

Dual recognition of chromatin and microtubules by INCENP is important for mitotic progression

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The chromosomal passenger complex (CPC), composed of inner centromere protein (INCENP), Survivin, Borealin, and the kinase Aurora B, contributes to the activation of the mitotic checkpoint. The regulation of CPC function remains unclear. Here, we reveal that in addition to Survivin and Borealin, the single α -helix (SAH) domain of INCENP supports CPC localization to chromatin and the mitotic checkpoint. The INCENP SAH domain also mediates INCENP's microtubule binding, which is negatively regulated by Cyclin-dependent kinase-mediated phosphorylation of segments flanking the SAH domain. The microtubule-binding capacity of the SAH domain is important for mitotic arrest in conditions of suppressed microtubule dynamics, and the duration of mitotic arrest dictates the probability, but not the timing, of cell death. Although independent targeting of INCENP to microtubules or the kinetochore/centromere promotes the mitotic checkpoint, it is insufficient for a robust mitotic arrest. Altogether, our results demonstrate that dual recognition of chromatin and microtubules by CPC is important for checkpoint maintenance and determination of cell fate in mitosis.

Introduction

Accurate chromosome segregation requires bipolar attachment of microtubules (MTs) to the kinetochore. Unattached kinetochores activate the mitotic checkpoint (or spindle assembly checkpoint [SAC]) to delay anaphase onset while erroneous kinetochore microtubule (kMT) attachments are being corrected (Foley and Kapoor, 2013). Both processes are promoted by the chromosomal passenger complex (CPC), composed of inner centromere protein (INCENP), Survivin, Borealin (also known as Dasra and CDCA8), and the kinase Aurora B (Carmena et al., 2012; Trivedi and Stukenberg, 2016).

The CPC regulates error correction and the SAC by phosphorylating multiple substrates at the kinetochore. First, Aurora B destabilizes kMT attachment by phosphorylating the MT-binding protein Hec1 (Ndc80; DeLuca et al., 2006; Welburn et al., 2010), generating unattached kinetochores that can signal the SAC (Etemad et al., 2015; Tauchman et al., 2015). Second, Aurora B promotes kinetochore recruitment of Mps1 (Saurin et al., 2011; van der Waal et al., 2012; Nijenhuis et al., 2013; Zhu et al., 2013), which stimulates the SAC by phosphorylating KNL1 (London et al., 2012; Sheppard et al., 2012; Yamagishi et al., 2012; Vleugel et al., 2015). Phosphorylated KNL1 further recruits the SAC proteins Bub1, Bub3, BubR1, Mad1, and Mad2 (Zich et al., 2012; Primorac et al., 2013; Tipton et al.,

2013; London and Biggins, 2014). Third, Aurora B promotes kinetochore recruitment of KNL1 and the Ndc80 complex by phosphorylating Dsn1, a subunit of the Mis12 complex (Yang et al., 2008; Akiyoshi et al., 2013; Kim and Yu, 2015). Finally, Aurora B antagonizes protein phosphatase 1 (PP1)-mediated silencing of the SAC by phosphorylating the PP1 binding motif on KNL1 to prevent PP1 localization (Liu et al., 2010; Rosenberg et al., 2011). Aurora B-dependent phosphorylation is high on unattached or erroneously attached kinetochores but low on bioriented kinetochores that are under MT-dependent tension (Knowlton et al., 2006; Liu et al., 2009; Welburn et al., 2010; DeLuca et al., 2011). How Aurora B-dependent kinetochore phosphorylation responds to kMT attachment status remains unclear.

Aurora B activation depends on its interaction with the C-terminal IN-box motif of INCENP and on autophosphorylation of Aurora B and INCENP (Adams et al., 2000; Bishop and Schumacher, 2002; Honda et al., 2003; Sessa et al., 2005). Because this autophosphorylation is facilitated by local enrichment of the CPC (Kelly et al., 2007), Aurora B activity is often coupled to its localization. During early mitosis, the CPC is enriched at the inner centromere through Survivin and Borealin (Gassmann et al., 2004; Sampath et al., 2004), which form a trimeric complex with the N-terminal CEN domain of INCENP (Klein et al., 2006; Jeyaprakash et al., 2007). Survivin

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Abbreviations used: CPC, chromosomal passenger complex; CSF, cytosolic factor; DoM, duration of mitosis; FL, full length; INCENP, inner centromere protein; IQR, interquartile range; kMT, kinetochore microtubule; MT, microtubule; MTBD, microtubule-binding domain; NEB, nuclear envelope breakdown; PP, protein phosphatase; PRD, phospho-regulatory domain; SAC, spindle assembly checkpoint; SAH, single α -helix.

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interacts directly with histone H3 phosphorylated at threonine 3 (H3T3ph; Kelly et al., 2010; Wang et al., 2010; Yamagishi et al., 2010), whereas Borealin indirectly binds histone H2A phosphorylated at threonine 120 (H2A T120ph; Tsukahara et al., 2010). However, the roles of CPC at the centromere in KMT regulation and SAC activation have been questioned in budding yeast (Campbell and Desai, 2013).

The CPC also interacts weakly with spindle MTs during early mitosis (Tseng et al., 2010). The interaction of Aurora B and EB1 at growing MT ends stimulates recruitment of the CPC to the inner centromere by promoting feedback between Aurora B and Bub1 (Banerjee et al., 2014). Ubiquitylated Aurora B also interacts with UBASH3B/MKLP2 on MTs and is required to concentrate the CPC at the inner centromere (Krupina et al., 2016). In addition, the CPC binds MTs directly through the single α -helix (SAH) domain (previously termed the putative coiled-coil domain) of INCENP (Mackay et al., 1993; Tseng et al., 2010; Samejima et al., 2015; van der Horst et al., 2015). The SAH domain is essential for viability in chicken DT40 cells, effective Dsn1 phosphorylation, and CPC relocation to the spindle midzone at anaphase in human cells (Samejima et al., 2015; van der Horst et al., 2015). It was also reported that deleting the SAH domain attenuates the SAC in taxol-treated human cells without affecting centromeric localization of the CPC (Vader et al., 2007). Here, we investigate the molecular mechanism by which the SAH domain regulates kinetochore phosphorylation by Aurora B and contributes to the SAC.

Results

The INCENP SAH domain is required for CPC localization to chromatin and the SAC in *Xenopus* egg extracts

The nocodazole-induced SAC depends on Aurora B in *Xenopus laevis* egg extracts (Kallio et al., 2002; Vigneron et al., 2004; Gadea and Ruderman, 2005). To determine whether the INCENP SAH domain is required for the SAC in this system, the CPC was depleted from M-phase extract and reconstituted with GFP-tagged full-length *Xenopus* INCENP (xINCENP) or xINCENP Δ SAH (Fig. 1 A). The SAC was assayed by adding a high concentration of sperm nuclei and nocodazole, followed by calcium to release the cytosolic factor (CSF)-mediated metaphase arrest (Minshull et al., 1994). Mock-depleted extract remained in M phase with high H3T3ph levels for the duration of the assay, indicative of a SAC arrest, whereas CPC-depleted extract (Δ CPC) exited into interphase within 30 min after calcium addition. This defect was fully rescued when the extract was reconstituted with CPC containing xINCENP but not xINCENP Δ SAH, indicating that the SAH domain is required for the SAC in *Xenopus* egg extract. Although xINCENP was broadly distributed along sperm chromatin with enrichment near the centromere, the level of xINCENP Δ SAH on chromatin was dramatically reduced (Fig. 1 B). A similar localization defect on preassembled CENP-A chromatin arrays in M-phase extract was also observed (Fig. 1 C).

When a GFP-tagged SAH domain fragment (xSAH) was expressed in M-phase extract, it localized weakly and broadly along sperm chromatin (Fig. 1 D). This localization was independent of endogenous INCENP, as 6Myc-tagged xSAH

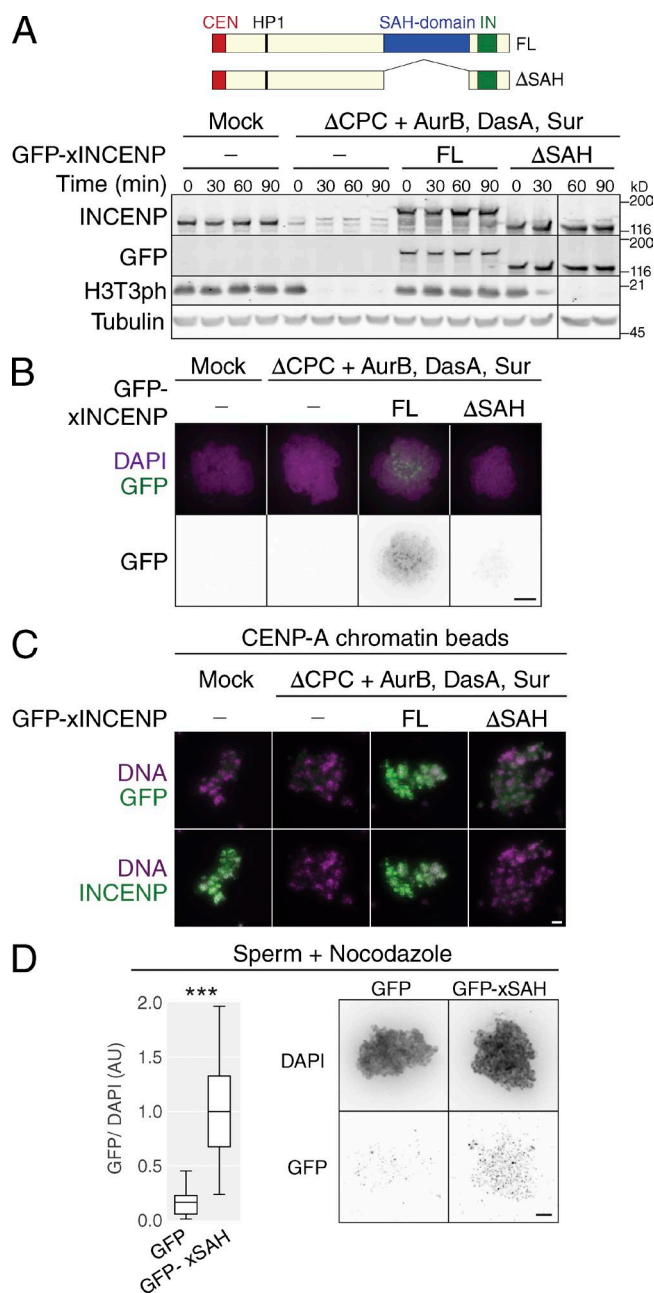


Figure 1. The SAH domain is required for CPC localization and the SAC in *Xenopus*. (A) Top, diagram of INCENP constructs. CEN, centromere-targeting domain; IN, IN-box; HP1, HP1-binding motif. Bottom, *Xenopus* egg extracts were either mock depleted (Mock) or depleted of CPC (Δ CPC); reconstituted with xAurora B, xDasra A, xSurvivin, and FL or Δ SAH GFP-INCENP; and maintained at M-phase with CSF. After addition of high concentrations of sperm and nocodazole, CSF was inactivated by calcium to monitor SAC-mediated maintenance of M phase. Time after calcium addition is indicated. Presence of H3T3ph, monitored by Western blot, marks M phase. (B) Immunofluorescence of GFP-xINCENP on sperm chromatin in M-phase extract treated with nocodazole. Green, GFP; purple, DAPI. Bar, 5 μ m. (C) Immunofluorescence of GFP-xINCENP (green) on preformed CENP-A chromatin arrays (purple) in M-phase extracts. Bar, 6 μ m. (D) Immunofluorescence of GFP or GFP-xSAH on sperm chromatin in M-phase extracts treated with nocodazole. Median, interquartile range (IQR; box), and $\pm 1.5 \times$ IQR (error bars) are shown. Two-tailed Mann-Whitney t test; $n = 20$ spreads per sample; ***, $P \leq 0.001$. Bar, 2 μ m.

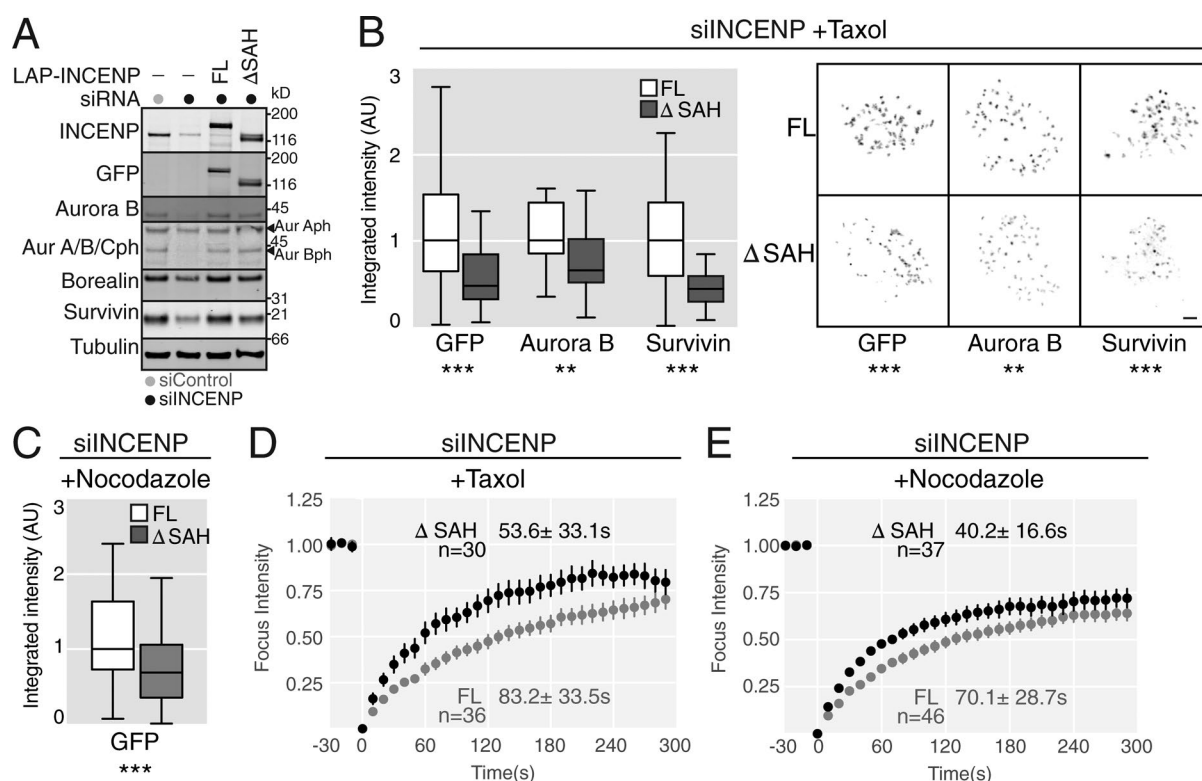


Figure 2. The SAH domain is required for CPC localization at the centromere in human cells. (A) Western blot of HeLa cell extracts reconstituted as indicated. Anti-Aurora A/B/Cph detects the phosphorylated T-loop of Aurora A and Aurora B. (B and C) Integrated intensity at the centromere was quantified for each epitope in taxol (B) or nocodazole (C). Data represent $n \geq 28$ cells in B and $n \geq 74$ cells aggregated from three independent experiments for C. Median, IQR (box), and $\pm 1.5 \times$ IQR (error bars) are shown. Two-tailed Mann-Whitney t test; **, $P \leq 0.01$; ***, $P \leq 0.001$. Bar, 2 μ m. AU, arbitrary units. (D and E) FRAP of centromeric foci in taxol-treated (D) or nocodazole-treated (E) cells reconstituted with hINCENP or hINCENP Δ SAH. The mean FRAP recovery curve with a 95% confidence interval is displayed with the mean $t_{1/2} \pm$ SD for each sample; data compiled from $n = 2$ independent experiments per condition, $n \geq 30$ centromeres for each condition.

localized to chromatin even after depletion of endogenous CPC (Fig. S1 A), and endogenous INCENP did not copurify with the 6Myc-tagged xSAH domain (Fig. S1 B). These results indicate that in addition to Survivin and Borealin, the SAH domain interacts with chromatin to support the SAC in *Xenopus* egg extracts.

The SAH domain supports CPC localization at the centromere in human cells

To test whether the SAH domain contributes to centromeric localization of the CPC in human cells, we generated doxycycline-inducible HeLa T-REx cell lines containing full-length human INCENP (hINCENP) or hINCENP Δ SAH. These constructs are resistant to an INCENP siRNA and N-terminally fused to the LAP tag, which contains GFP (Cheeseman and Desai, 2005; Vader et al., 2006). Treating these cells with INCENP siRNA reduced endogenous INCENP protein levels by 80%, whereas doxycycline induced LAP-hINCENP expression to near-wild-type levels (Fig. 2 A). As previously published (Honda et al., 2003), knockdown of INCENP reduced the abundance of the other CPC subunits, but it was rescued by expressing either hINCENP or hINCENP Δ SAH (Fig. 2 A). Coimmunoprecipitation of CPC subunits with hINCENP Δ SAH confirmed that the SAH domain is dispensable for CPC formation (Fig. S2 A).

hINCENP and hINCENP Δ SAH both showed the appropriate spatiotemporal distribution, including enrichment at the centromere during pro-/metaphase and transfer to the spindle

midzone during anaphase (Fig. S2, B and C). Compared with hINCENP, however, hINCENP Δ SAH was less abundant at the centromere and at the spindle midzone, consistent with our observations in *Xenopus* (Fig. 1 B) and the previously published role of the hINCENP SAH domain in midzone localization (Samejima et al., 2015; van der Horst et al., 2015), respectively. Quantitation confirmed that expression of hINCENP Δ SAH significantly reduced the abundance of all tested CPC components at the centromere relative to hINCENP without affecting the number of INCENP-positive centromeres in either nocodazole- or taxol-treated cells (Fig. 2, B and C, and Fig. S2 D). Reduced CPC abundance at the centromere was accompanied by a slight reduction in centromeric H3T3ph and the appearance of H2A T120ph on chromosome arms (Fig. S2 E). These defects are consistent with the idea that reduced CPC abundance causes defects in Aurora B-dependent activation of the H3T3 kinase Haspin and Aurora B-dependent recruitment of the H2A T120 kinase Bub1 to the kinetochore (Carmena et al., 2012).

To determine whether the SAH domain contributes to CPC stability at the centromere, we performed FRAP on hINCENP or hINCENP Δ SAH foci at the centromere in mitotic cells treated with taxol (Figs. 2 D and S2 F). hINCENP had a mean half-time of recovery ($t_{1/2}$) of 83.2 ± 33.5 s, whereas INCENP Δ SAH had a reduced $t_{1/2}$ (53.6 ± 33.1 s). hINCENP Δ SAH also showed a reduced $t_{1/2}$ in nocodazole-treated cells relative to full-length (FL) hINCENP. Additionally, the maximum intensity of recovery for hINCENP Δ SAH is higher than that of hINCENP

FL, suggesting that a larger fraction of the population is mobile (Figs. 2 E and S2 F). These results suggest that the SAH domain helps the CPC stabilize on centromeric chromatin in both *Xenopus* egg extracts and human somatic cells.

The SAH domain is required for SAC maintenance and mitotic cell death in taxol-treated human cells

A previous study investigating the role of the SAH domain in the SAC used flow cytometry to measure the relative abundance of mitotic cells (Vader et al., 2007). However, that analysis did not determine the duration of the mitotic arrest or how cells exit mitosis. Therefore, we measured the duration of mitosis (DoM) for single cells using live microscopy in the presence of 500 nM taxol, a concentration known to result in the longest DoM in HeLa cells (Yang et al., 2009). We defined the DoM as the time from nuclear envelope breakdown (NEB) to mitotic exit, which occurred by either slippage into interphase or death in mitosis. In taxol, all cells undergoing slippage produced a single, multinucleate cell because of an inability to complete cytokinesis, whereas cells that died in mitosis exhibited dramatic blebbing followed by compaction into a crenulated spherical mass (Fig. S3 A).

Cells treated with control siRNA exhibit a heterogeneous arrest time in taxol, with a median DoM of 18.6 h (Fig. 3 A). Consistent with the CPC being required for SAC maintenance, INCENP siRNA decreased the median DoM to 8.3 h. Although expression of hINCENP fully rescued the DoM (18.7 h), hINCENP Δ SAH did not (11.1 h), confirming that the SAH domain is required to maintain the SAC in taxol-treated human cells. Treating cells with monastrol, which inhibits spindle bipolarization and activates the SAC (Kapoor et al., 2000), demonstrated a similar requirement for the CPC and the INCENP SAH domain (Fig. 3 B).

To assess how manipulation of INCENP affects mitotic cell fate (slippage or death in mitosis), we aggregated data from at least five independent replicates of our taxol-induced SAC experiment by normalizing the DoM for each construct to the median DoM of hINCENP in the same replicate (Fig. 3 C). We then visualized the normalized DoM by cell fate for each construct. Although the majority of control cells (84%) or those expressing INCENP (71%) died in mitosis, the majority of cells lacking the CPC (75%) or containing INCENP Δ SAH (59%) underwent mitotic slippage. It is known that impairing the SAC shortens the duration of mitotic arrest in taxol, leading to an increased frequency of mitotic exit by slippage (Gascoigne and Taylor, 2008). Our data are consistent with this observation, as cells undergoing mitotic slippage arrested for half as much time in mitosis when expressing hINCENP Δ SAH relative to hINCENP.

To better correlate the DoM with cell fate, we performed a linear regression of the median DoM against cell survival for every construct in every experiment in this article, in addition to experiments not depicted (Fig. S3 B). We saw a strong, positive correlation between the median DoM of a sample and the percentage of cells dying in mitosis in that sample. The DoM of cells that die in mitosis was relatively constant, indicating that the probability of mitotic cell death, but not the timing of cell death, strongly correlates with the capacity of INCENP to hold the SAC. Thus, the CPC and the INCENP SAH domain ensure a robust SAC response in taxol, which promotes mitotic death.

The SAH domain is required for SAC protein recruitment and kinetochore phosphorylation in taxol-treated human cells

Next, we investigated the importance of the SAH domain in recruiting SAC regulators to the kinetochore. Although it was reported that BubR1 recruitment is qualitatively not affected by deleting the SAH domain in U2OS cells (Vader et al., 2007), HeLa cells expressing INCENP Δ SAH exhibited a quantitative reduction in the amount of Mad1, Bub1, and BubR1 at the kinetochore compared with hINCENP (Fig. 3 D), without affecting the number of Bub1/BubR1-positive kinetochores (Fig. S2 D). hINCENP Δ SAH showed no defect in recruitment of CENP-A or Hec1 (Fig. S3 C) or in Aurora B-dependent phosphorylation of CENP-A at the centromere or H3S10 on chromatin (Fig. 3 E). However, hINCENP Δ SAH had a reduced amount of phosphorylation on kinetochore substrates implicated in KMT stability and SAC maintenance, such as Dsn1 S100ph, KNL1 S24ph, and Hec1 S44ph (Hsu et al., 2000; Zeitlin et al., 2001; Welburn et al., 2010; DeLuca et al., 2011; Fig. 3, E and F; and Fig. S3 D). These results, together with a previous study (Samejima et al., 2015), suggest that the SAH domain can promote the SAC by supporting Aurora B-dependent kinetochore phosphorylation and SAC protein recruitment.

Neither cytoplasmic activation of Aurora B nor tethering of INCENP to the centromere/kinetochore is sufficient to support the SAC

A study in budding yeast has suggested that centromere targeting of the CPC is not required for its functions, including SAC activation, if Aurora B is activated by an alternative mechanism such as binding to MT (Campbell and Desai, 2013). Thus, we tested whether artificial activation of Aurora B by forced dimerization could bypass the requirement of the SAH domain for the SAC. As previously shown in *Xenopus* egg extracts (Tseng et al., 2010), replacing the SAH domain with the dimerization domain of GCN4 (hINCENP Δ SAH Δ VCN4) dramatically increased the overall level of Aurora B autophosphorylation in human cells (Fig. 4, A and B). However, this construct phenocopied hINCENP Δ SAH, failing to rescue the defect in INCENP centromeric localization (Fig. S3 E), kinetochore phosphorylation (Fig. S3 E), and the SAC (Fig. 4 C). These results suggest that cytoplasmic activation of Aurora B is insufficient to support the SAC in taxol.

We next tested whether tethering INCENP to the centromere or the kinetochore bypasses the requirement of the SAH domain. We generated HeLa cell lines in which the N-terminal CEN domain of INCENP was replaced with one of the following targeting modules: FL human Mis12 (Mis12-hINCENP Δ CEN) for kinetochore targeting, FL CENP-B (CB-hINCENP Δ CEN) for stable centromere targeting, or CENP-B DNA binding domain (CB^{DBD}-hINCENP Δ CEN) for more dynamic centromere targeting (Fig. 4 D; Liu et al., 2009; Wang et al., 2011). We then depleted endogenous INCENP, expressed the targeting chimeras, and monitored the DoM of these cells in taxol by live imaging (Fig. 4 E). Consistent with the requirement of the Survivin-H3T3ph interaction in SAC maintenance in human cells (Lens et al., 2006; Wang et al., 2010), hINCENP Δ CEN had a significantly shorter DoM than cells treated with siINCENP alone, likely because of hINCENP Δ CEN sequestering Aurora B from remaining endogenous INCENP protein. Surprisingly,

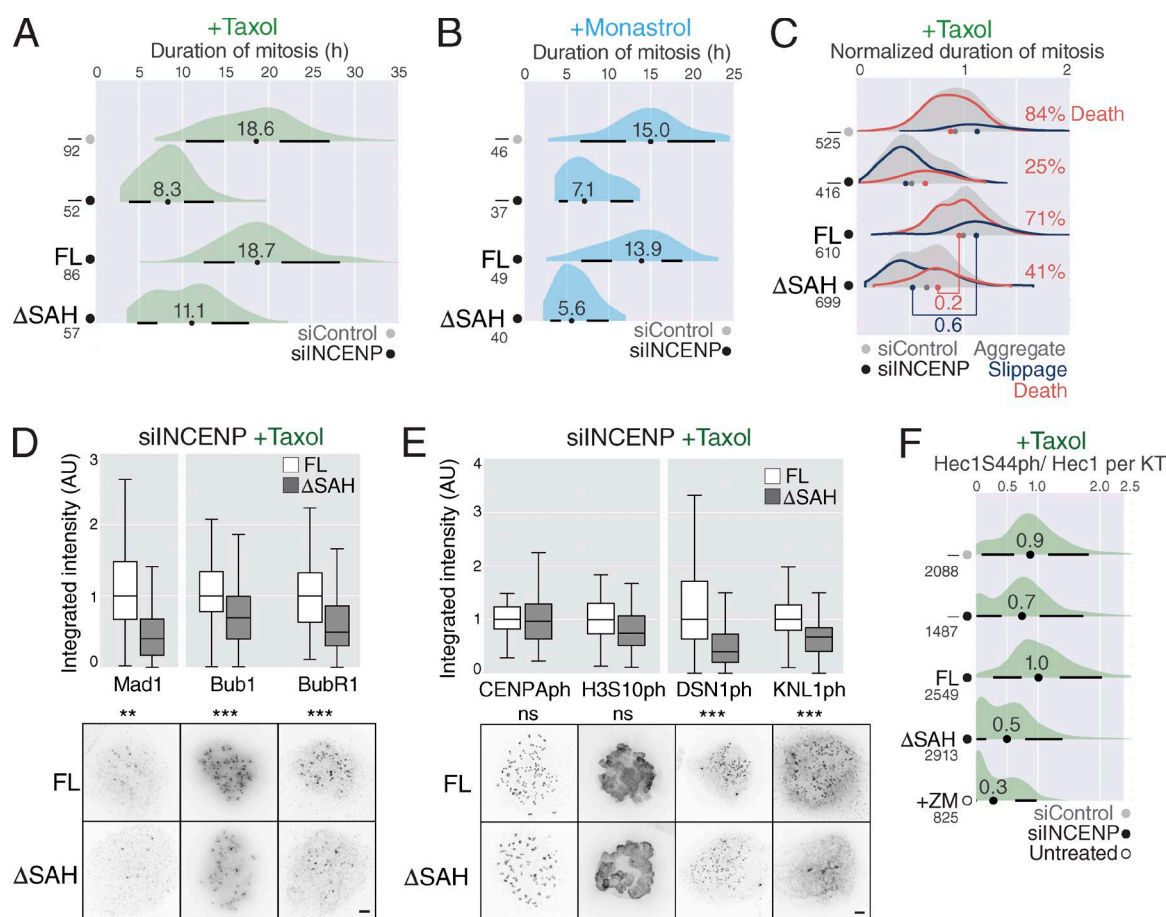


Figure 3. The SAH domain is required for the SAC and kinetochore phosphorylation in human cells. (A) DoM for HeLa cells with taxol, pretreated with control siRNA or INCENP siRNA and complemented with hINCENP FL or hINCENP-ΔSAH (ΔSAH). Data are displayed as a kernel density estimate of DoM for all cells in each condition. A box plot with a circle at the median and whiskers extending to $\pm 1.5 \times \text{IQR}$ is under each density, n values indicated under sample name, siRNA treatment indicated by gray (siControl) or black (siINCENP) circles adjacent to the sample name. (B) Same as A, except cells were treated with monastrol. (C) Normalized DoM for all cells (gray) and for cells undergoing death in mitosis (red) or mitotic slippage (blue). Data aggregated from at least five independent experiments. All values were normalized to the median of hINCENP in that experiment. Maximum height of each curve is proportional to percentage composition of that class in the aggregate. Red numbers on the right indicate the overall percentage of cells dying in mitosis. The difference in the normalized DoM between cells complemented with FL or ΔSAH hINCENP is also indicated. (D and E) Immunofluorescence quantification of SAC proteins (D) and Aurora B-dependent phosphorylation (E) in taxol-treated cells. In D, the data represent integrated intensity at the kinetochore from $n \geq 18$ cells for Mad1 and $n \geq 86$ cells aggregated from three independent experiments for Bub1 and BubR1. In E, the data represent integrated intensity at the centromere for $n \geq 52$ cells aggregated from two experiments for CENP-A S7ph, integrated intensity on chromatin for $n \geq 20$ cells for H3S10ph, and $n \geq 1,800$ individual kinetochore measurements for Dsn1 S100ph and Knl1 S24ph using a marker epitope. Representative images approximating the median are presented. Median, IQR (box), and $\pm 1.5 \times \text{IQR}$ (error bars) are shown. Two-tailed Mann-Whitney t test; ns, not significant; **, $P \leq 0.01$; ***, $P \leq 0.001$. Bars, 2 μm . (F) Immunofluorescence quantification of Hec1 S44ph at individual kinetochores standardized to total Hec1 at that kinetochore. Data are visualized as a kernel density estimate, n values indicated under the sample name. ZM, Aurora B inhibitor ZM447439. The maximum height of the density for ZM-treated cells was scaled independently.

none of the INCENP-targeting chimeras supported a robust SAC response. The DoM of cells in which INCENP was targeted to CENP-B chromatin or the kinetochores was longer than the DoM of cells expressing hINCENP ΔCEN but similar to the DoM of cells treated with siINCENP alone. These data demonstrate that targeting INCENP to the centromere or the kinetochore cannot fully bypass the requirement of the CEN domain.

Failure of these chimeric constructs to support the SAC was unexpected, because they are known to promote kMT destabilization and induce a mitotic delay when expressed on top of endogenous INCENP (Liu et al., 2009; Wang et al., 2011). Consistent with these published results, when expressed on top of endogenous INCENP in the absence of any MT poisons, Mis12-hINCENP ΔCEN caused the mitotic delay much more effectively than hINCENP ΔCEN (Fig. 4 F), whereas overexpression of hINCENP did not induce any mitotic delay (Fig. S3

F). Notably, the untagged version of hINCENP ΔCEN ΔSAH was not as potent as hINCENP ΔCEN, but the effect of Mis12-hINCENP ΔCEN ΔSAH was comparable to that of Mis12-hINCENP ΔCEN. These results indicate that the SAH domain of INCENP that is stably associated with the kinetochore is dispensable for SAC activation.

INCENP MT binding is negatively regulated by mitotic phosphorylation in the regions flanking the SAH domain

Aurora B is critical in human cells for the SAC in the presence of MTs (taxol, monastrol) but less important in the absence of MTs (nocodazole; Ditchfield et al., 2003; Hauf et al., 2003; Santaguida et al., 2011). Therefore, we wondered whether the MT-binding capacity of the SAH domain contributes to the SAC. As a first step, we tested whether a particular region of

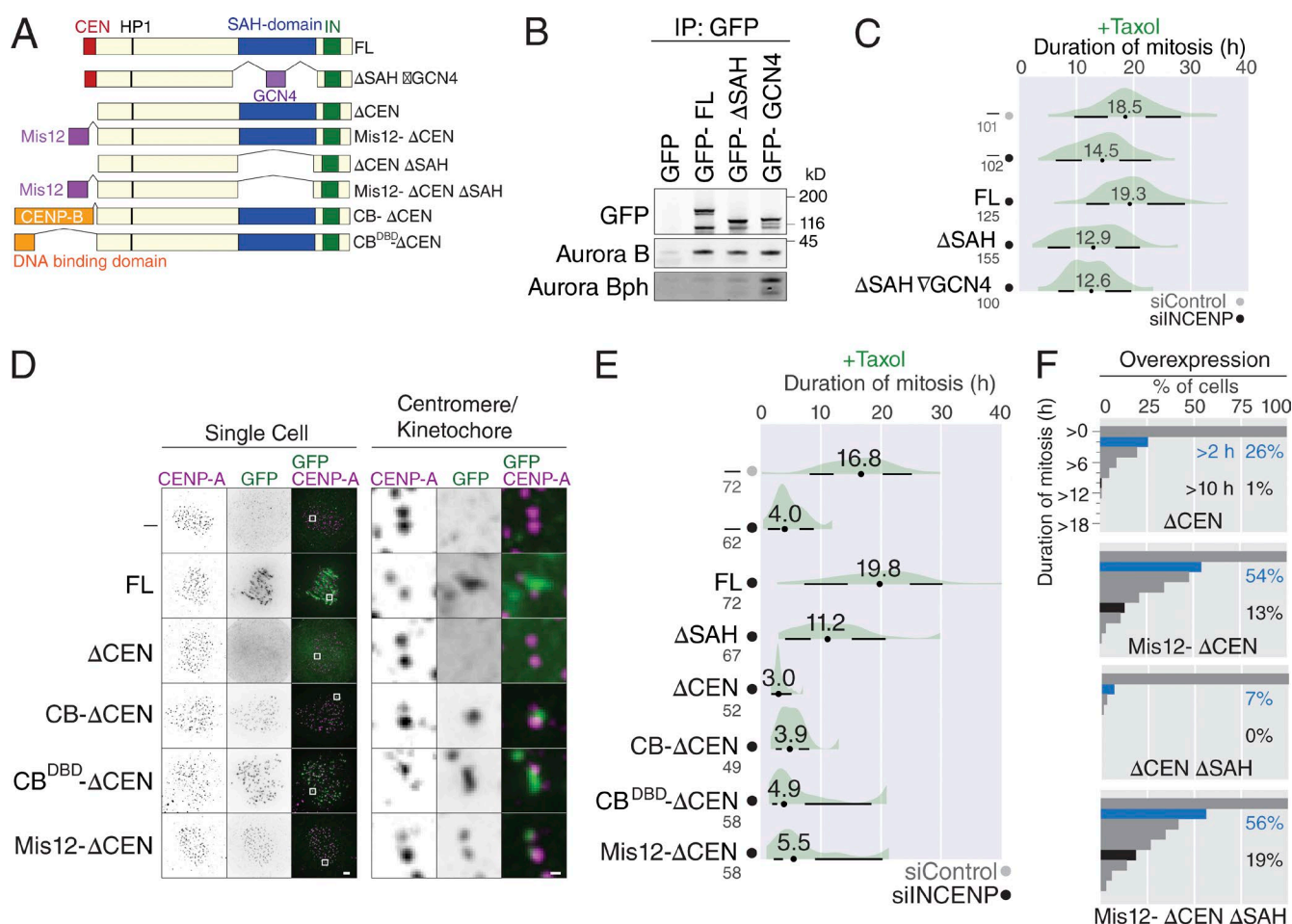


Figure 4. Clustering or kinetochore targeting of the CPC is insufficient to support the SAC. (A) Diagram of LAP-tagged hINCENP constructs. (B) Western blot of anti-GFP immunoprecipitation (IP) of the indicated constructs from mitotic HeLa cell extracts. (C) DoM for cells in taxol. Representative of $n = 2$ independent experiments. (D) Immunofluorescence of various INCENP constructs. Each construct was expressed over endogenous INCENP. For each construct, the left three panels are single-cell images; the right three are zoom-ins of a representative kinetochore pair indicated by white boxes. Bars: (single cell) 2 μ m; (centromere/kinetochore) 0.25 μ m. (E) DoM for cells in taxol. The maximum height for the density estimate of hINCENP ΔCEN was scaled independently. Representative of $n = 2$ independent experiments. (F) Each construct was expressed over endogenous INCENP and synchronized at the G2/M transition by treatment with the Cdk1-inhibitor RO-3306. After washout of the inhibitor, cells synchronously entered mitosis. Plots show the cumulative minimum DoM for each construct. Blue bars, arrested for at least 2 h; black bars, arrested for at least 10 h; $n \geq 90$ cells per sample.

the SAH domain contributes to MT binding using *Xenopus* egg extracts and found that 6Myc-tagged xSAH domain localized to the spindle with particular enrichment at the spindle poles (Fig. S4 A). Using a series of deletions, we mapped this spindle binding activity to the first ~100 aa of the SAH domain. We also identified a shorter 35-aa deletion in the N terminus of the xSAH domain (ΔSAH^N) that was unable to bind the spindle. Thus, the N terminus of the *Xenopus* SAH domain binds the spindle, consistent with reports for the SAH domain of human and chicken (Samejima et al., 2015; van der Horst et al., 2015).

Spindle pole enrichment of the xSAH domain is distinct from xINCENP, which localizes along the entire spindle (Tseng et al., 2010). Searching for elements flanking the SAH domain that might account for this difference, we identified four putative Cdk-dependent phosphorylation sites ([S/T]P) upstream of the SAH domain and two sites downstream of the SAH domain that are widely conserved among vertebrates species, and thus we named these flanking regions phospho-regulatory domain (PRD) 1 and 2, respectively (Figs. 5 A and S4 B).

To determine whether these putative Cdk sites are phosphorylated in *Xenopus* egg extracts, we prepared a fragment encompassing PRD1, SAH, and PRD2 (PRD-xSAH), as well as versions in which the six conserved putative Cdk sites, in addition to one SP site in PRD1 not conserved in humans, were made unphosphorylatable by mutation to alanine (PRD-xSAH 7A) or phosphomimetic by mutation to aspartate (PRD-xSAH 7D; Fig. 5 A). These fragments were incubated in M-phase or interphase extracts, followed by treatment with or without phosphatases (Fig. 5 B). Although the gel migration pattern of xSAH domain alone was not affected by cell-cycle stage or phosphatase treatment, the PRD-xSAH fragment in M phase ran as a smear that collapsed to a single band upon phosphatase treatment, indicative of phosphorylation. The mobility shift of this fragment was greatly reduced in interphase, suggesting that these [S/T]P sites are phosphorylated only in metaphase. Furthermore, the PRD-xSAH 7D fragment ran at the same size as the upper smear of PRD-xSAH, and the PRD-xSAH 7A mutant ran at a similar size to the PRD-xSAH upon phosphatase treatment.

To examine whether phosphorylation of the PRD regulates MT binding, the localization of the PRD-xSAH fragments

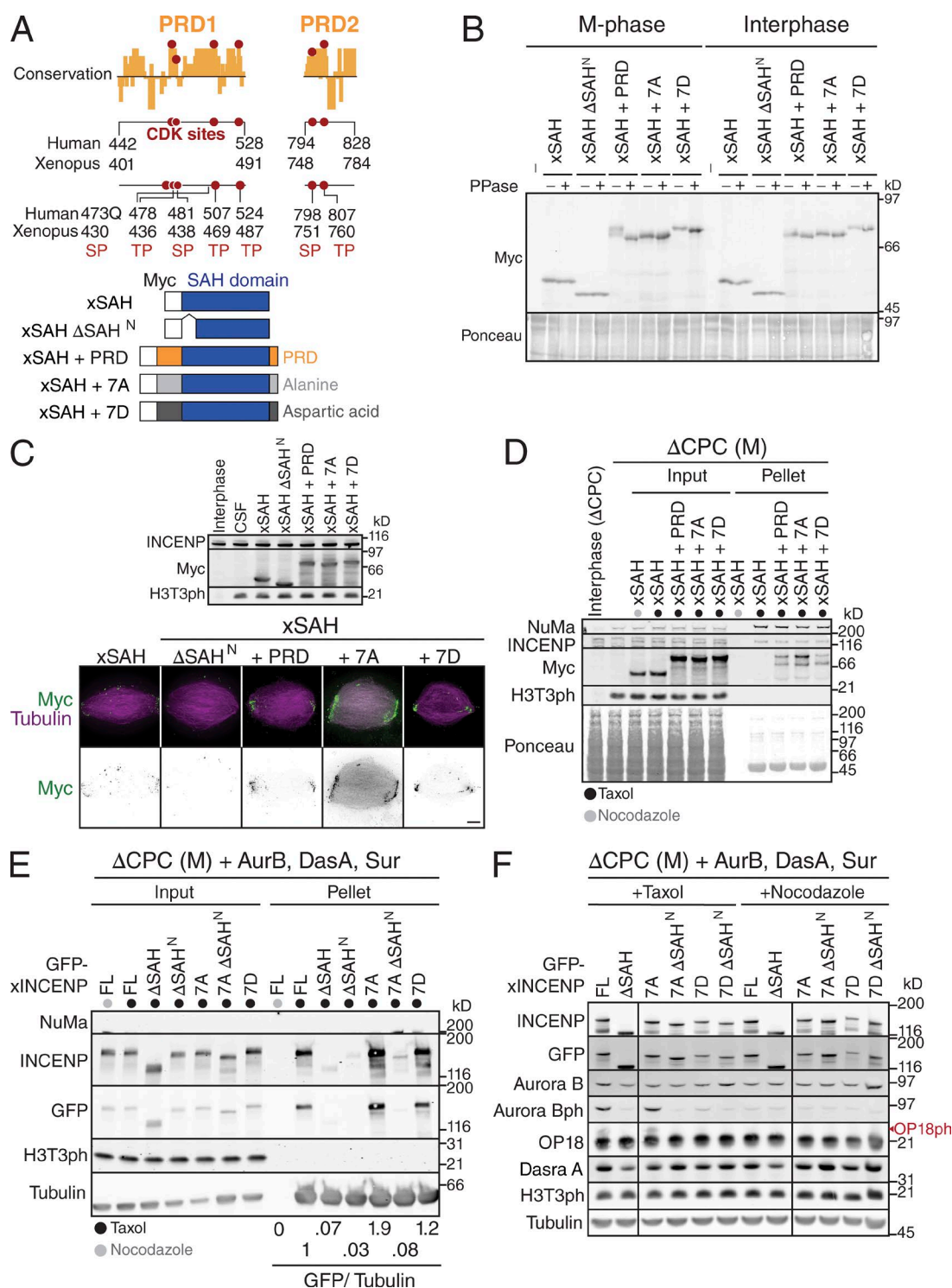


Figure 5. MT binding of the SAH domain is negatively regulated by phosphorylation. (A) Conservation of each residue in the PRDs relative to the mean conservation of residues in INCENP (top). Conservation was determined using conSURF (Landau et al., 2005) with 25 vertebrate INCENP homologs (Fig. S4 B). Putative Cdk sites conserved in at least 22 species are indicated (red circles). Diagrams of 6Myc-tagged constructs used in B, C, and D (bottom). (B) Western blot of various Myc-tagged xSAH domain constructs expressed in *Xenopus* egg extracts. Samples were either taken from extracts (–) or diluted in buffer containing lambda phosphatase and incubated at 30°C for 40 min (+). (C) Immunofluorescence of the indicated Myc-tagged constructs on the spindle in M-phase *Xenopus* egg extract (bottom). Western blot shows equal protein expression (top). Bar, 5 μ m. (D) MT pelleting assay with the indicated constructs after depletion of endogenous INCENP. M-phase egg extract was incubated with nocodazole (gray circle) or taxol (black circles) for 30 min, and MT pellets were analyzed by Western blot. (E) MT pelleting assay as in D in extract reconstituted with the indicated GFP-xINCENP construct, xAurora B, xDasra A, and xSurvivin. Quantitation of pelleted GFP-xINCENP constructs relative to pelleted tubulin was performed in ImageJ and is shown at the bottom. (F) Western blot visualizing the MT-dependent activation of Aurora B. M-phase extract was reconstituted with the indicated INCENP constructs and then incubated in the presence of the indicated drug for 40 min. Aurora B activation was assessed by the appearance of Aurora B autophosphorylation (Aurora Bph) and OP18 hyperphosphorylation (OP18ph).

on the spindle was assessed in M-phase egg extracts. Although the xSAH domain only weakly bound the spindle pole, the PRD-xSAH 7A mutant was highly enriched on the spindle and spindle pole (Fig. 5 C). The PRD-xSAH 7D mutant and the PRD-xSAH fragment localized to the spindle pole only slightly better than the xSAH domain alone. Furthermore, MT cosedimentation assays showed that the SAH domain exhibited little to no MT binding in the absence of the PRD (Fig. 5 D). The PRD-xSAH 7A fragment exhibited a twofold increase in sedimentation with MTs compared with PRD-xSAH. These results indicate that the PRD and SAH domain promote MT binding, which is negatively regulated by Cdk-dependent phosphorylation at the PRD.

To test whether the PRD regulates the MT binding of xINCENP FL, we generated versions of GFP- xINCENP containing the phospho-null (xINCENP 7A) and phospho-mimetic (xINCENP 7D) mutations either alone or in combination with the 35-aa deletion that abrogates the MT binding capacity of the SAH domain (Δ SAH^N). M-phase Δ CPC extracts were reconstituted with the CPC containing these INCENP constructs, and the MT cosedimentation assay was performed (Fig. 5 E). Although xINCENP demonstrated robust MT binding, this was dramatically reduced in xINCENP Δ SAH and xINCENP Δ SAH^N. xINCENP 7A showed an almost twofold increase in MT binding, whereas xINCENP 7D bound to MTs similarly to xINCENP. Incorporating Δ SAH^N into INCENP 7A abrogated MT binding, suggesting that the PRD is insufficient to support MT binding in the absence of the SAH domain.

We have previously shown that MTs assembled by taxol in *Xenopus* egg extracts activate Aurora B in a manner dependent on the SAH (Tseng et al., 2010). Indeed, the CPC reconstituted with xINCENP and xINCENP 7A exhibited robust Aurora B autophosphorylation (Fig. 5 F). However, deleting the MT-binding region of the SAH domain (xINCENP Δ SAH, Δ SAH^N, or 7A Δ SAH^N) or xINCENP 7D prevented MT-dependent activation of Aurora B. These results indicate that the SAH^N region and dephosphorylation of the PRD are required for MT-dependent activation of Aurora B.

The SAH domain binds MTs to maintain the SAC in taxol-treated cells

In *Xenopus* egg extracts, INCENP is still functional in spindle assembly when the SAH domain is replaced with an exogenous MT-binding domain (MTBD; Tseng et al., 2010). We adopted a similar strategy to determine whether the SAH domain binds MTs to sustain the SAC in taxol-treated HeLa cells. We replaced the SAH domain with the MTBD from MAP4 (hINCENP Δ SAH ∇ MAP4; Fig. 6 A) and confirmed that this construct localized to both MTs and chromatin (Fig. 6 B). Although the median DoM for hINCENP Δ SAH in taxol was 10.9 h, hINCENP Δ SAH ∇ MAP4 partially rescued this defect with a median DoM of 13.6 h and completely rescued the defect in mitotic cell fate and BubR1 localization (Fig. 6, C and D), suggesting that the SAH domain supports the SAC through MT binding.

Given that the SAH domain interacts with both chromatin and MTs, we sought to more precisely disrupt MT binding by deleting the 35 aa in the N terminus of the INCENP SAH domain required for MT binding in *Xenopus* (hINCENP Δ SAH^N). Although hINCENP had a median DoM of 16.6 h, hINCENP Δ SAH^N arrested for 12.9 h (Fig. 6 C). Several lines of evidence indicate that this defect was a result of attenuated

MT-binding activity, rather than a defect in chromatin binding or the length of the SAH domain. First, the levels of hINCENP Δ SAH^N at the centromere were indistinguishable from those of hINCENP (Fig. S3 E). Second, two larger deletions (47 and 107 aa) in the SAH domain downstream of the MT-binding region caused a relatively minor reduction in DoM compared with the 35-aa deletion in Δ SAH^N (Fig. S5 A). Third, replacing the N terminus of the SAH domain with the MAP4-MTBD (hINCENP Δ SAH^N ∇ MAP4) completely rescued the SAC defect (Fig. 6 C). Finally, although the SAH domain was required for BubR1 recruitment and Dsn1 S100ph at the kinetochore in taxol (Fig. 3, D and E), it became dispensable when MTs were depolymerized using nocodazole (Fig. 6 E). hINCENP Δ SAH also supported relatively higher levels of Hec1 S44ph at the kinetochore in nocodazole (Fig. 6 F) than in taxol (Fig. 3 F), whereas the levels of KNL1 S24ph and Bub1 recruitment were similarly defective in both conditions (Fig. 3, D and E, and Fig. 6 E). These observations are consistent with the idea that human INCENP binding to MTs through its SAH domain supports kinetochore phosphorylation and checkpoint activation in the presence of MTs.

INCENP PRD phosphorylation attenuates MT binding and the SAC in human cells

To test the functional significance of PRD phosphorylation, which suppresses MT binding of the SAH domain, we treated HeLa cells with siRNA against endogenous hINCENP and induced the expression of mutants in which Cdk sites in the PRD were mutated to alanine (hINCENP 6A) or the phosphomimetic aspartate (hINCENP 6D; Fig. 7 A). As predicted, hINCENP 6A localized prominently to MTs, whereas hINCENP and hINCENP 6D did not (Fig. 7, B–D). Expression of hINCENP 6A caused a mitotic delay without any MT poisons, driven by an increase in the time from metaphase alignment to anaphase onset, whereas constructs defective in MT binding (6D and Δ SAH) initiated anaphase in slightly less time than hINCENP (Figs. 7 D and S5 B). In the presence of taxol, the DoM of hINCENP 6A was significantly longer than that of hINCENP 6D, which phenocopied the SAC defect of hINCENP Δ SAH and Δ SAH^N (Fig. 7 E). These results indicate that the capacity of INCENP to support the SAC can be regulated by PRD phosphorylation status, such that PRD phosphorylation attenuates MT binding and promotes SAC silencing.

Finally, we asked whether hINCENP MT binding could support the taxol-mediated SAC independent of chromatin binding. We deleted the hINCENP CEN domain and mutated the PRD to alanine (hINCENP Δ CEN 6A), which enhances MT binding, or aspartate (hINCENP Δ CEN 6D; Fig. 7 A). As we saw previously (Fig. 4 E), expression of hINCENP Δ CEN reduced the median DoM relative to INCENP siRNA alone from 8.6 to 4.3 h (Fig. 7 F). hINCENP Δ CEN 6A interacted with the spindle and modestly increased the DoM to 5.8 h. This increase was dependent on MT binding, as the same construct with a deletion in the N terminus of the SAH domain (hINCENP Δ CEN 6A Δ SAH^N) bound weakly to the spindle and had a median DoM of 4 h. Furthermore, hINCENP Δ CEN 6D had a median DoM identical to that of hINCENP Δ CEN. Collectively, these results suggest that targeting INCENP to the mitotic spindle can contribute to SAC maintenance independently of binding to chromatin, though it is insufficient to fully support the SAC without the CEN domain of INCENP.

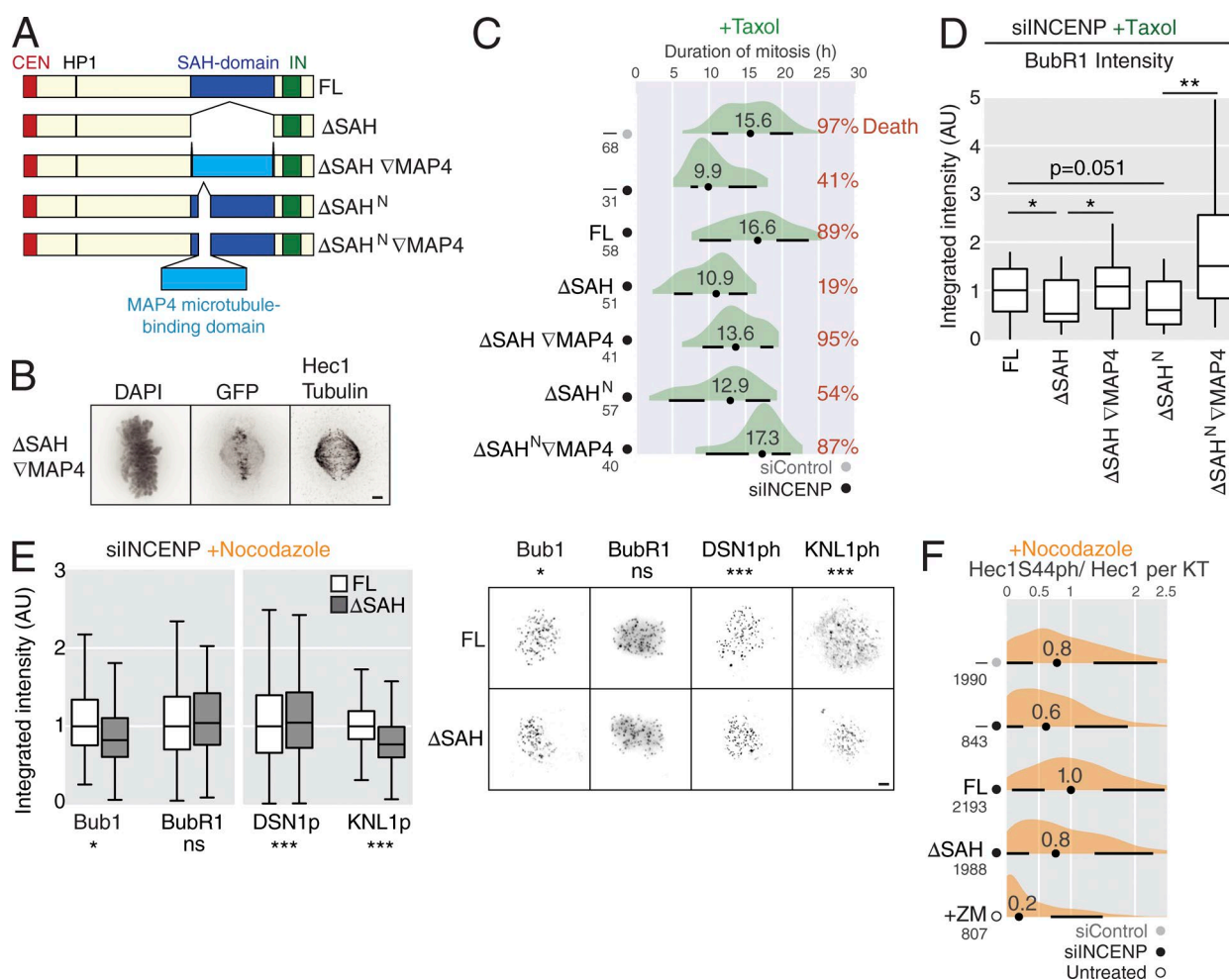


Figure 6. The SAH domain binds MTs to sustain the SAC. (A) LAP-tagged hINCENP constructs. (B) Immunofluorescence of hINCENP Δ SAH Δ MAP4 expressed over endogenous INCENP in asynchronous HeLa cells. Bar, 2 μ m. (C) DoM for cells in taxol. Red values are the percentage cell death for each construct. (D and E) Immunofluorescence quantification of BubR1 in taxol-treated cells (D) and SAC proteins and Aurora B-dependent phosphorylation in nocodazole-treated cells (E) expressing the indicated constructs. In D, the data represent integrated intensity at the kinetochore from $n \geq 27$ cells. In E, the data represent integrated intensity at the kinetochore from $n \geq 54$ cells aggregated from three independent experiments for Bub1 and BubR1, and $n \geq 1,720$ individual kinetochore measurements for Dsn1 S100ph and Knl1 S24ph using a marker epitope. Representative images approximate the median. Two-tailed Mann–Whitney t test; ns, not significant; *, $P \leq 0.05$; **, $P \leq 0.01$; ***, $P \leq 0.001$. Bar, 2 μ m. (F) Immunofluorescence quantification of Hec1 S44ph at individual kinetochores standardized to total Hec1 at that kinetochore for cells treated with nocodazole. The data are visualized as a kernel density estimate, n values indicated under the sample name. ZM, the Aurora B inhibitor ZM447439. The maximum height of the density for ZM-treated cells was scaled independently.

Discussion

The INCENP SAH domain maintains the CPC on chromatin

Here we demonstrate that in addition to Survivin and Borealin, which interact with the INCENP CEN domain, the INCENP SAH domain contributes to recruitment of the CPC to the centromere in both *Xenopus* egg extract and HeLa cells (Fig. 8). Deleting the SAH domain reduces CPC abundance and enhances its dynamics at the centromere even in the absence of MTs, indicating that this function is independent of the previously suggested role for the Aurora B–MT interaction in CPC enrichment at the centromere (Banerjee et al., 2014; Krupina et al., 2016). Given that the GFP-tagged xSAH domain is broadly distributed along sperm chromatin in *Xenopus* egg extract in the absence of MTs (Fig. 1 D), the SAH domain may enhance the affinity of the CPC for chromatin.

Although centromere targeting of the CPC is dispensable in budding yeast (Campbell and Desai, 2013), the INCENP CEN domain is required for robust SAC maintenance in human cells (Fig. 4 E), and the SAH domain promotes the SAC in nocodazole-treated *Xenopus* egg extracts (Fig. 1 A). However, artificially targeting INCENP Δ CEN to the centromere (CENP-B fusion) or the kinetochore (Mis12 fusion) did not bypass the requirement for the CEN domain in the taxol-induced SAC (Fig. 4 E). This may be caused by a defect in chromatin-dependent Aurora B activation, which may require a process beyond chromatin targeting, perhaps involving Borealin-mediated dimerization (Bourhis et al., 2009; Bekier et al., 2015; van der Horst et al., 2015). However, even though INCENP dimerization by GCN4 induced Aurora B autoactivation (Fig. 4 B; Tseng et al., 2010), this did not bypass the requirement for the SAH domain in the SAC maintenance, suggesting that activation of Aurora B is insufficient

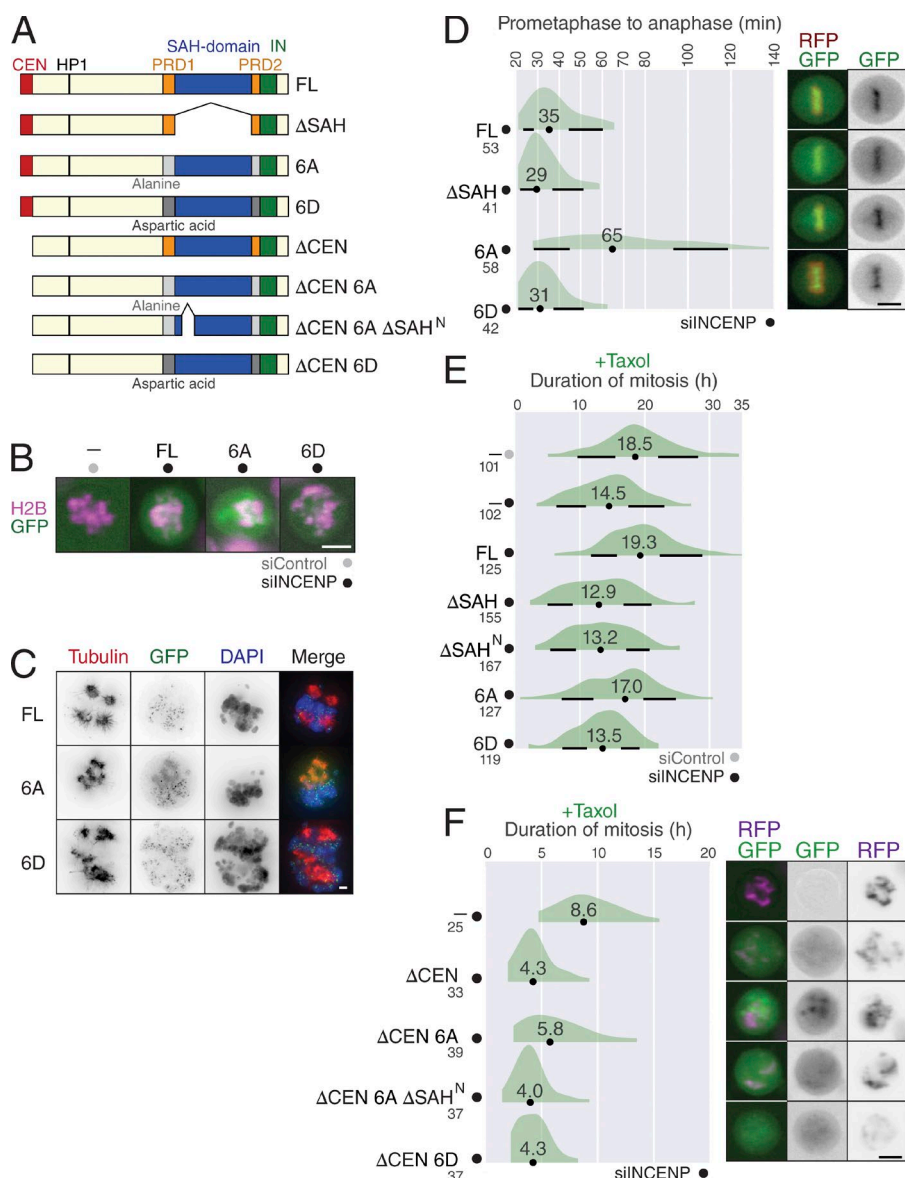


Figure 7. PRD phosphorylation regulates INCENP MT binding and the SAC. (A) LAP-tagged hINCENP constructs. (B) Fluorescence images of cells expressing the indicated hINCENP constructs in taxol after knockdown of endogenous INCENP. Taken from a live imaging experiment. Green, GFP-INCENP; purple, RFP-H2B. Bar, 10 μm. (C) Immunofluorescence of the indicated constructs expressed over endogenous INCENP in cells treated with taxol. Bar, 2 μm. (D) DoM for cells undergoing unperturbed mitosis (left) and representative fluorescence images from the live imaging experiment (right). Representative of *n* = 3 independent experiments. Green, GFP-INCENP; red, RFP-H2B. Bar, 10 μm. (E) DoM for cells in taxol. Data were collected from the same experiment as Fig. 4 C. Representative of *n* = 3 independent experiments. (F) DoM for cells in taxol (left) and representative fluorescence images from the live imaging experiment (right). Green, GFP-INCENP; purple, RFP-H2B. Data from a single experiment. Bar, 10 μm.

to support its function at the centromere/kinetochore in the absence of the SAH domain.

The INCENP SAH domain binds MTs to support the SAC in taxol-treated cells

Consistent with previous studies (Samejima et al., 2015; van der Horst et al., 2015), we have shown that the N-terminal segment

of the SAH domain is critical for MT binding. The MAP4 MTBD bypassed the requirement of this segment (Fig. 6 C), suggesting that the MT-binding capacity of the SAH domain is important for SAC activation/maintenance in response to the aberrant KMT attachments generated by taxol or monastrol.

The SAH–MT interaction could promote the SAC through several mechanisms. First, as previously reported

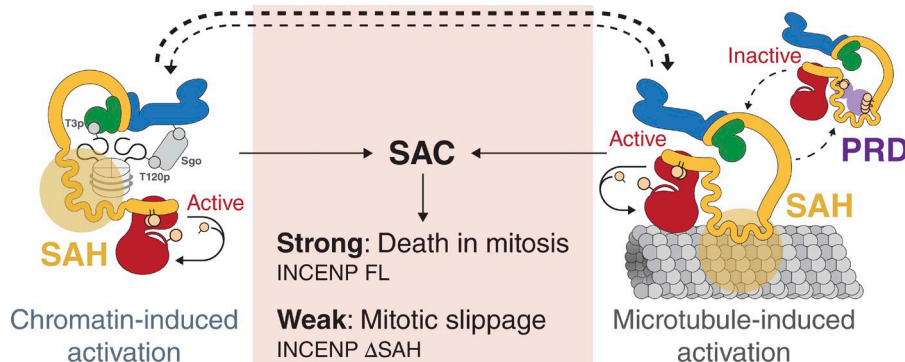


Figure 8. Model for the role of the CPC in the SAC. The SAH domain supports an interaction with chromatin to stabilize the CPC at the centromere (left). Similarly, the SAH domain mediates direct binding to MTs and MT-dependent activation of Aurora B (right). The MT–CPC interaction is suppressed by Cdk-dependent phosphorylation at the PRD of INCENP (top right). After activation, Aurora B may diffuse and phosphorylate substrates at the kinetochore/centromere/kMT (middle) to support SAC activation/maintenance. CPC subunits are Aurora B (red), INCENP (yellow), Survivin (green), and Borealin (blue).

(Banerjee et al., 2014; Krupina et al., 2016), CPC–MT interaction may facilitate CPC enrichment at the centromere. However, this is unlikely to be the sole mechanism, as a 35-aa deletion in the MT-binding region (hINCENP Δ SAH^N) attenuated the SAC without affecting localization to the centromere (Figs. 7 and S5 A). Second, the INCENP SAH domain may support MT-dependent activation of Aurora B (Fig. 5 F; Tseng et al., 2010). Centromeric enrichment of the CPC may enhance the accessibility of the CPC to local kMTs, where Aurora B is activated to phosphorylate kinetochore substrates. Third, SAH–MT binding may support effective phosphorylation of MT-binding proteins, such as Hec1, the Ska complex, mDia3, and EB2, to destabilize kMT attachment (Cheng et al., 2011; DeLuca et al., 2011; Chan et al., 2012; Iimori et al., 2016). Consistent with this model, Aurora B–dependent phosphorylation can be detected on the mitotic spindle (Tseng et al., 2010), and the INCENP–MT interaction preferentially promotes Aurora B–dependent phosphorylation of MT-bound substrates over unbound substrates *in vitro* (Banerjee et al., 2014). This may be related to the fact that Aurora B is a basophilic kinase (Alexander et al., 2011), and that the basic residues of MTBDs, such as those at the Hec1 N-terminal tail and the SAH, commonly interact with the negatively charged E-hooks of tubulin (Alushin et al., 2010; van der Horst et al., 2015). Because kMT attachments are stabilized by tension independently of Aurora B (Akiyoshi et al., 2010), the SAH domain may competitively displace MTBDs at the kinetochore from MTs to facilitate their phosphorylation by Aurora B preferentially at the kinetochore with no tension. Our observation that the SAH is dispensable for ectopic SAC activation by kinetochore targeting is consistent with the idea that the SAH increases Aurora B substrate accessibility, because stabilizing INCENP at the kinetochore would facilitate this in a manner independent of MT-binding (Fig. 4 F). However, the possibility that the SAH domain facilitates Aurora B substrate recognition independently of MT or chromatin binding may not be ruled out. Minor contributions to the SAC by the C-terminal SAH segment that is not supporting MT binding may reflect such functions (Fig. S5 A).

Although the SAH–MT interaction supports the SAC, it is not sufficient for a robust mitotic arrest in the absence of the CEN domain (Fig. 7 F). We propose that the CPC must interact with both centromeric chromatin and MTs to support a robust SAC arrest (Fig. 8).

Cdk-dependent regulation of the INCENP SAH domain

In budding yeast, phosphorylation of the INCENP homologue Sli15 by Cdk1 prevents binding to the mitotic spindle before anaphase (Pereira and Schiebel, 2003). These Cdk1 phosphorylation sites are also enriched in the region upstream of the putative coiled-coil (or SAH) domain in Sli15 (Pereira and Schiebel, 2003), similar to vertebrate INCENP homologs. However, there is a conflict in the literature as to whether phosphorylation of Sli15 supports or inhibits the SAC (Mirchenko and Uhlmann, 2010; Makrantonis et al., 2014). In human cells, Cdk1-dependent phosphorylation at T59 of INCENP indirectly inhibits CPC binding to the spindle by preventing it from binding to MKLP2 (Hümmer and Mayer, 2009). Here, we demonstrate that the PRD enhances SAH-dependent MT binding and that Cdk-dependent phosphorylation of the PRD reduces the binding of INCENP to MTs (Figs. 4 and 8). Furthermore, our data suggest that whereas the SAH–MT interaction contributes to SAC activation in taxol (Fig. 7 E),

suppressing the SAH–MT interaction by phosphorylation of the PRD is required to silence the SAC after metaphase alignment of chromosomes during an unperturbed mitosis (Fig. 7 D), although it is possible that INCENP 6A supports metaphase alignment without bipolar attachment (Kapoor et al., 2006).

Because Cdk1–Cyclin B activity is high in early mitosis, it may be counterintuitive that dephosphorylation of the PRD helps sustain the SAC. However, the INCENP PRD may be targeted for context-dependent dephosphorylation to locally regulate Aurora B activity. At the outer kinetochore, PP2A recruited to BubR1 could facilitate CPC-mediated regulation of kinetochore-proximal MTs (Suijkerbuijk et al., 2012; Kruse et al., 2013; Xu et al., 2013). Given that kinetochore localization of the PP2A B56 subunits are more pronounced in prometaphase than in metaphase (Foley et al., 2011), dephosphorylation of the INCENP PRD may be locally restricted to PP2A-enriched kinetochores during prometaphase. The centromeric PP2A–Sgo1, which inhibits Aurora B activation (Meppelink et al., 2015), may also potentiate MT-dependent activation of Aurora B. It would be interesting to see whether PRD phosphorylation levels change depending on kinetochore attachment status.

The CPC sustains the SAC to promote death in mitosis

Our live-imaging analyses show that the CPC and the INCENP SAH domain sustain the SAC in taxol to promote cell death. We see a strong, linear increase between the median DoM of a sample and the percentage of cells dying in mitosis (Fig. S3 B). The amount of time it took a cell to die in mitosis was similar between all of our constructs, suggesting that the timing of mitotic cell death can be largely uncoupled from the strength of the SAC. This is consistent with the competing-networks model, in which the decision to undergo slippage or death in mitosis is a winner-take-all race between cyclin B degradation and accumulation of pro-apoptotic signals (Gascoigne and Taylor, 2008; Huang et al., 2010; Topham and Taylor, 2013). Our data indicate that even a mild reduction in SAC strength can have a dramatic impact on cell fate during mitotic arrest (Fig. 8).

Materials and methods

Cell culture, siRNA, and drug treatments

HeLa cells were cultured at 37°C with 5% CO₂ in DMEM (Thermo Fisher Scientific) supplemented with 10% tetracycline-free FBS (Atlanta Biologicals) and nonessential amino acids (Thermo Fisher Scientific). HeLa T-Rex cell lines, which stably express the tetracycline repressor, were generated using the manufacturer's protocol from a parental line containing a single FRT integration site and stably integrated RFP-H2B (gift of A. Desai and R. Gassmann, University of California, San Diego, La Jolla, CA; Gassmann et al., 2010).

For live-cell imaging experiments, 300,000 cells were seeded onto glass-bottom dishes (no. 1.5 thickness; MatTek Corporation). After 12–16 h, 100 mg/ml doxycycline was added to induce construct expression, and siRNA was transfected using Lipofectamine RNAi-Max (Thermo Fisher Scientific) and 60 pmol of either ON-TARGET-plus control siRNA (GE Healthcare) or a previously described siRNA against the coding region of human INCENP (5'-UGACACGGAGAU UGCAAC-3'; Vader et al., 2006). After 24–36 h, cells were transferred to fresh culture media lacking phenol red before imaging.

For live-cell RO-3306 (RO) synchronization and release experiments (Figs. 4 F and S3 F), cells were plated as described earlier and

allowed to adhere in the presence of doxycycline for 6 h, before the addition of RO for 16–18 h. Cells were rinsed three times in Dulbecco's PBS and placed in fresh media before imaging.

For fixed-cell immunofluorescence, cells were seeded in 10-cm dishes, adhered for 12–16 h, treated with doxycycline and siRNA for 24–36 h, and split onto either poly-D-lysine-coated coverslips or Millicell EZ glass chamber slides (EMD Millipore) and a 6-cm dish (processed in parallel and used to prepare mitotic extracts for immunoblotting). After adhering for 12 h, cells were synchronized with RO for 12 h, followed by three washes in Dulbecco's PBS and addition of fresh media containing either nocodazole or taxol for 2–4 h before fixation. When used, ZM447439 was added 1 h before fixation.

Drug concentrations for human tissue culture cells, unless otherwise indicated, are as follows: 9 μ M RO (Sigma-Aldrich), 3.3 μ M nocodazole (Sigma-Aldrich), 500 nM taxol (EMD Millipore), 100 μ M monastrol (Tocris Bioscience), and 2 μ M ZM447439 (Chemietek).

Plasmids

Plasmids are listed in Table S1. All GFP-tagged xINCENP constructs and xSAH domain constructs were prepared from GFP-xINCENP FL (Tseng et al., 2010). FL siRNA-resistant hINCENP was a gift from S. Lens (University Medical Center Utrecht, Utrecht, Netherlands). All human constructs were cloned into a pcDNA5-FRT-TO plasmid containing the LAP tag (gift of R. Gassmann). FL hCENP-B and hCENP-B DNA-binding domain were gifts of M. Lampson (University of Pennsylvania, Philadelphia, PA; Wang et al., 2011). FL hMis12 and a plasmid containing the MAP4-MTBD were gifts of T. Kapoor (Rockefeller University, New York, NY; Maldonado and Kapoor, 2011; Tan and Kapoor, 2011). All human phospho-site mutants were generated using the QuikChange Multi Site-Directed Mutagenesis kit (Agilent Technologies) per the manufacturer's recommendations. *Xenopus* phospho-site mutants were generated using Gibson Assembly (Gibson et al., 2009) with synthesized DNA fragments corresponding to the desired mutations. All constructs either were cloned using QuikChange Site-Directed Mutagenesis (Agilent Technologies), a PCR-based digestion/ligation strategy using a pair of unique restriction sites in the pcDNA5-FRT-TO plasmid, or were generated using Gibson Assembly. The coding frame of all constructs was verified by DNA sequencing before insertion into HeLa T-REx cells.

Immunofluorescence

HeLa cells were fixed in 2% PFA in PBS, pH 7.4, or 1% PFA + 0.2% Triton X-100 (for DSN1 S100ph and KNL1 S24ph [gifts of I. Cheeseman, Massachusetts Institute of Technology, Cambridge, MA] and Hec1 S44ph [gift of J. Deluca, Colorado State University, Fort Collins, CO]) for 10 min at RT (Welburn et al., 2010; DeLuca et al., 2011). Cells were washed once in PBS, permeabilized in PBS + 0.2% Triton X-100 for 10 min, rinsed twice in PBS, and blocked for at least 30 min in 5% BSA + 0.2% Tween 20 (blocking). Cells were incubated in primary antibody diluted in blocking for at least 1 h, rinsed twice in PBS + 0.2% Tween 20 (PBST), and incubated in secondary antibody in blocking for at least 1 h. Cells were rinsed twice in PBST, incubated for at least 5 min in PBS plus 1 μ g/ml DAPI, rinsed once in PBS, and mounted with ProLong Gold antifade mountant (Thermo Fisher Scientific). All primary and secondary antibodies are listed in Table S2.

For *Xenopus* egg extract, immunofluorescence of spindles and sperm chromatin was performed as previously published (Desai et al., 1999; Tseng et al., 2010). In brief, extract was diluted in fixative (1 $\times$ BRB80, 2% formaldehyde) for 5 min, layered over a glycerol cushion, and pelleted onto a stage containing a circular coverslip for 15 min at 51,000 rpm in a Sorvall HS7 swinging-bucket rotor. The cushion was rinsed twice with 1 $\times$ BRB80 and aspirated away. The coverslip

was removed, fixed in ice-cold methanol for 5 min, rinsed twice in 10 mM TBS, pH 7.4, and 0.1% Triton X-100, and incubated with Abdil (10 mM TBS, pH 7.4, 150 mM NaCl, 0.1% Triton X-100, and 2% BSA) overnight at 4°C. Samples were incubated with primary and secondary antibodies diluted in Abdil for at least 1 h, with a 2 $\times$ PBST rinse after each incubation. Samples were washed once in PBS and mounted with VectaShield antifade mountant containing DAPI (Vector Laboratories).

Live cell imaging and analysis

Live cell imaging was performed at 37°C in 5% CO₂ in a LCV110U VivaView FL incubator microscope (Olympus) equipped with an X-Cite-exacte illumination source (Excelitas Technologies) and Orca-R2 CCD camera (Hamamatsu Photonics). For SAC experiments, spindle poisons diluted in media were added to each dish at least 20 min before imaging. For RO synchronization and release experiments, samples were imaged directly after the final washout. For taxol and RO synchronization and release experiments, images were acquired with a 20 $\times$ objective every 15 min for 48–72 h in the differential interference contrast (DIC), RFP, and GFP channels. For enhanced time resolution of unperturbed mitosis (Fig. 7 F), cells were imaged every 5 min. Individual cells were manually analyzed using the CellCognition browser. DoM and class at exit from mitosis were inferred from DIC and RFP morphology (Fig. S3 A). For SAC experiments, only cells observed transitioning from interphase to mitosis during the first 16 h of the video were analyzed. The DoM was calculated as the time from NEB until either slippage into interphase or cell death.

For RO synchronization and release experiments (Figs. 4 F and S3 F), each sample was imaged before washout to verify that >95% of cells were in interphase. After washout, cells in prophase/prometaphase at the beginning of the video or those seen transitioning into mitosis within the first 6 h of the video were analyzed. The DoM was calculated as the time from NEB until chromosome segregation or cell death.

Image acquisition and processing

Immunofluorescence images were acquired on a DeltaVision Image Restoration Microscope (Applied Precision Ltd.) using an inverted IX-70 stage (Olympus), pco.edge sCOS camera (PCO), Insight SSI solid-state illumination system (Applied Precision Ltd.), and 20 $\times$ /0.75 air and 100 $\times$ /1.4 oil objectives (Olympus) and running SoftWoRx v6.1 software (Applied Precision Ltd.). Images were acquired at RT using immersion oil with a refractory index of 1.516. The fluorophores DAPI, Dylight594, Alexa Fluor 488, and Cy5 were detected using filters for DAPI, Alexa Fluor 594, FITC, and Cy5, respectively. All images were acquired as z-stacks with 0.4- μ m optical sectioning and deconvolved using SoftWoRx constrained iterative deconvolution with default settings. All images within an experiment used identical acquisition settings.

For image quantification, deconvolved stacks were exported as 16-bit TIFFs and quantified using MetaMorph (Molecular Devices). For each sample, a maximum projection of the DAPI channel was used to create a DAPI region to select a single cell within a field. Epitopes that form foci at the centromere or kinetochore have the highest intensity in these regions; therefore, we empirically determined a threshold that primarily captured these intense foci, while excluding the signal outside these regions. This threshold was applied to each plane of the z-stack to quantify the integrated intensity of foci for the epitope. In a single experiment, each sample was quantified using the same threshold for a given channel. Images for which this threshold did not capture primarily centromere/kinetochore signal (for example, because of an artifact or high background) were discarded. A mean of ~10% or fewer cells from each sample were discarded for this reason.

Kinetochore intensity for phospho-specific kinetochore antibodies (Dsn1 S100ph, Knl1 S24ph) was quantified using a marker channel,

stained by anti-CENP-C antibody (a gift of H. Kimura, Tokyo Institute of Technology, Tokyo, Japan; Ando et al., 2002), to manually place a circle of defined size around each kinetochore. These kinetochore regions were then used to quantify the signal intensity from a maximum projection of the phospho-specific antibody channel. To calculate individual kinetochore intensity for Hec1 S44ph relative to Hec1, a threshold/watershed segmentation was performed on the Hec1 channel to create kinetochore regions. These regions were then used to quantify the intensity of Hec1 and Hec1 S44ph at each kinetochore from a maximum projection of each channel.

Automated foci counts were performed using MetaMorph. A mask generated from the DAPI channel was used to select a single cell within a field. For the channel of interest, a threshold/watershed segmentation was performed to separate individual foci. All foci larger than a minimum size were counted for each cell, yielding the number of foci per cell for a given epitope. The segmented regions were manually inspected for a subset of the total images.

Immunoblotting

Mitotic HeLa cell extracts were prepared by placing cells in 0.33 μ M nocodazole for 6–8 h followed by mitotic shake-off. Cells were pelleted at 200 g, rinsed twice in PBS, and resuspended in 2 $\times$ Laemmli buffer. Extracts were sonicated at 4°C in a Bioruptor water bath (Diagenode) at medium-intensity cycling 30 s on/60 s off for a total of 10 min. Samples were denatured, the protein concentration was determined using 260/280 nm on a NanoDrop 2000, and an equal amount of each sample was run on a precast 4–12% Bis-Tris gel (Thermo Fisher Scientific) followed by transfer to a nitrocellulose membrane overnight at 30 V. For *Xenopus* egg extract, samples were diluted in sample buffer, denatured, run on a precast gel as described earlier, and transferred to a nitrocellulose membrane overnight at 15 V.

Membranes were blocked in sterile 4% nonfat milk in PBS for at least 30 min, followed by incubation with primary antibody diluted in 5% BSA for 1–6 h at RT. After three 5-min washes in PBST, the membranes were incubated for 1 h with IRDye 800CW and 680LT secondary antibodies diluted in Odyssey Blocking Buffer with 0.1% Tween. Membranes were then washed three times for 5 min in PBST and detected using the Odyssey Infrared Imaging System (LI-COR Biosciences). All antibody information is listed in Table S2. Anti-NuMA antibody was a gift of R. Heald (University of California, Berkeley, Berkeley, CA).

FRAP

After 36- to 48-h siRNA treatment and construct induction, cells were treated with 3.3 μ M nocodazole or 500 nM taxol for 1 h then placed in a 37°C chamber mounted on a Deltavision Image Restoration System microscope (see Image acquisition and processing). FRAP was performed using SoftWoRx v6.1 software and a 488-nm QLM laser pulse to bleach three centromeric foci per cell. Images were acquired with a 60 $\times$ /1.42 oil objective from a single *z*-section every 10 s for 30 s before and 5 min after bleaching. Quantification was performed in ImageJ using a circle of defined size to quantify the integrated intensity of each bleached focus, a same-sized background focus, and the entire cell. Only foci remaining in the focal plane for the duration of the video were quantified. Full-scale normalization and single-exponential fitting were performed in Matlab using the easyFRAP module (Rapsomaniki et al., 2012) to determine the $t_{1/2}$ of each focus.

Immunoprecipitation

For immunoprecipitation from mitotic HeLa cell extracts, construct expression was induced by doxycycline in a confluent 15-cm dish for 8 h followed by the addition of 100 ng/ml nocodazole for 16 h. Mitotic

cells were collected by shake-off, rinsed in 1 $\times$ PBS, resuspended in 50 μ l of 1 $\times$ PBS, and incubated in lysis buffer (50 mM Tris, pH 8, 150 mM NaCl, 1% NP-40, 0.5 mM PMSF, 5 mM β -glycerol phosphate, phosphatase inhibitor cocktail [Roche]), and EDTA-free protease inhibitor [Sigma-Aldrich] for 30 min. Samples were sonicated in a water bath 5 $\times$ 10 s on/10 s off and passed through a 251/4 gauge needle. Samples were clarified by spinning for 10 min at maximum speed in a tabletop centrifuge. Clarified lysate was incubated with magnetic beads cross-linked to rabbit IgG or rabbit anti-GFP antibodies for 1.5 h with constant agitation. Beads were removed, rinsed once in lysis buffer, resuspended in sample buffer, and boiled. Half the sample was loaded for Western blot analysis. All steps were performed at 4°C.

Xenopus egg extract

A published protocol was used to prepare CSF extracts (Murray, 1991). Depletion of endogenous CPC and reconstitution was previously described (Tseng et al., 2010). For depletion, magnetic protein A Dynabeads (Invitrogen) were coupled and cross-linked to either rabbit anti-IgG or rabbit anti-xINCENP beads using BS₃ (Thermo Fisher Scientific) according to the manufacturer's recommendations. CSF extract was incubated with the appropriate beads in a 1:1 ratio for 1 h at 4°C, then beads were removed to yield depleted extract. The CPC was reconstituted in Δ CPC extract by the addition of 500 nM recombinant MBP-xAurora B, 0.3 μ g/ μ l of mRNA for the appropriate GFP-xINCENP construct, and 0.075 μ g/ μ l each of mRNA encoding xSurvivin and xDasra A. To translate these mRNA constructs, extract was cycled into interphase by the addition of 300 μ M calcium chloride for 1.5 h at 22°C, then driven back into metaphase by adding an equal volume of fresh CSF extract and incubating for 30–45 min at 22°C. mRNA was prepared using the SP6 mMessage mMachine RNA transcription kit (Ambion). All xSAH domain fragments were expressed from mRNA added to extract at 0.15–0.3 μ g/ μ l and cycled as described earlier. Immunoprecipitation of GFP and GFP-xSAH was performed similarly to CPC depletion using rabbit anti-GFP antibodies.

For immunofluorescence on nocodazole-treated sperm chromatin, 1,000–2,000 sperm/ μ l were added before calcium addition. After incubation with fresh CSF, samples were incubated with 33 μ M nocodazole for 30 min before processing for immunofluorescence.

The SAC assay was performed similarly to Minshull et al. (1994). In brief, samples were reconstituted as described earlier. CSF extract was then incubated with 10,000 sperm/ μ l and 33 μ M nocodazole for 45 min, followed by addition of 600 μ M calcium chloride. Western blot samples were taken at 30-min intervals after the addition of calcium chloride.

MT pelleting assays were modified from Xue et al. (2013). Extract was incubated for 30 min with 33 μ M nocodazole or 10 μ M taxol at 22°C. An input sample was taken, and 20 μ l of extract was diluted in fixative (1 $\times$ BRB80, 30% glycerol, 0.1% Triton X-100, and phosphatase inhibitor) and layered over a glycerol cushion (1 $\times$ BRB80 and 40% glycerol). Samples were spun in a tabletop centrifuge at RT for 15 min at maximum speed. The cushion was washed twice with 1 $\times$ BRB80 and aspirated away, and the pellet was resuspended in 1 $\times$ BRB80 and 0.1% Triton X-100 and transferred to a fresh tube. The sample was spun again for 15 min, and the pellet was resuspended in 40 μ l sample buffer. For phosphatase treatment, samples were diluted in buffer containing lambda phosphatase for 40 min at 30°C (New England Biolabs, Inc.).

CENP-A nucleosome arrays were assembled according to Guse et al. (2011). In brief, chromatin was assembled by salt dialysis of 3.6 μ M human CENP-A/*Xenopus* H4 dimers that had been copurified from bacteria using hydroxyapatite (Bio-Rad Laboratories) and HiTrap SP (GE Healthcare) columns, 4.0 μ M *Xenopus* H2A/H2B dimers generated by purifying his-tagged constructs separately from

inclusion bodies using Ni-NTA (QIAGEN) and dialyzed together after cleavage of the His tags using TEV protease (Zierhut et al., 2014), and 0.2 $\mu\text{g}/\mu\text{l}$ of a DNA fragment containing 19 copies of the 601 nucleosome positioning sequence, which had been biotinylated on both ends using Klenow (New England Biolabs, Inc.) in the presence of biotin-14-dATP, thio-dTTP, and thio-dGTP. Plasmids encoding this DNA template and for CENP-A/H4 coexpression were gifts from A. Straight (Stanford University, Stanford, CA). Plasmids for expression of His-H2A and His-H2B were gifts from C.D. Allis (Rockefeller University, New York, NY). Coupling to Dynabeads M-280 Streptavidin (Thermo Fisher Scientific) at a concentration in which 1 μg of DNA was coupled to 10 μl of bead slurry for 1 h was done just before addition to extract. To assay CPC localization, array-coupled beads were added to a final volume of 100 μl control (ΔIgG), CPC-depleted, or CPC-reconstituted extract. Extracts were then cycled to express RNA, and 33 μM nocodazole was added. Beads were removed from extract with a magnet, washed in ice-cold CSF-XB with 0.05% Triton X-100, fixed for 5 min in 2% formaldehyde, washed with Abdil, added to acid-washed, poly-L-lysine-coated coverslips, and processed for immunofluorescence in the same manner as sperm chromosomes.

Data presentation

Broken violin plots were generated using a custom script in Python 2.7. In brief, a kernel density estimate of the underlying data was determined using the `scipy.stats.gaussian_kde` module with a bandwidth determined by the Scott method. The integrated area underneath each density estimate is equal between all samples in a plot unless otherwise indicated. For visualization, density estimates were cut at the minimum and maximum value of the underlying data. Beneath each density is a minimalist boxplot with a lower whisker from $-1.5 \times \text{IQR}$ to the first quartile, a circle at the median, and an upper whisker from the third quartile to $1.5 \times \text{IQR}$. Data were organized using the Pandas module, programmatically visualized using matplotlib and seaborn modules, and labeled and formatted in Adobe Illustrator CC.

Representative images are no more than $\pm 20\%$ of the median value of the construct based on quantification. Representative images from qualitative experiments were chosen based on morphology. Graphs were generated using Python or Prism software. All box plots are Tukey box plots with whiskers extending $\pm 1.5 \times \text{IQR}$. For visual simplicity, points past the whiskers (considered statistical outliers) are omitted. All statistical analyses were conducted in Prism using a two-tailed Mann-Whitney test using all samples (even statistical outliers). Individual p-values and *n* values are indicated in figure legends. In several cases, data from multiple independent experiments were aggregated and used for statistics. Aggregation was performed by normalizing the intensity of each sample to hINCENP FL within that experiment, then pooling these normalized values from multiple experiments together. For these experiments, representative images were chosen from a single experiment.

Online supplemental material

Fig. S1 shows characterization of 6Myc-xSAH in *Xenopus* egg extracts. Fig. S2 shows CPC binding and localization analyses of GFP-INCENP ΔSAH in HeLa cells. Fig. S3 (A and B) shows mitotic fates upon taxol treatment, Fig. S3 (C–E) shows immunofluorescence of kinetochore markers in HeLa cells whose INCENP was replaced with INCENP SAH deletion mutants, and Fig. S3 F shows a control for Fig. 4 F. Fig. S4 A shows immunofluorescence of 6My-tagged xINCENP SAH domain constructs in *Xenopus* egg extracts, and Fig. S4 B shows sequence alignment of PRD1 and PRD2. Fig. S5 A shows the effect of C-terminal deletion of the SAH on DoM in taxol-treated HeLa cells, and Fig. S5 B shows the effect of expressing INCENP 6A on unperturbed HeLa cells. Tables S1 and S2 list plasmids and antibodies used in this study, respectively.

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Author contributions: M.S. Wheelock designed and executed all experiments and analyzed data, except for an experiment shown in Fig. 1C, which was performed by D.J. Wynne. D.J. Wynne and B.S. Tseng have generated preliminary data in *Xenopus* egg extracts, which guided this study. H. Funabiki supervised the entire study. M.S. Wheelock and H. Funabiki wrote the manuscript, which was edited by D.J. Wynne and B.S. Tseng.

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