Histone H3K4 methylation regulates deactivation of the spindle assembly checkpoint through direct binding of Mad2

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Histone H3 methylation on Lys4 (H3K4me) is associated with active gene transcription in all eukaryotes. In *Saccharomyces cerevisiae*, Set1 is the sole lysine methyltransferase required for mono-, di-, and trimethylation of this site. Although H3K4me3 is linked to gene expression, whether H3K4 methylation regulates other cellular processes, such as mitosis, is less clear. Here we show that both Set1 and H3K4 mutants display a benomyl resistance phenotype that requires components of the spindle assembly checkpoint (SAC), including Bub3 and Mad2. These proteins inhibit Cdc20, an activator of the anaphase-promoting complex/cyclosome (APC/C). Mutations in Cdc20 that block Mad2 interactions suppress the benomyl resistance of both *set1* and H3K4 mutant cells. Furthermore, the HORMA domain in Mad2 directly binds H3, identifying a new histone H3 “reader” motif. Mad2 undergoes a conformational change important for execution of the SAC. We found that the closed (active) conformation of both yeast and human Mad2 is capable of binding methylated H3K4, but, in contrast, the open (inactive) Mad2 conformation limits interaction with methylated H3. Collectively, our data indicate that interactions between Mad2 and H3K4 regulate resolution of the SAC by limiting closed Mad2 availability for Cdc20 inhibition.

[**Keywords:** chromatin; lysine methylation; spindle assembly checkpoint; Set1; H3K4; HORMA domain]

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Lysine methylation within histone H3 is commonly associated with gene regulation. H3K4 trimethylation in particular is associated with gene activation and is localized to gene promoters. In contrast, H3K4 dimethylation is present across the gene body of both active and repressed genes (Bernstein et al. 2002; Santos-Rosa et al. 2002). Lysine methylation also contributes to other cellular processes, such as DNA damage repair (Sollier et al. 2004) and telomere maintenance (Krogan et al. 2002). H3K4 methylation alters chromatin structure and also serves to recruit or exclude binding of nonhistone proteins to chromatin.

A single lysine methyltransferase, Set1, mediates all H3K4 methylation events in *Saccharomyces cerevisiae*

(Briggs et al. 2001). Set1 is incorporated into a multiprotein complex termed COMPASS (Miller et al. 2001; Roguev et al. 2001; Krogan et al. 2002) and is homologous to human MLL1, which also methylates H3K4 and is incorporated into a COMPASS-like complex (Shilatifard 2012). Translocations of MLL1 are associated with leukemogenesis (Zelevnik-Le et al. 1994) and a poor prognosis (Pui et al. 1994). However, how mutations in *MLL1* contribute to leukemogenesis and an unfavorable outcome is not clear. Identifying novel pathways regulated by *SET1* in yeast can help elucidate cellular functions regulated by *MLL1* in humans (Schneider et al. 2005; Milne et al. 2010; Shilatifard 2012).

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COMPASS also methylates Dam1, a kinetochore protein (Zhang et al. 2005). Like H3K4 methylation, Dam1 methylation is regulated in *trans* by histone H2B ubiquitination (Latham et al. 2011). Moreover, Dam1 dimethylation on K233 inhibits phosphorylation of surrounding serines by the Aurora kinase Ipl1. Accordingly, mutations that eliminate Set1 expression or activity suppress the effects of conditional mutations in Ipl1 (Zhang et al. 2005).

Ipl1 phosphorylates a number of kinetochore proteins in response to inappropriate microtubule–kinetochore interactions that lead to triggering of the spindle assembly checkpoint (SAC) (Biggins and Murray 2001). The SAC guards against aberrant chromosome segregation during mitosis by preventing progression to anaphase until defective microtubule–kinetochore attachments are resolved and mitotic spindle tension is established (Rieder et al. 1994; Li and Nicklas 1995).

S. cerevisiae has provided a useful model to identify several SAC components, including Mad1 (mitotic arrest defect 1), Mad2, Mad3 (Li and Murray 1991), Bub1 (budding uninhibited by benomyl 1), and Bub3 (Hoyt et al. 1991). The SAC involves an orchestration of protein–protein interactions and phosphorylation events that ultimately prevent activation of the anaphase-promoting complex/cyclosome (APC/C) (Jia et al. 2013). The APC is an E3 ubiquitin ligase that requires Cdc20 to recruit proteins for ubiquitination and subsequent degradation. SAC proteins bind directly to Cdc20, preventing proteolysis of key APC substrates such as Securin (Pds1).

As our previous studies indicated that Set1 opposes the functions of Ipl1 (Zhang et al. 2005), we reasoned that Set1 might have additional functions during mitosis. Here we report that the highly conserved HORMA domain in Mad2 is a novel H3K4 methyl reader and that this modification and Set1 play an important role in regulating the release of the SAC through Mad2 interactions.

Results

Loss of SET1 induces benomyl resistance

To further address how lysine methylation might regulate mitosis, we subjected cells bearing deletions in *SET1* or in other SET domain-encoding genes to growth in the presence of the microtubule depolymerizing agent benomyl, which interferes with mitotic spindle stability and antagonizes mitotic progression. Mutations in genes required for formation of the mitotic spindle, in components of the kinetochore, or in the activation and maintenance of mitotic checkpoints are characteristically sensitive to microtubule depolymerizing drugs (Spencer et al. 1990; Stearns et al. 1990; Hoyt et al. 1991; Li and Murray 1991).

We discovered that *set1* mutant cells are highly resistant, rather than sensitive, to high levels of benomyl (Fig. 1A; Supplemental Fig. S1A). *set1* mutant cells grow similarly to wild-type cells on rich medium or in the presence of dimethyl sulfoxide (DMSO) alone but display continued growth in levels of benomyl (30–40 μ g/mL) that completely block growth of wild-type cells. Benomyl resistance was not observed upon deletion of any other

Figure 1. Loss of COMPASS-mediated lysine methylation results in benomyl resistance. (A) A serial fivefold dilution assay of yeast strains with the indicated genotypes was performed to visualize growth phenotypes. Cells were placed onto YPD plates (YPD) with or without 30 μ g/mL benomyl, and these plates were placed for 2 d at 30°C. (B) Whole-cell extracts (WCEs) were isolated from strains with the indicated genotypes. The amount of total Set1 protein was assessed by immunoblot analysis using an antibody specific for Set1 (1:100; Santa Cruz Biotechnology, SC-101858). Pgk1 protein levels were used as a loading control. (C) Both wild-type Set1 and Set1G951S proteins similarly associate with the COMPASS component Bre2. Immunoprecipitation (IP) assays were used to pull down Bre2, and the associated levels of Set1 were assessed in the wild-type (WT) and *set1G951S* strains. Bre2-TAP (α -protein A) and Pgk1 (α -Pgk1) were used as loading controls. (D) *bre1Δ*, *set1Δ*, and the catalytically inactive mutant *set1G951S* were subjected to the same assay as in A. (E) COMPASS mutants were assessed for growth phenotypes using the same serial fivefold dilution assay as in A and D.

SET domain-containing gene, indicating a unique function for Set1 in responding to microtubule poisons (Fig. 1A). Interestingly, since *SET2* encodes the lysine methyltransferase required for histone H3K36 mono-, di-, and trimethylation, which is also associated with active transcription, our results suggest that defective transcription is not sufficient to confer benomyl resistance.

To address whether the benomyl resistance phenotype reflects loss of Set1 catalytic activity, we subjected cells bearing a mutation in the Set1 catalytic site, G951S (Nagy et al. 2002), to growth in the presence of the drug. Although some Set1 mutations that affect methyltransferase activity result in lower amounts of total Set1 protein (Soares et al. 2014), we detected equal amounts of wild-type Set1 and Set1G951S protein in immunoblots of whole-cell extracts (Fig. 1B). Moreover, we found equal incorporation of wild-type and mutant Set1 into the COMPASS complex, isolated via affinity purification of

Bre2, another COMPASS component (Fig. 1C). As expected, the G951S mutation abrogated the ability of Set1 to methylate histone H3K4 (Supplemental Fig. S1B). Importantly, cells bearing the *set1G951S* mutation displayed benomyl resistance equivalent to that caused by *SET1* deletion (Fig. 1D), indicating that Set1 catalytic activity is required for a normal response to microtubule depolymerization.

Our results are consistent with a previous large-scale screen that indicated that *BRE1* mutants display resistance to benomyl (Rieger et al. 1999). Bre1 is an E3 ubiquitin ligase and a targeting factor for the E2 enzyme Rad6. Both Bre1 and Rad6 are required for histone H2BK123 monoubiquitination (Nakanishi et al. 2009), an event required upstream of Set1-mediated methylation of both H3K4 and Dam1 K233 (Lee et al. 2007; Latham et al. 2011).

Mutations in different components of the COMPASS complex (Fig. 1E) affect distinct methylation states of Set1 substrates (Schneider et al. 2005). We found that deletion of genes encoding subunits required for all three methylation states, such as *SWD1* and *SWD3*, also results in benomyl resistance (Fig. 1E). Deletion of *BRE2* or *SDC1* inhibits dimethylation and trimethylation of H3K4 (Schneider et al. 2005; Latham et al. 2011), and these mutations also induce substantial resistance to benomyl, although not quite to the same degree as caused by *SET1* mutations. Deletion of *SPP1* eliminates trimethylation of H3K4 without affecting mono- and dimethylation (Schneider et al. 2005). Interestingly, *spp1* mutants display only a slight benomyl resistance, indicating that loss of trimethylation of COMPASS substrates is not sufficient to trigger resistance. Deletion of *SHG1* increases di- and trimethylation of H3K4 (Dehe et al. 2006) but does not affect benomyl resistance. Together, our results indicate that mono- and dimethylation of substrates by Set1, working within the COMPASS complex, are required for proper response to microtubule poisons.

Benomyl resistance of set1G951S cells requires an intact SAC and Cdc20 inhibition

Resistance to spindle poisons likely reflects failure in the regulation of the SAC, which maintains genome integrity by preventing anaphase progression in the presence of defective microtubule–kinetochore attachments (Fig. 2A).

Importantly, although the components and functions of the SAC are highly conserved across evolution, kinetochore–microtubule interactions are maintained throughout the yeast cell cycle, with only a short disruption during early S phase (Kitamura et al. 2007). Unlike mammalian cells, the SAC becomes essential in *S. cerevisiae* only upon active damage to the spindle, such as upon exposure to benomyl (Hoyt et al. 1991; Li and Murray 1991). Consistent with those findings, *set1Δ* cells do not display any cell cycle aberrations, as measured by flow cytometry (Supplemental Fig. S1C) or a budding index assay (Supplemental Fig. S1D). Our *set1Δ* cells did grow at a marginally slower rate than wild-type cells (Supplemental Fig. S1E), as previously reported (Miller et al. 2001), correlating with a slight reduction in colony-forming units (Supplemental Fig. S1F), but no abnormal cellular morphologies were observed (Supplemental Fig. S1G). Therefore, the benomyl resistance phenotype displayed by *set1* mutants is unlikely to be a downstream effect of other cell cycle anomalies.

To determine whether activation of the SAC is required for the benomyl resistance of *set1* mutants, we deleted genes encoding SAC components in the *set1G951S* strain. Deletion of *BUB1* and *BUB3* eliminated the resistance of *set1G951S* cells to benomyl (Fig. 2B). These data indicate that these SAC components are required for the resistance phenotype and also that the *set1* mutation does not prevent uptake of the drug, as the double mutants displayed a sensitivity to benomyl similar to that of *bub1* and *bub3* single mutants. Deletion of *MAD1*, *MAD2*, and *MAD3* also suppressed the benomyl resistance phenotype of *set1G951S* cells (Fig. 2C). Collectively, these data indicate that the SAC must be activated and intact for the benomyl resistance to occur in the *set1* mutant cells.

SAC component proteins inhibit Cdc20 to block entry into anaphase upon disruption of microtubule–kinetochore interactions. To further define the role of SAC activation in the benomyl resistance phenotype, we introduced a Cdc20 mutation that blocks inhibition into *set1G951S* cells. The dominant *cdc20-127* allele disrupts binding of Mad2 and Mad3, thereby inhibiting SAC-mediated cell cycle arrest upon exposure to microtubule poisons (Hwang et al. 1998). We found that *set1G951S cdc20-127* double mutants are sensitive to benomyl compared with *set1G951S* mutants alone, indicating that binding of SAC

Figure 2. The *set1G951S* mutant requires the SAC and Cdc20 inhibition for benomyl resistance. (A) An illustration of Cdc20 associated with the APC complex in both an active and an inactive state. Mad2 and Mad3 bind to Cdc20 to inhibit the ubiquitination of APC substrates. (B–D) Serial fivefold dilution assay of yeast strains with the indicated genotypes grown on normal YPD plates (YPD) or plates containing 30 μ g/mL benomyl for 2 d at 30°C.

components to Cdc20 is required for benomyl resistance (Fig. 2D) and that lysine methylation events mediated by Set1 function to limit Cdc20 inhibition.

set1 mutants display a thick mitotic spindle and a defect in the release of the SAC

One phenotype reported previously for *cdc20-1*, a temperature-sensitive mutant of Cdc20, is a thick mitotic spindle (Sethi et al. 1991), although this phenotype appears to be variable (Lim et al. 1998). Confocal analyses indicated that the mitotic spindles in *set1G951S* mutant cells are thicker than in wild-type cells (Fig. 3A). Indeed, although thick spindles were rarely observed in wild-type cells (e.g., three of 34 cells examined [8.8%]), they occurred in a majority (10 of 13 [76.9%]) of *set1* mutant cells (Fig. 3A). As Cdc20 recruits proteins to the APC/C for ubiquitination and degradation, the thick mitotic spindles in *cdc20* mutants likely reflect a failure in the degradation of microtubule-bundling proteins (Sethi et al. 1991; Juang et al. 1997). To determine whether the thick mitotic spindles in *set1G951S* mutants also reflect Cdc20 inhibition, we monitored destruction of a Cdc20–APC/C substrate, Pds1 (Securin), after SAC activation and release. Pds1 ubiquitination and proteolysis are required for the separation of sister chromatids and progression through anaphase. Both wild-type and *set1G951S* cells were treated with nocodazole to activate the SAC. Flow cytometry indicated that both wild-type and *set1G951S* cells arrested in G2/M, as expected. The cells were then released into fresh medium without drugs, and mitotic progression was monitored by assessing Pds1 stability coupled with cell cycle analysis. Immunoblots indicate that Pds1 is degraded at a slower rate in *set1* mutant cells (Fig. 3B, cf. lanes labeled 45' and 60'). These results indicate that *set1* mutants arrest normally after SAC activation but exhibit prolonged inhibition of the APC after checkpoint release.

Loss of H3K4 methylation results in benomyl resistance and a thick mitotic spindle

Two COMPASS substrates have been identified so far: H3K4 (Briggs et al. 2001) and Dam1 K233 (Zhang et al. 2005). We created genomic mutations in these methylation sites in order to determine whether either of these COMPASS substrates plays a role in regulating cellular responses to mitotic poisons. We found that *dam1K233R* cells are sensitive to benomyl, much like wild-type cells. However, cells bearing mutations in H3K4 (*H3K4R*) display benomyl resistance very similar to that seen in *set1G951S* cells (Fig. 3C). The yeast genome contains two alleles encoding H3: *HHT1* and *HHT2*. To determine whether partial loss of H3K4 methylation was sufficient to induce benomyl resistance, we compared growth of wild-type cells, cells bearing single mutations (*hht1K4R* or *hht2K4R*), and cells harboring mutations in both H3 alleles (labeled here as *H3K4R* for simplicity). We found that only the double mutation conferred benomyl resistance (Supplemental Fig. S2A). Moreover, the loss of *SET1* in *H3K4R* cells did not confer any further resistance, suggesting that H3K4 is the primary Set1 substrate regulating normal cellular response to mitotic poisons (Supplemental Fig. S2B). Importantly, we confirmed that Set1 protein levels and incorporation into the COMPASS complex were not affected by the *H3K4R* mutation (Supplemental Fig. S2C,D).

We also examined the mitotic spindle in wild-type, *dam1K233R*, *H3K4R*, and *set1G951S* cells (Fig. 3D). The mitotic spindle in *dam1K233R* cells appeared similar to that in wild-type cells. Only one in 16 (6.2%) displayed a thick spindle. In contrast, a majority (24 of 28 cells [85.7%]) of *H3K4R* cells displayed a thick mitotic spindle phenotype resembling that of *set1G951S* cells (Fig. 3D). These data indicate that methylation of H3K4 plays a novel role in regulating the response to microtubule

Figure 3. Loss of H3K4 methylation results in a thick mitotic spindle. (A) Cells bearing a *SET1* catalytic mutant allele (*set1G951S*) display a more robust mitotic spindle staining compared with wild-type cells (WT). Immunofluorescent confocal images of wild-type and *set1G951S* mitotic cells are shown. Tubulin (green) and DAPI (blue) stainings were used to identify cells undergoing mitosis in an asynchronous culture. Mitotic cells are outlined in white. (B) Cell cycle arrest and release experiments reveal that Pds1 is more stable in *SET1* mutants (*set1G951S*) than in wild-type cells. Cells were arrested in G2/M and released into fresh medium. Samples were taken at the indicated time points, and whole-cell extracts and immunoblots were prepared and probed with α -Myc antibody to assess the protein levels of Myc-tagged Pds1. Immunoblots of Pgk1 served as a loading control. Cells from the indicated time points were taken for analysis by flow cytometry. Histograms of DNA content reveal the cell cycle profile of the indicated strains at specific times after cell cycle arrest and release. (C) Serial fivefold dilution assays of wild-type and mutants with the indicated genotype placed onto either control plates (YPD) or plates containing 30 μ g/mL benomyl as in previous figures. (D) Immunofluorescent confocal images of wild type and mutants with the indicated genotypes. Staining and selection of cells were performed as in A.

depolymerizing drugs, related to increased amounts of polymerized tubulin or increased stability of the mitotic spindle, consistent with inhibition of Cdc20 function.

We next determined whether an intact SAC is required for benomyl resistance of H3K4R cells. As seen in *set1G951S* cells (Fig. 2), deletion of SAC component genes suppressed the benomyl resistance phenotype caused by H3K4R mutations (Fig. 4A,B). Immunoblots confirmed that loss of SAC components such as Mad2 had no effect on H3K4 methylation events, as H3K4 di- and trimethylation were unaltered in *mad2Δ* cells (Fig. 4C). Eliminating SAC function through deletion of *MAD2* in H3K4R cells also resulted in suppression of the thick mitotic spindle phenotype (two of 18 [11.1%] for the double mutant and one of 18 [5.5%] for the *MAD2* deletion alone, compared with 24 of 28 [85.7%] as noted above) (Fig. 4D; multiple panels in Supplemental Fig. S2E). These data further link the mitotic spindle phenotype with the benomyl resistance observed in the H3K4R cells. To test whether the H3K4R mutant also regulates the SAC through Cdc20 inhibition, we introduced the *Cdc20-127* mutation into the H3K4R strain. As in *set1G951S* cells, H3K4R cells require Cdc20 inhibition for benomyl resistance (Fig. 4E), supporting a model of an overactive SAC when H3K4 methylation is inhibited.

Since H3K4 methylation is involved in transcriptional regulation, we compared previously published microarray data (Venkatasubrahmanyam et al. 2007) with known

loss-of-function mutants that cause benomyl resistance (Brown et al. 2006). Loss of *SET1* results in 127 significantly down-regulated genes, none of which are associated with mitosis or the SAC. We confirmed that expression of SAC components is not down-regulated upon loss of H3K4 methylation by comparing levels of Bub1, Bub3, Mad1, Mad2, and Mad3 RNA in both wild-type and *set1Δ* cells (Supplemental Fig. S2F). We also evaluated H3K4 methylation (H3K4me) events at pericentric regions through chromatin immunoprecipitation (ChIP) followed by sequencing (ChIP-seq) but found little enrichment of H3K4 dimethylation (H3K4me2) or trimethylation (H3K4me3) in locations proximal to the centromere (Supplemental Fig. S2G). These results indicate that the presence of H3K4me2/me3, but not a pericentromeric location of this mark, is important for normal cellular response to benomyl.

Mad2 binds methylated histone H3K4 in a conformation-specific manner

A number of protein domains “read” the methylation state of H3K4, including PHDs, WD40 domains, and Tudor domains (Huyen et al. 2004; Wysocka et al. 2005; Kim et al. 2006; Shi et al. 2006). Mad2 lacks those domains but does contain a HORMA domain, which is found in several chromatin-associated proteins (Aravind and Koonin 1998; Muniyappa et al. 2014). To test whether Mad2 can bind directly to histones, we performed pull-down experiments using recombinant GST-yMad2 fusion proteins and calf thymus histones. We found that GST-yMad2 binds directly to H3, but not to H2A or H2B, in vitro (Fig. 5A). Since Mad2 is composed almost entirely of a single HORMA domain, our data indicate that the HORMA domain likely acts as an H3 interaction domain.

Mad2 adopts two conformations—an inactive, open state (O-Mad2) and an active, closed state (C-Mad2)—in both yeast and human cells (Luo et al. 2000; Sironi et al. 2002; Nezi et al. 2006). After SAC activation, Mad2 assumes the C-Mad2 conformation upon binding to Mad1 at the kinetochore. C-Mad2 then binds and inhibits Cdc20. Recombinant wild-type Mad2 can adopt both O-Mad2 and C-Mad2 conformations and can dimerize in solution (Fang et al. 1998; Luo et al. 2004). Specific mutations limit dimerization and structurally constrain Mad2 to the O-Mad2 form or promote the C-Mad2 conformation (Luo et al. 2004; Nezi et al. 2006). Mutations that delete the C-terminal tail of Mad2 result exclusively in a monomeric O-Mad2 conformation (Fang et al. 1998). In contrast, the Mad2^{RQEA} mutation, which also blocks dimerization, results in a high ratio of C-Mad2 to O-Mad2 monomers (Luo et al. 2004; Nezi et al. 2006). To determine whether Mad2 conformational changes affect binding to H3, we generated GST fusions to yeast wild-type, O-yMad2, and yMad2^{RQEA} proteins. We found that recombinant O-yMad2 interacted less well with H3 than did wild-type yMad2. In contrast, yMad2^{RQEA} pulled down more H3 than did either the wild-type or O-yMad2 conformations. None of the yMad2 proteins bound to H2A or H2B, confirming specificity for H3 interactions (Fig. 5A).

Figure 4. The H3K4R mutant requires the SAC and Cdc20 inhibition for benomyl resistance. (A,B) Serial fivefold dilution assay of wild-type (WT) and mutants with the indicated genotypes placed onto either control plates (YPD) or plates containing 30 μg/mL benomyl, as in previous figures. (C) Immunoblots of whole-cell extracts from wild-type, *set1Δ*, and *mad2Δ* cells to compare di- and trimethylation of histone H3 levels. Total H3 was used as a loading control. (D) Immunofluorescent confocal images of wild-type and mutants with the indicated genotypes. Tubulin (green) and DAPI (blue) were used to identify cells undergoing mitosis from an asynchronous culture. Mitotic cells are outlined in white, as in previous figures. (E) Serial fivefold dilution assay of wild-type and mutants with the indicated genotypes placed onto either control plates (YPD) or plates containing 30 μg/mL benomyl, as above.

Figure 5. Mad2 binds H3 methylated at K4 in a conformation-specific manner. (A,C) Colloidal staining of purified recombinant proteins. Pull-down assays of the indicated recombinant proteins to identify direct binding with calf thymus histones, assayed using immunoblots of membranes probed with antibodies against H2A, H2B, or H3. (B,D) Pull-down assay with recombinant GST-Mad2 fusion proteins and unmodified H3 (K4me0) or MLA full-length histones generated to mimic H3K4 monomethylation (K4me1), H3K4 dimethylation (K4me2), or H3K4 trimethylation (K4me3). Direct binding was assayed using immunoblots of membranes probed with an antibody recognizing histone H3. (E,F) GST pull-down assays using recombinant GST-yMad2 and GST-hMad2 with wild-type histone H3 or H3 with various point mutations. Direct binding was assessed by using immunoblots of membranes probed with an antibody against H3.

Importantly, O-yMad2 and yMad2^{RQEA} displayed strikingly different profiles binding to modified and unmodified forms of H3. yMad2^{RQEA} clearly bound to all methylation states of H3, whereas O-yMad2 bound best to H3K4me0 and displayed limited interaction with H3K4me1 (Fig. 5B). Both O-yMad2 and yMad2^{RQEA} are monomeric, and yMad2^{RQEA}–H3K4me binding indicates that C-Mad2, rather than O-Mad2, interacts with H3K4me. These findings identify Mad2 as a conformation-sensitive reader of lysine methylation states.

Yeast and human Mad2 structure and function are strikingly conserved. To determine whether human Mad2 (hMad2) also selectively binds H3 in a conformation-specific manner, we repeated the histone-binding assays with GST-hMad2 proteins. We purified GST fusions of wild-type hMad2, monomeric O-hMad2, and monomeric

hMad2^{RQEA}. Wild-type hMad2 did not bind well to any histone, but both O-hMad2 and hMad2^{RQEA} bound H3 but not H2A or H2B. Like yMad2, hMad2^{RQEA} pulled down more H3 than did O-hMad2 (Fig. 5C), indicating that the conformation sensitivity of Mad2 interactions with H3 is conserved from yeast to humans. Failure of the wild-type hMad2 to bind H3 suggests that hMad2 dimerization may limit its interactions with histones. Importantly, recombinant hMad2^{RQEA} interacted with methylated H3K4 much like yMad2^{RQEA} (Fig. 5D), indicating that closed hMad2 also serves as a reader of H3K4me states.

A sequence highly similar to a known consensus Mad2-binding site is found in amino acids 23–30 of H3. To test whether yMad2 requires this consensus sequence to directly bind histone H3, we generated two recombinant histone H3 mutant proteins K23A (H3.1K23A) and a second containing two mutations to both R26A and K27A (H3.1R26A/K27A). Wild-type histone H3 and the K23A mutant both bound to yMad2, while the R26A/K27A mutant H3 did not (Fig. 5E). Use of a GST-hMad2 construct confirmed that hMad2 also does not interact with the R26A/K27A H3 mutant (Fig. 5F). These results demonstrate that both yMad2 and hMad2 share a conserved H3-binding sequence.

The interactions of Mad2 with H3 suggest that the binding is bipartite, involving both the consensus sequence (amino acids 23–30) and the region around K4 in H3. To further test this idea, we tested previously described alanine scanning mutations in H3 (Dai et al. 2008) within the consensus Mad2 interaction motif (Luo et al. 2002; Sironi et al. 2002) for benomyl resistance (Supplemental Fig. S3A; Dai et al. 2008). As expected, H3K4A mutant cells displayed a striking benomyl resistance phenotype, as did H3R2A mutants. Previous reports indicated a functional link between R2 methylation and K4 methylation (Kirmizis et al. 2007). In addition, mutations to amino acids 23–30 also caused a moderate resistance to benomyl, consistent with the data above, indicating that this region also contributes to Mad2 interactions (Supplemental Fig. S3A).

set1G951S and the H3K4R mutations affect Mad2 localization in dividing cells

To further assess the consequences of loss of H3K4me on Mad2 function in vivo, we examined Mad2 localization in wild-type and *set1G951S* and *H3K4R* mutant cells. As we did not have an antibody for native Mad2, we sought to use established epitope-tagged Mad2 alleles. However, we found that C-terminal tags (both GFP and TAP tags) introduced into the *MAD2* locus suppressed the benomyl resistance of *set1Δ* cells, further emphasizing the importance of the Mad2 C-terminal region in interactions with H3K4 (Supplemental Fig. S3B). Therefore, we used an inducible, N-terminally tagged GST-Mad2 fusion protein construct to assess the subcellular localization of Mad2. As expected, we found that GST-Mad2 localizes to both the nucleus and the cytoplasm in wild-type and *set1G951S* and *H3K4R* mutant cells (Supplemental

Fig. S4A). However, in large budded G2/M cells, Mad2 was enriched around the bud neck (Supplemental Fig. S4A). This staining pattern is consistent with that reported previously for Kel2, a negative regulator of mitotic exit and Mad2-binding protein that localizes to the same region (Wong et al. 2007). Interestingly, this localization pattern was enriched roughly twofold in *set1G951S* and *H3K4R* mutant cells (Supplemental Fig. S4B), consistent with disruption of Mad2 localization upon loss of H3K4me.

We propose a model in which methylated H3K4 acts as a negative regulator of the SAC by sequestering active C-Mad2, thereby limiting Cdc20–Mad2 interactions and Cdc20 inhibition. In *set1* mutants, loss of H3K4me releases C-Mad2, allowing greater inhibition of Cdc20 and also possibly favoring binding of O-Mad2 to chromatin (Fig. 6).

Discussion

Our data reveal a new function for H3K4me in regulating the SAC and identify the Mad2 HORMA domain as a conformation-specific reader of H3K4me. These studies illustrate how specific chromatin states can provide critical regulatory scaffolds for regulation of essential cellular processes where the HORMA domain likely acts as a histone interaction domain in several important, evolutionarily conserved processes.

We found that Mad2^{RQEA} binds dimethylated and trimethylated H3K4, whereas O-Mad2 does not. In wild-type cells, O-Mad2 dimerizes to C-Mad2 at the kinetochore, where it undergoes a structural change resulting in an active form of Mad2 that binds to and inhibits Cdc20 (Nasmyth 2005; Yu 2006). Cells that cannot form an O-Mad2–C-Mad2 dimer or in which Mad2 is locked into an open conformation fail to arrest in mitosis after treatment with microtubule poisons (Nezi et al. 2006).

Mad2 binds to Cdc20 after SAC activation by adopting a conformation that inhibits ubiquitination of APC substrates, such as Pds1. Our data indicate that binding of C-Mad2 to H3K4me2 inhibits Cdc20 inhibition, likely by sequestering C-Mad2 and blocking O-Mad2 interactions with H3 (Figs. 4E, 5B). After tension is achieved by kinetochore attachments to the mitotic spindle, the SAC is deactivated, allowing APC activation and ubiquitination of APC substrates. We propose that loss of H3K4me allows continued inhibition of Cdc20 through

the C-Mad2–Cdc20 interaction, leading to a delay in the release of the SAC, as indicated by the prolonged stability of Pds1.

Our model is similar to that proposed for negative regulators that act to silence the SAC, such as p31^{comet} (MAD2L1BP), in vertebrate cells (Xia et al. 2004). Like H3K4me, p31^{comet} interacts directly with C-Mad2 but not O-Mad2. We propose that H3K4me2 and H3K4me3 may function similarly to p31^{comet} to sequester C-Mad2 and inhibit binding to Cdc20. Loss of this histone modification leads to excessive availability of active C-Mad2, which in turn leads to heightened inhibition of Cdc20 functions and delayed release of the SAC. p31^{comet} is not expressed in *S. cerevisiae*, so H3K4me may serve as the primary regulator of C-Mad2 and O-Mad2 interactions in yeast. Intriguingly, p31^{comet} is itself a HORMA domain protein, raising the question of whether H3K4me may regulate its function as well.

Multiple chromatin-associated proteins contain HORMA domains in mammals, including not only p31^{comet} and MAD2 but also REV7, HORMAD1, and HORMAD2. Like Mad2, both HORMAD proteins and REV7 are thought to undergo a conformational switch, although such structural changes have yet to be studied in detail (Hara et al. 2010; Kim et al. 2014). REV7 is required for proper DNA translesion synthesis and double-strand break repair (Prakash et al. 2005; Kolas and Durocher 2006; Sale 2015). Loss of REV7 results in PARP inhibitor resistance in cancer, highlighting the importance of its functions (Xu et al. 2015). HORMAD proteins align meiotic chromosomes for efficient homologous recombination (Handel and Schimenti 2010). It will be interesting to determine how binding of H3 by these proteins influences their functions. Perhaps, for example, loss of H3–REV7 interactions might also contribute to PARP inhibitor resistance.

H3 is subject to lysine methylation at a number of other sites, and lysines are subject to multiple types of post-translational modifications (PTMs) in addition to methylation. Lysine methylation and acetylation are also subject to regulation by other modifications, such as phosphorylation or ubiquitination, at other sites (Latham and Dent 2007). Given the multiple kinases and phosphorylation events required for proper execution of the SAC, it will be important to determine whether such cross-talk occurs between H3K4me and these events and how such cross-talk impacts binding to Mad2 with histone H3.

Misregulation of histone modifications or the SAC is associated with severe human disease conditions, including cancer (Sharma et al. 2010; Fang and Zhang 2011). Several MLL proteins highly homologous to yeast Set1 exist in humans. Translocations of Mll1, in particular, that result in fusion of MLL to various members of the superelongation complex are associated with human leukemias (Smith et al. 2011). Although Mll1 functions in both normal and disease cells are usually thought of in terms of transcription, our work indicates that these proteins may also be important to maintain normal chromosome ploidy through regulation of the SAC.

Figure 6. A model showing that methylated H3K4 binds to C-Mad2. An H3 sequence required for Mad2 binding, which is similar to the Mad2-binding consensus sequence, is underlined. Our in vivo and in vitro data indicate that H3K4me acts as a negative regulator of the SAC through direct binding of methylated H3K4 to an active C-Mad2, thereby limiting C-Mad2–Cdc20 interactions.

Materials and methods

Yeast strains, media, and culture conditions

All strains generated for this study are isogenic to BY4741 (Supplemental Table S1). Cells were grown in YPD medium at 30°C under standard conditions unless otherwise stated.

Benomyl resistance assays

Plates were prepared with the indicated amounts of benomyl dissolved in DMSO. Wild-type and mutant strains were grown overnight in rich medium and placed onto either control plates (YPD or YPD with DMSO) or plates containing increasing amounts of benomyl (10, 20, 30, and 40 µg/mL). Plates were placed for 2 d at 30°C unless otherwise stated.

Flow cytometry

Five milliliters of culture of mid-log phase cells was pelleted and resuspended in 3 mL of dH₂O. Seven milliliters of 95% ethanol was added to the culture and incubated overnight at 4°C. Cells were pelleted and washed in 5 mL of sodium citrate (pH 7.4). Cells were pelleted, resuspended in 1 mL 50 mM sodium citrate with 0.25 mg/mL RNase, and incubated for 1 h at 50°C. Proteinase K was then added for a final concentration of 1 mg/mL and incubated for an additional hour at 50°C. One milliliter of sodium citrate containing 16 µg/mL propidium iodide was added for a final concentration of 8 µg/mL and incubated for 30 min before flow cytometry analysis. Data were collected on a BD LSRFortessa (Becton Dickinson) and analyzed using FlowJo 7.6 software. The protocol was courtesy of Stephen P. Bell and can be found at <http://web.mit.edu/flowcytometry/www>.

Immunofluorescence

Immunofluorescence experiments were performed according to standard techniques (Burke et al. 2000). To visualize the mitotic spindle and nuclei, we used an anti-tubulin antibody (1:500; Millipore, CBL270) and 1 µM DAPI (Molecular Probes). To visualize GST-Mad2, cells were grown in YEP + 2% raffinose until mid-log phase. GST-Mad2 protein expression was induced by the addition of galactose to a final concentration of 2% for 30 min. We used an anti-GST antibody (1:500; GE, 27-4577-01) for immunofluorescence staining. Following staining procedures, images of wild-type and mutant yeast cells were acquired with a LSM510-Meta confocal microscope (Carl Zeiss, Germany) using a 100× PlanApoChromat/1.4 N.A. objective. Images were processed using ImageJ software (National Institutes of Health) and Adobe Photoshop.

Nocodazole arrest and release

Cells were diluted into YPD medium from overnight cultures and allowed to grow until a density of 10⁷ cells per milliliter. Nocodazole was added to a final concentration of 15 µg/mL, and the cells were allowed to arrest for 2 h. Cells were released from the G2/M arrest by washing twice in dH₂O, resuspended into fresh YPD medium, and collected at the indicated time points.

Recombinant protein generation and purification

Budding yeast genes were cloned into a pGEX4T1 vector to generate an N-terminally tagged GST fusion proteins. The plasmids used in this study are listed in Supplemental Table S2. Mutations

were generated by using the QuikChange XL kit (Stratagene) and sequenced. Proteins were expressed in the *Escherichia coli* strain Rosetta 2 (Novagen) by the addition of 0.2 mM IPTG for 20 h at 14°C. After protein expression, cells were collected and suspended in lysis buffer (50 mM Tris at pH 7.5, 150 mM NaCl, 0.05% NP-40, 1 mM PMSF, protease inhibitors, 0.5 mg/mL lysozyme solution). The cells were incubated for 45 min at 4°C and lysed using a EpiShear Probe sonicator (Active Motif). Lysed cells were centrifuged, and the supernatant was removed and incubated with glutathione sepharose 4B (GE). Recombinant GST fusion proteins were eluted by the addition of 15 mg/mL L-glutathione (Sigma) dissolved in 100 mM Tris (pH 8.0). Recombinant proteins were run on a 4%–12% BisTris gel and analyzed with colloidal staining.

Calf thymus histone-binding assays

Calf thymus histone-binding assays were performed by incubating 50 µg of calf thymus histones (Worthington) with 10 µg of recombinant GST fusion proteins in buffer (50 mM Tris at pH 7.5, 1 M NaCl, 1% NP-40) overnight. GST fusion protein histone interactions were assessed by adding glutathione sepharose 4B (GE) beads for 1 h and washing five times in buffer. The beads were resuspended in 5× SDS sample buffer, run on a 4%–12% Tris-Bis gel, and transferred to a nitrocellulose membrane followed by Western blot analysis. We used the following antibodies: anti-H3 (1:5000; Active Motif, 39163), anti-H2A (1:2000; Millipore, 07-146), and anti-H2B (1:2500; Millipore, 07-371).

MLA histone-binding assays

Similar to the calf thymus histone-binding assay, 2 µg of recombinant GST fusion proteins was incubated with 1 µg of full-length recombinant histone H3 with either no methylation on K4 or methylated lysine analogs (K4me1, K4me2, and K4me3) (Active Motif). GST-yMad2 fusion proteins were incubated in 50 mM Tris (pH 7.5), 1 M NaCl, and 1% NP-40, and GST-hMad2 proteins were incubated in 50 mM Tris (pH 7.5), 750 mM NaCl, and 1% NP-40 overnight. Glutathione sepharose 4B (GE) beads were added for 1 h, washed five times with the indicated buffer, and resuspended in 5× SDS sample buffer. Proteins were resolved on a gel, and interactions were assessed by transferring the proteins to a nitrocellulose membrane followed by Western blot analysis.

Histone H3.1 purification and binding assay

A gene encoding human H3.1 was cloned into the pET-30a(+) vector to create H3.1 C-terminally tagged with a 6xHIS epitope. Mutations were generated by using the QuikChange XL kit (Stratagene) and sequenced. Plasmids used in this assay are listed in Supplemental Table S2. Proteins were expressed in the *E. coli* strain Rosetta 2 (Novagen) by the addition of 0.2 mM IPTG for 16 h at 20°C. After protein expression, cells were collected and suspended in buffer (50 mM Tris at pH 8.0, 500 mM NaCl, 5% glycerol, 1 mM PMSF). The cells were disrupted using an EpiShear Probe sonicator (Active Motif). Cells were then pelleted and resuspended in denaturing buffer (50 mM Tris at pH 8.0, 500 mM NaCl, 5% glycerol, 6 M urea). The mixture was centrifuged, and the supernatant was removed and incubated with Ni-NTA agarose beads (Qiagen). Recombinant H3.1-6xHIS fusion proteins were eluted by the addition of 150 mM imidazole added to buffer containing 50 mM Tris (pH 8.0), 500 mM NaCl, and 5% glycerol. Eluates were run on a 4%–12% BisTris gel and evaluated with colloidal staining. Purified recombinant proteins were dialyzed overnight into buffer containing 20 mM Tris (pH 8.0), 5 mM DTT, 1 mM PMSF, 1 mM EDTA, 500 mM NaCl, and 5% glycerol

to remove urea. Purified proteins were then used in binding assays. Two micrograms of recombinant GST fusion protein was incubated with 2 μ g of full-length recombinant histone H3 in buffer (50 mM Tris at pH 7.5, 750 mM NaCl, 0.1% NP-40) overnight. Glutathione sepharose 4B (GE) beads were added to the binding assay for 1 h, washed five times with buffer, and resuspended in 5 $\times$ SDS sample buffer. Proteins were resolved on a gel, and interactions were analyzed by transferring the proteins to a nitrocellulose membrane followed by Western blot analysis.

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Histone H3K4 methylation regulates deactivation of the spindle assembly checkpoint through direct binding of Mad2

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