Cdc14 phosphatase directs centrosome re-duplication at the meiosis I to meiosis II transition in budding yeast [version 2; peer review: 4 approved]

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Abstract

Background: Gametes are generated through a specialized cell division called meiosis, in which ploidy is reduced by half because two consecutive rounds of chromosome segregation, meiosis I and meiosis II, occur without intervening DNA replication. This contrasts with the mitotic cell cycle where DNA replication and chromosome segregation alternate to maintain the same ploidy. At the end of mitosis, cyclin-dependent kinases (CDKs) are inactivated. This low CDK state in late mitosis/G1 allows for critical preparatory events for DNA replication and centrosome/spindle pole body (SPB) duplication. However, their execution is inhibited until S phase, where further preparatory events are also prevented. This “licensing” ensures that both the chromosomes and the centrosomes/SPBs replicate exactly once per cell cycle, thereby maintaining constant ploidy. Crucially, between meiosis I and meiosis II, centrosomes/SPBs must be re-licensed, but DNA re-replication must be avoided. In budding yeast, the Cdc14 protein phosphatase triggers CDK down regulation to promote exit from mitosis. Cdc14 also regulates the meiosis I to meiosis II transition, though its mode of action has remained unclear.

Methods: Fluorescence and electron microscopy was combined with proteomics to probe SPB duplication in cells with inactive or hyperactive Cdc14.

Results: We demonstrate that Cdc14 ensures two successive nuclear divisions by re-licensing SPBs at the meiosis I to meiosis II transition. We show that Cdc14 is asymmetrically enriched on a single SPB during anaphase I and provide evidence that this enrichment promotes SPB re-duplication. Cells with impaired Cdc14 activity fail to promote extension of the SPB half-bridge, the initial step in morphogenesis of a new SPB. Conversely, cells with hyper-active Cdc14 duplicate SPBs, but fail to induce their separation.

Conclusion: Our findings implicate reversal of key CDK-dependent phosphorylations in the differential licensing of cyclical events at the meiosis I to meiosis II transition.
Keywords
Meiosis, Cdc14, SPB, centrosome

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Introduction

Meiosis is a specialized cell division, which generates gametes. In the canonical mitotic cell cycle, ploidy is maintained by alternating S and M phases. In contrast, during meiosis, chromosome duplication in S phase is followed by two consecutive chromosome segregation phases, meiosis I and meiosis II, to generate gametes with half the ploidy of the parental cell. Therefore, in addition to distinct modifications to the chromosome segregation machinery, meiosis requires a re-wiring of cell cycle controls (reviewed in Duro & Marston, 2015). Progression through the cell cycle is driven by cyclin-dependent kinases (CDKs), in association with distinct cyclin subunits. CDK activity is low in G1, but upon cell cycle entry, activation of S-phase and mitotic CDKs in turn promote DNA replication followed by spindle assembly and chromosome segregation. Following completion of chromosome segregation, CDKs are inactivated, triggering spindle disassembly and the return to G1 (mitotic exit) (Stegmeier & Amon, 2004). This state of low CDK activity in G1 allows for the re-licensing of DNA replication origins and centrosomes/spindle pole bodies (SPBs), events that must be restricted to once per cell cycle. In budding yeast, the Cdc14 phosphatase triggers CDK inactivation through multiple mechanisms to promote exit from mitosis and return to G1 (Jaspersen et al., 1998; Visintin et al., 1998; Zachariae et al., 1998). Cdc14 is regulated by its localization: for the majority of the cell cycle it is sequestered in the nucleolus through association with its inhibitor, Cfi1/Net1 (Shou et al., 1999; Visintin et al., 1999). Upon chromosome segregation at anaphase onset, the Cdc Fourteen Early Anaphase Release (FEAR) network promotes Cdc14 release from the nucleolus into the nucleus; later in anaphase the Mitotic Exit Network (MEN) maintains Cdc14 release throughout the cytoplasm (Pereira et al., 2002a; Stegmeier & Amon, 2004; Yoshida et al., 2002). While FEAR-dependent Cdc14 release promotes successful completion of chromosome segregation, only MEN-dependent Cdc14 release is sufficient to trigger exit from mitosis, leading to spindle disassembly and entry into G1 (Yellman & Roeder, 2015).

The state of low CDK activity in G1 is permissive for the assembly of pre-replicative complexes, the later firing of which requires S phase CDKs (reviewed in Blow & Dutta, 2005; Drury & Diffley, 2009). This separation of pre-RC assembly and firing into differential CDK activity states ensures that DNA replication is strictly restricted to once per cell cycle. Similarly, centrosome/SPB duplication must occur exactly once in each cell cycle. Yeast SPBs are microtubule-organising centres, composed of at least 19 proteins, forming three layers that are assembled into a cylindrical organelle embedded in the nuclear envelope (reviewed in Jaspersen & Winey, 2004). SPB duplication is initiated in G1 by extension of the half-bridge (Byers & Goetsch, 1974; Vallen et al., 1994), which protrudes from the central SPB layer and consists of Kar1, Mps3, Sfi1 and Cdc31. The full bridge structure serves as the site for new SPB assembly (Jaspersen et al., 2002; Kilmartin, 2003; Spang et al., 1993; Spang et al., 1995), so that at the close of G1, two side-by-side SPBs are physically connected by a full-bridge (Byers & Goetsch, 1974). S phase CDK activity severs the bridge structure triggering SPB separation. Recent work has identified the bridge component Sfi1 as the target of S phase CDKs (Avena et al., 2014; Elserafy et al., 2014). S phase CDKs also activate the Cin8 and Kip1 motors that drive separation of the SPBs to enable bipolar spindle formation (Crasta et al., 2006; Crasta et al., 2008; Haase et al., 2001). In mitosis, M phase CDKs prevent SPB re-duplication through phosphorylation of Sfi1, thereby preventing the initiation of half-bridge elongation (Avena et al., 2014; Bouhlel et al., 2015; Elserafy et al., 2014; Haase et al., 2001). For SPBs to re-duplicate in the following G1-phase, Sfi1 must be dephosphorylated and Cib-CDKs inhibited. Recent findings have implicated Cdc14 as the phosphatase responsible for this dephosphorylation event in budding yeast (Avena et al., 2014; Elserafy et al., 2014).

The meiosis I to meiosis II transition requires specialized cell cycle controls. Uniquely, chromosome segregation at meiosis I exit is followed not by a DNA replication phase, but by a second chromosome segregation phase, meiosis II. Importantly, centrosome/SPBs must be re-licensed at meiosis I exit to permit this additional segregation event, yet DNA replication origins must not be re-set to avoid over-duplication of chromosomes. How this is controlled is not clear. One potential contributing factor is the retention of partial CDK activity between meiosis I and meiosis II. Indeed, there is evidence to suggest that CDKs are only partially downgraded between meiosis I and II in Xenopus oocytes and fission yeast (Iwabuchi et al., 2000; Izawa et al., 2005). However, how these global alterations in CDK activity impinge on the differential re-licensing of DNA replication origins and SPBs has not been investigated.

In budding yeast, the Cdc14 phosphatase plays a prominent role in the meiosis I to meiosis II transition (Buonomo et al., 2003; Marston et al., 2003). Following meiosis I chromosome segregation, cdc14-1 mutants disassemble the spindle, only to reassemble a single spindle that directs segregation of some chromosomes in a meiosis II-like manner (Bizzari & Marston, 2011). The result is binucleate, rather than tetranucleate, cells with a mixed complement of chromosomes (Sharon & Simchen, 1990). Furthermore, ectopic activation of Cdc14 is also detrimental to meiosis. Depletion of the regulatory subunit of protein phosphatase 2A, Cdc55, results in premature release of Cdc14 from the nucleolus in meiosis and a block to spindle assembly, so that nuclear division largely fails (Bizzari & Marston, 2011; Kerr et al., 2011; Nolt et al., 2011). Inactivation of Cdc14 in Cdc55-depleted cells enables spindle assembly and the production of binucleate cells, indicating that over-active Cdc14 is responsible for the block to spindle assembly (Bizzari & Marston, 2011; Kerr et al., 2011). Therefore, proper regulation of Cdc14 is critical to control spindle morphogenesis during meiosis, but not the dissolution of linkages between chromosomes. While the FEAR network plays a vital role in the release of Cdc14 in anaphase I, MEN is dispensable (Buonomo et al., 2000; Kamieniecki et al., 2005; Marston et al., 2003; Pablo-Hernando et al., 2007) and does not appear to be active until anaphase II (Attner & Amon, 2012).
Since FEAR-dependent Cdc14 release appears insufficient to trigger CDK inactivation (Stegmeier et al., 2002; Yellman & Roeder, 2015), it is likely that the critical role of Cdc14 at the meiosis I to meiosis II transition is to reverse the phosphorylation of key substrates.

Here we investigate the role of Cdc14 in executing the meiosis I to meiosis II transition. Our findings suggest that a critical role of Cdc14 at meiosis I exit is to re-license SPB duplication. This re-licensing ensures assembly of a pair of spindles for a second round of nuclear division at meiosis II. Conversely, premature Cdc14 activation prevents SPB separation. We provide evidence that Cdc14 associates with the SPB in meiosis and that this localization is important for permitting the duplication cycle. Our data suggest that the critical function of Cdc14 at the meiosis I to meiosis II transition is to reverse key phosphorylations to enable SPB re-duplication.

Materials and methods
Yeast strains and plasmids
Yeast strains used in this study were generated using standard genetic methods and are given in Table 1. pCLB2-3HA-CDC55

| Strain number | Genotype |
|---------------|----------|
| AM1835        | MATa/MATα Wild type |
| AM9319        | MATa/MATα CDC14-SZZ(TAP)::KanMX6/CDC14-SZZ(TAP)::KanMX6 cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55 |
| AM9434        | MATa/MATα CDC14-SZZ(TAP)::KanMX6/CDC14-SZZ(TAP)::KanMX6 |
| AM9459        | MATa/MATα cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55 |
| AM11443       | MATa/MATα SPC42-3FLAG-KANMX6/SPC42-3FLAG-KANMX6 cdc14::KanMX6/cdc14::KanMX6 trp1::cdc14-1::TRP1::LEU2/trp1::cdc14-1::TRP1::LEU2 |
| AM11444       | MATa/MATα SPC42-3FLAG-KANMX6/SPC42-3FLAG-KANMX6 |
| AM11517       | MATa/MATα CDC14-GFP-LEU2/CDC14-GFP-LEU2 SPC42-tdTomato::NAT/SPC42-tdTomato::NAT GAL-NDT80::TRP1/GAL-NDT80::TRP1 ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM13123       | MATa/MATα CDC14-GFP-LEU2/CDC14-GFP-LEU2 SPC42-tdTomato::NAT/SPC42-tdTomato::NAT cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55 GAL-NDT80::TRP1/GAL-NDT80::TRP1 ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM13989       | MATa/MATα SPC42-tdTomato::NAT/SPC42-tdTomato::NAT GAL-NDT80::TRP1/GAL-NDT80::TRP1 ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM15543       | MATa/MATα CDC14-GFP-LEU2/CDC14-GFP-LEU2 SPC42-tdTomato::NAT/SPC42-tdTomato::NAT bub2Δ::NAT/bub2Δ::NAT GAL-NDT80::TRP1/GAL-NDT80::TRP1 ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| Strain number | Genotype |
|---------------|----------|
| AM15984       | MATa/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM15985       | MATa/MATα  
his3::HIS3p-GFP-TUB1-HIS3/his3::HIS3p-GFP-TUB1-HIS3  
PDS1-tdTomato-KITRP1/PDS1-tdTomato-KITRP1  
spo12Δ::LEU2/spo12Δ::LEU2  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16019       | MATa/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
bfa1Δ::NAT/bfa1Δ::NAT  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16020       | MATa/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
slk19Δ::KANMX6/slk19Δ::KANMX6  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16064       | MATa/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
spo12Δ::LEU2/spo12Δ::LEU2  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16065       | MATa/MATα  
his3::HIS3p-GFP-TUB1-HIS3/his3::HIS3p-GFP-TUB1-HIS3  
PDS1-tdTomato-KITRP1/PDS1-tdTomato-KITRP1  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16066       | MATa/MATα  
his3::HIS3p-GFP-TUB1-HIS3/his3::HIS3p-GFP-TUB1-HIS3  
PDS1-tdTomato-KITRP1/PDS1-tdTomato-KITRP1  
cdc14::KanMX6/cdc14::KanMX6  
leu2::cdc14-1::LEU2::TRP1/leu2::cdc14-1::LEU2::TRP1  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16077       | MATa/MATα  
cdc14::KanMX6/cdc14::KanMX6  
leu2::cdc14-1::LEU2::TRP1/leu2::cdc14-1::LEU2::TRP1  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16079       | MATa/MATα  
CDC14-GFP-LEU2/CDC14-GFP-LEU2  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
bfa1Δ::NAT/bfa1Δ::NAT  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| Strain number | Genotype |
|---------------|----------|
| AM16080       | MATα/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
bub2Δ::NAT/bub2Δ::NAT  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16110       | MATα/MATα  
his3::HIS3p-GFP-TUB1-HIS3/his3::HIS3p-GFP-TUB1-HIS3  
PDS1-tdTomato-KIT/PR1/PDS1-tdTomato-KITPR1/slk19Δ::TRP/slk19Δ::TRP  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16163       | MATα/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
cdc14::KanMX6/cdc14::KanMX6  
trp1::cdc14::TRP1::LEU2/cdc14-1::TRP1::LEU2  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM16198       | MATα/MATα  
cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM17134       | MATα/MATα  
CDC14-GFP-LEU2/CDC14-GFP-LEU2  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
kin4Δ::NAT/kin4Δ::NAT  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM17341       | MATα/MATα  
CDC14-GFP-LEU2/CDC14-GFP-LEU2  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
bmh1Δ::NAT/bmh1Δ::NAT  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM17740       | MATα/MATα  
Bfa1-tdTomato::NAT/Bfa1-tdTomato::NAT  
SPC42-CFP::TRP1/SPC42-CFP::TRP1  
CDC14-GFP-LEU2/CDC14-GFP-LEU2  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
| AM17904       | MATα/MATα  
SPC42-tdTomato::NAT/SPC42-tdTomato::NAT  
cdc14::KanMX6/cdc14::KanMX6  
trp1::cdc14-1::TRP1::LEU2/trp1::cdc14-1::TRP1::LEU2  
cdc55::KanMX6::PCLB2-3HA-CDC55/cdc55::KanMX6::PCLB2-3HA-CDC55  
GAL-NDT80::TRP1/GAL-NDT80::TRP1  
ura3::pGPD1-GAL4(848).ER::URA3/ura3::pGPD1-GAL4(848).ER::URA3 |
(Clift et al., 2009), CLB1-9MYC (Buonomo et al., 2003), pCLB2-CD20 (Lee & Amon, 2003), GAL-NDT80 and pGPD1-GAL4(848), ER (Benjamin et al., 2003), cdc14-1, slk19Δ and spo12Δ (Marston et al., 2003) CDC14-GFP, PDS1-tdTomato and GFP-TUB1 (Matos et al., 2008) were as described. SPC42-tTomato, CDC14-SZZ(TAP), SPC42-3FLAG, SPC42-GFP, BFA1-tTomato, bub1Δ, bfa1Δ, kin4Δ and bmh1Δ were made using a one-step PCR method (Longtine et al., 1998). The SPC42-CFP strain was obtained by integrating the pHX144 plasmid at the SPC42 locus (He et al., 2000).

**Growth conditions**

To induce meiosis, diploid strains were removed from -80°C storage onto YPG (2% Bacto-peptone, 1% Bacto-yeast extract, 2.5% glycerol) plates and grown overnight (~16 h). The following day, cells were patched to 4%YPDA (2% Bacto-peptone, 1% Bacto-yeast extract, 4% glucose, 0.3 mM adenine) plates. On the third consecutive day, YPDA (2% Bacto-peptone, 1% Bacto-yeast extract, 2% glucose, 0.3 mM adenine) liquid cultures were inoculated with yeast strains and grown overnight. Cells were then diluted to OD₆₀₀ =0.2 in BYTA (2% Bacto-peptone, 1% Bacto-yeast extract, 1% Potassium acetate, 50 mM Potassium Phthalate) liquid culture and grown to OD₆₀₀ = 6-10. On the fifth day, cells were washed once in sterile water and resuspended in SPO liquid media (0.3% Potassium acetate, pH 7.0) at OD₆₀₀ = 1.8-3. Meiosis was performed at 30°C. For experiments with temperature sensitive cdc14Δ mutants, all steps were performed at room temperature prior to resuspension in SPO medium, upon which cultures were shifted to 30°C and incubated at this temperature for the remainder of the experiment.

**Fluorescence microscopy**

To visualize chromosomes and SPBs labelled with fluorescent proteins in fixed cells, 100 µl of meiotic culture was added to eppendorf tubes containing 10 µl of 37% formaldehyde and incubated for 8–10 mins at room temperature. Cells were spun down, washed with 1 ml of 80% ethanol and resuspended in 20 µl of 1 µg/ml DAPI before microscopy. Indirect immunofluorescence of meiotic spindles was carried out as previously described (Bizzari & Marston, 2011).

Imaging of live cells at isolated timepoints was performed on ~1mm deep 2% agarose slides. In total, 100 µl of meiotic culture was spun down and resuspended in 5 µl of sporulation (SPO) media (0.3% Potassium acetate) and 2–3 µl of cell suspension was added to each agarose pad. The slide was covered with a glass coverslip and sealed with a molten mixture of vaseline:lamarlin:paraffin (1:1:1) before microscopy.

Live-cell meiotic movies were generated using CellASIC® ONIX Y040D Microfluidics plates (Merck Millipore). All chambers on the plate were washed three times with 500 µl of SPO media before 200 µl of SPO media plus 1mM β-estradiol was added to chambers 1–6. Plates were then pre-incubated at 30°C 30 mins. After incubation, 200 µl of prophase I arrested cells (by Ndt80-depletion; Carlile & Amon, 2008) was loaded into chamber 8. A total of four different strains can be imaged on a single plate. The microfluidics plate was attached via a low-profile manifold to the CellASIC® ONIX Microfluidic Platform Control System, and the assembly was placed on a Deltavision Elite microscope. Cells were loaded, visualised, washed with β-estradiol-containing SPO media and imaged.

Imaging of fixed cells or live cells at isolated time-points was carried out using a Zeiss Axio Imager Z1 and a Photometrics EMCCD camera. Images were taken using Micro-Manager v1.4 (https://micro-manager.org/) and processed using ImageJ software v1.47 (https://imagej.nih.gov/ij/). For the generation of microfluidics movies, a Deltavision® Elite live cell imaging system was utilised with an Olympus IX-71 microscope and a Photometrics EMCCD Cascade II camera. Multi-point images were taken using SoftWoRx v5.5 (http://www.gelifesciences.com/), movies were assembled in Image-Pro Plus (Media Cybernetics) and processed in ImageJ v1.47.

**Quantification of fluorescence signal**

Quantification of fluorescence signal was performed as described in Hoffman et al., 2001. In brief, the following equations were used:

\[
FBk = (FO - Fi) \times (Ai / (AO - Ai))
\]

\[
F_x = Fi - FBk
\]

O and I represent outer and inner regions, respectively. The inner region (I) contained ~90% of the signal measured. The outer region (O) was at least twice the area of the inner region (Ai=16 pixels; AO=64 pixels), and was used to calculate the surrounding background (Bk) signal. F signifies integrated fluorescence signal, calculated from Raw Integrated Densities, and A is the area of the boxes. AO were 16 and 64 pixels, respectively.

**Electron microscopy**

For sample preparation, 3 ml of culture from meiotic cell cycle time-course was vacuum filtered through a 0.45 µm Millipore filter. The cell paste was rapidly frozen under high pressure in a Wohlwend Compact 02 High Pressure Freezer. Frozen cell pellets were then freeze substituted in acetone containing 2% (w/v) osmium tetroxide and 0.1% (w/v) uranyl acetate at -80°C. Samples were slowly warmed to room temperature over three days. After washing cells twice in acetone, samples were embedded in Epon 812 resin (Hexion) through multiple changes of diluted resin with acetone (1:3, 1:1 and 3:1). Three more changes using undiluted Epon 812 resin were then freeze substituted in acetone containing 2% (w/v) osmium tetroxide and 0.1% (w/v) uranyl acetate at -80°C. Samples were slowly warmed to room temperature over three days. After washing cells twice in acetone, samples were embedded in Epon 812 resin (Hexion) through multiple changes of diluted resin with acetone (1:3, 1:1 and 3:1). Three more changes using undiluted Epon 812 resin were carried out over two days before resin was polymerised at 60–70°C overnight. Epon blocks were serially sectioned at a thickness of 70 nm and stained with 2% (w/v) uranyl acetate in sterile water for 8 mins, and then in Reynolds’ lead citrate for 3 mins. Sections were viewed on a Philips CM120 transmission electron microscope, and images were collected with a Gatan Orius CCD camera and processed using ImageJ v1.47.

**Immunoprecipitation**

Meiotic cells were harvested and washed with sterile water by centrifugation at 4000 rpm for 6 mins. Cells were resuspended in 0.2x cell volume of sterile water before drop-freezing in liquid nitrogen. Cells were ground five times in a Retsch Mixer Mill MM400. For Cdc14-SZZ(TAP) purification, the yeast lysate...
was thawed in Hyman (50 mM Bis-Tris propane, pH7; 100 mM KCl; 5 mM EGTA; 10% (v/v Glycerol)) with inhibitors (5 µg/ml each chymostatin, leupeptin, antipain, pepstatin A, E-64; 4mM AEBSF (pefabloc); 2mM benzamidine, 2mM PMSF, 0.4 mM LR-microcystin, N-ethylmaleimide (NEM), sodium orthovanadate, b-glycerolphosphate, sodium pyrophosphate). For Spc42-3FLAG purifications, we adapted SPB buffer (Niepel et al., 2005) by addition of inhibitors as above. Following thawing, Triton X-100 was added to a final concentration of 1% (w/v) and samples were sonicated at 39% amplitude for 1 x 30 secs per 10 ml of lysate. Lysates were centrifuged at 4000 rpm for 5 mins at 4°C and the supernatant was transferred to a new 50 ml Falcon tube. Immuno-precipitation was performed by adding 5 µg of rabbit IgG-coupled Dynabeads or 18 mg of M2 αFLAG-coupled Dynabeads per 30 g lysed yeast, and the lysates were rotated at 4°C for 2 h. Lysates were then washed five times in cold buffer without inhibitors and then transferred to a 1.5 ml eppendorf tube with 1 ml buffer. Residual buffer was removed, 25 µl of 1x NuPAGE® LDS sample buffer was added, samples were boiled at 100°C for 5 mins before 5 µl of β-mercaptoethanol was added and samples were boiled for a further 5 mins, spun down at 13000 rpm for 5 mins and loaded onto a precast NuPAGE® 8–12% Bis-Tris gel (Novex). Bands were visualized after staining using the Pierce silver staining kit (Thermo Scientific).

Results

Spindle disassembly is only moderately delayed in cdc14-1 mutants

A hallmark of mitotic exit is spindle disassembly, an event that is critically dependent on Cdc14 in budding yeast mitosis (Stegmeier & Amon, 2004). Initial analysis of fixed temperature-sensitive cdc14-1 mutant cells undergoing meiosis at the restrictive temperature revealed an increased frequency of cells with long spindles characteristic of anaphase I, suggesting blocked spindle disassembly and impaired meiosis I exit (Marston et al., 2003). However, live-cell imaging revealed that meiosis I spindles frequently disassemble in cdc14-1 cells, only to reassemble at the presumptive time of meiosis II (Bizzari & Marston, 2011), suggesting that Cdc14 may be refractory for spindle disassembly and meiosis I exit.

To establish the importance of Cdc14 in spindle disassembly following meiosis I, we determined the time from anaphase I onset until spindle breakdown in live cells with impaired Cdc14 function. Securin (Pds1-tdTomato) degradation was used as a marker for anaphase I onset and the time taken for the meiosis I spindle (GFP-Tubulin) to completely disassemble after Pds1 proteolysis was measured in individual cells. In the wild type example (Figure 1A), spindle disassembly was observed 40 min after anaphase I onset, after which meiosis II spindles formed (note that Pds1-tdTomato is not visualised in meiosis II cells, presumably due to slow maturation of the fluorophore (Matos et al., 2008)). Spindle disassembly occurred 45.7 min after anaphase I, on average (Figure 1B) and was observed in 100% of wild type cells (Figure 1C). In ~82% cdc14-1 mutant cells, anaphase I spindles broke down and a new spindle did not assemble (Figures 1A and C), which is consistent with what we previously reported (Bizzari & Marston, 2011). We observed a modest, yet significant, increase (up to 52.2 min) in the time from anaphase onset to spindle disassembly in cdc14-1 cells (Figure 1B). In contrast, slk19Δ and spo12Δ cells, which retain Cdc14 in the nucleolus during meiosis I (Buonomo et al., 2000; Marston et al., 2003), disassembled anaphase I spindles with a timing comparable to wild type cells (43.8 and 41.9 min, respectively; Figures 1A and 1B). As reported previously for cdc14-1 mutants (Bizzari & Marston, 2011), spindle reassembly at the presumptive time of meiosis II was observed in a fraction of slk19Δ and spo12Δ cells, though the extent to which this occurred varied between the different mutants for reasons that are unclear. Taken together, these findings indicate that, while FEAR and Cdc14 appear to work together to ensure that two spindles are produced between the different mutants for reasons that are unclear. Taken together, these findings indicate that, while FEAR and Cdc14 appear to work together to ensure that two spindles are produced following meiosis I, suggesting blocked spindle disassembly and meiosis I exit. However, live-cell imaging revealed that meiosis I spindles frequently disassemble in cdc14-1 cells, only to reassemble at the presumptive time of meiosis II (Bizzari & Marston, 2011), suggesting that Cdc14 may be refractory for spindle disassembly and meiosis I exit.

Mass spectrometry

Protein bands were excised from Coomassie-stained NuPAGE® 8–12% Bis-Tris gels and washed alternatingly with 50 mM ammonium bicarbonate and acetonitrile solutions until Coomassie staining was removed. Gel pieces were treated with 10 mM DTT in 50 mM ammonium bicarbonate for 30 mins at 37°C, then DTT was removed and samples were washed with acetonitrile. A total of 55 mM iodoacetamide in 50 mM ammonium bicarbonate was added to the gel slices, and these were incubated at room temperature in the dark for 20 mins. After washing again with 50 mM ammonium bicarbonate and acetonitrile, gel pieces were incubated with trypsin for 15 mins on ice, and then samples were transferred to 37°C for overnight digestion. The following morning, digestion reactions were treated with 0.1% (w/v) trifluoroacetic acid and left for 15 mins to allow peptides to diffuse from the gel. Samples were then passed through an equilibrated StageTip consisting of two layers of Empore Disks C18 within a pipette tip (Rappsilber et al., 2007). A single StageTip was used per sample, as peptides within samples bind to StageTips. Peptides were later eluted for analysis via mass spectrometry (MS), performed as previously described (Sarangapani et al., 2014). MS data was compiled in MaxQuant v1.4.1.2 (http://www.coxdocs.org/doku.php?id=maxquant:start). Quantitative analysis was performed using Perseus v1.5.1.6 (http://www.coxdocs.org/doku.php?id=perseus:start). MaxQuant LFQ intensities were normalized to Spc42-3FLAG, filtered to remove contaminants, data was logarithmised to log2(x), then further filtered to only include proteins present in 5 out of the 6 groups (i.e. 2 wild type and 3 cdc14-1). The statistical test for significance was a t-test (FDR = 0.05, s0=1). No changes were found to reach significance.
Figure 1. Meiosis I spindle disassembly in the absence of functional Cdc14. Wild type (AM16065), cdc14-1 (AM16066), slk19Δ (AM16110) and spo12Δ (AM15985) cells carrying TUB1-GFP and PDS1-tdTomato were induced to sporulate, released from prophase I arrest and imaged at 10 min intervals for a total of 12 h in a microfluidics device. (A) Representative images are shown. The black arrow denotes Pds1 degradation, marking entry of cells into anaphase I. The white asterisk marks the time of spindle breakdown. (B) The time taken for complete spindle disassembly after Pds1 degradation was recorded for individual cells and plotted for a total of 67 wild type cells and 100 of each of cdc14-1, slk19Δ and spo12Δ cells. Mean rates of spindle breakdown are shown (red line), with error bars representing standard deviation. The two-tailed Student's t-test was used to calculate p values. (C) Inactivation of Cdc14 results in abnormal spindle behaviour. Cells were categorised based on spindle morphology as indicated in the legend.
cannot completely rule out retention of partial activity by the temperature sensitive Cdc14 proteins, together these findings suggest that Cdc14 is more critical for spindle disassembly in mitosis than meiosis.

Cdc14 associates with the SPB at the meiosis I to meiosis II transition

We took an unbiased approach to identify cellular processes targeted by Cdc14 to regulate the meiosis I to meiosis II transition. Following its release from the nucleus during anaphase I, Cdc14 is expected to associate with, and dephosphorylate, substrates that facilitate the transition to meiosis II. We reasoned that identification of Cdc14 interacting partners in both wild type cells and pCLB2-CDC55 cells, in which Cdc14 is ectopically released from the nucleus, would inform on the processes it regulates. Anti-FLAG immunoprecipitates from wild type and pCLB2-3HA-CDC55 cells harvested 4 h after induction of meiosis and carrying CDC14-3FLAG were analysed by mass spectrometry. Despite similar Cdc14 peptide counts in wild type and pCLB2-3HA-CDC55 samples, we observed a lower Cfi1/Net1 peptide count in the latter sample, consistent with premature release of Cdc14 from the nucleus in Cdc55-deficient cells (Figure 2A). Interestingly, however, the predominant class of proteins identified in both samples were components of the yeast centrosome/spindle pole body (SPB) (Figures 2A and B).

To determine the timing of Cdc14 association with the SPB during meiosis, we imaged live cells carrying CDC14-GFP and the SPB marker, SPC42-tdTomato undergoing meiosis. As previously reported, in wild type cells, Cdc14 is sequestered in the nucleolus throughout prophase I and metaphase I of meiosis and, accordingly, we did not observe co-localization with SPBs at these stages ((Buonomo et al., 2003; Marston et al., 2003; Matos et al., 2008); Figure 3A). During anaphase I, however, concomitant with its release from the nucleolus, Cdc14-GFP was detected at the SPB (Figure 3A, arrows). To confirm the timing of Cdc14 association with the SPB we determined the ratio of intensity of Cdc14-GFP and Spc42-tdTomato fluorescence (Figure 3B). This revealed the strongest association of Cdc14-GFP with the SPB in anaphase I, with a weaker association in anaphase II (Figure 3B). Interestingly, Cdc14-GFP localized asymmetrically, generally associating with just one of the two SPBs in anaphase I, or two of the four SPBs during anaphase II, with no detectable SPB association during metaphase I or metaphase II (Figure 3C).

We, and others, previously showed that ectopic release of Cdc14 prevents nuclear division in pCLB2-3HA-CDC55 cells (Bizzari & Marston, 2011; Kerr et al., 2011). To determine whether premature association of Cdc14 with the SPB could underlie this phenotype, we induced pCLB2-3HA-CDC55 cells carrying CDC14-GFP and SPC42-tdTomato to undergo meiosis and categorised cells based on the localization of Cdc14-GFP: nucleolar sequestration (class 1); partial release (class 2) or complete release (class 3) (Figures 3D and E). Cdc14-GFP was detected at the SPB in virtually all pCLB2-3HA-CDC55 cells where Cdc14-GFP was either completely or partially released from the nucleolus (Figure 3E). Note that the vast majority of pCLB2-3HA-CDC55 cells contain only a single Spc42-tdTomato foci, therefore it was not possible to address whether Cdc14-GFP remains asymmetric in these cells.

We sought to identify factors that are required for Cdc14 localization to SPBs during anaphase I. During mitosis, components of the MEN are associated with the SPB. However, the MEN is dispensable for the meiosis I to meiosis II transition and the majority of its components are not found at SPBs (Attner & Amon, 2012; Kamieniecki et al., 2005; Pablo-Hernando et al., 2007). An exception is the two component GAP, Bub2/Bfa1 which localizes symmetrically at SPBs during metaphase I, anaphase I but asymmetrically during metaphase II (Attner & Amon, 2012; Figures 4A and B). We found that Cdc14-GFP association with SPBs was abolished in both bub2Δ and bfa1Δ anaphase I cells (Figures 4C and D). We conclude that upon release from the nucleolus in anaphase I and anaphase II, Cdc14 associates asymmetrically at SPBs in a manner dependent on Bub2/Bfa1.

Generation of 4 Spc42-tdTomato foci during meiosis II depends on Cdc14

Recently, Cdc14 has been identified as a licensing factor that enables SPB duplication upon exit from mitosis (Avena et al., 2014; Elserafy et al., 2014). Taken together with our findings above, this suggests that a major function of Cdc14 during meiosis could be to license a second round of SPB duplication, thereby enabling the assembly of two spindles in meiosis II. To test this idea, we monitored SPB number in wild type cells where Cdc14 was inactivated (cdc14-1) by scoring Spc42-tdTomato foci as cells progressed synchronously through meiosis after release from a prophase I block (Figures 5A and B). As expected, wild type cells produced two, then four Spc42-tdTomato foci concomitant with the appearance of binucleate and tetranucleate cells (Figure 5A). In contrast, and consistent with the observed single nuclear division, cdc14-1 cells produced a maximum of two Spc42-tdTomato foci (Figure 5B), as did cells lacking the two FEAR activators, Spo12 and Silk19 (Figures 5C and D). Upon deletion of Cdc55, a single Spc42-tdTomato focus was observed in the majority of cells (Figure 5E), indicating a failure in the first round of SPB duplication or separation. Furthermore, this lack of SPB duplication/separation prior to meiosis I in Cdc55-depleted cells was a consequence of ectopic Cdc14 activation, since cdc14-1 pCLB2-3HA-CDC55 cells produced two Spc42-tdTomato foci, similar to cdc14-1 single mutant cells (Figure 5F). Consistent with a requirement for Cdc14 at SPBs, a large fraction of bub2Δ and bfa1Δ cells completed only a single meiotic division with only 2 Spc42-tdTomato foci (Figures 6A–C). During mitosis, Bub2/Bfa1 is asymmetrically localized on SPBs, but this asymmetry is broken in response to defective spindle positioning, in a manner dependent on Kin4 and Bmh1 (Gryaznova et al., 2016; Maekawa et al., 2007; Monje-Casas & Amon, 2009). We found, however that Kin4 and Bmh1 are dispensable for either the asymmetric SPB localization of Cdc14 (Figure 4D) or the execution of two meiotic divisions (Figures 6E–F), at least under normal conditions. These findings indicate that localization of Cdc14 at the SPB is important for the successful execution of the meiosis I to meiosis II transition.

Cdc14 is essential for SPB duplication at the meiosis I to meiosis II transition

To determine whether cdc14-1 and pCLB2-3HA-CDC55 mutants are defective in SPB duplication or separation, we initially used
Figure 2. The centrosome/SPB co-purifies with Cdc14 during meiosis. (A) Wild type (AM9434) and pCLB2-HA-CDC55 (AM9319) cells carrying CDC14-SZZ were harvested 4 h after induction of sporulation. Cdc14-SZZ-associated complexes, purified on IgG-coupled beads, were analysed by mass spectrometry. The number of identified peptides of the indicated proteins is given. See also https://osf.io/g5cmh/ (Marston, 2016). (B) Schematic diagram of the budding yeast spindle pole body.
Figure 3. Cdc14 localizes asymmetrically at SPBs during anaphase I. (A) Sporulating CDC14-GFP SPC42-tdTomato cells (AM11517) were imaged in a microfluidics chamber at 15 min intervals for a total of 12 h. Example of a cell in which Cdc14-GFP localizes asymmetrically to one SPB during anaphase I (arrows). (B and C) Wild type cells as in (A) were released from a prophase I arrest and imaged at 30 min intervals on agarose pads. (B) The ratio of Cdc14-GFP/Spc42-tdTomato signal per SPB is shown with error bars representing standard error. Cells were classified into different cell cycle stages (metaphase I, anaphase I, metaphase II, anaphase II) by scoring the number of SPBs, the distance between them and Cdc14 nucleolar sequestration. The two-tailed Student's t-test was used to calculate p values (* p<0.001) n=50 cells. (C) Co-localization of Cdc14-GFP with SPBs was scored in the indicated cell cycle stages. (D and E) Cdc14 localises prematurely to SPBs in the absence of Cdc55. pCLB2-3HA-CDC55 cells carrying CDC14-GFP and SPC42-tdTomato (AM13123) were induced to sporulate, released from pGAL-NDT80 block and imaged 3h later on agarose pads. Cdc14 localisation was classified into three categories: sequestered in nucleolus (1), partially released from nucleolus (2) and completely released from nucleolus (3). (D) Example images of with numbered arrows showing examples of each category. (E) Co-localisation of Cdc14 with SPBs was scored in 100 cells of each category.
Figure 4. Asymmetric localisation of Cdc14 to SPBs is Bfa1/Bub2-dependent. (A and B) Bfa1 localises symmetrically to SPBs during meiosis I. Wild type (AM17740) cells containing SPC42-CFP, BFA1-tdTomato and CDC14-GFP were induced to sporulate, released from pGAL-NDT80 block and imaged at 30 minute intervals on agarose pads. Cells were classified into different meiotic stages based on number of SPB foci, distance between SPBs and Cdc14 nucleolar sequestration. (A) Representative images of Bfa1 localisation. White arrows indicate Bfa1-tdTomato co-localization with Spc42-CFP foci. Grey arrowheads indicate Spc42-CFP foci where Bfa1-tdTomato is absent. (B) Co-localisation of Bfa1 with SPBs was scored throughout meiosis in 200 cells at each stage. Note that Spc42-CFP bleeds through to the GFP emissions channel so Cdc14-GFP signal is not shown, though its nucleolar sequestration and release was used to classify meiotic stages. (C and D). Wild type (AM11517), bfa1Δ (AM16079), bub2Δ (AM15543), kin4Δ (AM17134) and bmh1Δ (AM17341) cells containing CDC14-GFP and SPC42-tdTomato were induced to sporulate, released from a pGAL-NDT80 block and imaged at 30 minute intervals on agarose pads. (C) Co-localisation of Cdc14 with SPBs was scored in 100 anaphase I cells. (D) The ratio of Cdc14-GFP/Spc42-tdTomato signal per SPB per cell was quantified, with error bars representing standard error. Cells were classified as anaphase I based on distance between SPB foci and Cdc14 release. The two-tailed Student’s t-test was used to calculate significance. ** indicates p<0.001. n = 100 foci (50 cells).
Figure 5. Active Cdc14 is required for cells to form four distinct Spc42-tdTomato foci in meiosis. Wild type (AM13989; A), cdc14-1 (AM16163; B), cdc55mn (AM15984; C), slk19Δ (AM16020; D) spo12Δ (AM16064; E) and cdc14-1 cdc55mn (AM17904; F) cells carrying SPC42-tdTomato were induced to sporulate and released from a pGAL-NDT80 block. The percentages of binucleate and tetranucleate cells (upper graph) and the number of Spc42-tdTomato per cell were scored in 200 cells at each timepoint (lower graph).
Figure 6. Bfa1 and Bub2 are required for timely re-duplication of SPBs in meiosis. (A–C) Wild type (AM13989; A), bfa1Δ (AM16019; B) and bub2Δ (AM16080; C) cells containing SPC42-tdTomato were induced to sporulate and released from pGAL-NDT80 block. The percentages of binucleate and tetranucleate cells (upper graph) and the number of Spc42-tdTomato per cell were scored in 200 cells at each timepoint (lower graph). (D–F) Wild type (AM13989; D), kin4Δ (AM17134; E) and bmh1Δ (AM17341; F) cells containing CDC14-GFP and SPC42-tdTomato were induced to sporulate, released from a pGAL-NDT80 block and imaged at 30 min intervals on agarose pads. The percentages of binucleate and tetranucleate cells (upper graph) and the number of Spc42-tdTomato per cell were scored in 200 cells at each timepoint (lower graph).
intensity of Spc42-tdTomato foci did not increase, even though cells produced 2 Spc42-tdTomato foci (x1.07) (Figure 7D), suggesting that SPB re-duplication failed to occur prior to SPB separation. The single Spc42-tdTomato focus of pCLB2-3HA-CDC55 cells also did not increase in Spc42-tdTomato intensity throughout the timecourse (x0.98), suggesting that both SPB separation and re-duplication fail to occur upon Cdc55 depletion.

To gain further insight into how Cdc14 influences SPB morphogenesis during meiosis, we compared the composition of the SPB in wild type and cdc14-1 cells by quantitative mass spectrometry. SPBs were purified from wild type and cdc14-1 cells undergoing meiosis and carrying SPC42-3FLAG by immuno precipitation using anti-FLAG antibodies (Figure 7F). Comparison of relative peptide intensities for three biological replicates (Figure 7G).

Figure 7. Quantitative analysis of SPBs in meiosis indicates a requirement for Cdc14 in SPB duplication. (A) Wild type (AM13989) cells containing SPC42-tdTomato were induced to sporulate, released from pGAL-NDT80 block and imaged at 15 minute intervals for a total of 12 hours in a microfluidics chamber. Representative images of wild type cells are shown. (B) SPBs are duplicated at meiotic prophase. Wild type cdc14-1 (AM16163), and pCLB2-3HA-CDC55 (AM15984) cells containing SPC42-tdTomato and pGAL-NDT80 and were resuspended in sporulation medium in the absence of β-oestradiol and imaged immediately (t=0) or after 3 and 6h as they progress into the prophase I arrest due to the absence of Ndt80. Individual SPB foci were quantified and mean total SPB fluorescence intensity (F) per cell was plotted, with error bars (obscured by the markers) representing standard error, n=100 per timepoint. (C–E) SPB fluorescence was quantified as in B from movies of live cells after release from the prophase I arrest by addition of β-oestradiol as in (A). The first time point at which a cell contained 2 Spc42-tdTomato foci was defined as 1h and SPB fluorescence in the preceding 4 (1 focus) and following 4 time points was quantified (n=10 cells). Note that at the 1.5h time point 7 wild type cells carried 2 Spc42-tdTomato foci, while 3 wild type cells carried 4 Spc42-tdTomato foci. (F and G) LFQ Proteomic analysis of SPB composition and environment in wild type and cdc14-1 cells. Strains used were AM1835 (no tag), AM11444 (SPC42-3FLAG), AM9459 (cdc14-1) and AM11443 (cdc14-1 SPC42-3FLAG). After 4 h, cells were harvested and SPBs purified by anti-FLAG immunoprecipitation. Peptides were generated by in-gel trypsin digestion and LC-MS data sets for 3 biological replicas was analysed using MaxQuant software. (F) Example silver stained SDS-PAGE showing Spc42-3FLAG immunoprecipitates of one of three biological replicas used in LFQ proteomic analysis. BSA standards were used to estimate protein concentration. (G) Statistical analysis of relative LFQ intensity output was carried out using Perseus. Volcano plot shows –log of P values versus ratio of wild type/cdc14-1 for all 254 proteins in >5 columns. No significant change in composition and environment was observed between wild type and cdc14-1 SPBs. (FDR = 0.05, s0 = 1). Proteins of interest are highlighted as follows: red = SPB components; blue = Cdc14; green = Bfa1/Bub2. See also https://osf.io/g5cmh/ (Marston, 2016).
indicated significant depletion of Cdc14 on SPBs from *cdc14-1* cells, consistent with the idea that the mutant protein fails to associate with the SPBs. Though not reaching the stringent cut off for statistical significance (FDR=0.05), we further noticed that components of the SPB half bridge tended to be depleted on *cdc14-1* SPBs. Interestingly, the half bridge component, Sfi1, which was recently confirmed as a Cdc14 target in mitosis (Avena *et al*., 2014; Elserafy *et al*., 2014) showed the greatest change in abundance. These results are consistent with the idea that a major function of Cdc14 at the meiosis I to meiosis II transition is to enable half bridge extension, thereby allowing SPB re-duplication.

To examine SPB morphogenesis more directly, we analyzed *cdc14-1* and *pCLB2-3HA-CDC55* meiotic cells by electron microscopy. As predicted by the quantitative fluorescence microscopy, SPB re-duplication was not observed in *cdc14-1* mutants (n=8) and cells arrested with two unduplicated SPBs. We observed late meiosis II events in 3/8 cells. In the example shown (Figure 8A) a long spindle connects two unduplicated SPBs. Assembly of the outer plaque and vesicles are apparent at one of the SPBs (SPB 1, white arrow), indicating that the cell is in a late stage of meiosis II, though outer plaque formation has not been initiated at the other SPB (SPB 2, white arrow). In *pCLB2-3HA-CDC55* cells, also as predicted from our quantitative fluorescence microscopy, two side-by-side SPBs connected by a half bridge were invariably observed (n=6; Figure 8B). However, we found no evidence of over-duplication of SPBs in *pCLB2-3HA-CDC55* cells. This suggests that Cdc14 must be held inactive during early meiosis to allow SPB separation and acts in a licensing step, rather than as an assembly factor.

**Discussion**

The existence of two consecutive rounds of chromosome segregation without an intervening S phase is a characteristic feature of meiosis that underlies sexual reproduction. Unique, yet poorly understood, controls allow a second round of spindle formation, but prevent a second round of DNA replication. Our results implicate Cdc14 regulation as being central to this distinction. In mitosis, following chromosome segregation, MEN-dependent release of Cdc14 triggers CDK inactivation permitting both the re-licensing of SPBs and DNA replication origins. In contrast, following meiosis I, MEN is not active (Attner & Amon, 2012) and Cdc14 release is under the control of only the FEAR network (Buonomo *et al*., 2003; Kamieniecki *et al*., 2005; Marston *et al*., 2003), which is incapable of triggering mitotic exit (Yellman & Roeder, 2015). Here we show that FEAR-dependent Cdc14 is critical to initiate SPB duplication, thereby enabling assembly of two separate meiosis II spindles. We show that Cdc14 released by the FEAR network associates with the SPB in a Bub2/Bfa1-dependent manner and provide evidence that SPB-localized Cdc14 is critical to trigger SPB duplication. Based on recent findings in mitotic cells (Avena *et al*., 2014; Elserafy *et al*., 2014), we suggest that Cdc14 re-licenses SPBs through dephosphorylation of half-bridge components, in particular Sfi1 during anaphase I (Figure 9). Overall, our findings show that Cdc14 is required to re-license SPB duplication.

![Figure 8. cdc14-1 and pCLB2-3HA-CDC55 mutants arrest with unduplicated and unseparated SPBs, respectively.](https://example.com/figure8.png)
between meiosis I and meiosis II and that its retention in the nucleolus during early meiosis is required to allow SPB separation during meiosis I.

**SPB localization of Cdc14**

Bub2/Bfa1-dependent association of Cdc14 with the SPB is not unique to meiosis, indeed it has been observed in early mitosis (Pereira et al., 2002a; Yoshida et al., 2002), suggesting that it is a FEAR-triggered event here too. SPB-localized Cdc14 has been implicated both in MEN activation in early anaphase and, through Bfa1 dephosphorylation, in MEN inactivation in late anaphase (Pereira et al., 2002b). An attractive possibility, which remains to be tested, is that in meiosis I, Cdc14 at the SPB acts to maintain Bfa1/Bub2 in the dephosphorylated state, thereby preventing MEN activation. Our data also suggest another function for SPB-localized Cdc14 during meiosis I, to trigger SPB duplication.

Curiously, we observed Cdc14 at a single SPB during anaphase I. Asymmetric localization of Cdc14 and MEN components is observed in budding yeast mitosis, where the requirement to partition the nucleus through the bud neck imposes an intrinsic polarity on cell division. The asymmetric localization of MEN components, both on the SPB and within the bud contributes to the spindle position checkpoint which prevents mitotic exit in response to spindle alignment defects (Caydasi & Pereira, 2012). However, during meiosis I, MEN is not active (Attner & Amon, 2012) and meiotic yeast cells are not obviously polarized. Furthermore, Cdc14 activity is presumably required at both SPBs during anaphase I to trigger their duplication, thereby ensuring production of a pair of spindles in meiosis II. While Cdc14 is detectable by microscopy only on 1 SPB during anaphase I, we speculate that undetectable levels of Cdc14 on the other SPB are sufficient to trigger SPB duplication. This however, raises the question of how and why Cdc14 is more concentrated on a single SPB, particularly considering that Bub2/Bfa1 is itself symmetrically localized. The origin and significance of the asymmetric localization of Cdc14 at the SPB during anaphase I therefore remain unexplained.

**Control of the cell cycle at meiosis I exit**

Although Cdc14 is essential for mitotic exit, accumulating evidence suggests that Cdc14 plays a lesser role in CDK down-regulation at meiosis I exit. We found that spindle assembly is only slightly delayed in cells with impaired Cdc14 activity and cyclin destruction appears to occur on schedule in cdc14Δ cells (Bizzari & Marston, 2011; Kerr et al., 2011; Tibbles et al., 2013). Instead it is likely that cyclin degradation upon APC\(^{Cdc20}\) activation at anaphase I onset initiates meiosis I exit. Understanding how this is regulated to ensure step-by-step release of cohesion, spindle elongation and spindle disassembly at meiosis I is an important priority for the future.

**Data availability**

Source data for mass spectrometry results from Figures 2A (Fox_Data 1) and 7G (Fox_Data 2), example movies of spindle elongation (Figure 1), additional examples of electron
microscopy images, source data files for quantification of SPB fluorescence (Figure 7) and source data files for quantification of Cdc14-GFP localization at SPBs (Figure 3 and Figure 4) are available at: https://osf.io/g5cmh/ (doi: 10.17605/OSF.IO/G5CMH; Marston, 2016).

Author contributions
CF and AM conceived the study and designed the experiments. CF carried out the research. JZ and JR analysed mass spectrometry data. CF and AM prepared the manuscript. All authors have approved the final content.

Competing interests
No competing interests were disclosed.

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✔ Simonetta Piatti
Centre de Recherche en Biologie Cellulaire de Montpellier (CRBM), Montpellier, France

Authors have addressed all my comments and suggestions. Altogether it is a nice piece of work, carefully designed and executed.

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Reviewer Report 24 February 2017
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✔ Soni Lacefield
Department of Biology, Indiana University Bloomington, Bloomington, IN, USA

The authors have addressed my concerns and made appropriate changes to their manuscript in their revision.

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
The centrosome is a fascinating organelle; it serves as a scaffolding platform for numerous cell cycle regulators in addition to the major microtubule organising centre, which plays an essential role in chromosome segregation by acting as the spindle pole. During the mitotic cell cycle, a rigorous licensing mechanism that involves phosphorylation and dephosphorylation of the centrosome components restricts the number of centrosome duplications to exactly one per cell cycle. Physiological importance of this regulation is underlined by the fact that tumour cells are often associated with abnormally amplified centrosomes. The timing of the centrosome duplication in many organisms coincides with S phase, where DNA replication takes place. Common cell cycle regulators, such as CDKs, trigger both centrosome duplication and DNA replication events.

In meiosis, however, centrosome duplication must be uncoupled from DNA replication, because, no DNA replication must occur at the transition from meiosis I (MI) to meiosis II (MII). Although the mechanism of uncoupling of the centrosomal and DNA replication cycles in meiosis is mostly unclear, studies in *Xenopus* oocytes and the fission yeast found significant residual CDK activities at MI - MII transition, suggesting that these CDK activities may inhibit DNA replication, hence, decoupling the centrosome cycle from the DNA replication cycle. However, further supporting evidence for it or the molecular basis that directs the meiotic centrosome cycle has still been missing.

In this manuscript, Fox et al. used the budding yeast as a model and showed that Cdc14 phosphatase is one of the essential regulators of the meiotic centrosome duplication. By exploiting a conditional *cdc14-1* mutant, an absolute requirement of Cdc14 function in meiotic centrosome duplication was clearly demonstrated through extensive live and fixed cell imaging of the cells undergoing MI-MII synchronously. Observation using electron microscopy provided further supporting evidence.

The manuscript is very well written, the quality of the data (which are mostly very clearly presented) is high and sound data interpretations have been provided. Therefore, I would like to approve this study. A few minor points are listed below to help further clarify the authors’ message.

- **Culturing condition to inactivate Cdc14:**
  
  It was not too clear which temperature (and how long) was used to inactivate Cdc14 in the *cdc14-1* ts mutant. Or, does *cdc14-1* mutant show its meiotic phenotype at an intermediate temperature 30°C? It would be really helpful if the authors would clarify this point.
Fig. 3D and 3E:

In these figures, the Cdc14-GFP signal at the SPB is presented. If I understood correctly, in the pCLB2-3HA-CDC55 setting, SPB duplication does not occur and only one SPB can be found in each cell. If this would be the case, it would be helpful to clearly state so, thus, "asymmetric Cdc14 localisation" is not relevant in this experimental setting.

Page 8 final paragraph: “…but symmetrically during metaphase II” -> asymmetrically

Fig. 4A:

Bfa1-tdTomato localization is presented.

In the “Metaphase II” panel, it would be helpful to have additional small arrowheads in order to indicate SPBs (Spc42-CFP signal-positive dots) without Bfa1-tdTomato signal, so that asymmetric Bfa1-tdTomato localization is easily recognisable.

Page 12, 2nd paragraph:

It would be helpful to dub the SPB as “Spc42-tdTomato foci” in this paragraph. For example, instead of saying “two SPB” or “four SPB”, say “two Spc42-tdTomato foci” or “Four Spc42-tdTomato foci”. This is because until we see Fig. 7 and 8, we do not know whether a “Spc42-tdTomato focus” represents a non-duplicated SPB (I would call this one SPB) or non-separated SPBs (I would call this two SPBs).

Figure 7, figure legends:

Legends corresponding to panels (A-E) contain some inconsistencies, i.e., legends to panel (B) and (E) are missing, and C, D, and E are wrongly assigned.

In addition, would it be possible to indicate the size/band that corresponds to the Cdc14 protein in the gel in panel F?

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**Competing Interests:** No competing interests were disclosed.
I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 11 Feb 2017

Adele Marston, School of Biological Sciences, Edinburgh, UK

Thank you for your approval of our article, you kind comments and your constructive suggestions. Here we respond to the points made:

- **Culturing condition to inactivate Cdc14:**

  It was not too clear which temperature (and how long) was used to inactivate Cdc14 in the cdc14-1 ts mutant. Or, does cdc14-1 mutant show its meiotic phenotype at an intermediate temperature 30°C? It would be really helpful if the authors would clarify this point.

  *We used 30C, which we believe is efficient at largely inactivating Cdc14 (see comments to other reviewers) as vegetative cells fail to grow at this temperature. We have added this information to the methods.*

- **Fig. 3D and 3E:**

  In these figures, the Cdc14-GFP signal at the SPB is presented. If I understood correctly, in the pCLB2-3HA-CDC55 setting, SPB duplication does not occur and only one SPB can be found in each cell. If this would be the case, it would be helpful to clearly state so, thus, “asymmetric Cdc14 localisation” is not relevant in this experimental setting.

  *We added a statement to this effect in this results section*

  - Page 8 final paragraph: “…but symmetrically during metaphase II” -> asymmetrically

  *This has been corrected.*

- **Fig. 4A:**

  Bfa1-tdTomato localization is presented.

  In the “Metaphase II” panel, it would be helpful to have additional small arrowheads in order to indicate SPBs (Spc42-CFP signal-positive dots) without Bfa1-tdTomato signal, so that asymmetric Bfa1-tdTomato localization is easily recognisable.

  *Page 12, 2nd paragraph:*

  It would be helpful to dub the SPB as “Spc42-tdTomato foci” in this paragraph. For example, instead of saying “two SPB” or “four SPB”, say “two Spc42-tdTomato foci” or “Four Spc42-tdTomato foci”. This is because until we see Fig. 7 and 8, we do not know whether a “Spc42-tdTomato focus” represents a non-duplicated SPB (I would call this one SPB) or non-separated SPBs (I would call this two SPBs).

  *This is a very good point. We have changed the text and figure accordingly.*
Figure 7, figure legends:

Legends corresponding to panels (A-E) contain some inconsistencies, i.e., legends to panel (B) and (E) are missing, and C, D, and E are wrongly assigned. This has been corrected.

In addition, would it be possible to indicate the size/band that corresponds to the Cdc14 protein in the gel in panel F?

There is no obvious band corresponding to Cdc14, a 61 kd protein and as we are unable to confirm that any particular band is Cdc14 we prefer not to annotate this on the gel.

Competing Interests: None

Reviewer Report 19 January 2017

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Simonetta Piatti
Centre de Recherche en Biologie Cellulaire de Montpellier (CRBM), Montpellier, France

This is an interesting study addressing an important question, i.e. how yeast cells license SPB duplication at the MI/MII transition to be able to form bipolar spindles and support chromosome segregation during reductional division.

Previous work had shown that the Cdc14 phosphatase must be properly controlled to permit assembly of two MII spindles through an unknown mechanism. In addition, the SPB half-bridge protein Sfi1 has been recently identified as a critical target of Cdc14 in mitosis in order to license SPB duplication.

In this manuscript the authors show that Cdc14 associates with SPBs also in meiosis. Cdc14 localization at SPBs is asymmetric and requires the Bub2/Bfa1 complex. Cdc14, while not being strictly required for exit from MI (i.e. spindle disassembly), is required for SPB duplication and proper nuclear division in MII. Conversely, premature Cdc14 activation in cells lacking Cdc55 prevents SPB separation in MI. Finally, although SPBs do not duplicate upon Cdc14 inactivation, composition of the SPB remains mostly unchanged.

Overall, experiments are carefully executed and conclusions are mostly supported by the experimental data. I have, however, a couple of main issues and a few minor points that the authors might want to address.
Main points:

1. The incubation of *cdc14-1* mutant cells at 30°C, which might be a semi-permissive temperature, raises the question as to what extent Cdc14 is actually inactivated in these cells. Wouldn’t it have been better to use a tighter system to inactivate Cdc14 (e.g. expressing *CDC14* from the *CLB2* promoter)? It is even possible that the Cdc14-1 protein is prematurely released, thereby tempering the effects of temperature inactivation. Has nucleolar release of the Cdc14-1 protein ever been checked at 25°C/30°C?

This would also be relevant to interpret correctly the data in Fig. 1B and C, where *cdc14-1* cells do not behave exactly like FEAR mutants.

2. Quantification of SPB signals during meiotic progression (Fig. 7A-E) is an essential piece of data to support the conclusion that *cdc14-1* cells are defective in SPB duplication upon entry into MII. I wonder if a slow-folding fluorescent protein, such as Spc42-tdTomato, is actually a suitable marker for this kind of analyses.

A related issue concerns the way these data are presented, which I find a bit confusing. For instance, I do not understand why there are gaps in the plot of wild type cells at 0.75-1h and 1.5-1.75h (Fig. 7C), since fluorescence intensities are measured on movies.

Direct inspection of SPBs by EM, like the authors did, should greatly help understanding what is going on in the *cdc14-1* mutant. However, knowing how many MII cells display the phenotype shown in Fig. 8A is an essential piece of information that is currently missing. Another interesting information is whether SPB over-duplication was ever found in pCLB2-CDC55 cells.

Finally, if it were possible to inactivate Cdc14 before commitment to meiosis, would the authors expect a defect in SPB duplication also in MI? The likely possibility that Cdc14 is a crucial licensing factor in ALL kinds of divisions should perhaps be mentioned in the discussion.

3. Since the role of Cdc14 in licensing SPB duplication in mitosis has been linked to Sfi1 dephosphorylation, it would have been nice to check the phosphorylation state of Sfi1 in wild type and *cdc14-1* cells during meiosis. This kind of information might even be extracted from the mass spec data from Fig. 7F.

Along the same line, it would have been interesting to check if the published unphosphorylatable *sfi1* mutant can suppress the SPB duplication defect of *cdc14-1* cells. This would have offered strong experimental support to the model presented in Fig. 9.

Minor points

1. It is unclear why in the movies of *cdc14-1* cells (Fig. 1) spindles do not reassemble in the second meiotic division from the two unduplicated SPBs (like for example in Fig. 8A).

The abnormal spindle behaviour reported in Fig. 1C is not commented in the text. Surprisingly, *cdc14-1* have a milder phenotype than FEAR mutants in this respect, although *cdc14-1* has a more pronounced defect in spindle disassembly.
2. It is not clear why Spo21 is in the list of Tap-Cdc14 interacting proteins with 0 peptides identified.

3. Cdc14 is reported to localize at SPBs in most \( pCLB2\)-CDC55 cells where Cdc14 is partially or completely released from nucleolus. Yet, in the example cell #2 of Fig. 3D Cdc14 is not visible on the SPB. A better representative image could be selected.

4. Graphs in Fig. 5 and Fig. 6 look pixeled.

5. In Fig. 7C there is a label within the graph that belongs to the IPs underneath. Also, the graphs (C, D, E) are mislabeled relative to the legend.

6. In Fig. 7F two prominent bands appear in the Spc42-Flag IP from wt cells, while one of the two is much decreased in \( cdc14-1 \) cells. Which one of the two is Spc42?

**Competing Interests:** No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

**Author Response 11 Feb 2017**

**Adele Marston**, School of Biological Sciences, Edinburgh, UK

Thank you for your approval in principle of our article and you helpful suggestions, here we have addressed your reservations:

**Main points:**

1. The incubation of \( cdc14-1 \) mutant cells at 30°C, which might be a semi-permissive temperature, raises the question as to what extent Cdc14 is actually inactivated in these cells. Wouldn’t it have been better to use a tighter system to inactivate Cdc14 (e.g. expressing \( CDC14 \) from the \( CLB2 \) promoter)? It is even possible that the Cdc14-1 protein is prematurely released, thereby tempering the effects of temperature inactivation. Has nucleolar release of the Cdc14-1 protein ever been checked at 25°C/30°C?

This would also be relevant to interpret correctly the data in Fig. 1B and C, where \( cdc14-1 \) cells do not behave exactly like FEAR mutants.

*Please see response to comment by Soni Lacefield. We agree we cannot completely rule out partial inactivation of Cdc14.*

1. Quantification of SPB signals during meiotic progression (Fig. 7A-E) is an essential piece of data to support the conclusion that \( cdc14-1 \) cells are defective in SPB duplication upon entry into MII. I wonder if a slow-folding fluorescent protein, such as Spc42-tdTomato, is actually a suitable marker for this kind of analyses.

*Because of these and other concerns (see response to Soni Lacefield) we used EM as a completely independent way to examine SPB duplication in \( cdc14-1 \) mutants. We believe the EM data provides convincing evidence for our conclusions, while the fluorescence intensity measurements provide some supporting data.*
A related issue concerns the way these data are presented, which I find a bit confusing. For instance, I do not understand why there are gaps in the plot of wild type cells at 0.75-1h and 1.5-1.75h (Fig. 7C), since fluorescence intensities are measured on movies.

The data for individual cells was lined up so that the 1h timepoint is defined as the point at which 2 SPBs are first observed. The different coloured lines represent 1, 2 or 4 SPBs. We have clarified this point in the figure legend.

Direct inspection of SPBs by EM, like the authors did, should greatly help understanding what is going on in the cdc14-1 mutant. However, knowing how many MII cells display the phenotype shown in Fig. 8A is an essential piece of information that is currently missing. Another interesting information is whether SPB over-duplication was ever found in pCLB2-CDC55 cells.

Overall we observed SPBs in 8 cdc14-1 cells by EM. Out of these, evidence of late meiosis II events (i.e. spore formation/vesicles) was observed in 3 of these. The image shown in Figure 8A is one example of two in which a spindle could be followed between 2SPBs. The other example was spread over >8 sections (spore formation undetermined) so the image shown was favoured. The third example showed the spore wall forming around a nucleus so SPBs were not connected by spindles in that case.

No evidence of overduplication of SPBs was observed in pCLB2-CDC55 cells (n=6).

We have included this information in the relevant section in the paper. All EM images are available here: https://osf.io/g5cmh/ (Marston, 2016).

Finally, if it were possible to inactivate Cdc14 before commitment to meiosis, would the authors expect a defect in SPB duplication also in MI? The likely possibility that Cdc14 is a crucial licensing factor in ALL kinds of divisions should perhaps be mentioned in the discussion.

The licensing event that enables SPB duplication during meiosis I would be expected to occur in the last stages of vegetative growth i.e. at mitotic exit prior to commitment to meiosis. So yes, it would be required in the same way that Cdc14 is required to license SPBs during vegetative growth.

2. Since the role of Cdc14 in licensing SPB duplication in mitosis has been linked to Sfi1 dephosphorylation, it would have been nice to check the phosphorylation state of Sfi1 in wild type and cdc14-1 cells during meiosis. This kind of information might even be extracted from the mass spec data from Fig. 7F.

Unfortunately we did not detect Sfi1 phosphorylation in our quantitative mass spec analysis. We attempted to examine Sfi1 phosphorylation on western blots and phostag gels but since the time period where Sfi1 dephosphorylation is expected to be observed (anaphase I) is very short, we were unable to obtain sufficient time resolution to make firm conclusions about the effect of Cdc14 inactivation of Sfi1 phosphorylation status.

3. Along the same line, it would have been interesting to check if the published unphosphorylatable sfi1 mutant can suppress the SPB duplication defect of cdc14-1 cells. This
would have offered strong experimental support to the model presented in Fig. 9.

We agree that this would be a very interesting experiment but despite extensive efforts we were unable to generate yeast strains carrying the Sfi1 phosphonull mutants and were therefore sadly unable to do this experiment. Diploid cells carrying the mutations as the sole copy of Sfi1 were extremely sick and would not enter meiosis. Other approaches were also attempted, for example expression of Sfi1 only in meiosis but due to both biological and technical reasons we have so far been unable to address this important point.

Minor points

1. It is unclear why in the movies of cdc14-1 cells (Fig. 1) spindles do not reassemble in the second meiotic division from the two unduplicated SPBs (like for example in Fig. 8A). For reasons we do not understand, spindles reassemble in a fraction of cdc14-1 cells, depending on the growth condition. In live cell imaging (Fig. 1), about 10% of cells were observed to reassemble spindles, however, the fraction is likely to be higher in flasks (Fig. 8A).

The abnormal spindle behaviour reported in Fig. 1C is not commented in the text. Surprisingly, cdc14-1 have a milder phenotype than FEAR mutants in this respect, although cdc14-1 has a more pronounced defect in spindle disassembly.

We have included a sentence raising this point in the results section.

2. It is not clear why Spo21 is in the list of Tap-Cdc14 interacting proteins with 0 peptides identified.

For completeness, all SPB proteins were included in the list, regardless of whether they were identified in the mass spectrometer.

3. Cdc14 is reported to localize at SPBs in most pCLB2-CDC55 cells where Cdc14 is partially or completely released from nucleolus. Yet, in the example cell #2 of Fig. 3D Cdc14 is not visible on the SPB. A better representative image could be selected.

Cell #3 shows a cell where Cdc14 is on the SPB. Cell #2 is an example of where the diffuse released Cdc14 signal overlaps with the SPB. As these are the criteria that were scored, these examples are appropriate.

4. Graphs in Fig. 5 and Fig. 6 look pixeled.

We have uploaded new versions of these figures that should correct this problem.

5. In Fig. 7C there is a label within the graph that belongs to the IPs underneath. Also, the graphs (C, D, E) are mislabeled relative to the legend.

We have corrected this.

6. In Fig. 7F two prominent bands appear in the Spc42-Flag IP from wt cells, while one of the two is much decreased in cdc14-1 cells. Which one of the two is Spc42?
**The top band, which has similar intensity in the two samples, is Spc42-Flag**

**Competing Interests:** None

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**Reviewer Report 16 January 2017**

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**Fernando Monje-Casas**

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In their manuscript, Fox *et al.* demonstrate that Cdc14, a key phosphatase that promotes CDK inactivation and exit from mitosis in *Saccharomyces cerevisiae*, also plays a pivotal role in promoting the re-licensing of the SPBs at the meiosis I to meiosis II transition, thus ensuring the re-duplication of these structures between these two different meiotic phases. In order to fulfill this function, Cdc14 is loaded on a single SPB during anaphase I in a process that depends on the Bfa1/Bub2 complex. Once loaded on the SPB, Cdc14 likely promotes dephosphorylation of key substrates on this structure, thereby allowing the re-duplication of the SPBs in meiosis II. This newly suggested meiotic function of Cdc14 helps to shed light on how cells establish the particular pattern of chromosome segregation during meiosis, an essential cell division process by which gametes are generated. Overall, the experiments detailed in the manuscript are carefully designed, nicely presented and well executed. Also, and importantly, the final conclusions of the manuscript are properly sustained by the provided experimental data. Therefore, I support an approved status for this article. In any case, I also indicate some minor corrections that would need to be introduced, as well as some suggestions that might help the authors to improve the manuscript and strengthen some of the conclusions:

1. In page 8, when the authors indicate that *slk19Δ* and *spo12Δ* cells disassembled anaphase I spindles with a timing comparable to wild type cells, they only refer to Figure 1B, but Figure 1A should be also mentioned. Additionally, and regarding the data in Figure 1A, I would suggest to show images for other live cell experiments that are more representative of the previous statement. Although Figure 1B indeed shows that, as an average, time from anaphase I onset until spindle disassembly is similar for wild type, *slk19Δ* and *spo12Δ* cells, in the images shown in Figure 1A for the *slk19Δ* and *spo12Δ* cells the spindle disassembles, respectively, 20 min and 30 min after anaphase I onset, which is significantly earlier than for the wild type (40 min).

2. Also in page 8, it is later stated that “[…] the two component GAP Bfa1/Bfa1, […] localizes symmetrically at SPBs during metaphase I, anaphase I but symmetrically during metaphase II”. This sentence should be corrected, since, as shown in Figure 1B, Bfa1 localization is asymmetric during metaphase II.
3. The localization of Cdc14-GFP is difficult to assess in Figure 3A, due to the nucleolar background and the appearance of other GFP foci that do not co-localize with the SPBs and whose nature it is not indicated (e.g., 1:15 time point in Figure 3A). Since the authors show different proteins that interact with Cdc14 on the SPBs, they might consider to use a different approach if they plan to further analyze this localization in the future. As such, a bimolecular fluorescence complementation assay (BiFC) could be helpful to determine the exact timing and pattern of localization of Cdc14 on the SPBs. The BiFC assay could facilitate tracking Cdc14 localization exclusively to the SPBs by means of the reconstitution of a fluorescent signal when the phosphatase and a SPB component, both tagged with different fragments of a fluorescent protein, interact.

4. In page 16, when the authors show a 1.5-fold increase in Spc42-tdTomato intensity as wild type cells progressed from G1 into prophase I, they only refer to Figure 7A, but it is in Figure 7B where the quantification is shown.

5. A shocking observation is that, despite Bfa1/Bub2 being symmetrically localized in anaphase I and Cdc14 activity being presumably required at both SPBs to trigger their duplication in meiosis II, the phosphatase is asymmetrically localized to only one SPB during anaphase I. This is an obvious caveat and, although I do appreciate that it would require further extensive analysis for this question to be solved, the authors might have speculated a little more extensively about the reasons for this asymmetry in the discussion of the manuscript.

6. A prediction of the model proposed by Fox et al. is that re-duplication of the SPBs should be restored in a cdc14-1 mutant background by conditionally forcing Cdc14 loading on the SPBs at the appropriate time (e.g., by temporarily expressing a fusion between Cdc14 and a SPB component). This is a relatively straightforward experiment that the authors might consider to carry out at some point to further strengthen their model. Furthermore, and coming back to the differential localization of Cdc14 on the SPBs during anaphase I, this experiment could be informative to address the importance of this asymmetry in future studies. Cdc14 could be fused either to Bfa1 (asymmetrically localized) or an integral SPB component (symmetrically localized), and then analyze the consequences of expressing both Cdc14 fusions during meiosis.

References
1. Sung MK, Huh WK: Bimolecular fluorescence complementation analysis system for in vivo detection of protein-protein interaction in Saccharomyces cerevisiae. Yeast. 2007; 24 (9): 767-75 PubMed Abstract | Publisher Full Text

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
spindles with a timing comparable to wild type cells, they only refer to Figure 1B, but Figure 1A should be also mentioned. Additionally, and regarding the data in Figure 1A, I would suggest to show images for other live cell experiments that are more representative of the previous statement. Although Figure 1B indeed shows that, as an average, time from anaphase I onset until spindle disassembly is similar for wild type, \( slk19 \Delta \) and \( spo12 \Delta \) cells, in the images shown in Figure 1A for the \( slk19 \Delta \) and \( spo12 \Delta \) cells the spindle disassembles, respectively, 20 min and 30 min after anaphase I onset, which is significantly earlier than for the wild type (40 min).

We have now included the reference to Figure 1A. Under the conditions we observed them, the timing of spindle disassembly varied across all strains (Figure 1B), we tried to show representative images to reflect this.

2. Also in page 8, it is later stated that “[…] the two component GAP Bfa1/Bfa1, […] localizes symmetrically at SPBs during metaphase I, anaphase I but symmetrically during metaphase II”. This sentence should be corrected, since, as shown in Figure 1B, Bfa1 localization is asymmetric during metaphase II.

The typo has been corrected.

3. The localization of Cdc14-GFP is difficult to assess in Figure 3A, due to the nucleolar background and the appearance of other GFP foci that do not co-localize with the SPBs and whose nature it is not indicated (e.g., 1:15 time point in Figure 3A). Since the authors show different proteins that interact with Cdc14 on the SPBs, they might consider to use a different approach if they plan to further analyze this localization in the future. As such, a bimolecular fluorescence complementation assay (BiFC) \(^1\) could be helpful to determine the exact timing and pattern of localization of Cdc14 on the SPBs. The BiFC assay could facilitate tracking Cdc14 localization exclusively to the SPBs by means of the reconstitution of a fluorescent signal when the phosphatase and a SPB component, both tagged with different fragments of a fluorescent protein, interact.

It is presently unclear what the non-nucleolar, non-SPB Cdc14-GFP foci represent. Although outside the scope of the current manuscript, we appreciate the excellent suggestion to use a bimolecular complementation assay to probe the co-localization of Cdc14 with SPBs. It would be particularly interesting to determine whether Cdc14 can be detected at both SPBs using this system.

4. In page 16, when the authors show a 1.5-fold increase in Spc42-tdTomato intensity as wild type cells progressed from G1 into prophase I, they only refer to Figure 7A, but it is in Figure 7B where the quantification is shown.

This has been corrected.

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We agree, this is indeed a surprising finding that remains unexplained. We expanded our discussion as suggested, though ultimately this is indeed a very puzzling observation for which we currently do not have an explanation.

6. A prediction of the model proposed by Fox et al. is that re-duplication of the SPBs should be restored in a cdc14-1 mutant background by conditionally forcing Cdc14 loading on the SPBs at the appropriate time (e.g., by temporarily expressing a fusion between Cdc14 and a SPB component). This is a relatively straightforward experiment that the authors might consider to carry out at some point to further strengthen their model. Furthermore, and coming back to the differential localization of Cdc14 on the SPBs during anaphase I, this experiment could be informative to address the importance of this asymmetry in future studies. Cdc14 could be fused either to Bfa1 (asymmetrically localized) or an integral SPB component (symmetrically localized), and then analyze the consequences of expressing both Cdc14 fusions during meiosis.

We attempted this experiment using the GFP-GBP system. However, we were unable to obtain strains in which Cdc14 and SPB components were both tagged with these binding partners. Although other explanations are possible, these findings are consistent with the idea that forced constitutive targeting of Cdc14 to the SPB is lethal also in vegetative cells. Due to the lethality we were unfortunately unable to assess the effect on meiosis.

**Competing Interests:** None
of how SPBs duplicate between the two divisions. However, there are some points that need addressed:

Major considerations:
1. Does the Cdc14-1 allele result in a complete loss of function in meiosis? A discussion of whether this is a hypomorph or complete loss of function would be helpful, especially for thinking about the conclusion that "Cdc14 is not absolutely required for CDK down-regulation at meiosis I exit".

2. In Figure 1 C, why do only 50% of the cells form 2 spindles instead of all of the cells? The duration of the movies needs to be increased to ensure that there is enough time for meiosis II to occur.

3. I am having difficulty reconciling the results in Fig 7 with the model in Fig. 9. Why does the Spc42 fluorescence remain the same throughout meiosis I (metaphase I and anaphase I) when the SPB should be duplicating during the time? It would be helpful to have a graph showing the change in Spc42 fluorescence, tracking individual cells, also marking the time of anaphase I and anaphase II.

4. Does Sfi1 phosphorylation change throughout meiosis? Does Sfi1 remain phosphorylated in the Cdc14-1 cells?

5. A thought about the conclusions: The prematurely released Cdc14 in pCLB2-CDC55 cells also localized to SPBs but does not allow SPBs reduplication. Does this finding suggest that the Cdc14 SPB localization is required but not sufficient to promote SPB duplication if the SPBs have not first separated?

Minor points:
1. The last sentence in the abstract should end in "meiosis II transition." It currently says meiosis I.

2. The sentence in the last paragraph on page 8 should read, "An exception is the two component GAP Bub2/Bfa1 which localized symmetrically at SPBs during metaphase I, anaphase I by asymmetrically during metaphase II".

3. The reference to the white arrow in Figure 8 is right after talking about vesicle formation, but the arrow does not point at the vesicles.

4. On page 16, the reference to Fig. 7F is missing and one of the references to 7E should be to 7G.

**Competing Interests:** No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.
this is a hypomorph or complete loss of function would be helpful, especially for thinking about the conclusion that “Cdc14 is not absolutely required for CDK down-regulation at meiosis I exit”.

It is true that we cannot be sure that the cdc14-1 allele is completely inactivated during meiosis. Our attempts to deplete or degrade Cdc14 during meiosis have been unsuccessful and therefore temperature-sensitive alleles are the best tools we have. We find that spindle disassembly is impaired in cdc14-1 conditions in mitosis but not meiosis under similar temperature conditions, leading us to suggest a lesser requirement for Cdc14 in meiotic spindle disassembly. Recently, similar findings have been reported for meiosis II using a different temperature sensitive allele (Argüello-Miranda et al., 2017). We have revised the text to include the possibility that Cdc14 retains partial function using these alleles and included a reference to this recent study.

2. In Figure 1 C, why do only 50% of the cells form 2 spindles instead of all of the cells? The duration of the movies needs to be increased to ensure that there is enough time for meiosis II to occur.

The duration of the movies was 12h, however not all cells complete meiosis under the imaging conditions used for this experiment. Rather than bias the data by scoring only cells that complete meiosis II, all cells that enter meiosis were included in the analysis. This experiment was performed at a time where we were still establishing optimal conditions for live cell imaging of cells undergoing meiosis. Although we recognise that we could improve on this with our optimised conditions (where essentially all cells complete meiosis), we feel that the data presented clearly shows the effect of different mutants on meiosis I spindle disassembly.

3. I am having difficulty reconciling the results in Fig 7 with the model in Fig. 9. Why does the Spc42 fluorescence remain the same throughout meiosis I (metaphase I and anaphase I) when the SPB should be duplicating during the time? It would be helpful to have a graph showing the change in Spc42 fluorescence, tracking individual cells, also marking the time of anaphase I and anaphase II.

SPB fluorescence stays the same throughout meiosis I (metaphase I and anaphase I) because SPBs are already duplicated before metaphase I (i.e. in the ndt80D arrest – see Figure 7C). We do note, however, that a greater increase in fluorescence might be expected during the later stages of anaphase I in wild type cells. The source data for this experiment that would allow the reader to make traces of individual cells is available at https://osf.io/g5cmh/ (Marston, 2016). The fluorescence analysis of Spc42-tdTomato is complicated by two factors that mean we cannot make stand alone conclusions from it. First, in other experiments (not shown) we concluded that Spc42 incorporation into the SPB is continuous and exchange occurs. Second, the SPB undergoes a remodelling and maturation during meiosis II. As a result of these concerns we performed electron microscopy, which clearly show a failure in SPB re-duplication in cdc14-1 mutants. We believe that the EM data unequivocally supports our conclusions, while the fluorescence intensity data offer supportive, but not definitive evidence.

2. Does Sfi1 phosphorylation change throughout meiosis? Does Sfi1 remain phosphorylated in the Cdc14-1 cells?

Despite extensive efforts to examine Sfi1 phosphorylation in wild type and cdc14-1 mutants undergoing meiosis, the findings were inconclusive. Judging by the presence of multiple Sfi1 species on western blots, Sfi1 appears to be extensively phosphorylated during meiosis in wild type cells. Our model predicts transient dephosphorylation at the time of anaphase I, while phosphorylation should persist in cdc14-1 mutants. Unfortunately, using this method, we were
unable to reproducibly visualize Sfi1 dephosphorylation during anaphase I during wild type cells, likely due to the very short time window in which it occurs and variable cell synchronization, even using the best methods currently available (NDT80 block-release).

3. A thought about the conclusions: The prematurely released Cdc14 in pCLB2-CDC55 cells also localized to SPBs but does not allow SPBs reduplication. Does this finding suggest that the Cdc14 SPB localization is required but not sufficient to promote SPB duplication if the SPBs have not first separated?

SPBs undergo the first round of duplication in pCLB2-CDC55 cells but do not appear to be able to separate. The licensing model predicts that, following Cdc14 dephosphorylation of Sfi1, CDK activity is required to separate the SPBs. We suggest that Cdc14 at the SPB counteracts CDK activity, thereby preventing separation of SPBs. Therefore, we agree, SPB separation may be a pre-requisite for duplication.

Minor points:
1. The last sentence in the abstract should end in “meiosis II transition.” It currently says meiosis I.
   We have corrected this
2. The sentence in the last paragraph on page 8 should read, “An exception is the two component GAP Bub2/Bfa1 which localized symmetrically at SPBs during metaphase I, anaphase I by asymmetrically during metaphase II”.
   Corrected.
3. The reference to the white arrow in Figure 8 is right after talking about vesicle formation, but the arrow does not point at the vesicles.
   The arrows mark the SPBs, not the vesicles, as written in the figure legend. We made this clearer by writing “SPB1, white arrow”, “SPB2, white arrow”.
4. On page 16, the reference to Fig. 7F is missing and one of the references to 7E should be to 7G.
   We have corrected this.

Competing Interests: None