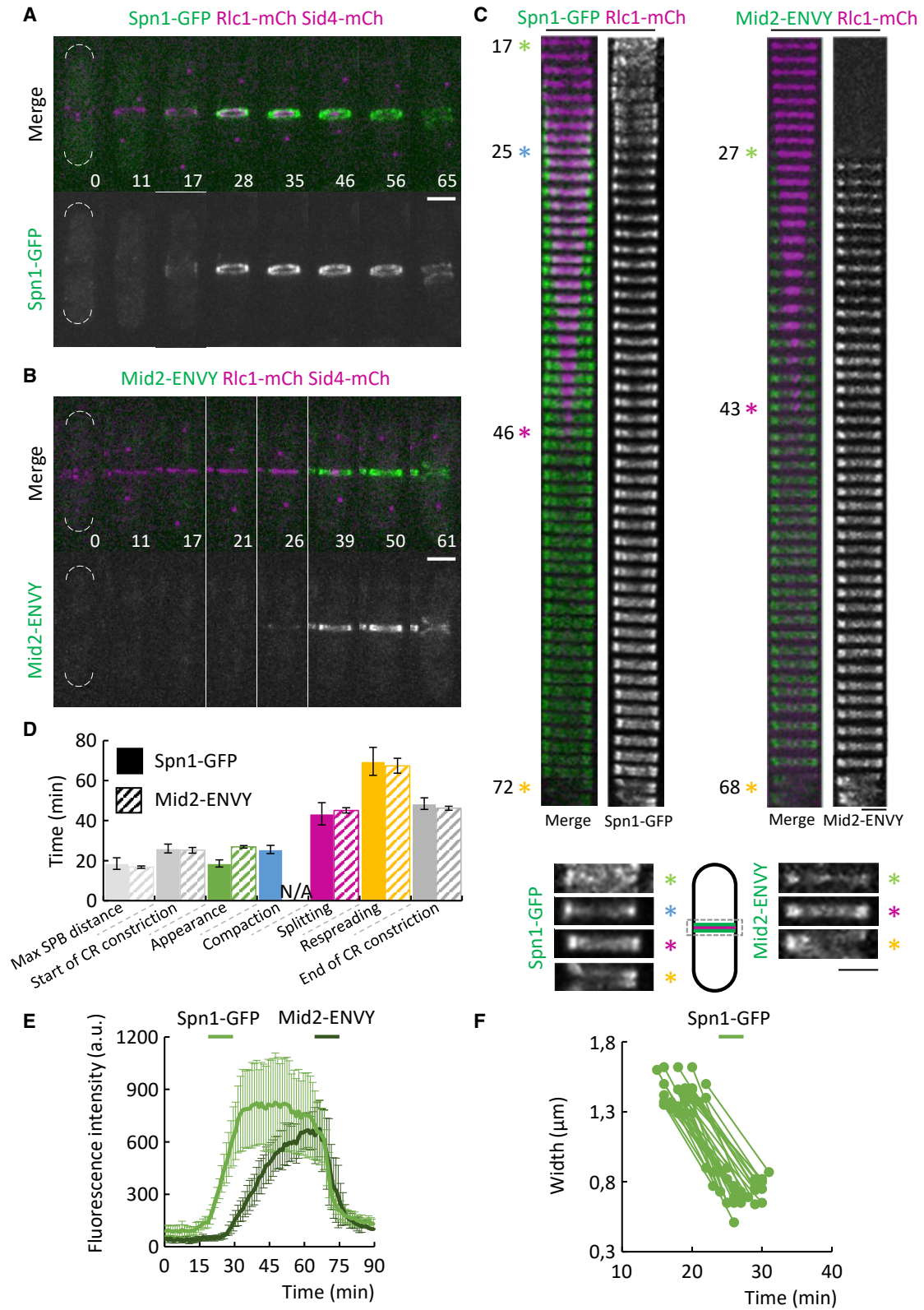
Septins are unique cytoskeletal components capable of self-assembling into high-order molecular structures of varied shapes such as filaments and rings. Their ability to interact with membranes through interaction with negatively charged phospholipids such as phosphatidylinositol 4,5-bisphosphate (PIP2) and with actin filaments and microtubules allows them to participate in many biological processes, working as diffusion barriers for protein compartmentalization and scaffolds for protein-protein interactions (Bertin et al., 2010; Bridges et al., 2014; Marquardt et al., 2019). These conserved GTP-binding proteins were initially discovered in budding yeast, based on their role in cytokinesis, a conserved function in animals (Hartwell, 1971; Neufeld and

Rubin, 1994). Septins in budding yeast serve as a scaffold for the sequential and ordered assembly of the acto-myosin-based CR (Juanes and Piatti, 2016; Bhavsar-Jog and Bi, 2017). Later on, when CR constriction occurs, septins act as cortical barriers to create confined zones at the cleavage furrow where membrane reorganization and septum formation happen during cell division (Dobbelaere and Barral, 2004).

In budding yeast, septins undergo multiple reorganizations during the cell cycle. In early G1, septins organize as a patch-like structure at the presumptive bud site. Upon bud emergence, this patch narrows to form a single ring marking the site of bud growth. Once the bud has formed, the septin ring extends into an hourglass-shaped collar present at the mother-bud neck until mitotic entry. At cytokinesis onset, the septin hourglass splits into two distinct rings that sandwich the CR (Gladfelter et al., 2001). Finally, after cell division, the septin rings disassemble and septin subunits are recycled in the new cycle (McMurray and Thorner, 2009).

One important factor in septin organization is the conserved protein anillin, whose disruption leads to defects in animal cytokinesis (Oegema et al., 2000; Straight et al., 2005; Zhao and Fang, 2005). *In vitro* studies have shown that anillin can bind septin and actin filaments, and indeed, anillin bridges septins and the CR during cell division (Kinoshita et al., 2002). Animal anillin interacts with septins and PIP2 through a C-terminal PH domain





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located after the anillin homology domain (Liu et al., 2012). The C-terminal region of Bud4, the budding yeast anillin ortholog, is important to direct septin organization during bud site selection and bud growth and to preserve the integrity of the septin ring during cytokinesis (Eluere et al., 2012). Thus, in *bud4Δ* mutants the septin double ring disassembles during cytokinesis (Kang et al., 2013), whereas the overexpression of Bud4 leads to the formation of ectopic septin structures. More recently, polarized fluorescence microscopy studies have described a key role of Bud4 in the reorientation of septin filaments in the hourglass to double ring transition (McQuilken et al., 2017).

The fission yeast *Schizosaccharomyces pombe*, a recognized model system for cytokinesis, is a rod-shaped organism that divides by the assembly and constriction of a medially placed actomyosin-based CR. *S. pombe* has two anillin-like proteins, Mid1 and Mid2, with non-overlapping functions (Berlin et al., 2003; Tasto et al., 2003). Mid1 accumulates at the medial cortex at mitosis onset to promote CR assembly (Pollard and Wu, 2010; Rincon and Paoletti, 2016). CR constriction and synthesis of the septum (the extracellular cell wall separating the two daughter cells) are triggered upon mitotic exit by the septation initiation network (SIN) pathway, related to the budding yeast mitotic exit network and the metazoan Hippo pathway (Hergovich et al., 2006). *S. pombe* septins were identified by sequence homology with the *S. cerevisiae* counterparts. Spn1–4 are expressed in vegetatively growing cells and localize to the division site. However, differently from *S. cerevisiae*, where septins are necessary for CR formation, *S. pombe* septins are recruited after the CR has fully assembled (Wu et al., 2003). Septins form first a single ring in late mitosis that splits in two during CR constriction and septum formation. This double ring does not constrict, but it remains at the base of the septum to delineate the boundaries of the cleavage furrow and dissociates after cell separation (Wu et al., 2003; An et al., 2004). Septin mutants have defects in cell-cell separation since septins have a key role, in concert with the exocyst complex, in the delivery of the septum hydrolytic enzymes to the area around the septum, thereby regulating the step that finishes cytokinesis (Martin-Cuadrado et al., 2005). Recently, it has also been shown that septins play a role in driving the proper recruitment and maintenance in the septum region of the septation initiation network (SIN) effector kinase Sid2 and of the glucan synthase enzymes Bgs1 and Ags1 to guarantee a successful cytokinesis (Zheng et al., 2018). Mid2 localizes to the division site after septin recruitment in a septin-dependent manner (Berlin et al., 2003). Fluorescence recovery after photobleaching analysis revealed that septins are more dynamic in

*mid2Δ* cells, indicating a role of Mid2 in septin organization (Berlin et al., 2003). Overexpression of a non-degradable Mid2 mutant let septin filaments persist through the next cell cycle, indicating that Mid2 is also involved in septin ring maintenance (Tasto et al., 2003).

How septins and Mid2 are precisely regulated to properly perform their function and how Mid2 contributes to septin ring organization remain elusive. Here, by using live cell imaging and precise timers for mitotic progression, we describe in detail septin and Mid2 behavior in fission yeast. Our approach identified a possible new step in septin ring assembly: septin filament compaction from a loose meshwork surrounding the CR into a tight ring when CR constriction is initiated. We show that this step requires Mid2, which has the properties of a bundler for septin filaments, promoting their compaction. Polarization microscopy shows that septin filaments transition from a disorganized state to a highly ordered state where they become parallel to the division plane during CR constriction, in a mechanism dependent on Mid2. Finally, we show that the process of septin compaction requires the SIN pathway signaling and CR constriction.

## RESULTS

### Septins form a diffuse network in the vicinity of the CR before compacting into a ring structure

Fission yeast septins localize to the division site where they form non-contractile rings defining the edge of the cleavage furrow while the CR constricts and the septum is built (Berlin et al., 2003; Tasto et al., 2003). However, septin ring assembly is poorly understood. We therefore decided to characterize by time-lapse imaging the dynamic organization of septins with respect to spindle assembly and elongation, used as a proxy for mitosis progression, and CR assembly and constriction.

To do so, we created a strain producing the septin Spn1 fused to GFP, the myosin II regulatory chain Rlc1 fused to mCherry to monitor the CR, and the spindle pole body (SPB) component Sid4 tagged with mCherry to monitor the mitotic spindle. Since septins function strictly as heteromeric complexes, imaging Spn1 is sufficient to have an overview of the whole septin network (An et al., 2004).

Septins appeared as a cortical band surrounding the CR about 18 min after SPB separation, by the time of maximum spindle elongation (Figures 1A and 1D,  $t = 18.6 \pm 2.3$  min). Later on, septins compacted into a tight ring structure ( $t = 25.6 \pm 2.8$  min), concomitant with the initiation of CR constriction ( $t = 26.1 \pm$

### Figure 1. Dynamics of septins and Mid2 during cytokinesis

(A and B) Time-lapse imaging of WT cells producing Rlc1-mCherry, Sid4-mCherry, and Spn1-GFP (A) or Mid2-ENVY (B). Maximum intensity projections of confocal images are shown. Scale bars: 4  $\mu$ m.

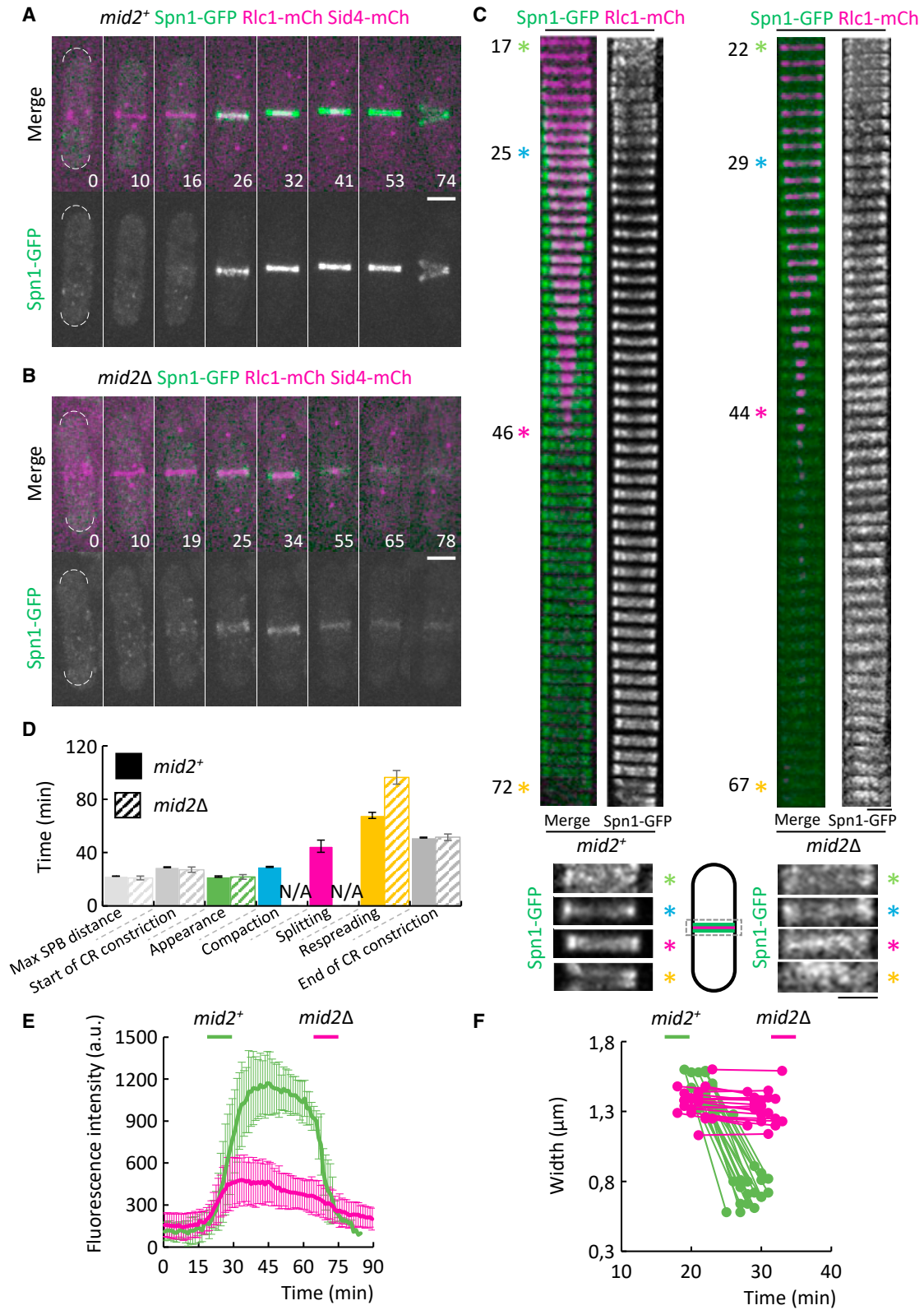
(C) Kymographs showing the medial region of the cell producing Spn1-GFP Rlc1-mCherry (left) or Mid2-ENVY Rlc1-mCherry (right). There is a 1-min interval between successive images. The major transitions in the behavior of Spn1 and Mid2 are indicated with the elapsed time shown in minutes and highlighted at the far right, in magnified panels with an asterisk (green for appearance, blue for compaction, magenta for splitting, and yellow for re-spreading). A scheme of a cell and the selected area are shown at the bottom. Scale bars: 2  $\mu$ m.

(D) Plot depicting the timing of Spn1-GFP (solid bars) and Mid2-ENVY (striped bars) transitions throughout cytokinesis from mitotic entry. Mean  $\pm$  SD are shown.  $n = 51$  cells for each strain.

(E) Analysis of Spn1-GFP (light green) and Mid2-ENVY (dark green) fluorescence intensities in the cell middle from mitotic onset throughout cytokinesis. The average line graphics  $\pm$  SD are displayed.  $n = 6$  cells in each case.

(F) Quantification of the extent of Spn1-GFP width compaction over time.  $n = 30$  cells.





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2.9 min). This event is best visualized on enlargements of the medial region of the cell recorded at 1-min intervals (Figure 1C left, green asterisk). 20 min later, by the end of CR constriction ( $t = 48.5 \pm 3.6$  min), septins were visualized as a double ring ( $t = 43.4 \pm 6.5$  min, magenta asterisk). Eventually, septins re-spread to the new cell tip generated when the two sister-cells separated from one another by cleavage of the septum (yellow asterisk).

Quantitative analysis of the septin signal further revealed that septin intensity increased for a period of about 15 min after they appeared on the medial cortex. The intensity then stayed roughly constant for 30 min before decreasing rapidly (Figure 1E). Furthermore, analyzing the width of the cortical region on which septins were distributed revealed a 2-fold compaction in width, from  $1.4 \pm 0.1$   $\mu\text{m}$  at their appearance to  $0.75 \pm 0.1$   $\mu\text{m}$  when CR constriction starts (Figure 1F).

This detailed live analysis of septins establishes the timing of the major transitions in septin organization and reveals a step in the process of septin ring assembly that we have named compaction.

### The anillin Mid2 is necessary for septin ring compaction

A similar imaging strategy was taken to define the dynamic organization of the anillin-like protein Mid2, known to co-localize with septins and modulate their dynamics (Berlin et al., 2003; Tasto et al., 2003). Using a strain expressing Mid2 C-terminally fused to the green fluorescent protein ENVY, we found that Mid2 appeared as a narrow band 8 min later than septins ( $t = 26.9 \pm 2.5$  min), coincident with the time of septin compaction (Figure 1C right, light green asterisk in the Mid2 kymograph, and Figure 1C left, blue asterisk in the Spn1 kymograph, Figure 1D) and CR constriction ( $t = 25.2 \pm 2.4$  min, Figures 1B–1E). The Mid2 domain transformed into two rings concomitantly to the double septin rings (Figure 1C right, magenta asterisk) and spread out on the newly formed cell tips at the end of cytokinesis (Figure 1C right, yellow asterisk).

Quantitative analysis of Mid2 signal confirmed the late recruitment of Mid2 compared to septins and showed that its intensity increased for about 30 min, stayed constant for 10 min, and decreased rapidly, similarly to septins (Figure 1E).

The coincidence between CR constriction, septin compaction, and Mid2 recruitment suggested that these events may be coupled. To determine if this was the case, we first tested whether Mid2 was involved in septin compaction. To do so, we analyzed septin ring assembly in a *mid2* $\Delta$  mutant. Time-lapse movies showed that in *mid2* $\Delta$  cells, initial septin recruitment

occurred normally (Figures 2A–2D), but septin intensity stopped increasing prematurely, about 8 min after septins appeared. As a consequence, septin maximum intensity was decreased by 3-fold in *mid2* $\Delta$  cells compared to control cells (Figure 2E). Furthermore, septin compaction was completely abolished (Figures 2C and 2F). Instead, septins retained a diffuse distribution around the CR and the septum-associated membrane (Figures 2C and S1A). As a consequence, splitting into two rings did not occur. Finally, whereas CR constriction took place normally (Figure 2D), cell separation was delayed, and the diffuse septin network persisted during the next cell cycle, as reported in a *mid2* $\Delta$  mutant (Figures 2A–2E; Berlin et al., 2003; Tasto et al., 2003).

### Mid2 may function as bundler for septin filaments

The fact that Mid2 is necessary for septin compaction suggested that Mid2 may function as a bundler for septin filaments, promoting their assembly into a ring structure. To perform this activity, Mid2 should be able not only to interact with septins, but also to dimerize or multimerize to bridge together two or more septin filaments.

In line with this hypothesis, Mid2 co-immunoprecipitated with Spn1 (see Figure S1B). To test if Mid2 self-associates, we designed co-immunoprecipitation assays between differentially tagged versions of Mid2 co-expressed in the same cells. These experiments were performed in the presence or in the absence of Spn1 to exclude the detection of differentially tagged molecules of Mid2 bridged by septin filaments. Our experiments show that Mid2-mEGFP co-immunoprecipitates with Mid2-13Myc regardless of the presence of Spn1 (Figure 3A). This interaction was also detected even in the absence of Spn1-5 (Figure S2A).

In order to map the Mid2-Mid2 interaction site, we performed co-immunoprecipitation assays with a truncated version of Mid2 lacking the PH domain (Mid2  $\Delta$ PH) or with the isolated PH domain (see Figure 3C for Mid2 domains organization). Interestingly, Mid2-mEGFP co-immunoprecipitated with Mid2- $\Delta$ PH-13Myc and Mid2 PH domain also showed the ability to self-interact, even in the absence of Spn1 (Figures 3A and 3B). These interactions were also detected in *spn1-5* $\Delta$  cells (Figures S2A and S2B), indicating that Mid2-Mid2 binding is septin independent. To strengthen this notion, a two-hybrid assay using Mid2 PH domain as a prey and as a bait was performed; the result suggests that Mid2 PH domain directly interacts with itself (Figure S2C). Furthermore, recombinant GST-Mid2-PH interacted with MBP-Mid2-PH in an *in vitro* binding assay (Figure 3D) showing that the interaction is direct.

### Figure 2. Defects in septin dynamics in *mid2* $\Delta$ cells

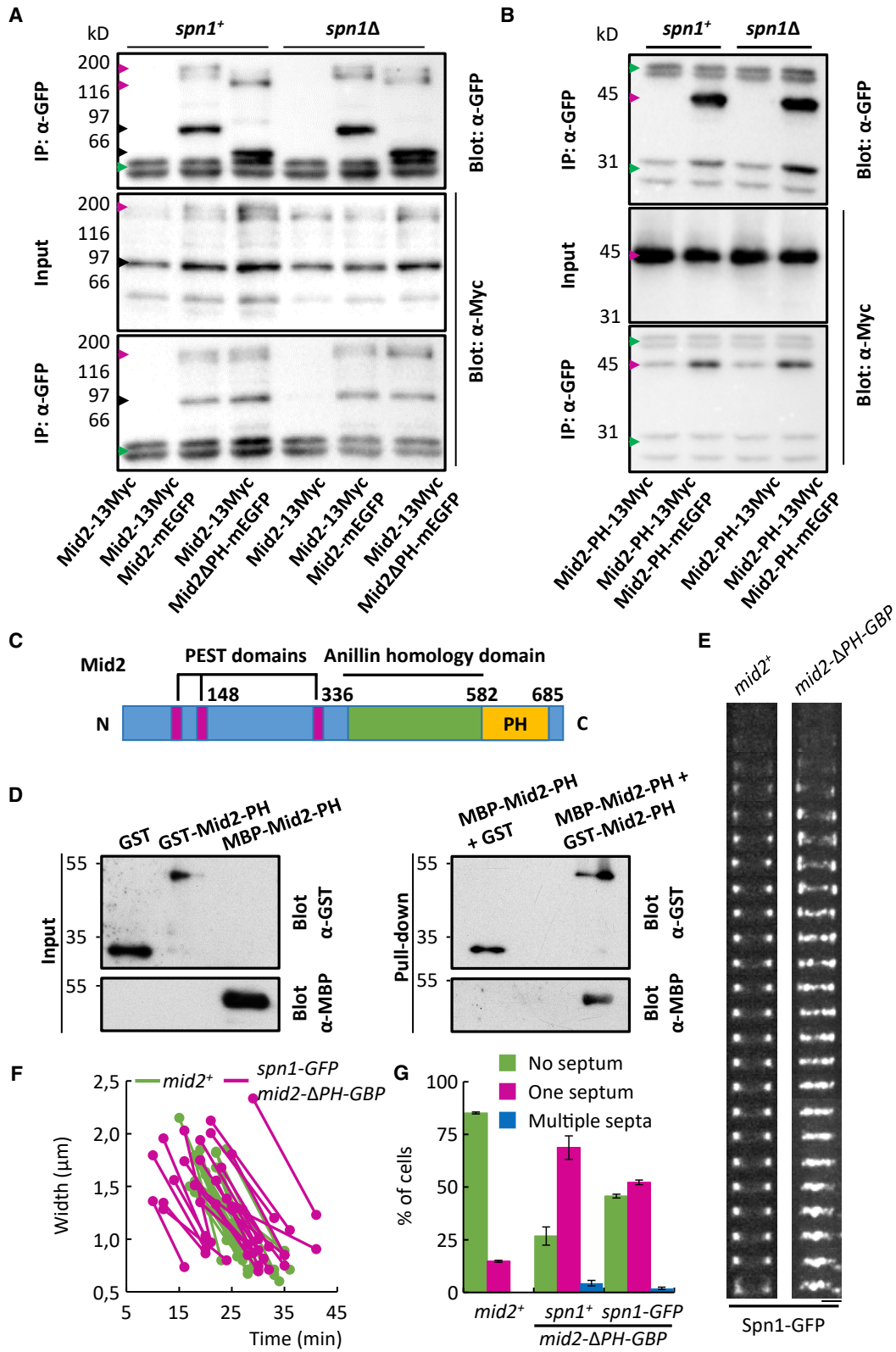
(A and B) Time-lapse imaging of WT (A) and *mid2* $\Delta$  cells (B) producing Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry. Maximum intensity projections of confocal images are shown. Scale bars: 2  $\mu\text{m}$ .

(C) Kymographs showing the medial region of WT (left) or *mid2* $\Delta$  cells (right) producing Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry, as displayed in the scheme of a cell at the bottom. There is a 1-min interval between successive images. The major transitions in Spn1-GFP dynamics and the defects observed in absence of *mid2* are indicated with the elapsed time shown in minutes and highlighted at the far right, in magnified panels with an asterisk (color code matches that in Figure 1C). Scale bars: 2  $\mu\text{m}$ .

(D) Plot displaying the dynamics of Spn1-GFP in WT (solid bars) and *mid2* $\Delta$  cells (striped bars) throughout cytokinesis. Mean  $\pm$  SD are shown.  $n = 52$  cells for each strain.

(E) Analysis of Spn1-GFP fluorescence intensity in the cell middle in WT (green) and in *mid2* $\Delta$  cells (magenta) from mitotic onset throughout cytokinesis. The average line graphics  $\pm$  SD are displayed.  $n = 6$  cells in each case.

(F) Quantification of the extent of Spn1-GFP width compaction over time in WT (green) and in *mid2* $\Delta$  cells (magenta).  $n = 20$  cells. See also Figure S1.



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To better define the contribution of Mid2 domains to septin ring compaction, we analyzed the functionality of a Mid2 truncation lacking the PH domain (Mid2- $\Delta$ PH). This protein does not localize to the division site, so it does not contribute to septin function (Tasto et al., 2003). To overcome this situation, Mid2- $\Delta$ PH was fused to the GFP-binding protein (GBP) and expressed as a Mid2 unique copy in a strain expressing Spn1-GFP to target it to septins by interaction between the GBP and GFP tags. The analysis of Spn1-GFP indicated that it appeared as a broad band that eventually compacted, despite the absence of Mid2 PH domain (Figures 3E and 3F). However, the septin rings collapsed as the CR constricted, suggesting that the PH domain contributes to septin ring stability. It should be noted, though, that in this situation, Spn1-GFP tends to form clumps that follow the CR during its constriction, and it remains longer at the division site. Nonetheless, expression of Spn1-GFP in the *mid2- $\Delta$ PH-GBP* background partially rescued the cell separation defect shown by *mid2 $\Delta$*  cells (Figure 3G).

This indicates that Mid2 has the ability to form structures of higher molecular order and that Mid2 lacking the PH domain is sufficient to drive septin compaction, when artificially bound to septins. Mid2 PH domain on the other hand may contribute to Mid2 association with septins and to septin ring stability at later stages. Since Mid2 arrival at the cortex also coincides with septin compaction and Mid2 is necessary for this step, our biochemical data support the hypothesis that Mid2 may promote septin compaction by bundling septin filaments.

### Septins and Mid2 form wide bands surrounding the CR when cells are blocked in mitosis

Our work shows that septin filaments appear precisely when the anaphase spindle length is maximum and compact into rings when Mid2 is recruited and the CR starts constricting. This provides evidence for a tight temporal regulation of septin dynamics. To get insight into how this is achieved, we first analyzed the localization of Spn1 and Mid2 in mutants blocked in mitosis with a high cyclin-dependent kinase (CDK) activity. We first used the cold-sensitive  $\beta$ -tubulin mutant *nda3-KM311*, which upon cold treatment, blocks in mitosis with an assembled CR. In these conditions, a large band of septins and Mid2 surrounding the CR formed as opposed to the septin or Mid2 ring detected in control cells (Figure 4A left). Interestingly, Spn1-GFP and Rlc1-mCherry intensity measurements by linescans along the cell axis showed a decrease in Spn1-GFP signal at the exact position of the CR (Figures 4A right, S3A and S3B). Furthermore, imaging these cells at 1-h intervals after the temperature shift showed that

the septin domain gradually spread along the cortex in *nda3-KM311* cells (from  $1.2 \mu\text{m} \pm 0.03$  after 1 h at  $18^\circ\text{C}$  to  $2.4 \mu\text{m} \pm 0.2$  after 7 h at  $18^\circ\text{C}$ ), as opposed to the tight ring present in control cells ( $0.9 \mu\text{m} \pm 0.03$ ; Figure S3C). A similar situation was observed for the Mid2 domain. Release from the mitotic block by transferring the cells back to  $25^\circ\text{C}$  for 1 h resulted in compaction of both septin and Mid2 domains with the domain widths decreasing to  $1.2 \mu\text{m} \pm 0.1$  and  $1.3 \mu\text{m} \pm 0.2$ , respectively (Figure S3C).

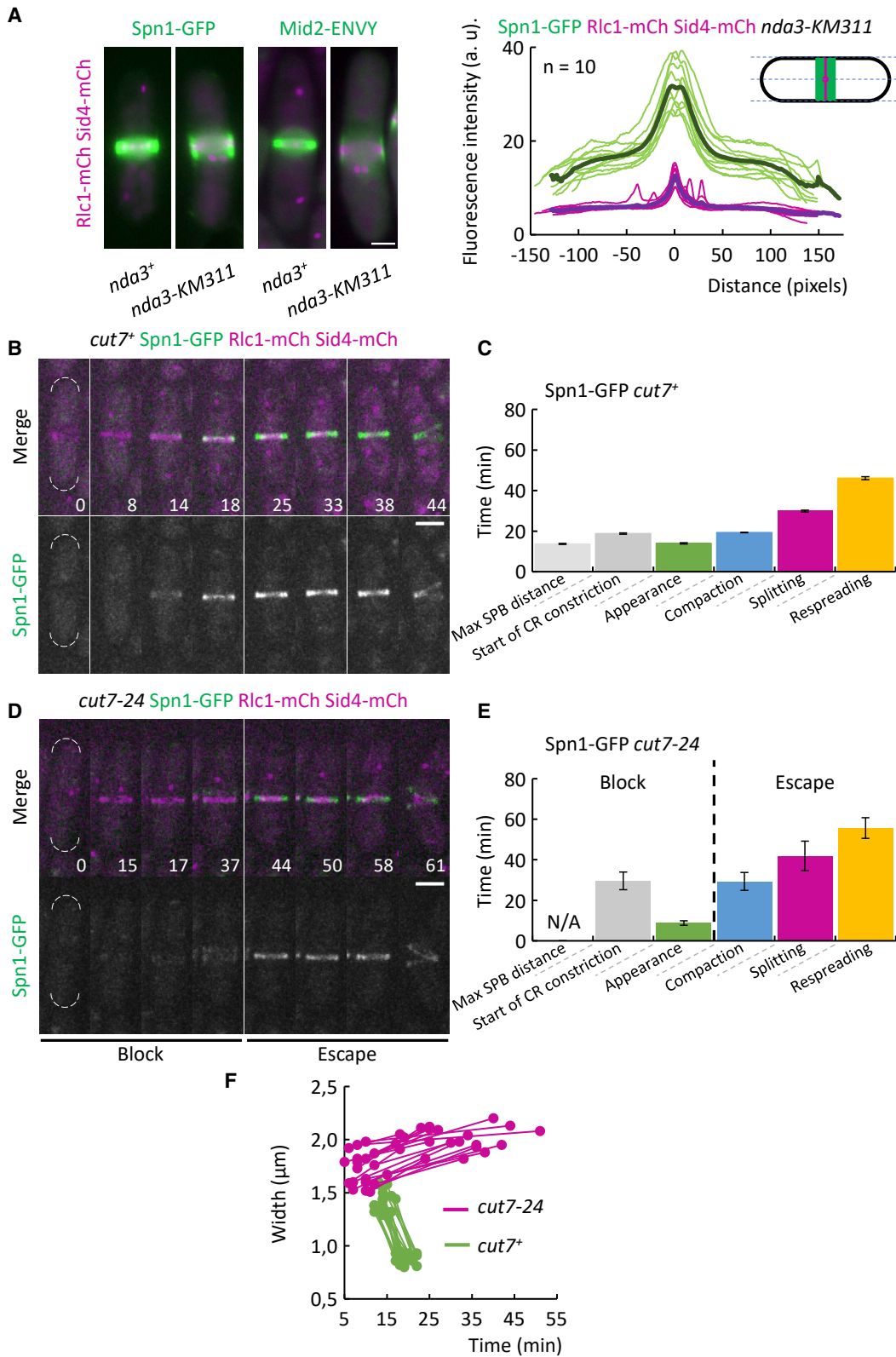
We also used the thermosensitive kinesin-5 *cut7-24* mutant, which forms a monopolar mitotic spindle at  $36^\circ\text{C}$ , producing a temporary cell-cycle arrest in metaphase with a high CDK activity (Hagan and Yanagida, 1990). Live cell imaging of the wild-type cells at  $36^\circ\text{C}$  slightly accelerated mitosis progression compared to  $25^\circ\text{C}$  (Figures 4B and 4C compared to Figures 1A and 1D), but septin appearance occurred again at the time of maximum spindle elongation, and septin compaction coincided with the initiation of CR constriction, like at  $25^\circ\text{C}$ . In the *cut7-24* mutant, since no SPB separation could be observed, the time of transfer to  $36^\circ\text{C}$  was defined as time 0 for this experiment. The timing measured in this background can therefore not be compared to those measured in the wild-type strain. In the *cut7-24* mutant, septins appeared 9 min after the temperature shift in a large band ( $1.7 \pm 0.2 \mu\text{m}$  compared to  $1.45 \pm 0.1 \mu\text{m}$  in control cells, Figure 4F) and were excluded from the CR as seen in *nda3-KM311* cells. Unlike in control cells, where septins compacted to  $0.9 \pm 0.05 \mu\text{m}$ , these bands spread along the membrane over time in *cut7-24* cells ( $2.0 \pm 0.1 \mu\text{m}$  Figures 4D–4F), until they escaped the cell-cycle arrest  $29.6 \pm 4.4$  min in average after transfer to  $36^\circ\text{C}$ , as visualized by the initiation of CR constriction, coincident with septin band compaction ( $t = 29.3 \pm 4.4$  min). Later on, septins were observed as two rings at  $t = 41.9 \pm 7.3$  min until cell separation at  $t = 55.6 \pm 5.1$  min (Figures 4D and 4E).

Live cell imaging of the Mid2-ENVY in the *cut7-24* mutant at  $36^\circ\text{C}$  showed that Mid2 also formed a faint band surrounding the CR that enlarged slightly over time (Figures S3D–S3H) until cells escaped mitosis and CR constriction initiated ( $t = 33.6 \pm 2.3$  min, Figures S3F–S3H). Ring splitting and re-spreading were strongly delayed ( $t = 45.7 \pm 1.2$  min and  $t = 66.4 \pm 5.2$  min, respectively; Figure S3G).

Overall, these data suggest that high CDK activity does not interfere with initial septin and Mid2 recruitment. However, later events, such as further septin accumulation at the division site and septin compaction and re-spreading may require a reduction in CDK activity. This in turn suggests that Mid2 bundling activity may be impaired upon high CDK activity.

### Figure 3. Mid2 interacts with other Mid2 molecules through different regions of the protein, contributing to septin compaction

- (A) Co-immunoprecipitation assay between Mid2-mEGFP and Mid2-13xMyc or Mid2  $\Delta$ PH-13xMyc in the presence or in absence of Spn1. Magenta arrowheads show the expected Mid2 bands; black arrowheads show degradation products; green arrowheads show the immunoglobulin bands.
- (B) Co-immunoprecipitation assay between the PH domain of Mid2 tagged with 13xMyc and the PH domain of Mid2 tagged with GFP in the presence or in absence of Spn1. Magenta, black, and green arrowheads show the expected Mid2 bands, degradation products, and the immunoglobulin bands, respectively.
- (C) Diagram showing the domain distribution of the anillin-like protein Mid2.
- (D) *In vitro* binding assay of recombinant MBP-Mid2-PH with GST-Mid2-PH or GST alone. Left panels show the inputs. Right panels show the GST pulldowns.
- (E) Kymographs showing the medial region of WT (left) or *mid2 $\Delta$ PH-GBP* cells (right) producing Spn1-GFP. There is a 2-min interval between successive images. Scale bar:  $2 \mu\text{m}$ .
- (F) Quantification of the extent of Spn1-GFP width compaction over time in WT (green) and in *mid2 $\Delta$ PH-GBP* cells (magenta).  $n = 20$  for WT cells and 25 for *mid2- $\Delta$ PH-GBP* cells.
- (G) Percentage of septating cells in the indicated strains at log phase.  $n < 500$  cells per strain and experiment. Mean  $\pm$  SD are shown. See also Figure S2.



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This prompted us to investigate if Mid2 still had the ability to self-interact in this context. To do so, we performed co-immunoprecipitations between Mid2-13Myc and Mid2-GFP, Mid2  $\Delta$ PH-GFP, or the single PH domain of Mid2 tagged with GFP in the control or *nda3-KM311* mutant cells, after 8 h of incubation at 18°C (Figure S2D). These experiments did not reveal any modification in the ability of Mid2 to self-interact.

We then tested if high CDK activity could reduce the avidity of Mid2 for septins. To test this hypothesis, co-immunoprecipitation assays between Spn1-GFP and Mid2-13Myc were performed in control and *nda3-KM311* cells after 8 h of incubation at 18°C or after 8 h at 18°C followed by 1 h at 25°C to release cells from the mitotic block (Figure S1B). Although the amount of Mid2 in mitotic cells is higher because its expression peaks at this stage of the cell cycle (Alonso-Nunez et al., 2005), these experiments did not reveal significant differences in the ability of Mid2 to interact with septins.

Altogether, these data suggest that, although Mid2 is necessary for septin compaction, it is not proficient for this function when CDK activity is high, independently of its ability to self-interact or to interact with septins. Since Mid2 is hyperphosphorylated during mitosis (Tasto et al., 2003), one hypothesis is that Mid2 phosphorylation prevents septin compaction. Or else, CDK activity regulates septin compaction independently of Mid2.

#### Mid2 controls the orientation of septin filaments

To solidify our hypothesis that Mid2 functions as a septin filament bundler, we decided to study the role of Mid2 in septin filament organization using polarization-resolved fluorescence microscopy (Vrabiou and Mitchison, 2006; DeMay et al., 2011a, 2011b; Kress et al., 2013; Wang et al., 2013). Polarimetry measurements with the regular Spn1-GFP fusion revealed randomly oriented GFP dipoles at all stages of cytokinesis (septin recruitment, compaction, splitting, and re-spreading) consistent with the flexibly linked GFP exploring orientations in all directions relative to Spn1 orientation (Figure S4A).

Thus, to use dipole orientation measurements of the GFP as a readout of septin filament orientation, the mobility of GFP relative to Spn1 had to be minimized. We therefore screened for Spn1-GFP fusions in which several amino acids between the C terminus of Spn1 and the N terminus of GFP were deleted without impairing Spn1 localization and function nor GFP fluorescence, and we identified a fusion of Spn1 to rotationally constrain GFP (conGFP), Spn1-conGFP (see STAR Methods for details).

The Spn1-conGFP construct was produced from the genomic *spn1* locus, under the control of its own promoter, giving rise to

cells with a wild-type phenotype indicating that the fusion was functional. Accordingly, Spn1-conGFP localized and redistributed similarly to Spn1-GFP (Figures S4B and S4C).

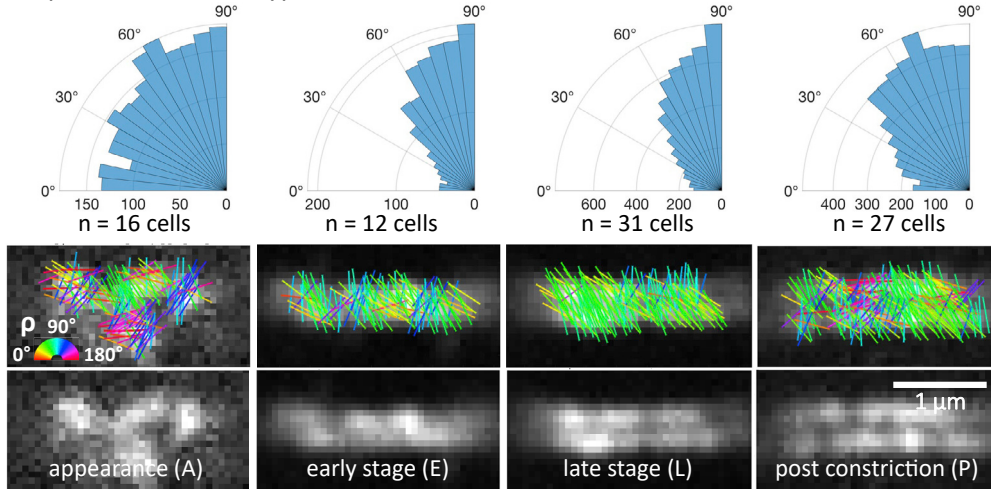
We measured GFP dipole orientation at four different stages during cytokinesis, namely septin recruitment (appearance), septin compaction (early), CR constriction (late), and CR disassembly (post-constriction) in this strain. We found that GFP dipoles were highly disordered when septins are first recruited to the division site, as evidenced by the random distribution of angles between the GFP dipoles and the axis of the CR (Figure 5A). However, when septins compacted into tight rings and throughout CR constriction, the distribution of angles between GFP dipoles and the CR axis became narrow, with GFP dipoles mostly oriented perpendicular to the CR axis (Figure 5A). Their distribution eventually started becoming broader again after CR disassembly, when the septin rings start to collapse. The narrowing of the GFP dipole orientation distributions upon and throughout CR constriction, which also persists post-constriction, indicates that the associated septin filaments are highly organized during these stages.

To determine the orientation of the GFP dipoles relative to septin filaments and be able to deduce septin filament orientation with respect to the CR axis, we took advantage of a Spn4-conGFP fusion (Figures S4D and S4E; see STAR Methods for details), which in addition to its normal localization to the septin ring (white asterisks in Figure S4D), also assembled into ectopic elongated cytoplasmic bars (arrows in Figure S4D). These ectopic bars most likely represent bundles of tightly packed parallel septin filaments, similar to elongated bar-like septin structures induced in *Ashbya gossypii* upon treatment with the septin filament stabilizing drug forchlorfenuron (DeMay et al., 2011a) and very similar to the ectopic bar-like structures detected in *spn2 $\Delta$*  cells containing both Spn1 and Spn4 (An et al., 2004). GFP dipoles were perpendicular to these septin bars (Figure S4D), suggesting that the septin filaments are oriented perpendicular to the GFP dipole in the Spn4-conGFP fusion. Importantly, GFP dipoles were also perpendicular to the CR axis during CR constriction and disassembly in these cells (Figure S4E), suggesting that septin filaments run parallel to the CR axis of constriction. Given that Spn4 is in a complex with Spn1 (An et al., 2004), we deduce that the septin filaments in the Spn1-conGFP construct are also oriented perpendicular to the GFP dipoles, and we propose that initially randomly organized septin filaments orient parallel to the plane of CR constriction upon compaction and keep this orientation throughout CR constriction, progressively losing their organization after CR constriction as septin rings progressively collapse (Figure 5D).

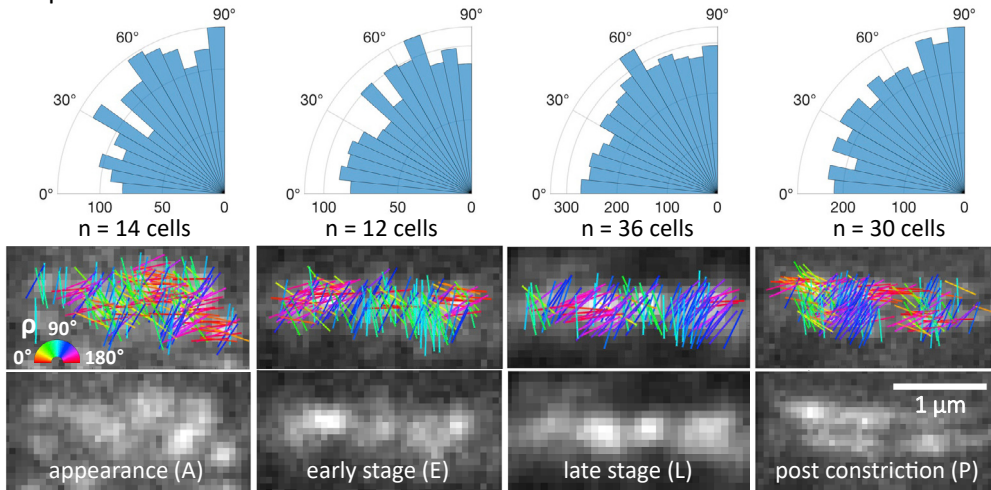
#### Figure 4. Septins accumulate as a broad band at the cell middle cortex in mitotic-blocked cells

(A–E) Epifluorescence images of cells producing Rlc1-mCherry, Sid4-mCherry, and Spn1-GFP (left panels) or Mid2-ENVY (right panels) in WT or *nda3-KM311* cells growing at 18°C for 4 h. Fluorescence intensity analysis of Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry signals along the entire cell length of *nda3-KM311* cells incubated during 7 h at 18°C. Individual lines are shown in light green (Spn1-GFP) and magenta (Rlc1-mCherry), while average lines are shown in dark green (Spn1-GFP) and purple (Rlc1-mCherry).  $n = 10$  cells (far right panel). A scheme of a cell displaying the line scan used for quantification of fluorescence intensity is shown. Time-lapse imaging of WT (B) or *cut7-24* cells (D) producing Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry growing at 36°C. Maximum intensity projections of confocal images are shown. Scale bars: 2  $\mu$ m. Plot showing Spn1-GFP transitions throughout cytokinesis in the WT (C) or the *cut7-24* mutant (E). Mean  $\pm$  SD are shown.  $n = 45$  cells in each case. The graph showing the behavior of the *cut7-24* cells was obtained by the analysis of the cytokinetic events of those cells that escaped the cytokinetic arrest (termed “Escape”). The backward analysis of previous cytokinetic events in the same cells is shown in the “Block” portion of the plot. (F) Quantification of the extent of Spn1-GFP width compaction over time in WT (green) and *cut7-24* cells (magenta).  $n = 20$  cells in each case. See also Figure S3.

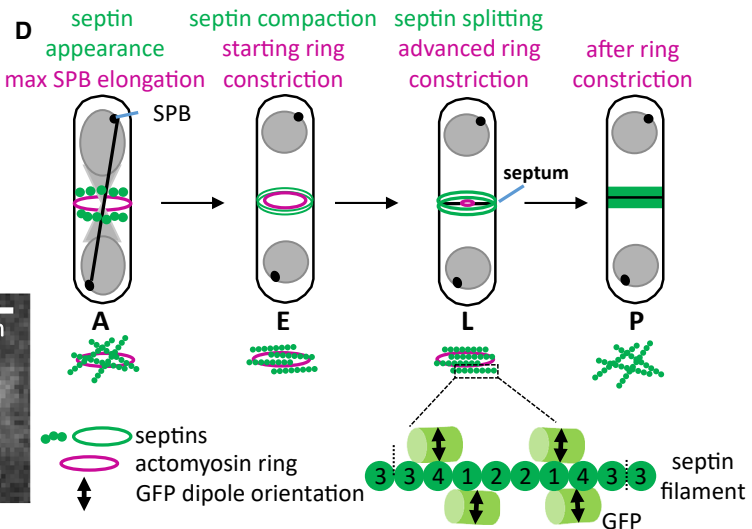
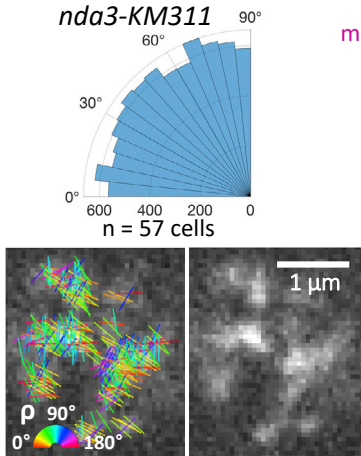
**A** Spn1-conGFP wild-type



**B** Spn1-conGFP *mid2Δ*



**C** Spn1-conGFP *nda3-KM311*



(legend on next page)



Septin filament orientation was then analyzed in *mid2Δ* cells. Similar to the wild-type, septin filaments appeared in a very disorganized manner in *mid2Δ* cells (Figure 5B). However, as CR constriction took place, GFP dipoles failed to reorient perpendicularly to the CR (Figure 5B). Septin filaments also failed to align in *nda3-KM311* cells where CDK activity is high (Figure 5C).

Overall, these data indicate that Mid2 contributes to the orientation of septin filaments parallel to the CR axis of constriction, supporting the hypothesis that Mid2 functions as a septin bundler (Figure 5D) during the septin compaction phase. Moreover, Mid2 bundling activity is negatively regulated by CDK activity in a direct or indirect manner, preventing septin orientation and compaction in early mitosis.

### SIN activity and CR constriction are required for normal septin accumulation and compaction at the division site

Since septin compaction is coincident with the initiation of CR constriction, an event known to be triggered by the SIN pathway, we next wondered if septins and Mid2 were also under the control of this pathway.

To test this hypothesis, we analyzed the behavior of septins in the temperature-sensitive hypo-active SIN mutant *sid2-250* at the non-permissive temperature of 36°C (Balasubramanian et al., 1998). Live cell imaging of *sid2-250* cells producing Spn1-GFP revealed that in the majority of cells, septins appeared as a diffuse cortical band at the normal timing compared to control cells (Figure 6A). However, the intensity of the septin signal was diminished by ~4.5-fold (Figure 6B), and compaction failed in 75% of cells, while a loose compaction was observed in the remaining 25%. Indeed, quantification of the extent of compaction showed that, while in control cells the width of the septin domain decreased from  $1.5 \pm 0.1 \mu\text{m}$  to  $0.9 \pm 0.1 \mu\text{m}$  after compaction, in the majority of *sid2-250* mutant cells, the initial width of the septin domain increased instead from  $1.6 \pm 0.1 \mu\text{m}$  to  $1.9 \pm 0.1 \mu\text{m}$  in a similar period of time, although it decreased in a minority of cells (from  $1.6 \pm 0.1 \mu\text{m}$  to  $1 \pm 0.1 \mu\text{m}$ ) (Figure 6C).

The lack of septin compaction suggested a similar defect in the function of Mid2. Indeed, *sid2-250* cells displayed a strong defect (3.5-fold) in the accumulation of Mid2 (Figures S5A and S5B), similar to the one observed in Spn1 accumulation.

Altogether, these data suggest that the SIN pathway is not responsible for the initial recruitment of septins or Mid2, but it

is required for their accumulation. The defects observed in septin compaction point toward an important role of the SIN in this event.

Since the SIN pathway triggers CR constriction,  $\beta(1,3)$ -glucan synthase Bgs1 localization, and septum synthesis initiation (Cortés et al., 2018), we next wondered if the SIN acts directly on septin compaction or indirectly by inducing CR constriction.

To test this, we decided to use the Bgs1 mutant *cps1-191*, which halts cytokinesis progression due to the inability to construct a septum, with an assembled CR that cannot constrict. Interestingly, the SIN pathway is active in this mutant and required for the stability of the CR (Liu et al., 2000), which provides the best scenario to answer this question. Analysis of Spn1-GFP and Mid2-ENVY behavior in a *cps1-191* mutant at restrictive temperature showed a very defective accumulation in both cases (Figures 6D, 6E, S5C and S5D). Moreover, septins did not compact (Figure 6F) and eventually dissociated from the division site, even when the CR remained stable (Figures 6D and S5C).

Altogether, these data strongly suggest that the CR itself is not sufficient to promote septin accumulation, and that the SIN pathway by inducing CR constriction and/or septum synthesis initiation allows septin filament accumulation. Following Mid2 recruitment, septin filaments further accumulate, compact, and align parallel to the axis of CR constriction to assemble the septin rings in a process strictly regulated by CR constriction and septum ingression.

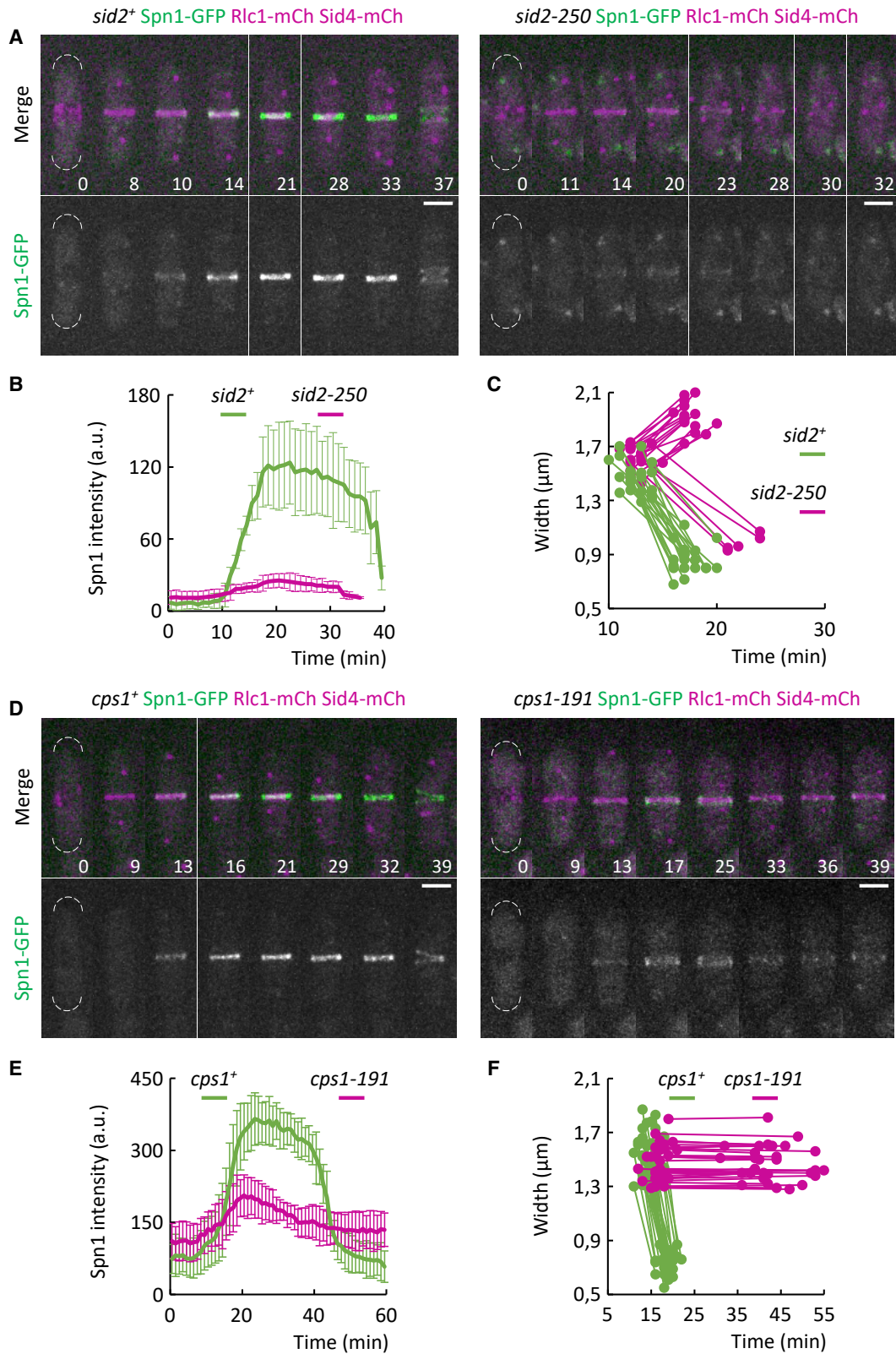
## DISCUSSION

The assembly of macromolecular structures is a distinctive feature of cytoskeletal elements. Septins are GTP-binding proteins with the ability to form non-polar filaments, which can assemble into a variety of membrane-associated structures such as filament networks or rings (Ong et al., 2014). A number of factors influence the ability of septins to get assembled into those structures, such as the septin monomers themselves, the lipid bilayer composition, post-translational modifications, or regulatory proteins.

Here, we investigated the role of the septin-binding protein, Mid2 in the assembly of supramolecular septin filament structures, using fission yeast as a model system. Mid2 is an anillin-like protein, previously involved in the stability of the septin ring assembled during cytokinesis (Berlin et al., 2003; Tasto et al., 2003). Similar to

### Figure 5. Septin filament organization changes during CR constriction and septum synthesis depend on Mid2

(A) Polar histograms of GFP dipole orientations of Spn1-conGFP filaments from WT cells at different stages of cytokinesis (top panels). The number of analyzed cells per stage is indicated below each polar histogram (typically 120–200 analyzed pixels per cell). Confocal images of Spn1-conGFP from selected WT cells and an overlaid color-coded stick representation of the measured orientations (angles  $\rho$ ) per analyzed pixel at each cytokinetic stage (bottom panels). Scale bar: 1  $\mu\text{m}$ . (B) Polar plots of GFP dipole orientations of Spn1-conGFP filaments from *mid2Δ* cells at different stages of cytokinesis (top panels). The number of analyzed cells per stage is indicated below each polar histogram. Confocal images of Spn1-conGFP from selected *mid2Δ* cells and an overlaid color-coded stick representation of the measured orientations per pixel, as in (A), at each cytokinetic stage (bottom panels). Scale bar: 1  $\mu\text{m}$ . (C) Polar histogram of GFP dipole orientations of Spn1-conGFP filaments from 57 *nda3-KM311* cells incubated 7 h at 18°C (top panel). Confocal images of Spn1-conGFP from *nda3-KM311* cells and an overlaid color-coded stick representation of the measured orientations per pixel, as in (A) (bottom panels). Scale bar: 1  $\mu\text{m}$ . (D) Schematic representation of septin localization related to major CR and septum transitions (top) and detail of septin filament orientation at each stage (bottom). The cartoon on the bottom right depicts the orientation of a septin filament with respect to the orientation of GFP dipoles (double arrowheads), as deduced from Spn4-conGFP fusions (see Figures S3D and S3E). The GFP barrels orient with their long axes parallel to the septin filament axis, i.e., with the GFP dipoles perpendicular to the septin filament axis. The numbers correspond to the septin subunits Spn1–4, and the dashed lines delimit a putative octameric septin complex within the filament. See also Figure S4.



(legend on next page)

other anillin-like proteins, such as budding yeast Bud4, Mid2 is recruited at the end of cytokinesis. In contrast to budding yeast, in which septins are first recruited to the presumptive bud site at the onset of G1 (Juanes and Piatti, 2016), in fission yeast, septins localize to the cortex during cell division. Indeed, septin recruitment requires the assembly of the CR prior to their localization.

Our data indicate that septins localize to the division site as a diffuse band that compacts into a tight ring by the time of CR constriction. The intensity of this band increases as septin filament compaction takes place. This might explain the differences found with Wu and collaborators, who established that septins appeared around 22 min after SPB separation (Wu et al., 2003). This compaction phenomenon is coincident with the appearance of Mid2, suggesting a cause-effect relationship. Accordingly, the lack of Mid2 results in the inability of septins to compact.

How might Mid2 trigger septin band compaction? One possibility is that Mid2 serves as a bridge connecting septin filaments. In fact, anillin is a scaffold protein, able to interact with many other proteins (Piekny and Glotzer, 2008). Anillin (and Mid2) contains a C-terminal septin-interacting domain. If Mid2 were to bundle septin filaments, it should be able to interact with itself so that each molecule could bind to a different septin filament. Our biochemical analysis of Mid2 self-association shows that Mid2 can directly interact with other Mid2 molecules via its C-terminal PH domain. The Mid2 fragment lacking the PH domain can also self-associate independently of septins (although we cannot rule out that other proteins may bridge this interaction) and is sufficient to trigger septin compaction if an interaction with septins is artificially triggered using the GBP system. This suggests that at least two molecules of Mid2 could bridge two septin filaments together. Importantly, in the absence of Mid2, septin accumulation is strongly affected; since Mid2 recruitment to the division site is strongly dependent on septins (Berlin et al., 2003; Tasto et al., 2003), we propose that septins and Mid2 establish a positive feedback loop to accumulate at the division site.

Septin filaments show a dynamic organization pattern during cell cycle progression. Our data based on polarization microscopy let us conclude that septin filaments are not organized when they first appear at initial stages of cytokinesis. Coincident with the arrival of Mid2, septin filaments become aligned parallel to the CR axis of constriction, a pattern that is absent upon deletion of *mid2*. This strongly supports the notion that Mid2 functions to bridge septin filaments and organize them. In budding yeast, after the initial recruitment to the bud site, septins show a “fluid” state, followed by a “frozen” state once the ring trans-

forms into a collar (Caviston et al., 2003; Dobbelaere et al., 2003); after ring duplication, septin dynamics increase again. At the same time, during the collar stage, septin filaments are parallel to the mother-bud axis (Vrabiou and Mitchison, 2006), while they reorganize to become parallel to the plane of CR constriction in later stages (DeMay et al., 2011a) in a process that might implicate disassembly and reassembly of septin filaments (Ong et al., 2014). This process might be different in fission yeast, where septin filaments only transition from a disordered to an ordered organization, parallel to the plane of CR constriction. In any case, anillin-like proteins seem to play the conserved role of septin filament reorganization and septin ring stabilization, as is also the case for budding yeast anillin Bud4 (McQuilken et al., 2017).

What signals trigger the dynamic behavior of septins? Our data are consistent with a tight control from the cell cycle machinery. In wild-type cells, septin recruitment to the division site takes place during late anaphase, while Mid2 appears later, by the time of detectable CR constriction and septum synthesis initiation. Analysis of Spn1 and Mid2 behavior in *nda3-KM311* or *cut7-24* mutants blocked at early stages of mitosis demonstrates that both strongly accumulate at the division site when CDK activity is high. Although we have no proof of a direct phosphorylation by CDK, Mid2 is highly phosphorylated during septation (Tasto et al., 2003), and our non-published data show a number of CDK consensus sites phosphorylated in *nda3-KM311*-blocked cells. This would not be surprising since both septins and anillin-like proteins have been shown to be direct targets of CDK in other model systems (Sinha et al., 2007; Eluere et al., 2012).

Analysis of septins and Mid2 in mitotic-blocked cells opens two questions. First, septin and Mid2 initial meshworks are physically excluded from the precise CR position on the cortex, raising the question on the molecular mechanisms at work to allow such an exclusion and whether steric hindrance on the plasma membrane surface might be sufficient. Whatever the mechanisms, from this observation it can be inferred that the compaction of the two septin domains on each side of the CR might produce a double septin ring, although it may not be detectable until the septum has grown enough to push them apart. Initial septin recruitment observation by super-resolution microscopy will contribute to clarify this point. Second, even though septins and Mid2 co-localize and interact in this context, this is not sufficient to trigger septin ring compaction. This suggests that another mechanism contributes to septin ring compaction. It might involve post-translational modifications of Mid2, additional

**Figure 6. Septin recruitment and compaction are defective in the SIN mutant *sid2-250* and the Bgs1 mutant *cps1-191***

- (A) Time-lapse imaging of WT (left) and *sid2-250* cells (right), producing Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry incubated at 36°C. Maximum intensity projections of confocal images are shown. Scale bars: 2 μm.
- (B) Analysis of Spn1-GFP fluorescence intensity from mitotic onset throughout cytokinesis in WT (green) and *sid2-250* cells (magenta). The average line graphics ± SD are displayed. n = 6 cells in each case.
- (C) Quantification of the extent of Spn1-GFP width compaction over time in WT (green) and *sid2-250* cells (magenta). Note that some *sid2-250* cells escape the SIN phenotype. n = 20 cells.
- (D) Time-lapse imaging of WT (left) and *cps1-191* cells (right), producing Spn1-GFP, Rlc1-mCherry, and Sid4-mCherry incubated at 36°C. Maximum intensity projections of confocal images are shown. Scale bars: 2 μm.
- (E) Analysis of Spn1-GFP fluorescence intensity from mitotic onset throughout cytokinesis in WT (green) and *cps1-191* cells (magenta). The average line graphics ± SD are displayed. n = 6 cells.
- (F) Quantification of the extent of Spn1-GFP width compaction over time in WT (green) and *cps1-191* cells (magenta). n = 20 cells in each case. See also Figure S5.

proteins working together with Mid2 to promote septin filament bundling, or yet another mechanism to be identified.

Importantly, septin meshwork compaction is coincident with the onset of CR constriction triggered by the SIN pathway, and we found the SIN plays a critical role in septin ring assembly. Although septin accumulation is inhibited in latrunculin A-treated cells, in which CR assembly is abolished (Tasto et al., 2003), we think the CR is not sufficient for septin accumulation and compaction, given that in SIN mutants, the actin ring begins to assemble (Hachet and Simanis, 2008), but septin recruitment is strongly impaired. The poor accumulation of Mid2 in the absence of SIN activity may arise from the defective accumulation of septin filaments, which could also affect their reorganization. This suggests that the SIN, directly or indirectly, regulates Mid2 to control septin accumulation and compaction.

Moreover, we also found that the Bgs1 mutant allele *cps1-191*, in which the CR assembles and is stable and the SIN pathway is active but CR constriction and initial septum synthesis are defective (Liu et al., 2000), phenocopies the SIN mutant *sid2-250* in terms of septin defects. This suggests a role for the septum structure to trigger septin and Mid2 accumulation and compaction. Although we cannot rule out the hypothesis that the SIN may target Mid2, this demonstrates that SIN activity on its own is not directly responsible for triggering the full accumulation of septins and Mid2 to promote septin meshwork compaction. Rather, CR constriction and septum formation by themselves might cooperate with Mid2 to promote septin filament accumulation and reorientation. Indeed, Bgs1 is the only glucan synthase localized to the division site at the time of CR constriction onset (Cortés et al., 2018). One possibility to explore in the future is that CR constriction or the initial deposition of septum material by the  $\beta(1,3)$ -glucan synthase Bgs1 may modify the plasma membrane locally at the ingression furrow, favoring Mid2 and septin accumulation on the cortex and septin filament reorientation.

### Limitations of the study

First, our data indicate that Mid2 can crosslink septin filaments by interacting with itself and septins, but we are still lacking an understanding on the high molecular order structures of Mid2 that are necessary to endow Mid2 with such properties. Second, Mid2-dependent septin compaction is halted when cells are blocked in mitosis showing that Mid2 activity is cell-cycle regulated. The molecular mechanisms at play remain unknown. Third, we are still missing information on how septin meshwork assembly is initiated specifically at the division site, in the vicinity of the CR. Anillin is known to bridge actin and septin cytoskeletons during cytokinesis in animal cells, but there is no report of Mid2 interaction with actin structures in fission yeast. Further biochemical and structural analysis are required to address these questions.

In summary, we have shown that the crosslinking activity of the anillin-like protein Mid2 promotes the alignment septin filaments parallel to the division axis, promoting their compaction from an initial broad band into tight rings at the onset of CR constriction and septum formation. The activity of the SIN is necessary but not sufficient to trigger this effect, and CR constriction, Bgs1, and/or septation may be the real trigger. It

will be interesting to explore whether the cleavage furrow acts on the properties of other anillin-like proteins in a similar manner in other systems.

### STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **RESOURCE AVAILABILITY**
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- **EXPERIMENTAL MODEL AND SUBJECT DETAILS**
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  - Septin-GFP fusions for orientation measurements
  - Spinning-disk polarization-resolved fluorescence microscopy
- **QUANTIFICATION AND STATISTICAL ANALYSIS**

### SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.celrep.2022.110722>.

### ACKNOWLEDGMENTS

We thank Vincent Fraisier and Carmen Castro for maintenance of the microscopes for imaging performed at the Institut Curie, Paris (PICT-IBISA, part of France-Bioimaging national research infrastructure), and at the IBFG (University of Salamanca/CSIC, Salamanca, Spain), respectively, and Cristel Chandre (I2M, CNRS, Marseille, France) for the Polarimetry Analysis application and MATLAB codes used for polarimetry data analysis. We are grateful to Henar Valdivieso and Jurg Bähler for sharing strains and plasmids. We also wish to thank Pilar Pérez and Carlos Rodríguez for their very helpful comments on the manuscript. This work was supported by a grant from Fondation ARC to A.P. (PJA 20171206550). F.A. received a fourth-year PhD fellowship from Fondation ARC (DOC20180507063). A.P. is a member of the Labex CeTisPhyBio (ANR11-LABX-0038). S.A.R. is a Ramon y Cajal fellow from the University of Salamanca (RYC-2016-20652). This work was also supported by the grants PGC2018-094090-B-I00 and PGC2018-098924-B-I00 (Ministerio de Ciencia e Innovación, MICINN, Spain, and ERDF, EU), and CSI150P20 and “Escalera de Excelencia” CLU-2017-03 (Junta de Castilla y León 14–20, Spain and ERDF, EU) and ANR-17-CE13-0014; SEPTIMORF (Agence Nationale de la Recherche).

### AUTHOR CONTRIBUTIONS

Conceptualization, F.A., A.P., and S.A.R.; Methodology, F.A., M.M., S.B., A.P., and S.A.R.; Investigation, F.A., M.M., M.H., and S.A.R.; Writing – Original Draft, F.A., A.P., and S.A.R.; Writing – Review & Editing, M.M., S.B., J.C.R., S.B., A.P., and S.A.R.; Funding Acquisition, F.A., M.M., J.C.R., S.B., A.P., and S.A.R.; Supervision, A.P. and S.A.R.

### DECLARATION OF INTERESTS

The authors declare no competing financial interests.



**INCLUSION AND DIVERSITY**

One or more of the authors of this paper self-identifies as a member of the LGBTQ+ community. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list.

Received: June 1, 2021  
Revised: February 10, 2022  
Accepted: March 30, 2022  
Published: April 19, 2022

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
Monoclonal anti-GFP	Sigma	Cat No. 11814460001; RRID: AB_390913
Monoclonal anti-myc	Sigma	Cat No. 11667149001; RRID: AB_390912
Monoclonal anti-GST	Thermo	Cat No. MA4-004; RRID: AB_10979611
Monoclonal anti-MBP	New England Biolabs	Cat No. E8032S; RRID: AB_1559730
Polyclonal anti-GFP	Thermo	Cat No. A-6455; RRID: AB_221570
Goat anti-mouse IgG-HRP conjugated	Bio-Rad	Cat No. 1706516; RRID: AB_11125547
Goat anti-mouse IgG-HRP conjugated	Jackson ImmunoResearch	Cat No. 115-035-008; RRID: AB_2313585
<b>Experimental models: Organisms/strains</b>		
<i>spn1-GFP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP5501
<i>mid2-ENVY:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP5678
<i>spn1-GFP:kanMX6 mid2Δ::kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP5582
<i>mid2-13xMyc:Nat ade6-M210 ura4-D18 leu1-32</i>	This study	AP5823
<i>leu1+:Pmid2-mid2-mEGFP mid2-13xMyc:Nat ade6- ura4-D18 leu1-32</i>	This study	AP5831
<i>leu1+:Pmid2-mid2ΔPH-mEGFP mid2-13xMyc:Nat ade6- ura4-D18 leu1-32</i>	This study	AP5829
<i>spn1Δ::ura4+ mid2-13xMyc:Nat ade6- ura4-D18 leu1-32</i>	This study	AP5918
<i>spn1Δ::ura4+ leu1+:Pmid2-mid2-mEGFP mid2-13xMyc:natMX6 ade6-</i>	This study	AP5920
<i>spn1Δ::ura4+ leu1+:Pmid2-mid2ΔPH-mEGFP mid2-13xMyc:natMX6 ade6- ura4-D18 leu1-32</i>	This study	AP5927
<i>leu1+:Pmid2-mid2-PH-13xMyc ade6- ura4-D18 leu1-32</i>	This study	AP5814
<i>mid2-PH-mEGFP:kanMX6 leu1+:Pmid2-mid2-PH-13xMyc ade6- ura4-D18 leu1-32</i>	This study	AP5932
<i>spn1Δ::ura4+ leu1+:Pmid2-mid2-PH-13xMyc ade6- ura4-D18 leu1-32</i>	This study	AP5930
<i>spn1Δ::ura4+ mid2-PH-mEGFP:kanMX6 leu1+:Pmid2-mid2-PH-13xMyc ade6- ura4-D18 leu1-32</i>	This study	AP5943
<i>mid2Δ::mEGFP:kanMX6 ade6::Pmid2-mid2-PH-13xMyc ade6- ura4-D18 leu1-32</i>	This study	SR1370
<i>mid2-ΔPH-GBP:kanMX6 spn1-GFP:hphMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	SR1216
<i>mid2-ΔPH-GBP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	SR1210
<i>spn1-GFP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP5515
<i>spn1-GFP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 cut7-24 ade6- ura4-D18 leu1-32</i>	This study	AP6268

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
<i>mid2-ENVY:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP5514
<i>spn1-msfGFPΔN12:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP6313
<i>spn1-msfGFPΔN12:kanMX6 mid2::hphMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP6334
<i>spn1-msfGFPΔN12:kanMX6 nda3-KM311 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP6324
<i>spn1-GFP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 sid2-250 ade6- ura4-D18 leu1-32</i>	This study	AP5585
<i>spn1-GFP:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 cps1-191 ade6- ura4-D18 leu1-32</i>	This study	AP5633
<i>mid2-13xMyc:natMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP6274
<i>spn1-GFP:KanMX6 mid2-13xMyc:natMX6 ura4-D18 leu1-32</i>	This study	AP6276
<i>spn1-GFP:KanMX6 mid2-13xMyc:natMX6 nda3-KM311 ura4-D18 leu1-32</i>	This study	AP6280
<i>mid2-13xMyc:natMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP6274
<i>leu1+:Pmid2-mid2-mEGFP mid2-13xMyc:natMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP6069
<i>leu1+:Pmid2-mid2ΔPH-mEGFP mid2-13xMyc:natMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP6073
<i>leu1+:Pmid2-mid2-PH-mEGFP mid2-13xMyc:natMX6 nda3-KM311 ade6- ura4-D18 leu1-32</i>	This study	AP6074
<i>mid2-mEGFP:kanMX6 ade6:mid2-13xMyc ade6- ura4-D18 leu1-32</i>	This study	SR1344
<i>spn1-5Δ::ura4+ mid2-mEGFP:kanMX6 ade6:mid2-13xMyc ade6- ura4-D18</i>	This study	SR1234
<i>mid2-ΔPH-mEGFP:kanMX6 ade6:mid2-13xMyc ade6- ura4-D18 leu1-32</i>	This study	SR1346
<i>spn1-5Δ::ura4+ mid2-ΔPH-mEGFP:kanMX6 ade6:mid2-13xMyc ade6- ura4-D18</i>	This study	SR1236
<i>PH-mid2-mEGFP:kanMX6 ade6: PH-mid2-13xMyc ade6- ura4-D18 leu1-32</i>	This study	SR1348
<i>spn1-5Δ::ura4+ PH-mid2-mEGFP:kanMX6 ade6: PH-mid2-13xMyc ade6- ura4-D18</i>	This study	SR1237
<i>mid2-ENVY:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 cut7-24 ade6- ura4-D18 leu1-32</i>	This study	AP6271
<i>spn4ΔC6-msfGFPΔN7:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 ade6- ura4-D18 leu1-32</i>	This study	AP6234
<i>mid2-ENVY:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 sid2-250 ade6- ura4-D18 leu1-32</i>	This study	AP5564
<i>mid2-ENVY:kanMX6 rlc1-mCherry:natMX6 sid4-mCherry:hphMX6 cps1-191 ade6- ura4-D18 leu1-32</i>	This study	AP5855

**Oligonucleotides**

See Table S1	N/A	N/A
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REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Continued</b>		
<b>Recombinant DNA</b>		
pJK148- <i>Pmid2-mid2-mEGFP-Tnmt1</i>	This study	pAF15
pJK148- <i>Pmid2-mid2ΔPH-mEGFP-Tnmt1</i>	This study	pAF16
pJK148- <i>Pmid2-mid2-PH-mEGFP-Tnmt1</i>	This study	pAF17
pFA6a- <i>Pmid2-mid2-mEGFP-kanMX6-Tmid2</i>	This study	pSRP75
pFA6a- <i>Pmid2-mid2ΔPH-mEGFP-kanMX6-Tmid2</i>	This study	pSRP76
pFA6a- <i>Pmid2-mid2-PH-mEGFP-kanMX6-Tmid2</i>	This study	pSRP77
pBP-Ade6- <i>Pmid2-mid2-13Myc-Tmid2</i>	This study	pSRP81
pBP-Ade6- <i>Pmid2-mid2ΔPH-13Myc-Tmid2</i>	This study	pSRP82
pBP-Ade6- <i>Pmid2-mid2-PH-13Myc-Tmid2</i>	This study	pSRP83
pFA6a- <i>Pmid2- mEGFP-kanMX6-Tmid2</i>	This study	pSRP78
pJK148- <i>Pmid2-mid2-PH-13Myc-Tnmt1</i>	This study	pAF18
pFA6a- <i>spn1-msfGFPΔN12-kanMX6</i>	This study	pAF32
pFA6a- <i>spn4ΔC6-msfGFPΔN7-kanMX6</i>	This study	pAF37
pFA6a- <i>Pmid2-mid2ΔPH-GBP-kanMX6-Tmid2</i>	This study	pSRP86
pJC20- <i>mid2-PH</i>	This study	pSRP96
pMALp2- <i>mid2-PH</i>	This study	pSRP90
pGADT7- <i>mid2-PH</i>	This study	pSRP68
pGBK7- <i>mid2-PH</i>	This study	pSRP71
<b>Software and algorithms</b>		
MetaMorph 7.7.8	Molecular Devices	<a href="https://www.moleculardevices.com">https://www.moleculardevices.com</a>
Fiji	National Institutes of Health	<a href="https://imagej.nih.gov/ij">https://imagej.nih.gov/ij</a>
MatLab	MathWorks	<a href="https://www.mathworks.com/products/matlab.html">https://www.mathworks.com/products/matlab.html</a>
<b>Other</b>		
Micro-Slide 8 Well Uncoated	IBIDI	AI-80821
Dynabeads protein A conjugated	Thermo	10006D
PureCube Gluthatione MagBeads	Cube Biotech	32225
CellLytic B Cell Lysis Reagent	Sigma	C8740
Amylose Resin	New England Biolabs	E8021S
Centricon YM-30	Millipore	4307

## RESOURCE AVAILABILITY

### Lead contact

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Sergio Rincon ([sarpadilla@usal.es](mailto:sarpadilla@usal.es)).

### Materials availability

All yeast strains and plasmids used in this study will be made available upon request without any restriction.

### Data and code availability

The data reported in this paper will be shared by the lead contact upon request.

This paper does not report original code.

Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

### Yeast genetics and culture conditions

Standard *S. pombe* media and genetic manipulations were used as described in (Moreno et al., 1991). All strains used in the study were isogenic to wild type h<sup>-</sup> 972 (see Key Resources Table). Strains from genetic crosses were selected by random spore germination and replica in plates with the appropriate supplements or drugs.

Plasmids and DNA transformations were performed by using the lithium acetate-DTT method. Briefly, 20 mL of exponentially growing cells (optical density 0.5–0.8 at 600 nm) were harvested by centrifugation and washed with 10 mM Tris HCl pH 7.4. Then, these cells were re-suspended in 100 mM lithium acetate with 10 mM DTT and incubated on an orbital wheel at room temperature for 40 min. 100  $\mu$ L of these cells were mixed with 80  $\mu$ L of 100 mM lithium acetate, 10  $\mu$ L of single stranded DNA from salmon testes (D9156-5ML, Sigma) and 2  $\mu$ g of the desired plasmid or the purified PCR product. After 10 min of incubation at room temperature on an orbital wheel, 300  $\mu$ L of PEG 4000, previously diluted 1:1 in 100 mM lithium acetate, were added. After a second round of 10 min at room temperature on the wheel, 15  $\mu$ L of DMSO were added and the cells were subjected to heat shock at 42°C for 20 min in a water bath. Cells were then washed, resuspended in 100  $\mu$ L of sterile water and plated on the appropriate selection plates.

## METHOD DETAILS

### Production of mutant and tagged strains

Wild type *mid2* was fused to the DNA sequence coding for the green fluorescent tag ENVY (Slubowski et al., 2015) or for the 13Xmyc tag at its 3' end by PCR amplification of a cassette with the corresponding fusion from the plasmid pFA6a-ENVY-*kanMX6* or pFA6a-13Xmyc-*natMX6*, respectively, as described in (Bahler et al., 1998).

To create the strains used for the co-immunoprecipitation experiments shown in Figures 3A, S1B and S2D, constructs containing *mid2*<sup>+</sup>, or the *mid2* fragments coding for Mid2- $\Delta$ PH (amino acids 1–581) or for the PH domain of Mid2 (amino acids 582–685) were cloned using XhoI and NotI sites into a pJK148-derived plasmid (Keeney and Boeke, 1994) containing *mEGFP* followed by the terminator of *nmt1*<sup>+</sup>, to be expressed under the control of the *mid2*<sup>+</sup> promoter cloned between KpnI and XhoI sites. The resulting plasmids were digested with NruI or Tth111 for plasmid linearization to allow integration in the endogenous *leu1-32 locus* upon yeast transformation.

To create the strains used for the co-immunoprecipitation experiments shown in Figures 3B, S2A and S2B, an integration plasmid for Mid2 constructs was made by insertion of 1 kb of the *mid2*<sup>+</sup> promoter and terminator into pFA6a-*mEGFP-kanMX6* (Bahler et al., 1998) between Sall and BamHI sites and SacI and SpeI sites, respectively. The *mid2*<sup>+</sup> open reading frame was then inserted upstream of *mEGFP* between BamHI and PacI. The *mid2* fragments coding for Mid2- $\Delta$ PH (amino acids 1–569) or for the PH domain of Mid2 (amino acids 569–706) were cloned into this plasmid, replacing full-length *mid2*<sup>+</sup>. pFA6a-*GFP-kanMX6*-derived plasmid was digested with NotI to release the linear DNA fragment that directs homologous recombination upon yeast transformation.

For production of Mid2-13Myc truncations in Figures S2A and S2B, 1 kb of the *mid2*<sup>+</sup> promoter was cloned into pBP-*ade6* (Beaudoin et al., 2006) using the sites Sall and XmaI. The resulting plasmid was used to clone a fragment coding for 13xmyc and 1 kb of *nmt1*<sup>+</sup> terminator obtained from pSR98 (Rincon et al., 2014) using the sites NotI and SacI. Finally, the *mid2* truncations were cloned in this plasmid using the sites XmaI and NotI. The resulting plasmids were digested with AatII for plasmid linearization to allow integration in the endogenous *ade6*<sup>+</sup> locus upon yeast transformation.

For production of the PH domain of Mid2 fused to 13Myc in the co-immunoprecipitation experiment shown in Figures 3B and a construct containing the *mid2* fragment coding for the PH domain of Mid2 (amino acids 582–685) was cloned using XhoI and NotI into a pJK148-derived plasmid containing 13xMyc followed by the terminator of *nmt1*<sup>+</sup>, to be expressed under the control of the *mid2*<sup>+</sup> promoter cloned between KpnI and XhoI. The resulting plasmid was digested with NruI for plasmid linearization and integration in the endogenous *leu1-32 locus* of a strain containing the PH domain of Mid2 (amino acids 582–685) followed by *mEGFP* expressed from *mid2*<sup>+</sup> endogenous locus.

To create the strains producing Spn1 fused to monomeric superfolder GFP lacking the 12 N-terminal amino acids (Spn1-conGFP, *spn1*<sup>+</sup>-*msfGFP $\Delta$ N12*), the *msfGFP* sequence was first amplified from plasmid pEX-A128-*msfGFP* and cloned into the pFA6a-*EGFP-kanMX6* plasmid (Bahler et al., 1998), replacing the original EGFP for *msfGFP* using the Gibson Assembly cloning strategy (Gibson et al., 2009). *msfGFP $\Delta$ N12* and *spn1*<sup>+</sup> were amplified and fused in a two-round PCR and the PCR product was used to replace *msfGFP* in the pFA6a-*msfGFP-kanMX6* plasmid. This construct was amplified with long oligos containing 80 bases from *spn1*<sup>+</sup> promoter and terminator to direct the homologous recombination, producing the integration module which was used to transform the wild type, *mid2 $\Delta$*  and *nda3-KM311* strains and replace *spn1*<sup>+</sup> with *spn1*<sup>+</sup>-*msfGFP $\Delta$ N12*. A similar strategy was used to create the strains producing Spn4 lacking the 6 C-terminal amino acids fused to GFP lacking the 7 N-terminal amino acids (Spn4-conGFP, *spn4- $\Delta$ C6-mEGFP- $\Delta$ N7*).

To create the strains producing Mid2- $\Delta$ PH-GBP, a DNA fragment coding for the GFP binding protein was obtained from pFA6a-*GBP-kanMX6* (Chen et al., 2017) by digestion with PacI-AscI, and cloned into the plasmid pFA6a-*mid2- $\Delta$ PH-mEGFP-kanMX6*, replacing the *mEGFP* for GBP. The obtained plasmid was linearized by digestion with NotI before yeast transformation.

All plasmids were confirmed by PCR and restriction enzyme digestion, and the DNA fragments amplified by PCR were confirmed by DNA sequencing.

### Microscopy and image analysis

Cells were incubated at 25°C in YE5S except in the case of cell cycle temperature-sensitive mutants (*cut7-24*, *sid2-250*, *cps1-191*), which were incubated for 3 h at 36°C or *nda3-KM311* cold sensitive mutant, which was incubated up to 7 h at 18°C.

Epifluorescence images were taken on a DMRXA2 upright microscope (Leica Microsystems), equipped with a 100 $\times$ /1.4NA oil immersion PlanApo objective and a Coolsnap HQ CCD camera (Photometrics). The exposure time was 2 s for GFP, 1 s for mCherry.

Quantification in the *nda3-KM311* mutant of Spn1-GFP fluorescence intensity with respect to the red fluorescence signal was performed by analysis along a line drawn throughout the cell length (dashed lines in [Figures 4A](#), [S3A](#) and [S3B](#)). The values obtained were background-subtracted using an equivalent region outside the cell. The line graphics obtained were then centered with respect to the position of the ring.

For time-lapse imaging, 1 mL of exponentially growing cells were harvested by centrifugation for 60 s at 800 g in a Minispin Eppendorf centrifuge equipped with an F-45-12-11 rotor, most of the supernatant was discarded and 1  $\mu$ L of the cells was deposited in a 2% YE5S agar pad at the center of polydimethylsiloxane slide chamber prepared as described in ([Costa et al., 2013](#)).

Time-lapse movies were performed on an inverted spinning disk confocal microscope (Roper/Nikon Instruments), equipped with a Plan ApoChromat 100 $\times$ /1.4 NA objective (Nikon Instruments), a PIFOC (perfect image focus) objective stepper, and a charge-coupled device camera (EMCCD 512 $\times$ 512 QuantEM; Photometrics). The microscope was equipped with a thermostatic methacrylate chamber to perform imaging at 25 $^{\circ}$ C (or 36 $^{\circ}$ C when analyzing thermosensitive mutants). To analyze Spn1-GFP Rlc1-mCherry Sid4-mCherry wild type as well as Mid2-ENVY Rlc1-mCherry Sid4-mCherry in the respective cell cycle thermosensitive mutants, z-stacks of 7 planes spaced by 1  $\mu$ m between planes were acquired at 1 min interval for 3 h (binning 1, 300 EM gain; 200 ms exposure for each channel).

Fluorescent images for each channel were scaled equivalently to their respective control and analyzed with MetaMorph software 7.7.8.

Analysis of Spn1-GFP and Mid2-ENVY fluorescent signals were quantified in cells at mitotic onset (SPBs separation) by measuring the average fluorescence intensity of the medial region with the linescan tool (MetaMorph software 7.7.8) of 5  $\mu$ m in length and 13  $\mu$ m in width perpendicularly oriented to the long axis of the cell on maximum intensity projection images. Background values, measured as the average intensity of an equivalent region outside the cell, were then subtracted.

To compare the width of Spn1-GFP and Mid2-ENVY at the two different time points indicated in the text in control and mutant cells, the linescan tool (MetaMorph software 7.7.8) was oriented parallel to the long axis of the cell. The values obtained are the average of 3 measurements over the cell width within the bands corresponding to Spn1-GFP or Mid2-ENVY.

### Co-immunoprecipitation experiments

Co-immunoprecipitation experiments were performed as described in ([Guzman-Vendrell et al., 2013](#)) with few modifications. In particular, for the experiments shown in [Figures 3A](#), [3B](#), [S1B](#) and [S2D](#), 200 mL of early log phase cells were incubated to an optical density of 1.0 at 600 nm at 25 $^{\circ}$ C in YE5S medium concentrated twice compared to the regular YE5S medium (YE5S2X), harvested in cold STOP buffer (150 mM NaCl, 50 mM NaF, 10 mM EDTA, 1 mM NaN<sub>3</sub>) and resuspended in 300  $\mu$ L Lysis buffer (50 mM Hepes, 100 mM NaCl, 1 mM EDTA, 1% NP-40, 50 mM NaF, 20 mM  $\beta$ -glycerophosphate) with 1 mM PMSF and a tablette of protease inhibitor cocktail (Thermo, 11697498001). Extracts were incubated with protein A-conjugated magnetic beads (Thermo, 10006D), previously coupled to 6  $\mu$ g of anti-GFP mAb (Roche). For the experiments shown in [Figures S2A](#) and [S2B](#), 200 mL of early log phase cells were incubated to an optical density of 0.7 at 600 nm at 25 $^{\circ}$ C in YE5S medium harvested in cold STOP buffer, and finally resuspended in 300  $\mu$ L Lysis buffer with proteases inhibitors (1 mM PMSF, 0.5  $\mu$ M aprotinin, 0.5  $\mu$ M leupeptin, 0.5  $\mu$ M pepstatin A, 0.1  $\mu$ M NaVO<sub>3</sub>). Extracts were incubated with protein A-magnetic beads (Thermo, 10006D) previously conjugated with 5  $\mu$ L of polyclonal anti-GFP antibody (Thermo A-6455).

Western blots were probed with mouse anti-GFP mAb (1/500 dilution, Sigma Cat No. 11814460001), mouse anti-myc mAb 9E10 (1/666 dilution, Sigma Cat No. 11667149001), mouse anti-GST, (1/1000 dilution, Thermo Cat No. MA4-004) and mouse anti-MBP (1/10000 dilution, NEB, Cat No. E8032S). Secondary goat anti-mouse antibodies were coupled to horseradish peroxidase (1/10000 dilution, Jackson Immunoresearch, Cat No. AB\_2313585 and Bio-Rad, Cat No., 1706516).

For *nda3-KM311* synchronization experiments, 1 L of cell culture was incubated overnight to an optical density of 0.6 at 600 nm. Then, the cells were synchronized by an 8 h incubation at 18 $^{\circ}$ C before being collected and processed.

### In vitro binding assay

DNA fragments encoding Mid2 PH domain (amino acids 569–706) were PCR-amplified and inserted into the GST-producing vector pJC20 or the MBP-producing vector pMAL-p2 (New England Biolabs Inc.) between the sites NdeI - BamHI or BamHI - Sall, respectively. The resulting plasmids were transformed into the E. coli strain BL21. 200 mL of LB cultures supplemented with ampicillin at an O.D. of 2.0 at 600 nm after induction of protein production with 0.1 mM IPTG for 3 h at 28 $^{\circ}$ C were harvested. For GST fusions, cells were resuspended and incubated 30 min at room temperature in 10 mL of CellLytic B Cell Lysis Reagent (Sigma-Aldrich, C8740) containing 50 u/mL Benzonase (Sigma-Aldrich, E1014), 1 mg/mL lysozyme and protease inhibitors (1 mM PMSF, 0.5  $\mu$ M aprotinin, 0.5  $\mu$ M leupeptin, 0.5  $\mu$ M pepstatin A). After a centrifugation at 10.000  $\times$  g for 30 min at 4 $^{\circ}$ C, the lysate was incubated with 100  $\mu$ L glutathione-conjugated magnetic beads (Cube Biotech) in rotation overnight at 4 $^{\circ}$ C. The beads were then PBS-washed three times and resuspended in 200  $\mu$ L of binding buffer (20 mM Tris-HCl pH 7.5, 0.2 M NaCl, 1 mM EDTA, 0.5% NP-40). For MBP fusions, cells were resuspended and incubated 10 min in rotation at room temperature in mM Tris-HCl pH 8.0, 20% sucrose, 1 mM EDTA (80 mL of buffer per gram of cells). After centrifugation, the pellet was incubated for 10 min in ice in 80 mL of 5 mM MgSO<sub>4</sub> buffer per gram of cells, frequently shaking gently. After a centrifugation at 8.000  $\times$  g for 20 min at 4 $^{\circ}$ C, 800  $\mu$ L of 50% of amylose resin (NEB, E8021) per 40 mL of lysate and Tris-HCl pH 7.5 to a final concentration of 20 mM were added and incubated in rotation at 4 $^{\circ}$ C overnight. The resin was then washed three times in washing buffer (20 mM Tris-HCl pH 7.5, 200 mM NaCl, 1 mM EDTA) and the MBP fusions were eluted

by adding a total of 6 mL 10 mM maltose. Samples were concentrated using a YM-30 centricon (Millipore) after washing five times with binding buffer. 30  $\mu$ L of GST or GST-Mid2-PH were mixed with 30  $\mu$ L MBP-Mid2-PH in a total volume of 100  $\mu$ L and incubated at 4°C for 2 h. Magnetic beads were washed three times with binding buffer, resuspended in 35  $\mu$ L of laemli buffer and analyzed by SDS-PAGE and western blotting.

### Septin-GFP fusions for orientation measurements

The Spn1-GFP fusion used throughout this study includes a flexible linker (RIPGLI) between the Spn1 and the GFP sequences rendering this fusion not appropriate for orientation measurements (Figure S3A). To generate fusions of septins with rotationally constrained GFP (conGFP) we generated and screened C-terminal GFP fusions of Spn1 and Spn4 with no linker sequence between the septin and the GFP, and with the C-termini of septins and the N-terminus of GFP truncated to different extents, indicated as  $\delta$ C and  $\Delta$ N, respectively. The coiled-coil prediction algorithm COILS was used to identify the ends of the predicted coiled-coils in Spn1 and Spn4 to guide the C-terminal truncations. We started with Spn1 fusions to monomeric EGFP (mEGFP) deleting C-terminal Spn1 and N-terminal GFP amino acids, generating Spn1 $\Delta$ C5-mEGFP $\Delta$ N5 (Spn1 1–464; mEGFP starting at EELF), Spn1 $\Delta$ C5-mEGFP $\Delta$ N6, Spn1 $\Delta$ C5-mEGFP $\Delta$ N7 and Spn1 $\Delta$ C5-mEGFP $\Delta$ N11; the first three fusions behaved normally localization- and septation-wise, but the GFP was still flexible, whereas the last one was not fluorescent. Further trimming of Spn1 C-terminus to generate Spn1 $\Delta$ C12-mEGFP $\Delta$ N5, Spn1 $\Delta$ C12-mEGFP $\Delta$ N6 and Spn1 $\Delta$ C12-mEGFP $\Delta$ N7 led to ectopic cytoplasmic bar-like structures (like the ones shown in Figure S3D). We thus turned to monomeric superfolder GFP (msfGFP) which tolerates more extensive N-terminal truncations (Pédrelacq et al., 2006), and generated a fusion of full-length Spn1 to msfGFP without using a linker and additionally removing residues before the first stave of the GFP  $\beta$ -barrel to obtain Spn1-msfGFP- $\Delta$ N12 (Spn1 1–469; msfGFP starting at VPILV). Spn1-msfGFP $\Delta$ N12, designated as Spn1-conGFP in this study, was rotationally constrained and was used for orientation measurements. We also generated Spn4 fusions to mEGFP deleting C-terminal Spn4 and N-terminal GFP amino acids to obtain Spn4 $\Delta$ C6-mEGFP- $\Delta$ N5 (Spn4 1–374; EGFP starting at EELF), Spn4 $\Delta$ C6-mEGFP $\Delta$ N6, Spn4 $\Delta$ C6-mEGFP $\Delta$ N7 and Spn4 $\Delta$ C6-mEGFP $\Delta$ N11. The first two constructs behaved normally, the third one formed ectopic cytoplasmic bar-like structures in addition to its localization in septin rings (Figure S3D), whereas the last one was not fluorescent. The GFP in Spn4 $\Delta$ C6-mEGFP $\Delta$ N7 was rotationally constrained and this fusion, designated as Spn4-conGFP in this study, was used to infer the orientation of septins with respect to the GFP dipole.

### Spinning-disk polarization-resolved fluorescence microscopy

Yeast cells were prepared for live polarimetry measurements as described above for live microscopy. Polarization-resolved fluorescence images were acquired on a custom-built optical setup employing a confocal spinning disk unit (CSU-X1-M1 from Yokogawa) connected to the side-port of a Perfect Focus System-equipped inverted microscope (Eclipse Ti2-E, Nikon Instruments), using a Nikon Plan Apo  $\times$ 100/1.45 NA oil immersion objective lens, 488- and 561-nm Sapphire laser lines (Coherent) and an iXon Ultra 888 EMCCD camera (1024  $\times$  1024 pixels, 13  $\times$  13  $\mu$ m pixel size, Andor, Oxford Instruments), resulting in an image size of 1 pixel = 65 nm. Polarimetry images were obtained by varying the incident linear polarization with a Pockels Cell electro-optic modulator (No 28-NP Quantum Technology) synchronized to the EMCCD camera, followed by a quarter waveplate (WQP05M-488, Thorlabs) (Wang et al., 2013). The quality of the polarization state at the sample plane was ensured by using a compensating dichroic beamsplitter similar to the one present in the spinning disk unit (405/488/561/635 nm Quad Dichroic mirror) but placed at 90° reflection to exchange s and p polarization state phase-shifts. A polarization stack is made of 18 polarized images using an incident linear polarization varying from 0° (corresponding to the horizontal direction in the image) to 170° with steps of 10°. The recorded stack is used to obtain the polarization response of the sample at each pixel, from which GFP orientation information can be deduced. Three to five large field-of-view images (66  $\times$  66  $\mu$ m) typically containing 10–20 cells during cytokinesis per image, were collected for wild-type and *nda3-KM311* cells; seven to eight such images were collected for *mid2* $\Delta$  cells to account for the smaller number of *mid2* $\Delta$  dividing cells per field of view. An exposure time of 0.5 s was used per image. Before each polarimetry measurement, a two-color z stack was acquired ( $\Delta$ z = 1.0  $\mu$ m) to image both septin-GFP fusions and myosin/spindle bodies; these images were used for staging the measured cells and correlating the measured orientation distributions with the different stages of cytokinesis. A polarization stack was then recorded for each position within a z stack ( $\Delta$ z = 1.0  $\mu$ m) for the septin-GFP channel, and thus allowed to obtain polarimetry images throughout the septin rings, containing both tangential-most views where the entirety of the ring can be seen parallel to the xy plane, and more equatorial views showing cross-sections that appear as spots on either side of the ring. To minimize bias in the measured orientations due to the contribution of off-plane orientations we focused on the tangential-most views, as shown in Figures 5 and S3.

Polarization stack images were first processed with the open-source image processing software ImageJ/Fiji. Images within each polarization stack were systematically registered using the StackReg plugin to correct for drift during the acquisition. The proper z planes, i.e. the ones containing the tangential-most views of the ring, were then identified for each cell, and polarimetry data were analyzed according to the framework defined by (Kress et al., 2013) to obtain the mean GFP dipole orientation (angle  $\rho$ ) per pixel. Analysis and data representation, including color-coded stick representations of the measured orientations per pixel and polar-plot histograms, were done by using the Polarimetry Analysis software which is a Matlab App Designer standalone application. The source code is available at [github.com/cchandre/Polarimetry](https://github.com/cchandre/Polarimetry), and the desktop app can be freely obtained at <https://www.fresnel.fr/polarimetry> under a BSD license.



The regions of interest for analysis were selected by a combination of intensity thresholding and manual selection of the features to analyze. Each region of interest contained typically 120–200 analyzed pixels i.e. 120–200 orientation color-coded sticks in the tangential-most view of the septin ring per cell (Figures 5 and S3). The mean orientation per pixel is represented as a stick whose orientation and color are the measured angle  $\rho$  within the resolution limit of the microscope ( $\sim 200$  nm). The distribution of mean orientations was represented in 0–180° polar-plot histograms. To assess the extent to which the measured orientations were more parallel or more perpendicular with respect to the CR axis, the ring axis in each cell was used to normalize the angle distributions from 0–180° to 0–90° and generate 0–90° polar plots (Figures 5 and S3), with 0° and 90° defining orientations parallel and perpendicular to the ring axis, respectively. To facilitate the visual comparison of orientation distributions between cells and also relate directly to the 0–90° polar plots, representative cells in Figures 5 and S3 have been rotated using the Polarimetry Analysis software so that their CR axes correspond to the horizontal orientations (0°). 0–90° normalized polar-plot histograms from all cells in a given condition were concatenated to represent all analyzed data in each condition, then they were rescaled to their respective maximum value to facilitate comparisons, and orientations were sorted in bins of 6°. The normalization of orientation distributions with respect to the CR ring axis, the normalization of 0–180° to 0–90° polar plots and the rescaling of histograms were done with custom-generated Matlab codes, the source codes of which are available at [github.com/cchandre/Polarimetry](https://github.com/cchandre/Polarimetry).

### QUANTIFICATION AND STATISTICAL ANALYSIS

The number of samples analyzed ( $n$ ) is defined in each figure and were derived from 3 independent experiments, except for the co-immunoprecipitations assays in Figures S1B and S2D, which were repeated twice. The error bars correspond to standard deviation (sd) between experiments and are specifically indicated in each figure.

**Cell Reports, Volume 39**

**Supplemental information**

**Septin filament compaction into rings  
requires the anillin Mid2 and  
contractile ring constriction**

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## SUPPLEMENTAL FIGURE LEGENDS

**Figure S1: Septin compaction is defective in *mid2Δ* cells, and Mid2 and septins interact in *nda3-KM311*-blocked cells and Mid2 self-interaction is septin independent, Related to Figure 2**

**A:** Kymographs of the lateral region of the medial cortex (indicated in the scheme at the bottom) of wild type (left) or *mid2Δ* cells (right) producing Spn1-GFP, Rlc1-mCherry and Sid4-mCherry. Elapsed time is shown in minutes. The major transitions in the behavior of Spn1 from mitosis onset and the defects observed in the absence of Mid2 are highlighted in magnified panels at the far right and indicated with a color-coded asterisk that matches that of the kymograph in figure 1C. **B:** Co-immunoprecipitation assay between Mid2-13XMyc and Spn1-GFP in control and *nda3-KM311* cells incubated at 18°C for 7 hours (lanes 1, 2, 3 and 4) and released for 1 hour at 25°C (lanes 5, 6, 7 and 8). Magenta arrowheads show the expected Spn1 or Mid2 bands; black arrowheads show degradation products; green arrowheads depict the immunoglobulin bands.

**Figure S2: The PH domain of Mid2 self-interacts in two hybrid assays and Mid2 co-immunoprecipitates with itself in *nda3-KM311*-blocked cells, Related to Figure 3**

**A:** Co-immunoprecipitation assay between full length Mid2-13Myc and Mid2-mEGFP or Mid2 ΔPH -13Myc and Mid2ΔPH-GFP either in the presence (wt) or in the absence of septins (*spn1-5Δ*). Magenta arrowheads show the expected Mid2 bands; black arrowheads show degradation products. **B:** Co-immunoprecipitation assay between the PH domain of Mid2 tagged with 13XMyc and the PH domain of Mid2 tagged with GFP either in the presence (wt) or in the absence of septins (*spn1-5Δ*). **C:** Two-hybrid assay showing the interaction of the PH domain of Mid2 with itself. The serially diluted growth of the indicated transformants is shown under non-selective (left panel) or selective conditions (right panel). **D:** Co-immunoprecipitation assay between Mid2-13XMyc and full length Mid2-GFP, Mid2ΔPH-GFP or the PH domain of Mid2 tagged with GFP in *nda3-KM311* cells incubated at 18°C for 8 hours. Magenta arrowheads

show the expected Mid2 bands; black arrowheads show degradation products; green arrowheads depict the immunoglobulin bands.

**Figure S3: Under high CDK activity, septins and Mid2 appear as a broad band at both sides of the CR, Related to Figure 4**

Fluorescence intensity analysis of Spn1-GFP, Rlc1-mcherry and Sid4-mcherry signal measured along the entire cell in epifluorescence images of the *nda3-KM311* mutant incubated at 18°C for 8 hours in a representative cell where the SPBs are aligned (**A**) or not (**B**) with respect to the ring position. **C**: Quantification of Spn1-GFP and Mid2-ENVY width over time in wild type (light green for Spn1-GFP and dark green for Mid2-ENVY) and *nda3-KM311* mutant cells (magenta for Spn1-GFP and purple for Mid2-ENVY) incubated at 18°C during 8 hours. The last data point has been taken after switching the cells to 25°C for 1 hour. The average curves  $\pm$  sd are displayed.  $n > 500$  cells. Time lapse imaging of wild type (**D**) or *cut7-24* cells (**F**) expressing Mid2-ENVY, Rlc1-mcherry and Sid4-mcherry incubated at 36°C for 3 hours. Maximum intensity projections of confocal images are shown. Time 0 corresponds to mitotic entry. Scale bars: 2  $\mu$ m. Plot displaying Mid2-ENVY dynamics throughout cell division in control (**E**) and *cut7-24* cells (**G**). Mean  $\pm$  sd are shown.  $n = 45$  cells in each strain. **H**: Quantification of the extent of Mid2-ENVY width compaction over time in wild type (dark green) and *cut7-24* cells (purple).  $n = 20$  cells in each case.

**Figure S4: Localization and orientation measurements of Spn1-conGFP and Spn4-conGFP in wild type cells, Related to Figure 5**

**A**: Polar-plot histograms of GFP dipole orientations of Spn1-GFP filaments from wild type cells at different stages of cytokinesis (top panels). The number of analyzed cells per stage is indicated below each polar-plot histogram. Confocal grayscale images of Spn1-GFP from selected wild type cells and an overlaid color-coded stick representation of the measured orientations (angles  $\rho$ ) per analyzed pixel at each cytokinetic stage (bottom panels). The mean orientation of GFP dipoles per pixel is represented as a stick whose orientation and color are the measured angle  $\rho$ . Scale



bar: 1  $\mu\text{m}$ . **B:** Maximum intensity projection images of Spn1-GFP, Rlc1-mCherry and Sid4-mCherry in wild type cells (left) and Spn1-conGFP, Rlc1-mCherry and Sid4-mCherry (right) in wild type, *mid2* $\Delta$  and *nda3-KM311* cells at each stage of cytokinesis. Representative large field-of-view maximum intensity projection images of wild type *rlc1-mCherry sid4-mCherry* cells producing Spn1-conGFP (**C**) or Spn4-conGFP (**D**) used for binning cells in each cytokinetic stage before polarimetry measurements. White asterisks in the insets in (D) point to septin rings in selected cells (outlined rectangles), while white arrows point to ectopic cytoplasmic bar-like structures upon Spn4-conGFP production. Scale bars: 10  $\mu\text{m}$ . An example of a polar plot of dipole orientations in a Spn4-conGFP bar-like structure is shown on the right. A confocal grayscale image of the bar-like structure and an overlaid color-coded stick representation of the measured orientations in this structure are shown below the polar-plot histogram. Scale bar: 1  $\mu\text{m}$ . **E:** Examples of cells with Spn4-conGFP in septin rings during CR constriction (left) and during early stages of post constriction (right). Confocal grayscale images of selected cells are shown together with the respective overlaid color-coded stick representation of the measured orientations and the corresponding polar-plot histograms.

**Figure S5: Mid2 recruitment to the division region is compromised under low SIN activity or in the  $\beta(1,3)$ -glucan synthase Bgs1 mutant *cps1-191*, Related to Figure 6**

**A:** Time lapse imaging of wild type (left series) and *sid2-250* cells (right series) producing Mid2-ENVY, Rlc1-mCherry and Sid4-mCherry incubated at 36°C for 3 hours. Maximum intensity projections of confocal images are shown. Time 0 corresponds to mitotic entry. Scale bars: 2  $\mu\text{m}$ . **B:** Analysis of Mid2-ENVY fluorescence intensity from mitotic onset throughout cell division in control (dark green) and *sid2-250* cells (purple). The average line graphics  $\pm$  sd are displayed. n=6 cells in each case. **C:** Time lapse imaging of wild type (left series) and *cps1-191* cells (right series) producing Mid2-ENVY, Rlc1-mCherry and Sid4-mCherry incubated at 36°C for 3 hours. Maximum intensity projections of confocal images are shown. Time 0 corresponds to mitotic entry. Scale bars: 2  $\mu\text{m}$ . **D:** Analysis of Mid2-ENVY fluorescence intensity from mitotic

onset throughout cell division in control (dark green) and *cps1-191* cells (purple). The average line graphics  $\pm$  sd are displayed. n=6 cells in each case.

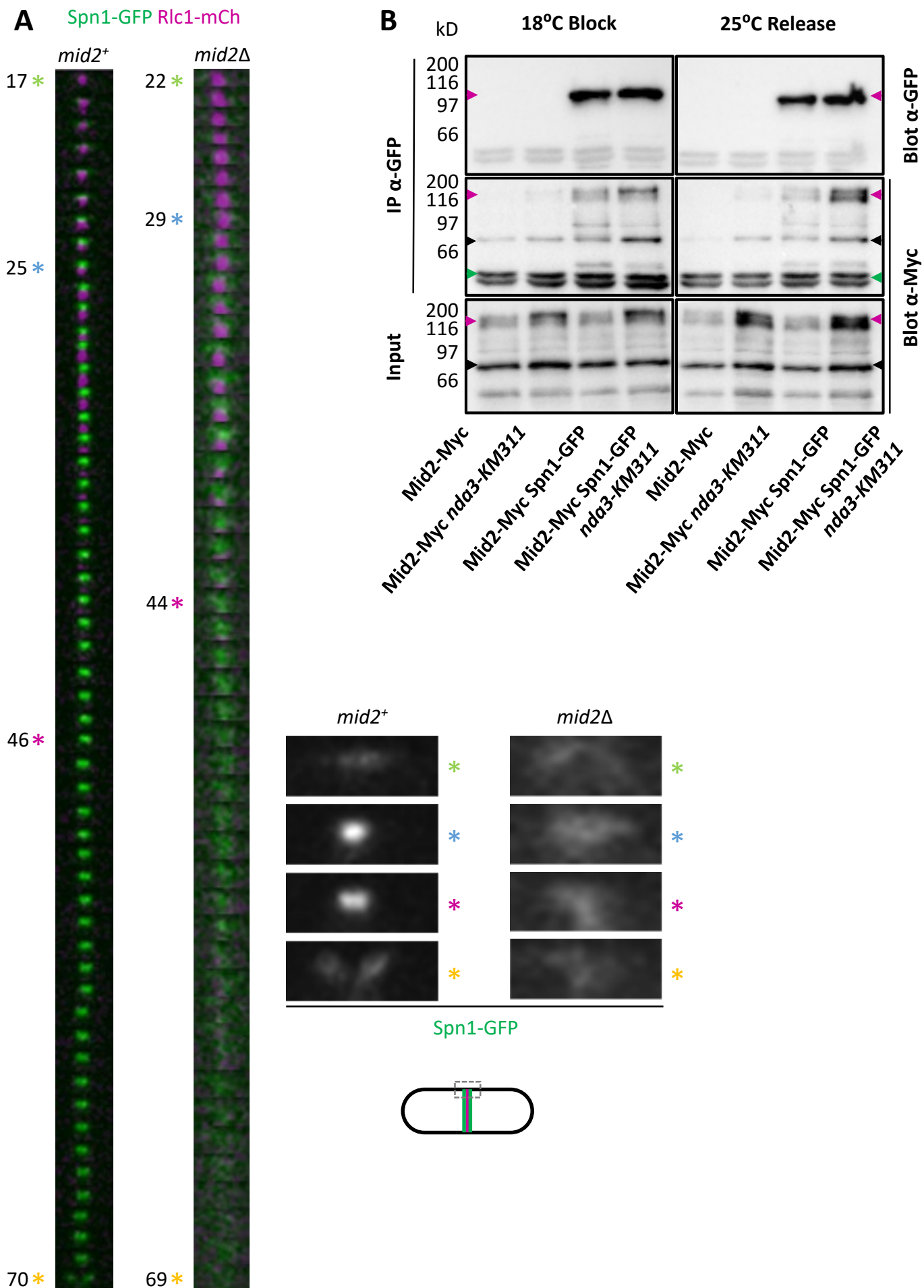


Figure S1, Related to Figure 2

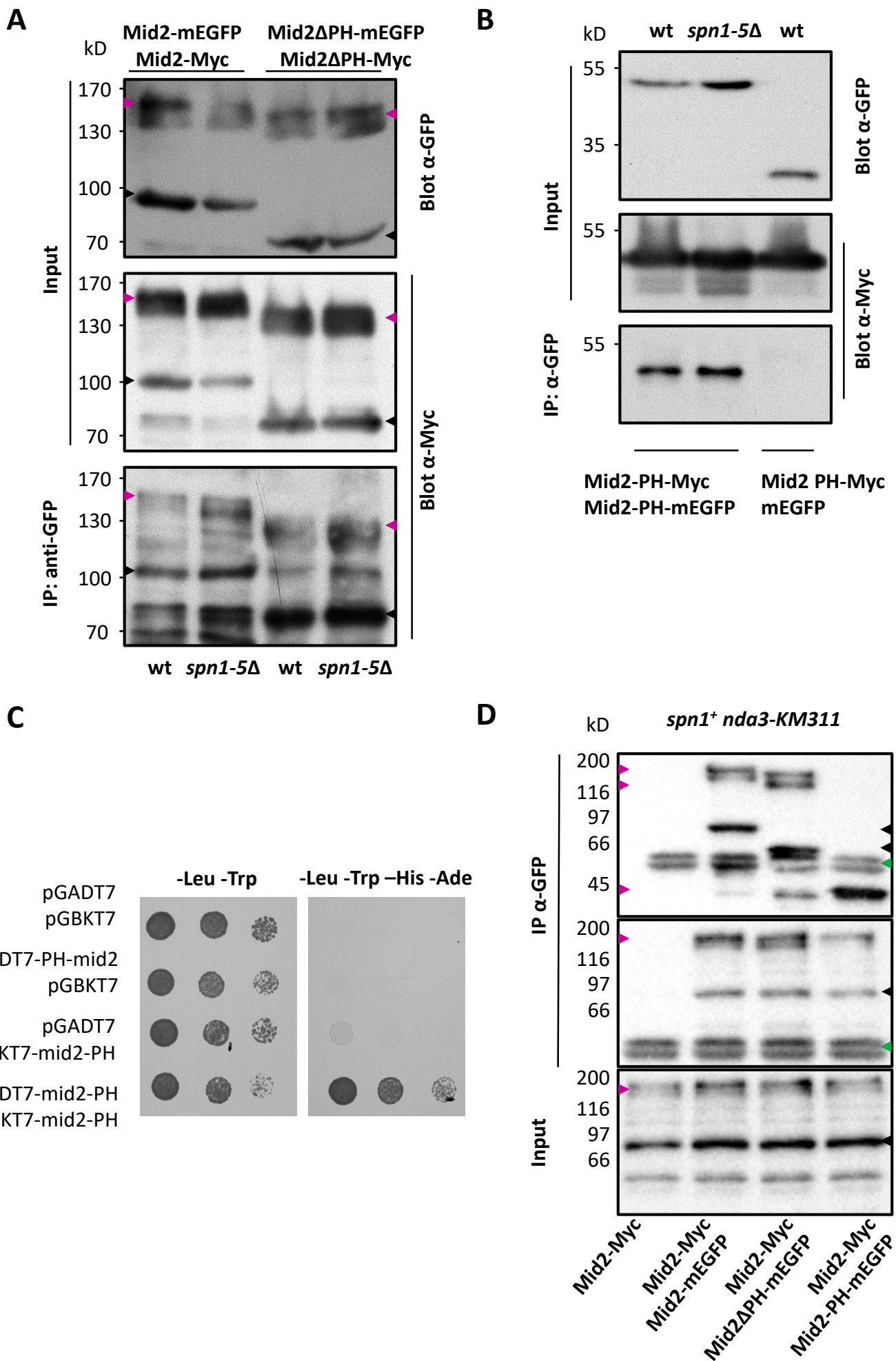
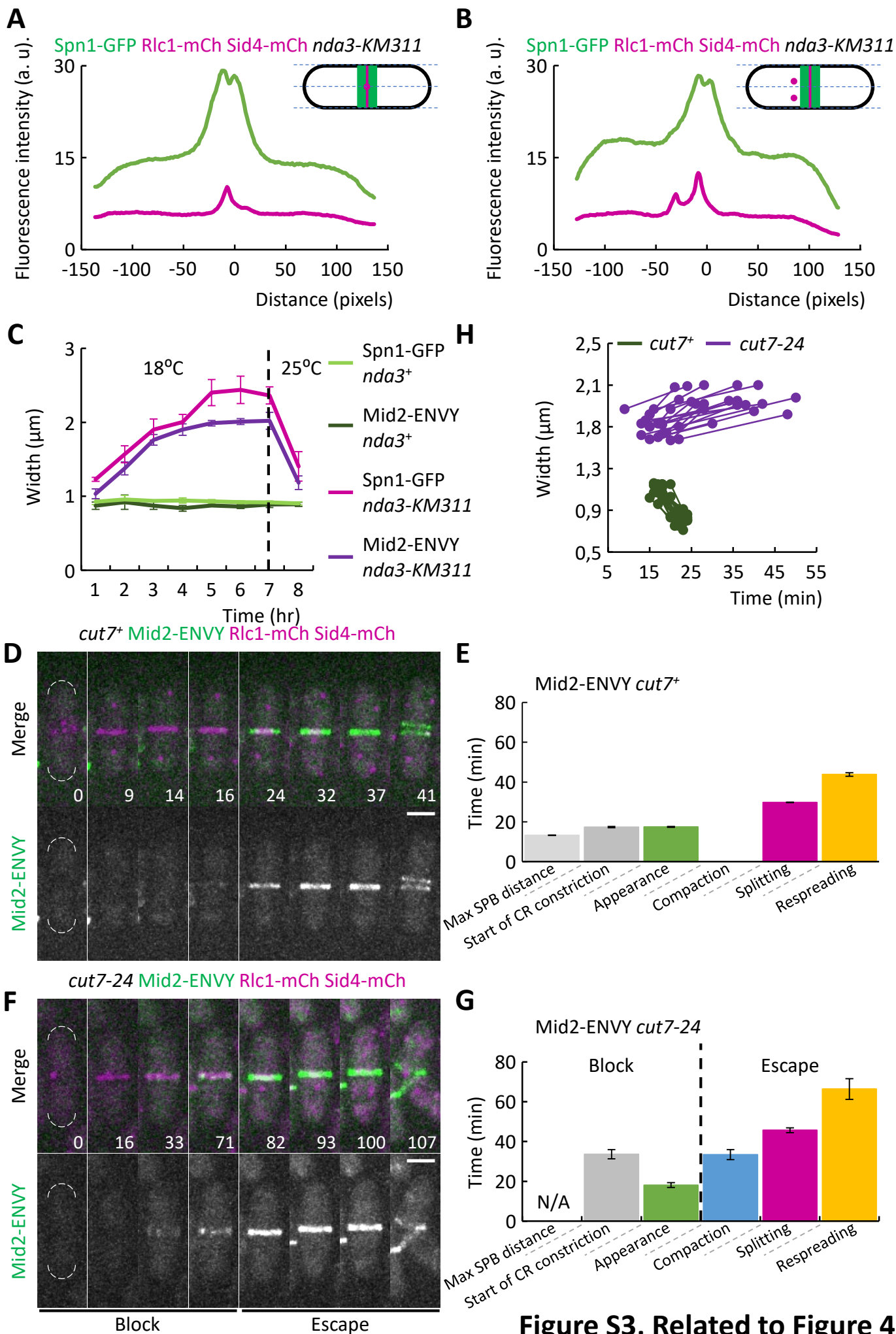
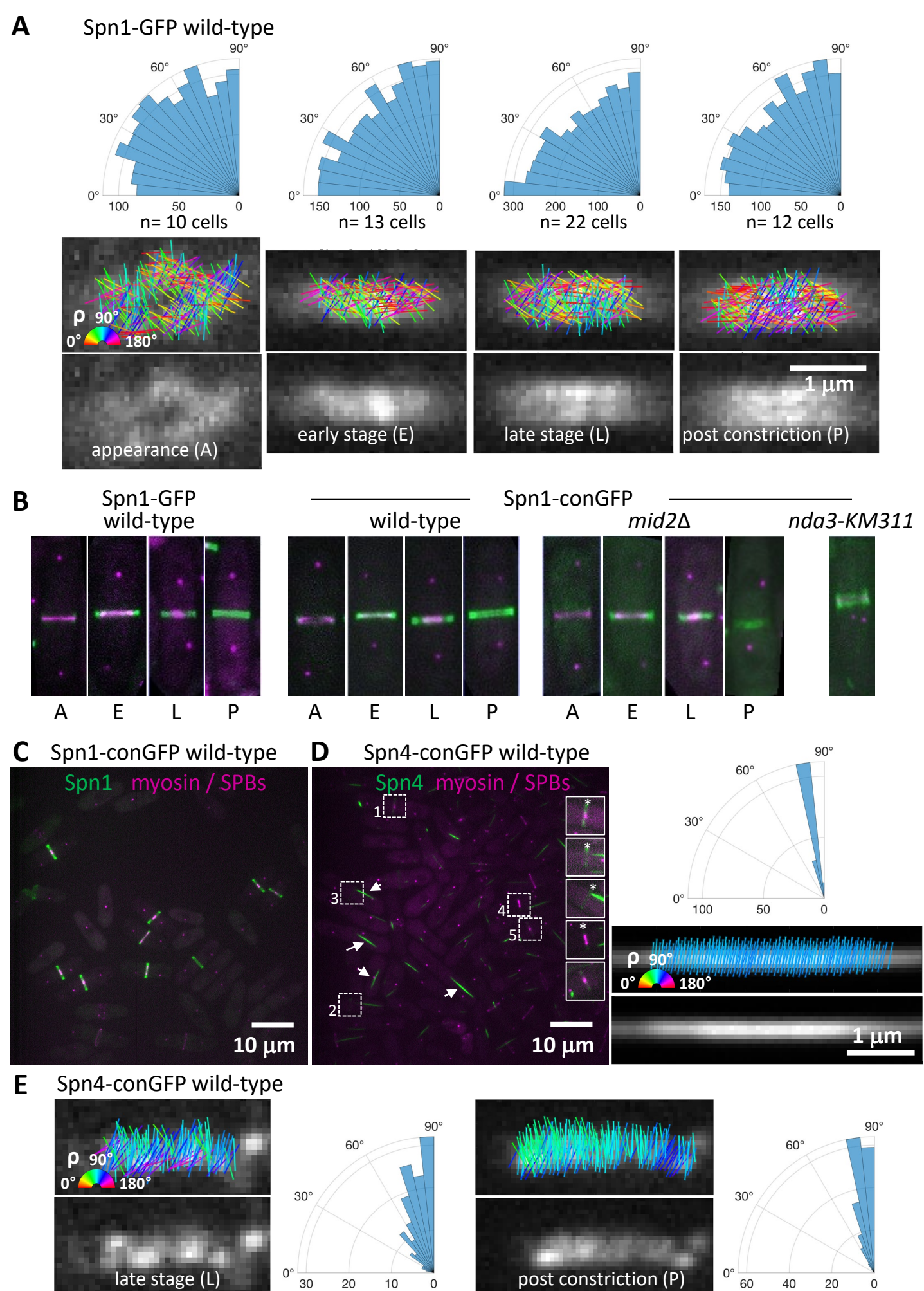


Figure S2, Related to Figure 3



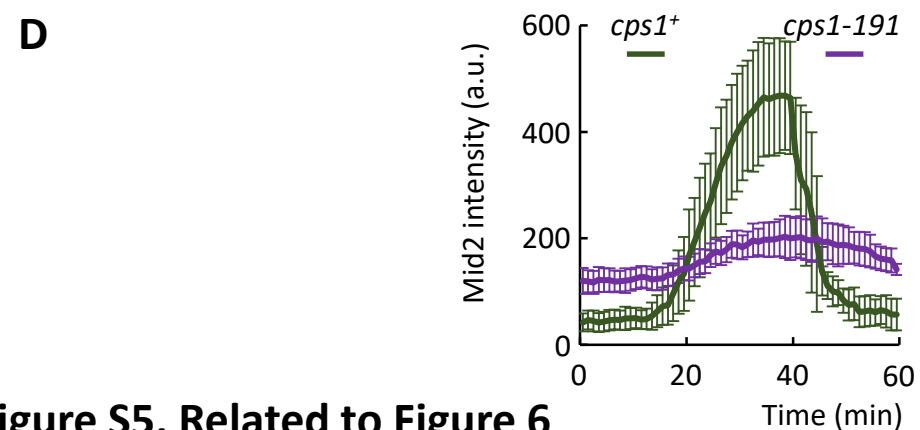
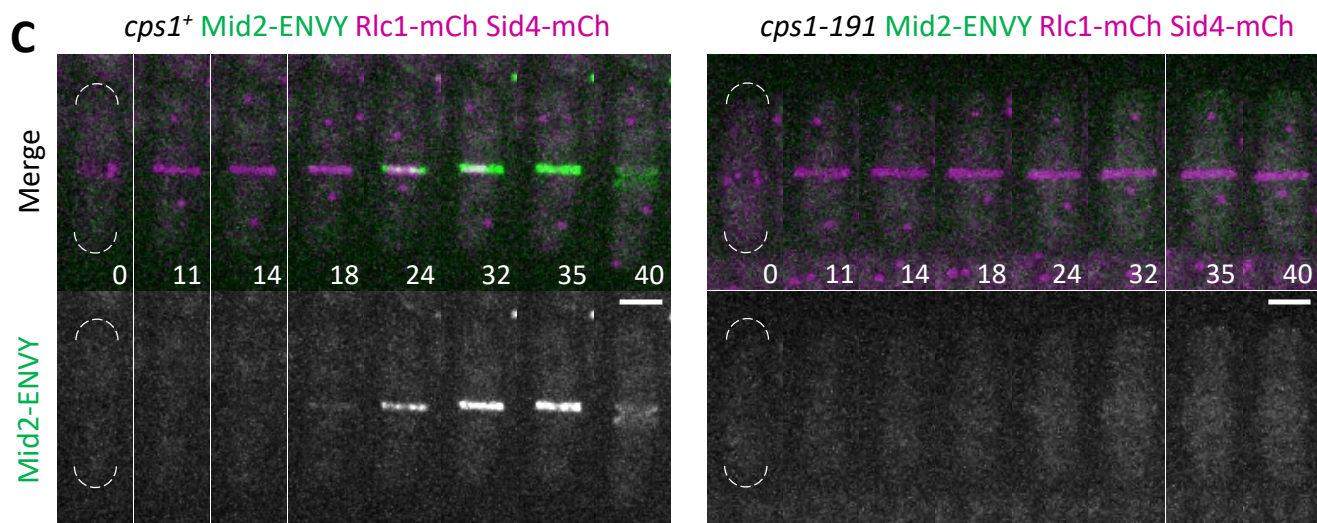
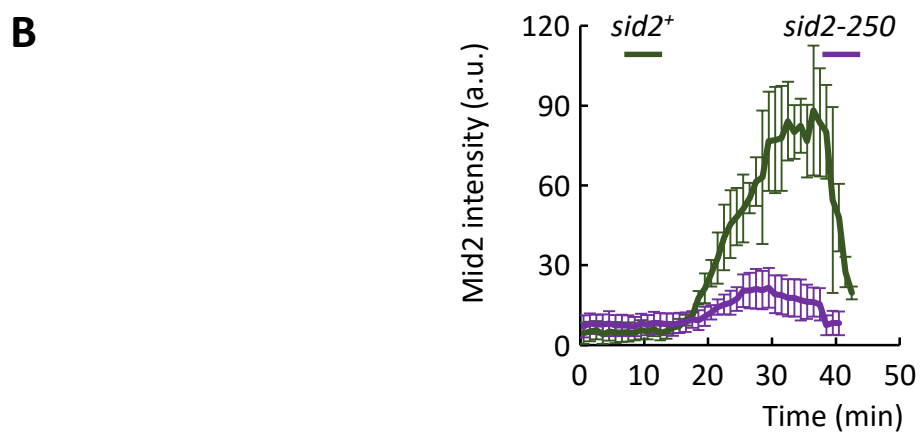
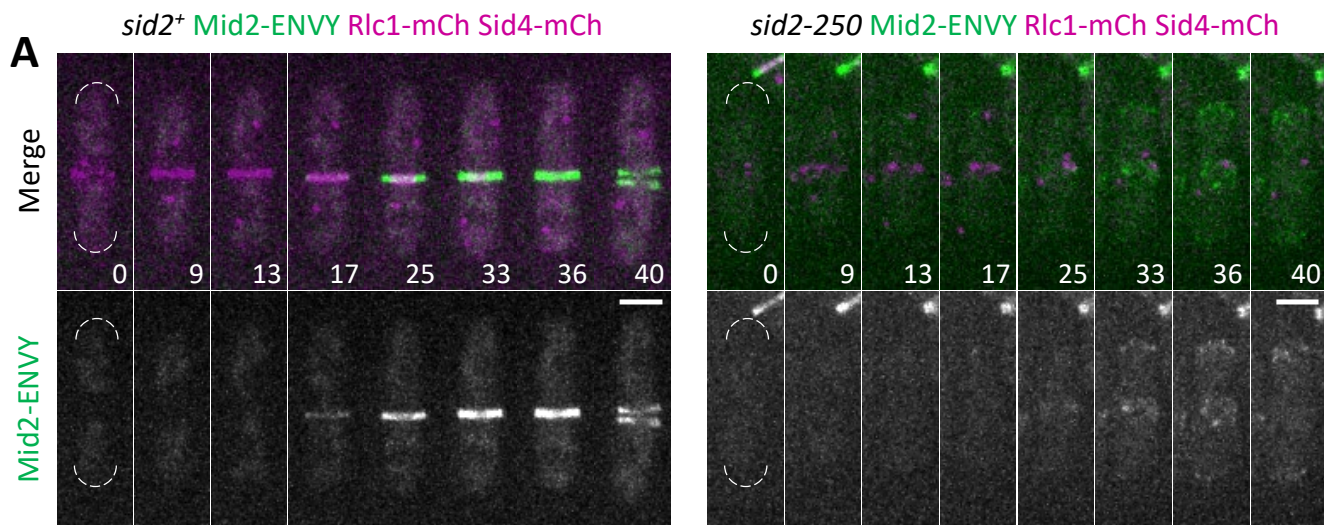


**Figure S3, Related to Figure 4**



**Figure S4, Related to Figure 5**





**Figure S5, Related to Figure 6**