The intrinsically disordered region of the cytokinetic F-BAR protein Cdc15 performs a unique essential function in maintenance of cytokinetic ring integrity

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ABSTRACT Successful separation of two daughter cells (i.e., cytokinesis) is essential for life. Many eukaryotic cells divide using a contractile apparatus called the cytokinetic ring (CR) that associates dynamically with the plasma membrane (PM) and generates force that contributes to PM ingression between daughter cells. In Schizosaccharomyces pombe, important membrane–CR scaffolds include the paralogous F-BAR proteins Cdc15 and Imp2. Their conserved protein structure consists of the archetypal F-BAR domain linked to an intrinsically disordered region (IDR). Functions have been assigned to the F-BAR and SH3 domains. In this study we probed the function of the central IDR. We found that the IDR of Cdc15 is essential for viability and cannot be replaced by that of Imp2, whereas the F-BAR domain of Cdc15 can be swapped with several different F-BAR domains, including that of Imp2. Deleting part of the IDR results in CR defects and abolishes calcineurin phosphatase localization to the CR. Together these results indicate that Cdc15’s IDR has a nonredundant essential function that coordinates regulation of CR architecture.

INTRODUCTION Cytokinesis is the final step in the cell division cycle when daughter cells separate. In Amoebozoa and Opisthokonta, cytokinesis employs an actin- and myosin-based proteinaceous apparatus termed the cytokinetic ring (CR; Mangione and Gould, 2019). The CR assembles at the future division plane and eventually constricts coincident with cleavage furrow ingression. Of the many proteins that compose the CR, membrane-bound scaffolds are crucial for transmitting CR-generated tension to the plasma membrane (PM) and for maintaining CR placement at the division site (Glotzer, 2017). F-BAR proteins are conserved membrane–CR scaffolds that function during cytokinesis as well as other processes involving coordination between the dynamic cytoskeleton and membranes (Lippincott and Li, 2000; Ahmed et al., 2010; Fricke et al., 2010; Roberts-Galbraith and Gould, 2010). The namesake F-BAR domain binds phospholipids and oligomerizes into a multivalent platform that can recruit proteins to membranes through interactions with the F-BAR domain itself or using additional functional domains (McDonald and Gould, 2016).

In the fission yeast Schizosaccharomyces pombe, the F-BAR protein Cdc15 is an essential protein that promotes CR stabilization and cell wall deposition (i.e., septation) that occurs coincident with CR constriction (Fankhauser et al., 1995; Wachtler et al., 2006; Huang et al., 2007; Pinar et al., 2008; Roberts-Galbraith et al., 2009; Arasada and Pollard, 2014; Cortes et al., 2015; Ren et al., 2015; Willet et al., 2015b; Sethi et al., 2016; Onwubiko et al., 2019). Cdc15 has an N-terminal F-BAR domain linked by a central intrinsically disordered region (IDR) to a C-terminal SH3 domain. The F-BAR domain directly binds the PM (McDonald et al., 2015) and the essential CR actin nucleator/elongator formin Cdc12 (Carnahan and Gould, 2003; Willet et al., 2015a). Additionally, the Cdc15 F-BAR domain oligomerizes into a multivalent scaffold with high avidity for the PM at the division site (McDonald et al., 2015). Disrupting the Cdc15-Cdc12 interaction...
RESULTS

The Cdc15 IDR is essential for viability

We tested the ability of various cdc15 mutants to support viability by integrating them at one cdc15 locus of diploid cells to make cdc15+/cdc15 mutant heterozygotes. Next, the diploids were sporulated and tetrads dissected to determine whether the cdc15 mutants could support viability when present as the sole cdc15 allele.

We found that the Cdc15 F-BAR domain alone was not sufficient to support cell viability (Figure 1, A and B). An allele encoding the Cdc15 F-BAR domain juxtaposed to the SH3 domain (cdc15-1/IDR) was also unable to support viability, verifying the necessity of Cdc15’s IDR (Figure 1, A and C). Deletion of the SH3 domain was previously determined to be viable (Roberts-Galbraith et al., 2009) and further C-terminal truncation was tolerated to aa 710 but not to aa 700 (Figure 1A). These results, and those from previous studies (McDonald et al., 2015), indicate that the F-BAR domain and part of the IDR are required to perform Cdc15’s essential function.

F-BAR domains can be exchanged

We next asked whether these features could be replaced by the comparable regions of other F-BAR proteins. We first generated a series of chimeras by fusing various F-BAR domains from human and yeast proteins to the Cdc15 C-terminus and then tested them for viability as previously described. Five of the 15 chimeras were viable: the fusions with F-BAR domains of the human proteins Fer, Gas7, PACSIN2, and PSTPIP1 and S. pombe Imp2 (Figure 2A; Supplemental Table S1). We tagged the viable chimeras with the fluorophore mNeonGreen (mNG) and they all localized to the CR (Figure 2B). The chimeras have septation defects (Figure 2B and Supplemental Figure S1, A and B) and are cold-sensitive (Supplemental Figure S1C). These defects are comparable among chimeras, suggesting a similar level of rescue.

In contrast, replacement of only the Cdc15 IDR with that of Imp2 (cdc15-imp2 chimera) did not support viability (Figure 2A). Additionally, a mutant in which the Cdc15 F-BAR was fused to the Imp2 IDR duplicated (to match the length of Cdc15’s IDR) plus Imp2’s SH3 domain (cdc15-imp2 extended chimera) did not support viability (Figure 2A). These results indicate that the Cdc15 IDR is essential in a sequence-specific manner.

Partial loss of the IDR results in cytokinesis defects

To separate the function of the IDR from that of the SH3 domain, which coordinates CR constriction and septum deposition through multiple binding interactions (Tajadura et al., 2004; Morrell-Falvey et al., 2005; Pinar et al., 2008; Roberts-Galbraith et al., 2009; Cortes et al., 2015; Ren et al., 2015; Sethi et al., 2016), we divided the IDR into three segments of 175 aa (Figure 3A) and deleted each
segment individually. Each of the three deletion mutants was viable (Figure 3B). However, the three mutants had different phenotypes: cdc15-∆1 cells were morphologically indistinguishable from WT cells, while cdc15-∆2 and cdc15-∆3 cells had defects in morphology and cell division, as evidenced by an increased number of nuclei and septa per cell (Figure 3, B and C). We considered the possibility that the cytokinetic defects observed in cdc15-∆2 and cdc15-∆3 resulted from simultaneously removing key aa residues and shortening the separation between the F-BAR and SH3 domains. Therefore, we tested whether replacing the IDR with duplicated or triplicated segment 2 (aa 503–677) would rescue the morphological defects of cdc15-∆3 (Supplemental Figure S1D). However, both of these strains resembled the phenotype of cdc15-∆3 cells: a high percentage of cells were septated and length at septation was increased, indicating growth initiation before separation was accomplished (Supplemental Figure S1E). This points to a specific function of aa 503–854.

As described above, the shortest viable C-terminal truncation of Cdc15 was at aa 710 (Figure 1A). Therefore, although segment 3 (aa 678–854) could be deleted when the SH3 domain was present (Figure 3B), cells required part of segment 3 in the absence of the SH3 domain. This result suggested that there are likely multiple functional motifs along the length of Cdc15 and that loss of some combinations, but not others, might be tolerated. To expand on this idea, we combined deletions of IDR segments and/or the SH3 domain. While cdc15-∆1∆3 and cdc15-∆1∆SH3 are viable, cdc15-∆1∆2, cdc15-∆2∆3, and cdc15-∆2∆SH3 are not (Figure 3D). These results indicate that region 2 mediates a particularly important function such that its loss is barely tolerated.

Segments 2 and 3 of Cdc15’s IDR contain multiple short stretches that have similar sequence identity among Schizosaccharomyces species (Supplemental Figure S2B). Many of these conserved motifs also have high prediction for protein binding (ANCHOR; Dosztanyi et al., 2009; Meszaros et al., 2009; Supplemental Figure S2A).
Therefore, deletions of these short conserved motifs were tested to better define important functional motifs within the IDR. However, none of the smaller deletions resulted in the cytokinetic phenotypes observed for cdc15Δ-2 or cdc15Δ-3 (Supplemental Figure S2C).

Cdc15 IDR mutants localize to the division site and are phosphoregulated

We confirmed by SDS–PAGE and immunoblotting that Cdc15-Δ1, Cdc15-Δ2, and Cdc15-Δ3 were produced at levels comparable to WT Cdc15 (Supplemental Figure S3A). Additionally, the three single segment–deletion mutants localized to nodes and CRs (Supplemental Figure S3B).

Next, we examined whether the Cdc15 segment deletions were still subject to cell cycle–regulated phosphoregulation, as the IDR is the primary site of Cdc15 phosphorylation (Wilson-Grady et al., 2008; Roberts-Galbraith et al., 2010; Koch et al., 2011; Chen et al., 2013; Carpy et al., 2014; Kettenbach et al., 2015; Swaffer et al., 2016, 2018; Lee et al., 2018). During interphase, Cdc15 is highly phosphorylated, which is evidenced by slower mobility on SDS–PAGE (Fankhauser et al., 1995; Roberts-Galbraith et al., 2010). During mitosis and cytokinesis, Cdc15 is dephosphorylated (Fankhauser et al., 1995; Roberts-Galbraith et al., 2010). Mutants that abolish phosphorylation exhibit defects in cytokinesis and localize abnormally to the cell cortex during interphase (Roberts-Galbraith et al., 2010). Lambda phosphatase collapse revealed that Cdc15-Δ1, Cdc15-Δ2, and Cdc15-Δ3 were still phosphorylated (Supplemental Figure S3C). Additionally, we determined that Cdc15-Δ1, Cdc15-Δ2, and Cdc15-Δ3 maintained cell cycle changes in phosphorylation (Supplemental Figure S3D). Furthermore, mEGFP-Cdc15-Δ2 and mEGFP-Cdc15-Δ3 showed diffuse localization during interphase (Supplemental Figure S3E), suggesting that the division defects of cdc15Δ-2 and cdc15Δ-3 are not mimicking Cdc15 phospho-ablating mutants. In contrast, mEGFP-Cdc15-Δ1 accumulated in cortical puncta during interphase despite the fact that cdc15Δ-1 have normal morphology (Supplemental Figure S3E). These findings suggest that phosphorylation events throughout the IDR may have differential effects on Cdc15 function.

The Cdc15 IDR is important for normal cytokinesis dynamics and CR integrity

To better understand the defects observed in cdc15Δ IDR segment deletions, we measured the cytokinesis dynamics of mutants expressing spindle pole body (SPB) protein Sid4-mNG and CR protein Rlc1-mNG, which mark progression through mitosis and cytokinesis, respectively. As expected from the morphological analysis, the cytokinesis dynamics of cdc15Δ-1 were like that of WT. In contrast, the CR of cdc15Δ-2 cells took longer to constrict, while cdc15Δ-3 cells showed increased duration of both CR maturation (defined as the time between CR assembly and constriction) and CR constriction (Figure 3E).

We were particularly interested in the cytokinesis defects of cdc15Δ-2 cells, since segment 2 was the minimal IDR that supported viability (i.e., cdc15Δ-1Δ3 was viable; Figure 3D). Interestingly, the increased length of CR constriction in cdc15Δ-2 mutants correlated with a loss of CR integrity upon constriction initiation and subsequent asymmetric deposition of cell wall material (Figure 3F). More precisely, the Rlc1-mNG signal in z-projections became asymmetric and accumulated at one point of the cell, appearing to collapse and lose circularity (Figure 3F, double arrowhead). Cdc15-Δ2 and actin colocalized with Rlc1, indicating that the entire CR collapsed to this single point (Supplemental Figure S4, A and B). After CR collapse, septum formation became visible at the point(s) where the CR proteins remained (Figure 3F, yellow arrowheads). This was confirmed by imaging Rlc1-mCherry (mCh) to mark the CR and glucan synthase GFP-Bgs4 to mark the position of cell wall deposition in cells oriented vertically (Wang and Tran, 2014) (Figure 4A) and in the standard xy plane (Supplemental Figure S4C). A Z-series of images of Rlc1-mCh and the membrane marker Acyl-GFP also provided a clear view of asymmetric furrowing in cdc15Δ-2 cells (Figure 4B). The observation that the position of septum formation corresponded to the asymmetric localization of Rlc1 is consistent with a previous report that the CR or actin remnants of the CR locally stimulate cell wall synthesis (Zhou et al., 2015). Asymmetric septation has not been observed in any previously characterized Cdc15 mutant with defective membrane association or oligomerization or reduced abundance (Huang et al., 2007; Arasada and Pollard, 2014; McDonald et al., 2015). Vice versa, the CR unraveling observed in the absence of both Cdc15's and Imp2's SH3 domains (Roberts-Galbraith et al., 2009) and the CR sliding observed in F-BAR oligomerization and membrane-binding mutants were not observed in cdc15Δ-2 cells during time-lapse live-cell imaging (e.g., Figure 3F), nor did cdc15Δ-2 cells have off-center septa (as measured by long-to-short cell ratio; Supplemental Figure S4D). These results suggest that the function of segment 2 is distinct from that of the F-BAR and SH3 domains.

Cell wall enzymes are trafficked to the division site and active in cdc15Δ-2

The septum is a trilaminar structure. The Bgs1-dependent primary septum is deposited first and sandwiched by secondary septa deposited by Bgs4 and Ags1; enzymatic digestion of the primary septum leads to daughter cell separation (reviewed in Willet et al., 2015b; García Cortés et al., 2016; Perez et al., 2016). We have already demonstrated that Bgs4 is recruited to the division site (Figure 4A and Supplemental Figure S4C). However, the Cdc15 IDR has been implicated in the recruitment of Bgs1 (Arasada and Pollard, 2014). Therefore, to test whether cdc15Δ-2 cells failed to recruit Bgs1 adequately, we monitored GFP-Bgs1 localization in cdc15Δ-2 cells. GFP-Bgs1 was recruited to the division site with similar temporal kinetics and abundance in WT and cdc15Δ-2 (Figure 5). However, consistent with the asymmetric localization of GFP-Bgs4 (Figure 4A) and the deposition of septum material (Figures 3E and 4), GFP-Bgs1 distribution was not always circumferential in cdc15Δ-2 cells (Figure 5A). Taken together, these data indicate that cell wall synthesis is activated at the proper time but not necessarily in the proper location in cdc15Δ-2. In fact, cdc15Δ-2 was synthetic lethal with bgs1-ts/cps1-191 (Supplemental Figure S5B), indicating that robust cell wall deposition is crucial for division and viability of cdc15Δ-2.

CR composition is altered in cdc15Δ-2 cells

The results described above indicate that the asymmetry of CR constriction in cdc15Δ-2 is not due to a defect in septation but rather results from a CR-intrinsic defect that leads to loss of a ring structure upon constriction onset and instead CR proteins remain at the division site as arcs or strands. Indeed, cdc15Δ-2 mutants were sensitive to a low dose of latrunculin A (Supplemental Figure S5A) and were synthetically sick or lethal with numerous regulators of the actin cytoskeleton (Supplemental Figure S5B), indicating lower tolerance to further perturbations of the CR. Therefore, we used fluorescence microscopy to screen actin-binding proteins, actin regulators, Cdc15-binding partners, and other CR proteins for changes in their localization to the division site in cdc15Δ-2 (Supplemental Figure S5C). The calcineurin phosphatase, which comprises catalytic subunit Ppb1 and regulatory subunit Cnb1, has
FIGURE 3: Cdc15 IDR deletion mutants have cytokinesis defects. (A) Schematic of Cdc15 indicating the aa boundaries of IDR regions. (B) BF images of the indicated strains. Bar, 5 µm. (C) Quantification of the number of nuclei (left) and septa (right) per cell of the indicated strains, determined from cells stained with DAPI and methyl blue to visualize DNA and septum, respectively. Results are the means from three biological replicates, \( n > 198 \) cells per replicate. Error bars are SD. *, \( p < 0.0001 \). Ordinary two-way analysis of variance (ANOVA) with Tukey's multiple comparisons test. (D) Table of cdc15 alleles, schematic of gene product, and ability to rescue cdc15 null. Schematics are drawn to scale. Black lines indicate deletion. (E) Duration of cytokinesis stages determined from imaging of strains expressing the indicated cdc15 allele plus rlc1-mNG and sid4-mNG to label the CR and SPB, respectively. Results are the means from three biological replicates.
been found to regulate Cdc15 (Martin-Garcia et al., 2018). Also, the morphologies of cdc15-Δ2 and ppb1Δ (Yoshida et al., 1994) are similar. This prompted us to examine Ppb1’s localization in cdc15-Δ2. We found that Ppb1 was greatly reduced at the CR of cdc15-Δ2 cells, but it localized normally in other cdc15 segment deletions (Figure 6A). This was further confirmed by two-color imaging of Ppb1 and the CR marker Rlc1 (Figure 6B). The localization of Ppb1 to the CR was previously found to depend on the CR protein Pxl1 (Martin-Garcia et al., 2018), which associates with Cdc15 by Co-IP (Pinar et al., 2008; Roberts-Galbraith et al., 2009). Given that Ppb1 was essentially absent from the CR, it was surprising to find that Pxl1 localized normally to the division site in cdc15-Δ2 (Figure 6C). This result suggests that Pxl1 is not sufficient for normal Ppb1 localization to the CR.

**DISCUSSION**

It was unknown what protein feature(s) distinguish the essential Cdc15 from its nonessential paralogue Imp2. Using domain-swapping experiments, we determined that the essential function of the F-BAR domain can be replaced by Imp2’s F-BAR domain as well as other human F-BAR domains, whereas Cdc15’s IDR is uniquely essential and cannot be replaced by that of Imp2. Deletions of segments of the IDR exhibit either lengthened periods of CR matura

CR strands or fragments direct primary septum deposition

In cdc15-Δ2, the CR appears to form normally and the glucan syntheses are normally recruited, but upon constriction, the CR loses its structure and Bgs1 then localizes with the CR-protein strands/ar- clike structures that remain. As a result, the septum is deposited— albeit inefficiently—from that point. This behavior of the CR and Bgs1 in cdc15-Δ2 cells is consistent with the idea that the primary purpose of the S. pombe CR may be to direct efficient cell wall deposition, which in turn provides the force for division (Proctor et al., 2012; Stachowiak et al., 2014; Thiyagarajan et al., 2015; Willet et al., 2015b; Zhou et al., 2015). The CR has been previously proposed to direct vesicular traffic (Vjestica et al., 2008), and the colocalization of glucan syntheses with CR remnants throughout constriction in cdc15-Δ2 indicates that the CR is indeed a landmark for glucan syn

In terms of the F-BAR-Cdc15 chimeras’ defects or lethality, multiple explanations can be imagined. In addition to the strong possibility that some fusion proteins were not structurally sound, some F-BAR domains bind directly to other proteins (Shoham et al., 2003; Hansen et al., 2011; Senju et al., 2011; Begonja et al., 2015; Garabedian et al., 2018; Liu et al., 2019), including Cdc15’s F-BAR domain that binds the formin Cdc12 (Willet et al., 2015a). However, a mutant that disrupts the Cdc15-Cdc12 interaction is viable (Willet et al., 2015a), so this is unlikely to be the only explanation for de

The exchangeability of F-BAR domains

Five of the 15 F-BAR domains tested replaced the essential function of Cdc15’s F-BAR domain: the F-BAR domains of S. pombe Imp2 and human Gas7, PSTPIP1, PACSIN2/syndapin2, and Fer (Supplemental Table S1). Given that the proteins in which these domains normally reside are involved in a variety of apparently unlinked functions and the F-BAR domains themselves have distinct biochemical characteristics and structures, we conclude that there is significant plasticity among F-BAR domains. This is perhaps not surprising, since all F-BAR domains share the ability to interact electrostatically with negatively charged phospholipids and to oligomerize (Supplemental Table S2; Tsujita et al., 2006; Shimada et al., 2010; Bai et al., 2012; Bai and Zheng, 2013; Goh et al., 2012; McDonald et al., 2015, 2016; McDonald and Gould, 2016). Although there are reported differences among F-BAR proteins in phospholipid or curvature specificity, F-BAR domains themselves interact relatively nonspecifically with negatively charged phospholipids, and it is only protein features adjacent to the F-BAR domain, such as the FX(C) domain (Itoh et al., 2009; Yamamoto et al., 2018), that convey phospholipid or curvature specificity to the core F-BAR domain (reviewed in Itoh and De Camilli, 2006). Thus, in the context of the full-length protein, an F-BAR domain acts as a module that concentrates the protein on membranes where other linked modules (e.g., SH3 domains, RhoGAP domains) provide the bulk of specific functions.

replicates, n > 5 cells per replicate, and the total number of cells analyzed for each strain is indicated on the graph. Error bars are SD. * p < 0.0001. Ordinary one-way ANOVA with Tukey’s multiple comparisons test. (F) Representative time-lapse series of the indicated strains. Images were acquired every 2 min. A single BF slice and max intensity z-projections of deconvolved mNG images are shown. Numbers indicate min from SPB separation. Bar, 5 µm. Double arrow indicates first frame when CR loses ringlike structure. Yellow arrowheads point out first and last frame with visible asymmetric septum deposition.
of their F-BAR domains and adjacent protein regions, including globular domains and IDRs. The IDR of *S. cerevisiae* Hof1 mediates its localization to the CR and is important for CR contraction; it also binds septins (Meitinger et al., 2011, 2013; Oh et al., 2013). The IDR in FCHO2 binds and allosterically activates AP-2 (Hollopeter et al., 2014; Umasankar et al., 2014). The IDR in Fer undergoes a disorder-to-order transition to convey a curvature sensing function (Yamamoto et al., 2018). In the case of Cdc15’s IDR, we found that it

FIGURE 4: Characterizing the CR defect in *cdc15Δ2*. (A) End-on imaging of the indicated strains. Numbers indicate minutes elapsed since the start of the series. Bar, 5 µm. (B) Z-slices of two representative cells of the indicated genotype. The pink dashed lines mark the middle slice. Z-slices were taken every 0.2 µm through the volume of the cell. Fluorescence images are deconvolved. Bar, 5 µm.
plays a key role in recruiting the phosphatase calcineurin. Calcineurin localization at the CR depends on the second region of Cdc15’s IDR (aa 503–677) and Pxl1 (Martin-Garcia et al., 2018), and determining how these two proteins collaborate for proper calcineurin localization will be an important next step. Cdc15-∆2 is dephosphorylated with timings similar to WT, which indicates that Cdc15’s role scaffolding calcineurin at the CR would be separate from its identity as a calcineurin substrate (Martin-Garcia et al., 2018), and furthermore that either other phosphatases are redundant with calcineurin or cytoplasmic calcineurin dephosphorylates Cdc15. Finally, because cdc15-∆3 also has cytokinesis defects, but is not required for calcineurin recruitment to the CR, it will be interesting to determine the other functions performed by Cdc15’s IDR.

MATERIALS AND METHODS

Yeast methods

*S. pombe* strains (Supplemental Table S3) were grown in yeast extract with supplements (YES), YES without additional adenine, or Edinburgh minimal media (EMM) plus selective supplements (Moreno et al., 1991; Forsburg and Rhind, 2006). Transformation of yeast with plasmid or linear DNA was accomplished using electroporation (Forsburg and Rhind, 2006) and lithium acetate methods (Keeney and Boeke, 1994; Forsburg and Rhind, 2006), respectively.

For integration of *cdc15* alleles at the endogenous locus, *cdc15*+/cdc15::ura4+ was transformed with pIRT2 vector with the desired *cdc15* allele plus *cdc15*5′ and 3′ noncoding regions (Roberts-Galbraith et al., 2009). Spores were germinated on selective medium to select for *cdc15::ura4*+ haploids that have the vector. Haploid integrants were recovered based on resistance to 5-fluoroorotic acid (FOA) and integration plus loss of vector was verified by growth on selective media, PCR and/or microscopy, and finally sequencing.

To confirm the lethality of *cdc15* mutants, the *ura4* cassette in a *cdc15*/cdc15::ura4* diploid strain was replaced with a cassette consisting of a 500-base-pair 5′ *cdc15* flanking region, the mutant *cdc15* coding sequence, *kan*5, and a 500-base-pair 3′ *cdc15* flanking region. Integrants resistant to 5-FOA and G418 were selected and confirmed by whole-cell PCR before sporulation and tetrad dissection on YES agar. Plates were grown at 32°C. At least 10

FIGURE 5: Cell wall synthases are properly trafficked in *cdc15-Δ2*. (A) Representative time-lapse series of the indicated strains. Sum z-projections are shown. Images were acquired every 2 min. Numbers indicate minutes from SPB separation. Bar, 5 μm. (B) Timing of GFP-Bgs1 arrival to the division site in the indicated strains also expressing Sid4-mNG and Rlc1-mCh. Error bars are SD, *p* = 0.67. Student’s *t* test. (C) Average peak fluorescence of GFP-Bgs1 in the indicated strains also expressing Sid4-mNG and Rlc1-mCh. Error bars are SD, *p* = 0.85. Student’s *t* test. (D) Timing of Rlc1 (×) and Bgs1 (●) peak intensity relative to SPB separation for the indicated strains determined from strains imaged as in (A). Results in A–D are the means from two biological replicates, *n* > 9 cells per replicate, and total number of cells analyzed for each strain is indicated on the graph.
tetrads were dissected for each integrant. Production of the gene product was confirmed using lysates from the diploid and/or by expressing from the plasmid. In some cases, the experiment was repeated and tetrad plates were grown at alternative temperatures of 25 and/or 29°C; in all cases, the results were the same.

Genes were tagged at the 3' ends of their ORFs with mNG:kan<sup>R</sup>, mNG:hyg<sup>R</sup>, or mCh:nat<sup>R</sup> using pFA6 cassettes as previously described (Wach et al., 1994; Bahler et al., 1998; Willet et al., 2015a). Integration of tags was verified using whole-cell PCR and/or microscopy.

The mNG-pxl1:kan<sup>R</sup> strain was generated by transforming KGY16851 with a cassette consisting of a 500-base-pair 5' pxl1<sup,+</sup> flanking region, the mNG-pxl1 coding sequence, kan<sup>R</sup>, and a 500-base-pair 3' pxl1<sup,+</sup> flanking region. Integrants resistant to G418 were selected and confirmed by whole-cell PCR.

Introduction of tagged loci or cdc15 mutants into other genetic backgrounds was accomplished using standard <i>S. pombe</i> mating, sporulation, and tetrad dissection techniques.

For growth assays, cells were grown to log phase in YES medium at 32°C for Supplemental Figure S2C and 25°C for Supplemental Figure S5, A and B. A suspension of 40 × 10<sup>6</sup> cells/ml YES was serially diluted 10-fold and 3 µl of each dilution was spotted on YES plates for growth at indicated temperatures. All spot assays were performed in duplicate.

**FIGURE 6:** Calcineurin is not recruited to CR in cdc15-∆2. (A–C) Representative sum projections of the indicated strains. Bar, 5 µm. (A) At least 100 cells from 3 d of imaging were examined. (B, C) At least 50 cells were examined.
Molecular biology
cdc15 plasmids were created by amplifying cdc15 fragments by PCR from pKG4456 (Roberts-Galbraith et al., 2009) and subsequent cloning into piRRT2 for integration at the endogenous locus (Roberts-Galbraith et al., 2009). All constructs were sequenced for verification.

Microscopy methods
Except for Figures 4A and 5, images of S. pombe cells were acquired using a Personal DeltaVision (GE Healthcare, Issaquah, WA) that includes an Olympus IX71 microscope, 60 × NA 1.42 Plan Apochromat and 100 × NA 1.40 U Plan S Apochromat objectives, fixed and live-cell filter wheels, a Photometrics CoolSnap HQ2 camera, and softWoRx imaging software (Applied Precision).

For live-cell fluorescence imaging, strains were grown overnight to log phase in YES media at 25°C in a shaking water bath, except for the chimeras in Figure 2B, which were grown at 32°C. For the brightfield (BF) images in Figure 3B and Supplementary Figure S1E, cells were grown at 32°C.

For Figures 2B, 3B, 4B, and 6 and Supplementary Figures S1E, S3B, S4C, and SSC, cells were imaged in YES media at 23°–29°C.

For Supplementary Figure S4A, time-lapse imaging was performed on a 2% MAUL agarose pad. Images were acquired every 2 min. For Figure 3, E and F, time-lapse imaging was performed using an ONIX microfluidics perfusion system (CellASIC ONIX; EMD Millipore). A suspension of 50 µl of 40 × 10^6 cells/ml YES was loaded into Y04C plates for 5 s at 8 psi. YES medium was flowed into the chamber at 5 psi throughout imaging. Time-lapse images were obtained at 2-min intervals. Representative fluorescence images are maximum-intensity or sum projections of z sections spaced at 0.2–0.5 µm that have been deconvolved with 10 iterations.

For Figure 4A, end-on images were acquired with a spinning-disk system, which includes a Zeiss Axiovert200m microscope, Yokogawa CSU-22, 63X NA 1.46 planApochromat and 100X numerical aperture (NA) 1.40 PlanApo oil immersion objectives, and a Hamamatsu ImageEM-X2 camera. Cells were grown to log phase in EMM plus adenosine, uracil, and leucine, loaded into vertical chambers in a 4% MAUL agarose pad, and imaged every 2 min. Six z-slices at 0.5-µm intervals around the middle of the cell were acquired at each time point.

For Figure 5, time-lapse images were acquired with a Nikon spinning-disk system, which incorporates a Yokogawa CSU-X1 spinning disk head, Andor DU-897 EMCCD, a high-speed piezo [2] stage, a four-line high-powered solid state laser laser, and a Plan Apo Lambda (oil) 60 × 1.40 NA WD 0.13 mm. Cells were mounted on 2% MAUL agarose pads. Images were acquired every 2 min. Fifteen z-slices at 0.4-µm intervals were acquired at each time point. Sum projections were analyzed and used for representative images. Sum projections were corrected for background fluorescence before quantitation.

For Figure 3C and Supplementary Figures S1, A and B, and S4D, to visualize nuclei and septa, cells were grown to log phase at 32°C and fixed with ice-cold 70% ethanol at 4°C for at least 30 min. Approximately 0.5 OD of fixed cells were washed once with PBS, pH 7.5, and then incubated in 50 µl 1-mg/ml mCherry (MB; Sigma; M6900) for 30 min at RT. MB-stained cells were centrifuged and mixed with 5 µg/ml DAPI (4′,6-diamidino-2-phenylindole) (Sigma; D9542) at a 1:1 ratio immediately before imaging.

For Supplementary Figure S4B, cells were fixed in formaldehyde, permeabilized for 2 min in PBS with 0.1% NP-40, washed with PBS, and then stained with Alexa-Fluor-488-phalloidin for 60 min before imaging.

Image analysis was performed using ImageJ software (Schindelin et al., 2012).

Protein methods
Cell pellets were snap-frozen in dry ice–ethanol baths. Denatured lysates were prepared by bead disruption (Fastprep cell homogenizer; MP Biomedicals) in NP-40 buffer containing SDS as previously described (Gould et al., 1991), except with the addition of 0.5 mM diisopropyl fluorophosphates (Sigma-Aldrich).

For immunoprecipitation and lambda phosphatase collapse, lysates were incubated with anti-Cdc15 (Roberts-Galbraith et al., 2009) and Protein A magnetic beads or sepharose (GE Healthcare; 17-5280-04) for 2 h at 4°C. Beads were washed three times with NP-40 buffer and twice with phosphatase buffer (150 mM NaCl, 50 mM HEPES, pH 7.4) before being split in two and added to either lambda phosphatase reaction or control. Lambda phosphatase collapse was performed according to the manufacturer’s protocol (New England Biolabs; P0753). Reactions were stopped by the addition of sample buffer.

Protein samples were resolved by SDS-PAGE and transferred to a polyvinylidene fluoride membrane (Immobilon P, EMD Millipore). Anti-α-tubulin (B512) mouse monoclonal antibody (Sigma-Aldrich), anti-Cdc2 (PSTAIRE) mouse monoclonal antibody (Sigma-Aldrich; P7962), or anti-Cdc15(1-405) rabbit polyclonal antibody (Roberts-Galbraith et al., 2009) was used in immunoprecipitations and/or as a primary antibody in immunoblotting. Secondary antibodies were conjugated to IRDye800 or IRDye680 (LI-COR Biosciences). Blotted proteins were detected via an Odyssey machine (LI-COR Biosciences).

Statistical analysis
Statistical analysis was performed using Prism (GraphPad).

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