

The fission yeast nucleoporin Alm1 is required for proteasomal degradation of kinetochore components

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Kinetochores (KTs) are large multiprotein complexes that constitute the interface between centromeric chromatin and the mitotic spindle during chromosome segregation. In spite of their essential role, little is known about how centromeres and KT are assembled and how their precise stoichiometry is regulated. In this study, we show that the nuclear pore basket component Alm1 is required to maintain both the proteasome and its anchor, Cut8, at the nuclear envelope, which in turn regulates proteostasis of certain inner KT components. Consistently, *alm1*-deleted cells show increased levels of KT proteins, including CENP-C^{Cnp3}, spindle assembly checkpoint activation, and chromosome segregation defects. Our data demonstrate a novel function of the nucleoporin Alm1 in proteasome localization required for KT homeostasis.

Introduction

Centromeres are specialized regions of chromatin that constitute the site of kinetochore (KT) assembly required for mitotic spindle attachment. Fission yeast centromeres are structurally complex, similar to those of metazoa, and contain repetitive elements (outermost repeats [otr]) flanking a central domain composed of a central core (cnt) surrounded by innermost repeats (imr; Takahashi et al., 1992; Pidoux and Allshire, 2005). Centromeric chromatin is transcriptionally silenced. However, whereas transcriptional silencing at imr/otr depends on the canonical heterochromatin marks (H3K9me2) and HP1/Swi6 binding, transcriptional silencing at cnt depends on the centromeric histone H3 variant CENP-A^{Cnp1} (Pidoux and Allshire, 2005; Castillo et al., 2007). Cnp1, together with the non-canonical nucleosome complex CENP-S-T-W-X (SpMhf1, SpCnp20, SpNew1, and SpMhf2), defines the landmark of centromeric chromatin structure (Stoler et al., 1995; Takahashi et al., 2000, 2005). Cnp1 directly interacts with CENP-C^{Cnp3}, which in turn recruits several factors, including CENP-L^{Fta1}, CENP-K^{Sim4}, or the monopolin complex Pcs1-Mde4, required for KT assembly and function (Cohen et al., 2008; Tanaka et al., 2009; Carroll et al., 2010; Folco et al., 2015).

KTs can be functionally divided into two domains: the inner KT, which constitutes the interface with centromeric

chromatin, and the outer KT, which constitutes the microtubule (MT)-binding interface (Santaguida and Musacchio, 2009). During mitosis, each KT must establish correct interactions with spindle MTs. Erroneous KT-MT attachments are detected by the spindle assembly checkpoint (SAC) and corrected by aurora B kinase. If not corrected, erroneous attachments might result in chromosome missegregation and aneuploidy, which contributes to chromosomal instability in cancer cells (Cimini et al., 2003, 2006). Both centromeric chromatin and proper KT structure are required for correct chromosome segregation (Goshima et al., 1999; Bernard et al., 2001; Nonaka et al., 2002; Pidoux et al., 2003; Pidoux and Allshire, 2005; Gregan et al., 2007; Tanaka et al., 2009). Similarly, the appropriate balance of KT proteins must be strictly regulated to ensure proper KT assembly and function (Castillo et al., 2007).

The 26S proteasome is a large, evolutionary conserved protease complex responsible for the ATP-dependent degradation of polyubiquitinated proteins. This proteolytic system plays a pivotal role in cellular quality control and protein homeostasis. Proteasomal degradation takes place in the cytoplasm and also inside the nucleus, where the proteasome is enriched (Wilkinson et al., 1998; Voges et al., 1999). It is known that the levels and distribution of centromeric CENP-A^{Cnp1} are regulated by proteasomal degradation, restricting its localization to centromeric chromatin (Collins et al., 2004; Moreno-Moreno et al., 2006; Ranjitkar et al., 2010; Kitagawa et al., 2014). Interestingly, Cut8, a nuclear envelope (NE) protein required for proteasome function and localization to the NE, is also required for Cnp1 centromeric localization (Tatebe and Yanagida, 2000; Takeda

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Abbreviations used: ChIP, chromatin immunoprecipitation; CC, coiled-coil; CHX, cycloheximide; cnt, central core; EMM, Edinburgh minimal media; GO, gene ontology; imr, innermost repeats; KT, kinetochore; MBC, methyl benzimidazol-2-yl carbamate; MT, microtubule; NE, nuclear envelope; NPC, nuclear pore complex; otr, outermost repeats; qPCR, quantitative PCR; SAC, spindle assembly checkpoint; SGA, synthetic genetic array; SPB, spindle pole body; TBZ, thiabendazole; TPR, translocated promoter region; ts, thermosensitive.

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and Yanagida, 2005; Kitagawa et al., 2014), suggesting that proteasome localization at the NE is functionally linked to proper centromere–KT composition. The NE further contributes to the quality control and protein homeostasis of KT components through the ubiquitin–proteasome pathway (Furth et al., 2011; Kriegenburg et al., 2014). However, to date, it is still unknown how the NE influences proteasome localization and regulation to ensure proper KT–centromere stoichiometry and function.

The NE is spanned by the nuclear pore complexes (NPCs), composed of ~30 different types of proteins called nucleoporins (Asakawa et al., 2014; Kabachinski and Schwartz, 2015). The classical view of the NPCs as mediators of nucleocytoplasmic transport has been further extended to many other cellular functions such as regulation of gene expression and genome stability (Palancade et al., 2007; Capelson et al., 2010; Kalverda and Fornerod, 2010; Yang et al., 2015). Associated with the NPCs are the translocated promoter region (TPR) nucleoporins, which are evolutionary conserved large coiled-coil (CC) proteins that localize at the nucleoplasmic side of the NPC, forming a specific structure called the nuclear basket (Cordes et al., 1997; Frosst et al., 2002; Asakawa et al., 2014). TPRs form filaments that connect adjacent NPCs and project into the nucleoplasm, serving as a binding platform for chromatin, transcription factors, and cell cycle regulators (Galy et al., 2000; Niepel et al., 2013; Umlauf et al., 2013; Rajanala et al., 2014). In human cells, TPR contributes to a variety of nuclear processes, such as small ubiquitin–related modifier (SUMO) homeostasis (Schweizer et al., 2013), spatiotemporal regulation of SAC components (Rodríguez-Bravo et al., 2014), and the exclusion of heterochromatin from NPCs (Krull et al., 2010). In addition, in *Saccharomyces cerevisiae*, the TPR proteins Mlp1 and Mlp2 are involved in anchoring transcriptionally active genes at the NPC (Luthra et al., 2007; Tan-Wong et al., 2009) and mRNA export (Niepel et al., 2013) and are required for maintenance of spindle pole body (SPB) integrity and telomere structure (Hediger et al., 2002; Zhao et al., 2004; Niepel et al., 2005). In *Schizosaccharomyces pombe*, there are two members of the TPR family of nucleoporins, Nup211 and Alm1 (Bae et al., 2009; DeGrasse et al., 2009). Whereas Nup211 plays an essential role in the regulation of mRNA export, the function of Alm1 has not been characterized to date.

Here, we show that the TPR nucleoporin Alm1 is required to maintain the proper localization of the proteasome anchor and regulator Cut8 and the proteasome subunits Mts2 and Mts4 to the NE. Cells lacking *alm1* show an altered stoichiometry of KT proteins, including Cnp3, and DNA segregation defects. Together, these results strongly suggest that Alm1 is required for proper 26S proteasome localization at the NE, and this regulation is critical to maintain correct KT stoichiometry.

Results

Alm1 localizes to the NE

TPR nucleoporins are large CC proteins that localize at the nucleoplasmic side of the NPC, forming the nuclear basket. Alm1 showed extensive regions of predicted CCs (Fig. S1 A). Moreover, Alm1-Tomato localized as bright dots at the nuclear periphery (Fig. S1 B) and partially colocalized with the nucleoporin Nup107-GFP (Fig. S1 C). This partial colocalization suggests that Alm1 also occupies areas that are devoid of NPCs, similar to what has been shown for *S. cerevisiae* Mlp1 and Mlp2

(Strambio-de-Castillia et al., 1999; Niepel et al., 2013). The localization of Alm1-Tomato mostly overlapped with Nup211-GFP (Fig. 1 A). Furthermore, in the absence of Alm1, a fraction of Nup211 lost its NE localization and accumulated in the nucleoplasm (Fig. 1 B), suggesting that both proteins interact at the NPC. It has previously been reported that Alm1 localizes at the SPBs and the medial ring (Jiménez et al., 2000). Careful analysis of the localization of Alm1-Tomato expressed from its own promoter showed only a transient colocalization of Alm1 with the SPB, and we never observed any Alm1-Tomato signal at the medial ring (Fig. S1 D).

Thus, as it has been reported for members of the Mlp/TPR family from yeast to human (Zimowska et al., 1997; Strambio-de-Castillia et al., 1999; Xu et al., 2007), the fission yeast TPR nucleoporin Alm1 localizes to the NPC and the nuclear rim.

alm1 deletion results in chromosome missegregation

The localization of Alm1 at the NE prompted us to analyze NE morphology in *alm1Δ* cells. Whereas we did not find severe alterations in nuclear morphology in *alm1Δ* cells, we did observe that 40% of mitotic *alm1Δ* cells presented daughter nuclei of different sizes, which was never observed in WT cells (Fig. 1 C, arrow). We then monitored DNA segregation by in vivo microscopy using histone H3 (*hht2*) tagged with GFP. We observed that 31.8% of mitotic *alm1Δ* cells ($n = 22$) showed lagging masses of DNA during anaphase B, of which approximately half of them resulted in aneuploidy, a phenotype that was never observed in WT cells ($n = 25$; Fig. S2 A). Chromosome missegregation defects in *alm1Δ* cells were further confirmed by minichromosome loss assays (Fig. 1 D).

The appearance of lagging chromosomes during anaphase B is indicative of erroneous KT–MT attachments (Gregan et al., 2007; Rumpf et al., 2010). Thus, we monitored KT behavior in cells coexpressing the KT marker Mis6-GFP and the SPB marker Sid2-Tomato. In WT cells, KTs appeared as two to six dots that oscillated between the two SPBs during prophase–metaphase. Just before anaphase onset, the KTs achieved bipolar attachment and simultaneously segregated to the spindle poles (Fig. 1 E; Nabeshima et al., 1998; Tournier et al., 2004; Courtheoux et al., 2007). However, in *alm1Δ* cells, KTs appeared more extended along the metaphase spindle compared with WT cells, and their pole to pole oscillations seemed to be more dynamic, which suggested that MT–KT interactions might be unstable in *alm1Δ* (Fig. 1 E, dashed boxes). We also noticed that segregation of sister chromatids at anaphase frequently occurred in an uncoordinated manner. In WT cells, all of the KTs were segregated to the spindle poles simultaneously, whereas in *alm1Δ* cells, after the segregation of most of the KTs, we still could observe one to two KTs in the spindle that are usually segregated 1–2 min later (Fig. 1 E, asterisk). We further confirmed the presence of lagging KTs in *alm1Δ* cells during anaphase B (Fig. 1 E, arrow).

To analyze the behavior of a single KT pair in vivo, we used a strain that harbors a tandem array of lacO DNA repeats at the *lys1* gene located close to the centromere of chromosome I (*cen1*). This lacO array allows binding of a GFP-LacI fusion protein (Nabeshima et al., 1998). In *alm1Δ* cells, we observed a prolonged metaphase in which the distance between the two *lys1*-GFP signals is transiently increased compared with WT cells (Fig. S2 B, arrows and dashed boxes).

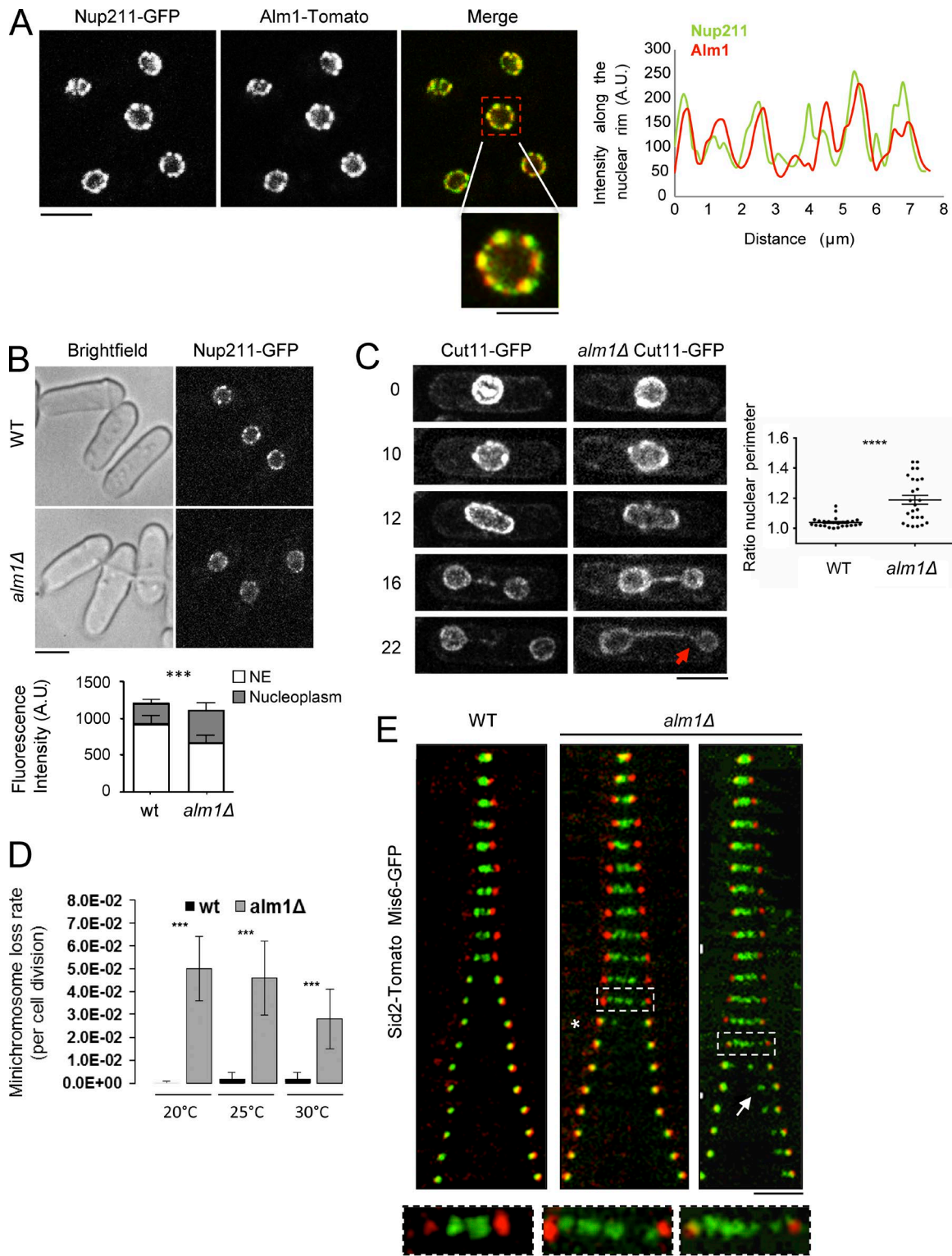


Figure 1. **Alm1 localizes to the NE and NPC, and its absence results in chromosome segregation defects.** (A) Alm1-Tomato colocalizes with Nup211-GFP. Maximal projection of three z sections is shown. (Inset) Magnification of the indicated nucleus. Bars, 2.5 μm . Fluorescence intensity levels along the nuclear rim of the selected nucleus from an arbitrary starting point. (B) Images of WT and *alm1* Δ cells expressing Nup211-GFP. Mean fluorescence intensity of Nup211-GFP at the NE and in the nucleoplasm. $n = 30$. (C) WT and *alm1* Δ cells expressing the NE marker Cut11-GFP. Arrow indicates smaller daughter nuclei. Time is indicated in minutes. Ratio of the perimeter of sister nuclei. $n = 25$. (D) Minichromosome loss assay. Three independent experiments. $n > 1,500$. (E) Images of WT and *alm1* Δ mitotic cells expressing Sid2-Tomato and Mis6-GFP. Time between frames is 1 min. Magnifications of the regions indicated by dashed boxes are shown below. Bar, 2.5 μm . Asterisk marks asynchronous KT segregation. Arrow indicates lagging KT. Bars, 5 μm . A.U., arbitrary units. Error bars represent SD. ***, $P < 0.001$; ****, $P < 0.0001$.

Thus, collectively, these results show that *alm1Δ* cells display abnormal KT behavior during metaphase and are affected in chromosome segregation, suggesting that, in the absence of *alm1*, either KTs or spindle MTs are altered.

To test whether the mitotic defects in *alm1Δ* cells are the result of a defective mitotic spindle, we followed spindle formation and behavior by expressing the α -tubulin subunit Atb2 tagged with GFP. In *S. pombe*, mitotic phases can be distinguished by the spindle dynamics. Phase I initiates with spindle formation (prophase). During phase II, the spindle maintains a constant length of ~2.5–3 μ m (metaphase) and culminates with KT segregation to the spindle poles (anaphase A). Finally, during phase III, the spindle elongates, segregating the sister nuclei (anaphase B; Fig. S2, C and D; Nabeshima et al., 1998). We did not observe any significant alteration in spindle formation and function in *alm1Δ* compared with WT cells. We found, however, that 66.7% of *alm1Δ* cells ($n = 30$) presented a delay at the metaphase to anaphase transition (phases I and II; 10.9 ± 0.1 min in WT cells vs. 15.1 ± 0.9 min in *alm1Δ*; Fig. S2, C–E).

***alm1+* deletion leads to SAC activation**

The SAC is a mitotic surveillance mechanism that monitors correct attachment between KTs and spindle MTs. This pathway delays the metaphase to anaphase transition until all KTs are properly attached to MTs of the mitotic spindle. The SAC is composed of the kinases Bub1 and Mps1 (Mph1 in *S. pombe*) and the Mad1, Mad2, Mad3, and Bub3 proteins. The inhibitory signal originates at unattached or tensionless KTs. Here, SAC proteins accumulate and prevent anaphase onset by inhibiting the ubiquitin ligase anaphase-promoting complex (APC)/cyclosome through binding to the APC activator Cdc20 (Musacchio and Salmon, 2007). *alm1Δ* showed a sustained localization of both Mad2-GFP and Bub1-GFP at KTs during mitosis (Fig. S3, A and B), which is indicative of SAC activation.

A conserved function of the TPR proteins is the spatio-temporal regulation of the SAC components Mad1 and Mad2 (Lee et al., 2008; De Souza et al., 2009; Schweizer et al., 2013). During interphase, both Mad1 and Mad2 localize to the NE. In mitosis, however, Mad1/Mad2 are released from the NE and relocalize to the nucleoplasm and KTs, where they are essential for SAC activation. Mlp/TPR proteins are required for correct localization of Mad2 to the NE during interphase and to KTs during mitosis. In addition, in human cells, TPR is required for full SAC activation in unperturbed mitosis and in response to MT damage (Schweizer et al., 2013; Rajanala et al., 2014). We found that during interphase, Mad2-GFP was not recruited to the NE in *alm1Δ*, which shows that this function is conserved in *S. pombe* (Fig. S3 C). The localization of Mad2-GFP at KTs during mitosis, however, was not affected (Fig. S3 A). Furthermore, we found that *alm1Δ* cells were able to respond to MT damage, as treatment with the MT inhibitor methyl benzimidazol-2-yl carbamate (MBC) induced a delay at the metaphase to anaphase transition and the accumulation of Mad2-GFP at KTs, as seen in WT cells (Fig. S3 E). Moreover, double mutants of *alm1Δ* and either *bub1Δ* or *mad2Δ* showed a negative genetic interaction, indicating that Mad2 and Bub1 become critical for *alm1Δ* survival (Fig. S3 F).

Aurora B kinase is required for the correction of erroneous attachments and for SAC functionality (Petersen et al., 2001; DeLuca et al., 2006; Knowlton et al., 2006; Zhang et al., 2007; Vanoosthuysen and Hardwick, 2009). To examine whether the appearance of lagging chromosomes in *alm1Δ* might result

from deregulation of aurora kinase (Ark1), we analyzed its localization and activity. We found that both localization to KTs and the kinase activity of Ark1 were unaffected in *alm1Δ* relative to WT cells (Fig. S3, G and H). Importantly, we found a synergistic phenotype in the double mutant comprising the *alm1Δ* and *ark1-t7* thermosensitive (ts) allele at the permissive temperature (Fig. S3 I), with increased frequency of lagging chromosomes and DNA missegregation events (Fig. S3 J).

Thus, the presence of an apparently normal mitotic spindle and the strong dependency of SAC functionality for survival suggest that *alm1Δ* cells might be defective in KT–MT capture or stability.

Analysis of genetic interactions of *alm1Δ* by synthetic genetic array (SGA)

Mutants affecting centromere or KT function usually display DNA segregation defects and are sensitive to MT perturbation (Takahashi et al., 1994; Ekwall et al., 1999; Pidoux et al., 2003). We found that *alm1Δ* is sensitive to MT perturbation, as growth was inhibited in the presence of 15 μ g/ml of the MT-depolymerizing drug thiabendazole (TBZ) compared with the WT strain (Fig. 2 A).

To gain further insights into the cellular function of Alm1, we screened for genetic interactions by performing an SGA assay (Tong et al., 2001). To enrich for genetic interactions related to KT function, we performed the SGA in conditions of MT perturbation (i.e., in the presence of TBZ; Fig. 2 B and see the SGA assay section of Materials and methods). The analysis of the sensitivity of single and double mutants toward TBZ revealed several mutant clusters that displayed increased sensitivity in combination with *alm1Δ* (Fig. 2 C and Fig. S4 A). Gene ontology (GO) enrichment analysis of these synthetic clusters showed an enrichment in genes encoding for KT components like *spc19*, *dis2*, or *ask1*; SAC components like *bub1*, *mad1*, and *mad2*; proteins involved in ubiquitin-dependent proteolysis like the ubiquitin-protease *upb2*; and chromatin regulators (Fig. 2 D). Given the enrichment in KT-encoding genes and the fact that most of the components of this structure are essential, we analyzed genetic interactions of *alm1Δ* with several KT ts alleles, which were not present in the deletion library (Fig. 2 E). After tetrad dissection, we indeed found viable spores only for the WT and single mutants, whereas *alm1Δ* showed synthetic lethality with mutants of centromere–KT factors, such as *cnp3*, *mis6*, *mis12*, or *mhf1* (Fig. 2 E). Furthermore, we also observed a strong genetic interaction between *alm1Δ* and the GFP-tagged versions of the outer KT proteins Ndc80 and Nuf2 (Fig. 2 E). Together, these results point to a functional relationship between *alm1* and centromere and KT genes.

***alm1Δ* mutant shows abnormal accumulation of centromere–KT proteins**

The synthetic lethality between *alm1Δ* and mutants in centromere and KT components prompted us to analyze the localization and intensity levels of several of these proteins in *alm1Δ* by video microscopy (Fig. 3, A–C). Although the levels of Cnp1 were unaffected in *alm1Δ* cells (Fig. 3, A and B), we found increased levels of the centromere–KT proteins Mhf2, Cnp20, Mde4, Fta1, and especially Cnp3, with a 1.8-fold increase in *alm1Δ* compared with the WT (Fig. 3, A–C). The increased signal of Cnp3 at KTs in the *alm1Δ* mutant correlated with an increase in total Cnp3 protein (Fig. 3 D). The mRNA levels, however, were comparable to the WT strain (Fig. 3 E),

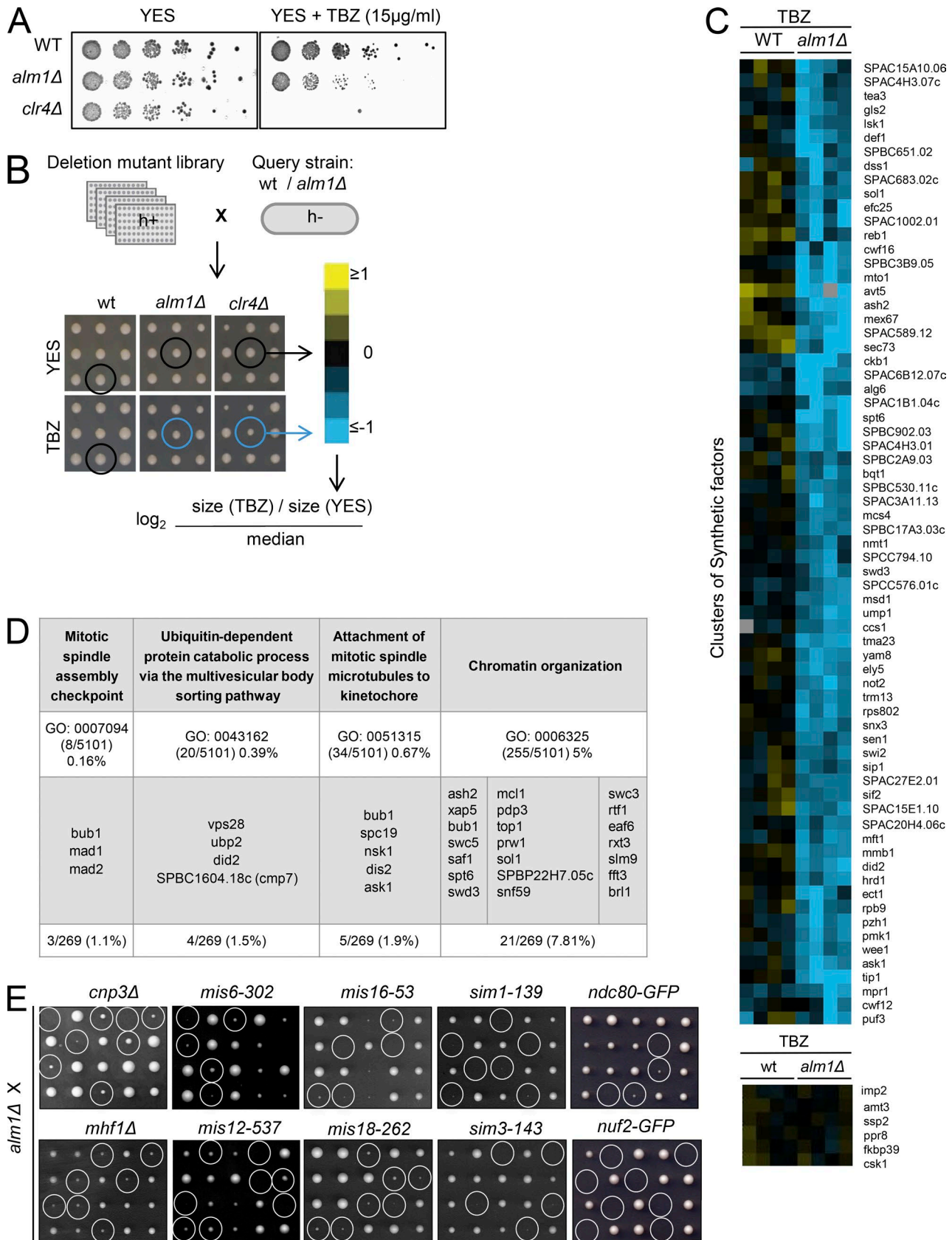


Figure 2. **SGA assay based on TBZ sensitivity identifies genetic factors that contribute to maintain KT structure and functionality.** (A) TBZ sensitivity assay of *alm1Δ* mutant. *clr4Δ* mutant was used as a positive control (Bernard et al., 2001). (B) Flow-through of the SGA to screen for genetic interactors of *alm1Δ* mutant in TBZ. WT and *alm1Δ* query strains were crossed with the *Bioneer* haploid deletion mutant library (v. 3) and spotted on YES- and TBZ-containing plates. Examples of colony growth on YES and YES + TBZ plates. The growth of the single and double mutants was quantified in both media, and we

suggesting that the increased protein levels of Cnp3 in *alm1Δ* cells were not the result of transcriptional deregulation but instead to altered protein stability or turnover.

Overexpression of Cnp1 was previously shown to cause its spreading from the centromeric *cnt* to the adjacent heterochromatin, resulting in altered pericentromeric silencing and chromosome missegregation (Castillo et al., 2007). We therefore analyzed the chromatin state at centromeres by measuring transcript levels of heterochromatin repeats (*imr*, *dg*, and *dh*) and *cnt* by reverse transcriptase assays combined with quantitative PCR (qPCR; Fig. 3 F). This analysis revealed a reduction in transcripts originating from the *imr* and *otr* repeats in *alm1Δ* cells compared with WT cells. Surprisingly, there was a threefold increase in transcripts derived from the *cnt* region of centromere I (Fig. 3 F), indicating that this chromatin region is derepressed in *alm1Δ*. We performed chromatin immunoprecipitation (ChIP) experiments using endogenously expressed Cnp3-GFP and found that Cnp3 distribution at the centromeric region in *alm1Δ* was indistinguishable from the WT cells (Fig. 3 G), indicating that the excess of Cnp3 protein in *alm1Δ* cells does not seem to increase the chromatin-bound fraction.

Collectively, these experiments demonstrate that *alm1Δ* shows abnormal accumulation of several KT proteins, including Cnp3. We further found that the absence of Alm1 alters the expression of centromeric transcripts; however, these changes are not caused by lateral spreading of Cnp3 into pericentromeric chromatin.

Cnp3 proteostasis is regulated by the proteasome

Previous studies have shown that centromeric levels and distribution of CENP-A^{Cnp1} are regulated locally through proteasomal degradation (Collins et al., 2004; Moreno-Moreno et al., 2006; Ranjitkar et al., 2010; Kitagawa et al., 2014). To test whether Cnp3 stability is regulated by proteasomal degradation, we first analyzed Cnp3 protein levels in a *ts* mutant of the non-ATPase Mts4, a subunit of the 19S regulatory particle of the proteasome (Wilkinson et al., 1997). Microscopic analysis of KT Cnp3-GFP levels in the *mts4* mutant revealed that the Cnp3-GFP signal is increased compared with WT cells, even at the permissive temperature, reaching similar levels to those found in the *alm1Δ* mutant (Fig. 4 A and Fig. S4 B). Consistently, the *mts4* mutant showed a twofold increase of lagging chromosomes compared with WT cells at the permissive temperature (Fig. 4 C). The double mutant *alm1Δ mts4* showed a further increase in Cnp3 levels at KTs at the permissive temperature (Fig. 4 A and Fig. S4 B). These results are consistent with the total Cnp3-GFP protein levels observed by Western blot analysis (Fig. 4 B).

In *S. pombe*, the 26S proteasome is enriched within the nucleus as seen in animal cells (Amsterdam et al., 1993; Enenkel et al., 1998; Wilkinson et al., 1998; Russell et al., 1999). Cut8 is required to anchor the proteasome to the NE in the

fission yeast (Tatebe and Yanagida, 2000; Takeda and Yanagida, 2005; Takeda et al., 2011), and *cut8* mutant phenotypes are similar to those of proteasome mutants such as *mts4*. We therefore analyzed the levels of centromeric Cnp3-GFP in the *cut8-563-ts* mutant. Although this mutant showed elevated centromeric Cnp3 levels already at permissive temperature, we observed a further twofold increase at restrictive temperature (Fig. 4 D and Fig. S4 C), which is consistent with total Cnp3-GFP protein levels (Fig. 4 E). We were not able to generate the double mutant of *alm1Δ* and *cut8-563*, suggesting that the combination of both deficiencies is synthetically lethal.

Next, we analyzed Cnp3 levels in WT and *alm1Δ* cells in the presence of the protein synthesis inhibitor cycloheximide (CHX). As a control, we analyzed the levels of cyclin B/Cdc13, a known substrate of the proteasome. We found that Cnp3 is unstable in WT cells, although it has a significantly longer half-life than cyclin B, suggesting that only a pool of Cnp3 is degraded (Fig. 4 F and Fig. S4 D). Importantly, the Cnp3 decay was delayed in *alm1Δ* cells compared with WT cells (Fig. 4 F). We did not observe significant changes in the turnover of cyclin B between WT and *alm1Δ* cells under these experimental conditions (Fig. S4 D).

To examine whether Cnp3 is ubiquitinated, we immunoprecipitated native Cnp3 from *mts4* mutant cells and probed the immunoblot with an antibody against ubiquitin, which revealed the typical smear of polyubiquitinated proteins (Fig. 4 G and see the Ubiquitin pull-down section of Materials and methods). Using a complementary approach, we coexpressed Cnp3-GFP and 6His-tagged ubiquitin in *mts4* mutant cells and performed pull-down experiments for His-tagged ubiquitin under denaturing conditions. We analyzed the fraction enriched in ubiquitinated proteins by anti-GFP immunoblotting, in which we detected a smear of slower migrating ubiquitinated forms of Cnp3-GFP (Fig. 4 H). Thus, both experiments are consistent with Cnp3 being ubiquitinated *in vivo*. Altogether, these data indicate that Cnp3 is regulated by ubiquitin-dependent proteasomal degradation.

Alm1 is required for proper localization of the 26S proteasome to the NE

Both Cut8 and Alm1 localize to the NE (Fig. 5 A; Takeda and Yanagida, 2000; Takeda and Yanagida, 2005; Takeda et al., 2011). To test whether the absence of Alm1 affects Cut8 localization, we analyzed Cut8 distribution in the *alm1Δ* background by live-cell microscopy. Intriguingly, we found that *alm1Δ* displays reduced mean levels of Cut8-GFP at the NE (Fig. 5 B). The total levels of Cut8 protein, however, were similar in *alm1Δ* and WT cells (Fig. 5 C). As it was known that Cut8 is required for proper proteasome nuclear localization, we tested whether the absence of Alm1 also affects the localization of the 26S proteasome. To this end, we analyzed the localization of the proteasomal subunits Mts2 and Mts4 by immunostaining. Notably, we

compared the ratio with the median ratio. Blue indicates small colony size, and yellow indicates large colony size (see Materials and methods). (C) Clustering analysis showing one of the clusters of deletion mutants that have a synthetic effect on TBZ sensitivity with *alm1Δ*. For each experiment, four replicates were performed. Clustering analysis showing one group of genes with no sensitivity to TBZ, neither in the single nor the double mutant with *alm1Δ*. Blue indicates synthetic interaction, yellow indicates suppressive interaction, black indicates no interaction, and gray indicates the absence of data. (D) GO enrichment analysis for biological processes of all of the genes identified in the three synthetic clusters. The table contains the percentage between the total number of genes of the indicated GO group and the total number of genes of *S. pombe* (upper value), and the percentage between the number of gene deletion mutants of the synthetic clusters belonging to the mentioned GO group and the total number of gene deletion mutants of the synthetic clusters (lower value). (E) Tetrad dissection analysis of crosses between *alm1Δ* and strains with the indicated genotypes. Double mutants between *alm1Δ* and the indicated mutant backgrounds are encircled.

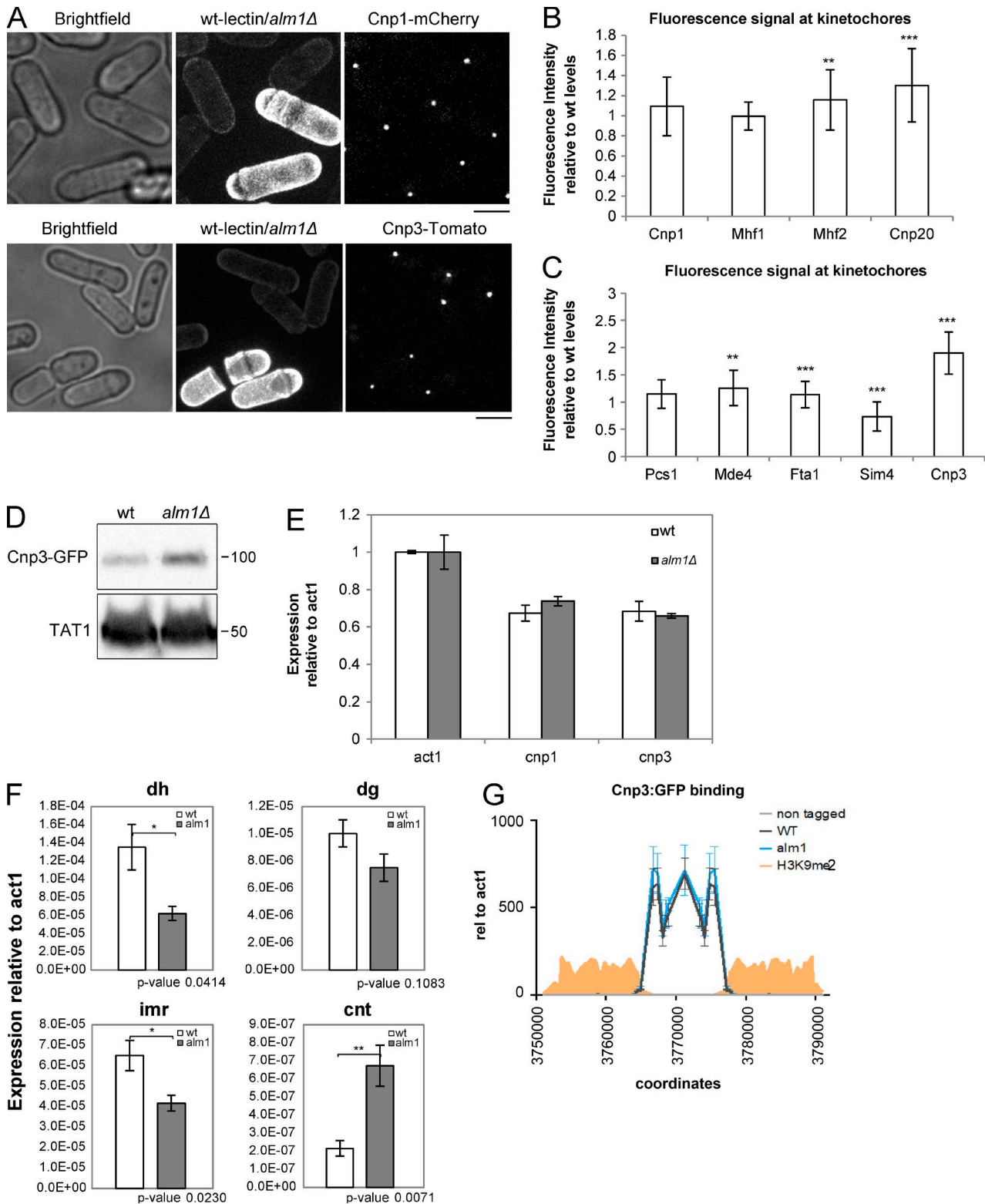


Figure 3. *alm1Δ* mutant shows altered accumulation of centromere and KT proteins. (A) WT (green lectin stained) and *alm1Δ* cells expressing Cnp1-mCherry (top) or Cnp3-Tomato (bottom). Bars, 5 μ m. (B and C) Mean fluorescence intensity in *alm1Δ* cells relative to WT cells of the indicated proteins (Cnp1-mCherry, Mhf1-GFP, Mhf2-GFP, Cnp20-GFP, Pcs1-GFP, Mde4-GFP, Fta1-GFP, Sim4-Tomato, and Cnp3-Tomato) at KTs during interphase. $n = 50$. (D) Western blot analysis of total Cnp3-GFP protein from WT and *alm1Δ* cells using anti-GFP mAb to detect Cnp3-GFP (top) and TAT1 as a loading control (bottom). Positions of molecular mass markers are indicated in kilodaltons. (E) RT-qPCR analysis of *act1*, *cnp1*, and *cnp3* mRNA levels in WT and *alm1Δ* cells. Three biological repeats were performed. (B, C, and E) Error bars represent SD. (F) RT-qPCR analysis of centromere I (*dh*, *dg*, *imr*, and *cnt*) transcript levels in WT and *alm1Δ* mutant (normalized to *act1* transcript levels). $n = 5$. Error bars represent SEM. (G) ChIP-qPCR analysis of Cnp3-GFP and H3K9me2 levels. ChIP data have been normalized to *act1* and are shown relative to the maximal enrichment in WT cells at the *cen1* region. *, $P < 0.05$; **, $P < 0.001$; ***, $P < 0.001$.

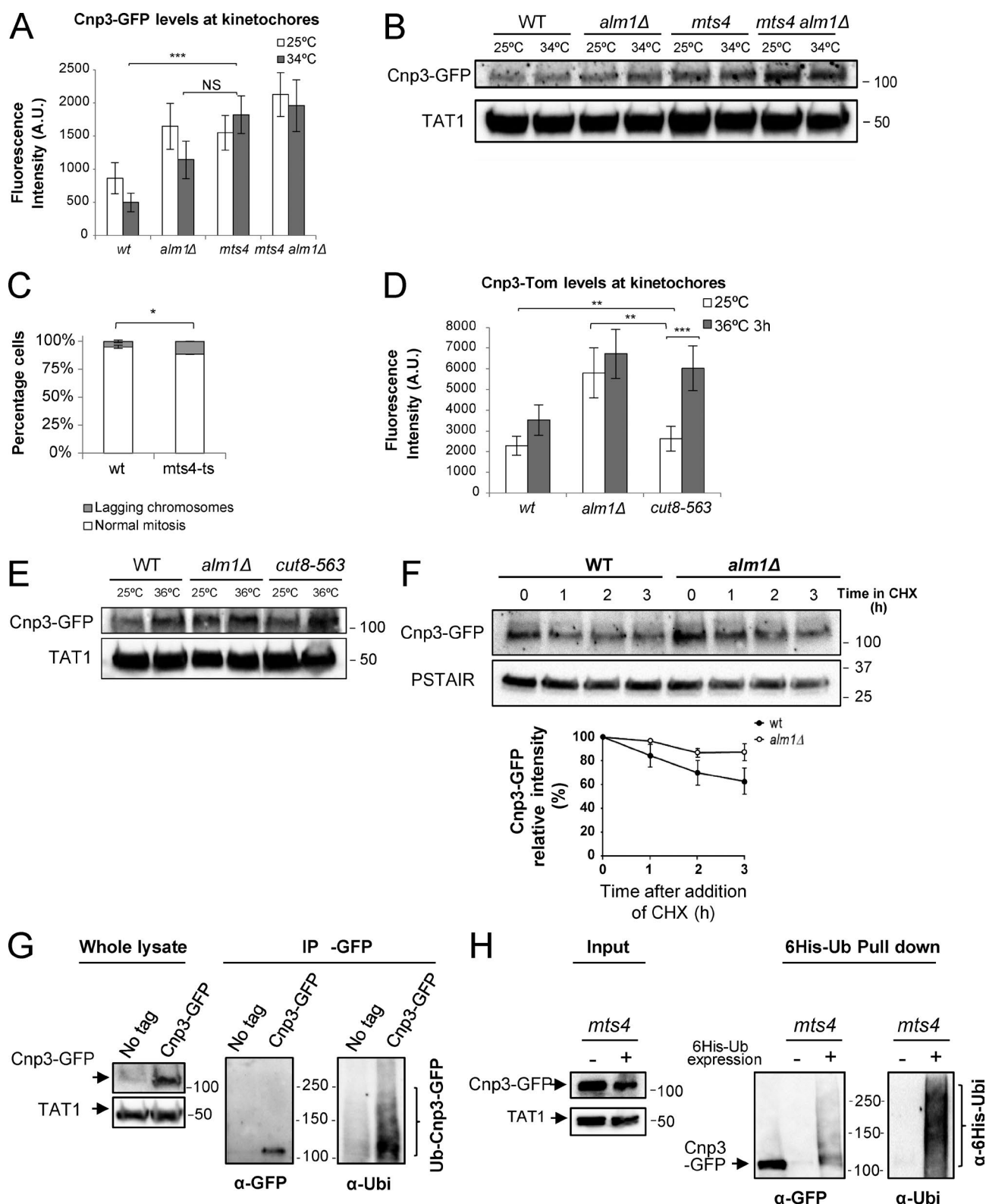


Figure 4. **Proteasome function and localization are required for stoichiometric accumulation of Cnp3 at KTs.** (A) Centromeric Cnp3-GFP levels in the indicated backgrounds and temperatures. $n = 50$. (B) Western blot analysis of total Cnp3-GFP, using anti-GFP mAb to detect Cnp3-GFP and TAT1 as a loading control. (C) Quantification of mitotic defects (lagging KTs during anaphase) of WT and *mts4* cells at 25°C by in vivo fluorescence microscopy, using Cnp3-GFP as a KT marker. $n > 140$. (D) Quantification of centromeric Cnp3-Tomato levels in the indicated backgrounds and temperatures. $n = 50$. (E) Western blot analysis of total Cnp3-GFP protein in the indicated strains, using anti-GFP mAb to detect Cnp3-GFP and antitubulin (TAT1) mAb as a loading control. (F) Cnp3 protein stability in WT and *alm1Δ* cells in the presence of CHX. Cnp3-GFP was detected by immunoblotting with anti-GFP mAb, and anti-PSTAIR mAb was used as a loading control. Quantification of Cnp3-GFP protein stability. Cnp3-GFP band intensities were quantified using ImageJ and normalized to PSTAIR signals. Relative intensity at time 0 was set up as 100% in each case. Error bars represent SD from three independent experiments. (G) *mts4* expressing Cnp3-GFP and untagged *mts4* cells were grown to midlog phase at 25°C and then shifted to 36°C for 3 h. Samples were collected and

found that both proteasome subunits fail to properly localize to the NE in the *alm1Δ* background (Fig 5, D–G).

Together, these results show that proteasome localization to the NE is altered in *alm1Δ* cells, which might affect the degradation of specific substrates.

An excess of Cnp3 impairs chromosome segregation

We reasoned that if the chromosome missegregation defects observed in *alm1Δ* cells are caused by abnormal accumulation of Cnp3 at KTs, the overexpression of this protein in WT cells would result in similar chromosome segregation defects. To test this hypothesis, we used a strain in which an additional copy of Cnp3-GFP was ectopically expressed under the control of the medium strength *nmt41* promoter (pINT-cnp3-GFP) in WT and *alm1Δ* Cnp3-GFP strains (Fennessy et al., 2014). Western blot analysis of total Cnp3-GFP protein in conditions of moderate overexpression (rich medium) showed increased levels of Cnp3-GFP in both WT and *alm1Δ* backgrounds (Fig. 6 A). Consistently, using live-cell microscopy, we observed a 2.4- and 1.4-fold increase in centromeric Cnp3-GFP intensity signal in WT and *alm1Δ* strains, respectively (Fig. 6, B and C). Furthermore, using the same Cnp3 overexpression conditions, we also found an increase in chromosome segregation defects, especially an increase in lagging chromosomes for both WT and *alm1Δ* cells (Fig. 6, D and E, asterisk and arrows).

We next tested whether a decrease in Cnp3 levels would rescue *alm1Δ* phenotypes. To this end, we inserted the thiamine-repressible *nmt81* promoter at the endogenous *cnp3* promoter locus in the *alm1Δ* background. Under repressed conditions (+ thiamine), this promoter rendered reduced Cnp3 levels (Fig. 6, F–H). This, however, only partially suppressed the chromosome segregation defects of *alm1Δ* (Fig 6 I). Thus, these data suggest that an excess of Cnp3 is one of the reasons for aberrant chromosome segregation seen in the absence of Alm1.

Discussion

The nucleoporins of the TPR family are the main structural element of the nuclear basket (Cordes et al., 1997; Bangs et al., 1998; Kosova et al., 2000; Frosst et al., 2002). As described for orthologues in other organisms, the fission yeast TPR Alm1 shows extensive regions of predicted CC domains, and we demonstrate that Alm1 indeed localizes at the nuclear rim, where it partially colocalizes with the NPC (Fig. S1, A–C). Alm1 colocalizes with Nup211, the other member of the TPR family of nucleoporins, in *S. pombe* (Fig. 1 A). Furthermore, Nup211 localization at the NE is decreased in the absence of Alm1 (Fig. 1 B), which suggests that they likely form a complex at the nuclear basket, as has been demonstrated for Mlp1 and Mlp2 in *S. cerevisiae* (Strambio-de-Castillia et al., 1999; Niepel et al., 2005).

In this work, we show that the absence of Alm1 causes a significant delay in the metaphase to anaphase transition because of SAC activation and the appearance of lagging chromosomes during anaphase B, which eventually leads to chromosome missegregation (Fig. 1, C–E; Fig. S2; and Fig. S3, A and B). A conserved function of the TPR proteins is the spatiotemporal regulation of the SAC through recruitment of Mad1 and Mad2 to the NE during interphase (Lee et al., 2008; Lince-Faria et al., 2009; Schweizer et al., 2013; Rodriguez-Bravo et al., 2014). We find that Alm1 is also required for the recruitment of Mad2 to the NE during interphase (Fig. S3 C). This indicates that this particular function of TPR nucleoporins has been conserved during evolution. In animal cells, depletion of TPR results in an accelerated anaphase onset, with decreased localization of Mad2 at KTs, weakened SAC response, and the appearance of lagging chromosomes (Lee et al., 2008; Lince-Faria et al., 2009; Rodriguez-Bravo et al., 2014). However, the *alm1Δ* mutant shows a delayed anaphase onset as a result of SAC activation, and Mad2 accumulates at KTs in unperturbed mitosis as well as in the presence of the MT-depolymerizing drug MBC (Fig. S3, A and E). This suggests that Alm1 is not required for SAC functionality, but that instead *alm1Δ* cells are dependent on this pathway for survival (Fig. S3 F). Thus, the chromosome missegregation observed in *alm1Δ* might result from dysfunctional KTs, as the mitotic spindle is apparently normal in *alm1Δ* cells (Fig. S2, C–E).

Proper centromere and KT structure and function are crucial to avoid erroneous MT–KT attachments (Nonaka et al., 2002; Gregan et al., 2007). For instance, inactivation of Cnp1 affects the loading of other KT proteins, such as Cnp20 or Mis6 (Castillo et al., 2007; Tanaka et al., 2009; Folco et al., 2015), and overexpression of Cnp1 leads to increased levels of centromeric Cnp3, Mal2, and Sim4 (Castillo et al., 2007), resulting in chromosome missegregation. The *alm1Δ* mutant shows increased levels of several KT proteins, including of Cnp3 (Fig. 3, A–C), which is not the result of deregulated transcription (Fig. 3 E). The increased level of Cnp3 at KTs observed in *alm1Δ* is not likely the result of its direct binding to centromeric DNA, as we did not observe an increase of centromere-associated Cnp3 by ChIP (Fig. 3 G). The C-terminal domain of CENP-C^{Cnp3} contains two regions that are highly conserved from yeast to humans that are called Mif2p homology domain II (also known as CENP-C motif) and III. The Mif2 homology domain II is responsible for CENP-C binding to centromeric DNA (Trauzzi et al., 2002, 2009), whereas the Mif2 homology domain III displays multiple activities, including recruitment of other KT components. Importantly, the Mif2 homology domain III is able to mediate CENP-C self-association *in vivo*, allowing the formation of higher order structures such as homodimers and homo-oligomers (Sugimoto et al., 1997; Cohen et al., 2008; Trauzzi et al., 2009). Therefore, it seems conceivable that the excess of Cnp3 protein in *alm1Δ* cells may cause its ectopic accumulation at KTs through binding to other Cnp3 molecules

subjected to anti-GFP immunoprecipitation (IP). Whole lysate (left) was immunoblotted with anti-GFP mAb to detect Cnp3-GFP and antitubulin (TAT1) mAb as a loading control. Immunoprecipitated proteins were immunoblotted with anti-GFP mAb to detect Cnp3-GFP and antiubiquitin pAb to detect ubiquitinated proteins. (H) *mts4* cells overexpressing Cnp3-GFP or Cnp3-GFP and His₆-ubiquitin were grown in EMM to midlog phase at 25°C and then shifted to 36°C for 3 h. Samples were collected, and polyubiquitinated proteins were purified with Ni²⁺-nitrilotriacetic acid beads in denaturing conditions. Ubiquitinated proteins were detected by immunoblotting with antiubiquitin pAb (right), and ubiquitinated forms of Cnp3 were detected by immunoblotting using anti-GFP mAb (left). A fraction of the whole cell extract (left) with an equal amount of total protein was immunoblotted with anti-GFP mAb to detect Cnp3-GFP and antitubulin (TAT1) mAb as a loading control. Positions of molecular mass makers are indicated in kilodaltons. Graphs represent mean and SD. A.U., arbitrary units. *, P < 0.05; **, P < 0.001; ***, P < 0.001.

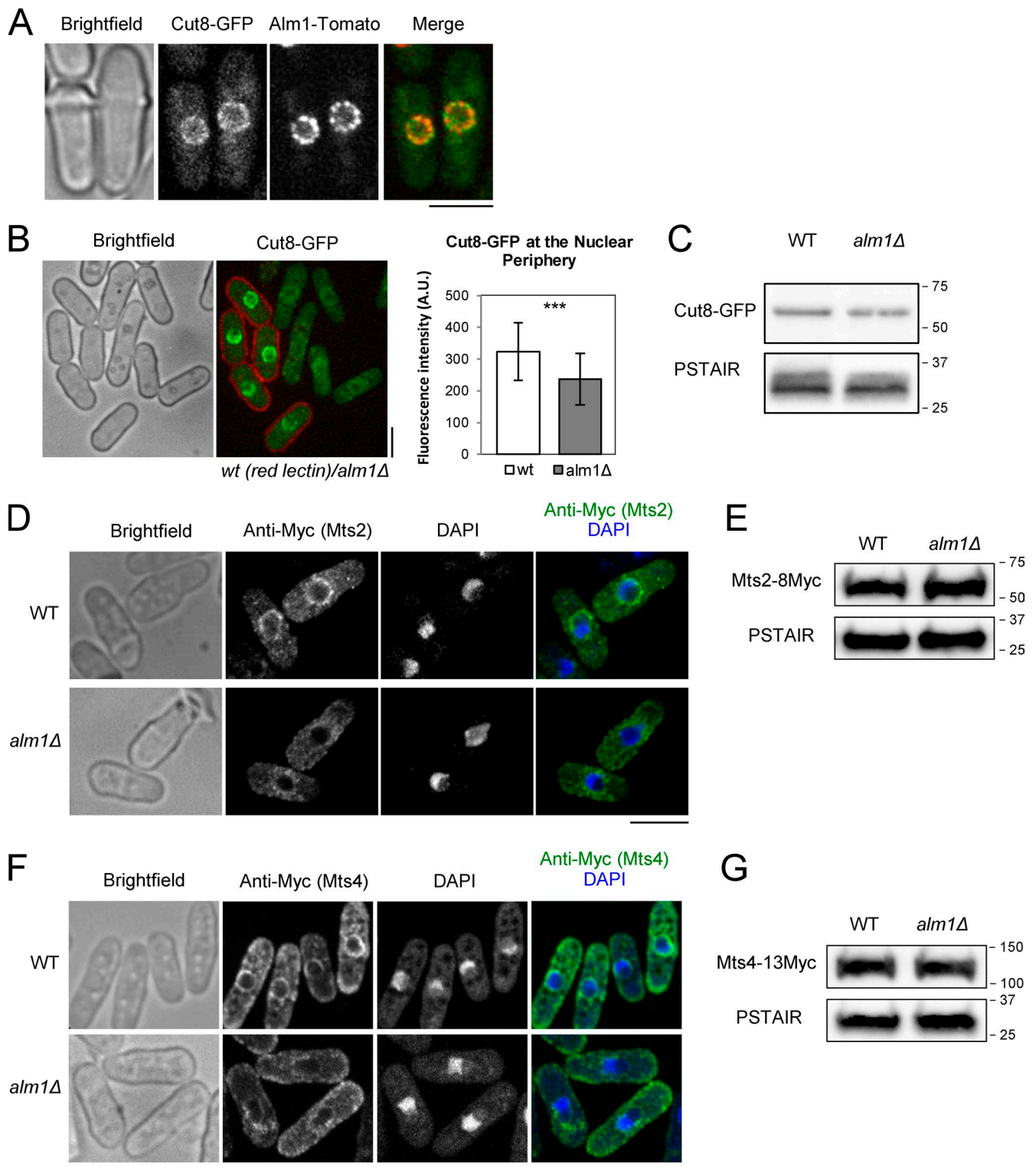


Figure 5. **Alm1 is required for proper accumulation of Cut8 and the proteasome to the NE.** (A) WT cells expressing Cut8-GFP and Alm1-Tomato. Images are maximal projections of three z sections. (B) WT (red lectin stained) and *alm1Δ* cells expressing Cut8-GFP. Cut8-GFP intensity levels at the NE in WT and *alm1Δ* cells. Graphs represent mean and SD. $n = 69$. (C) Western blot analysis of total Cut8-GFP protein of WT and *alm1Δ* cells using anti-GFP mAb to detect Cut8-GFP and anti-PSTAIR mAb as a loading control. (D) Brightfield and immunofluorescence images of WT and *alm1Δ* cells expressing Mts2-8Myc, using anti-Myc antibodies against Mts2-8Myc (green) and DAPI to stain DNA (blue). (E) Western blot analysis of total Mts2-8Myc protein in WT and *alm1Δ* cells using anti-Myc antibodies to detect Mts2-8Myc and anti-PSTAIR antibodies as a loading control. (F) Brightfield and immunofluorescence images of WT and *alm1Δ* cells expressing Mts4-13Myc, using anti-Myc antibodies against Mts4-13Myc (green) and DAPI to stain DNA (blue). (G) Western blot analysis of total Mts4-13Myc protein in WT and *alm1Δ* cells using anti-Myc antibodies to detect Mts4-13Myc and anti-PSTAIR antibodies as a loading control. Positions of molecular mass markers are indicated in kilodaltons. Bars, 5 μm . A.U., arbitrary units. ***, $P < 0.001$.

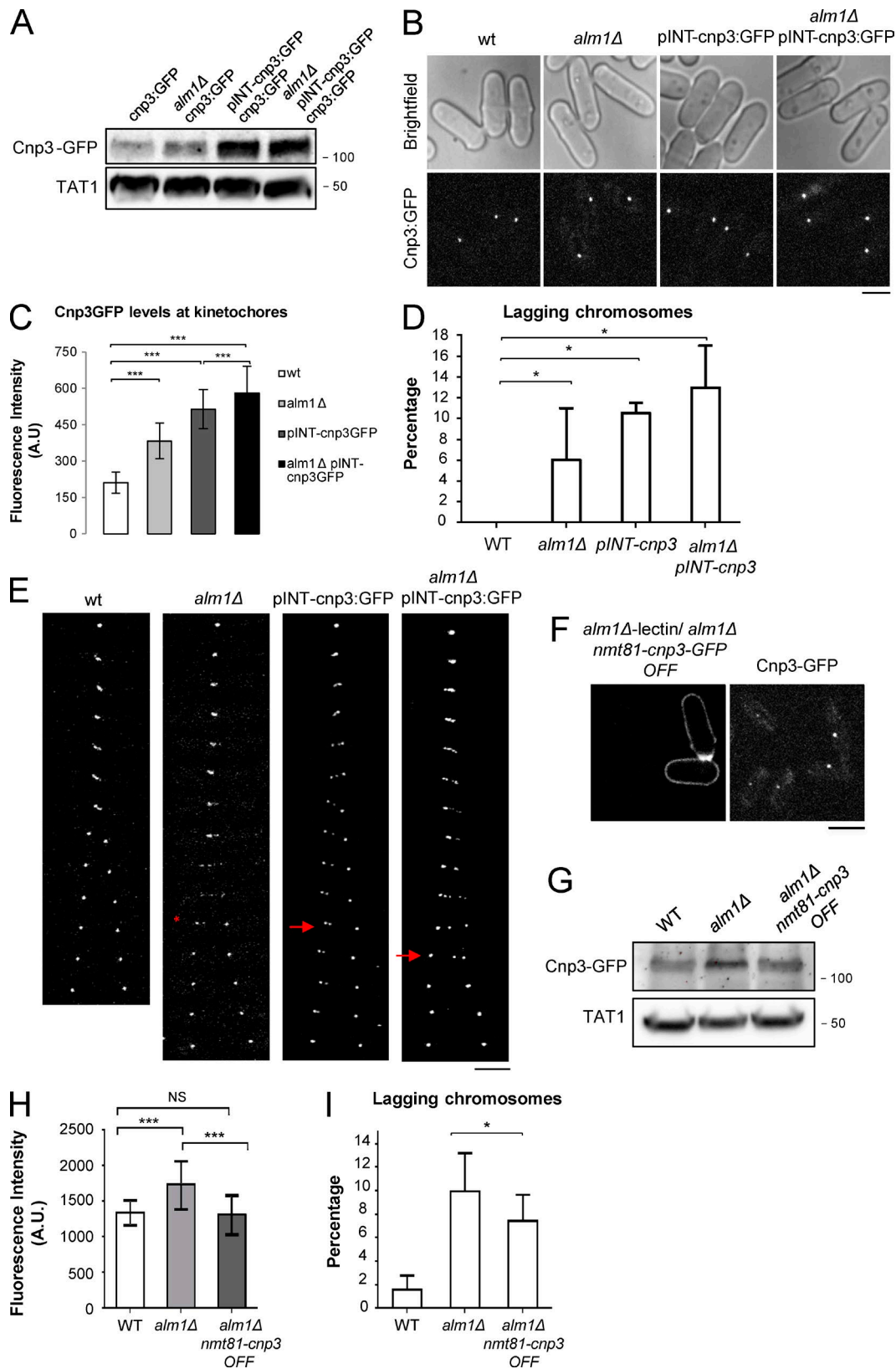


Figure 6. **Cnp3 overexpression phenocopies *alm1Δ* segregation defects.** (A) Western blot analysis of total Cnp3-GFP using anti-GFP mAb or antitubulin (TAT1) mAb as a loading control. (B) Images of WT and *alm1Δ* cells expressing Cnp3-GFP from the endogenous locus and WT and *alm1Δ* cells expressing an additional copy of Cnp3-GFP from the medium-strength promoter *nmt81* (pINT-Cnp3-GFP) in repressed conditions (YES media). (C) Centromeric Cnp3-GFP intensity levels in the indicated strains. $n = 50$. (D) Quantification of mitotic defects (lagging KTs during anaphase B). $n = 50$. (E) Time-lapse fluorescence images of the indicated strains. Time between frames is 2 min. Red asterisk denotes asynchronous KT segregation, and arrows denote lagging KTs. (F) Fluorescence images of *alm1Δ* cells expressing Cnp3-GFP from the endogenous promoter (marked with red lectins) and *alm1Δ* cells expressing

mediated by the Mif2 homology domain III. In *S. pombe*, Cnp3 binds through the N-terminal domain to Fta1, a component of the Sim4 complex required for proper KT function (Tanaka et al., 2009). Consistently, we observed increased levels of centromeric Fta1-GFP in the *alm1Δ* background (Fig. 3 C), suggesting that an excess of Cnp3 might drive the ectopic accumulation of other KT components.

Cnp3 is one of the substrates degraded in an Alm1- and proteasome-dependent manner (Fig. 4). However, we found that in WT cells, the Cnp3 turnover is less prominent than that of Cdc13 in the presence of CHX (Fig. 4 F and Fig. S4 D), suggesting that only a fraction of Cnp3 is degraded by the proteasome, similar to what has been shown for Cnp1 regulation by proteasomal degradation at centromeres (Kitagawa et al., 2014). In our experimental conditions, the absence of Alm1 did not significantly affect Cdc13 degradation (Fig. S4 D). Whether the different requirement of Alm1 for Cnp3 and Cdc13 degradation is the result of spatial specificity in proteasomal degradation or a consequence of different kinetics of degradation of both proteins remains an open question.

Overexpression of Cnp3 is sufficient to cause chromosome missegregation in WT cells (Fig. 6, D and E). However, a decrease of Cnp3-GFP levels only partially rescues chromosome segregation defects of *alm1Δ* cells (Fig. 6 I), suggesting that the imbalance of other centromere or KT factors (Fig. 3, B and C) may also contribute to the impaired KT function.

Among the synthetic genetic interactions obtained from the SGA analysis, we found that mutants with defects in KT components and SAC regulators are enriched (Fig. 2 D). This finding is in agreement with our observation of aberrant KT behavior, SAC activation and dependency, and chromosome missegregation in *alm1Δ* cells. We also obtained genes with roles in chromatin organization. These include *eaft6*, which encodes a subunit of the Mst2 histone acetyltransferase complex that regulates histone turnover at centromeres and prevents heterochromatin spreading into euchromatic sites (Wang et al., 2015). We further found *fft3*, which encodes an ATP-dependent chromatin remodeler required to prevent euchromatin formation at centromeres (Strålfors et al., 2011). Interestingly, Fft3 has also been involved in heterochromatin maintenance and NE attachment of subtelomeric regions (Steglich et al., 2015). Intriguingly, we found that transcription at centromeric chromatin is affected in *alm1Δ* cells (Fig. 3 F). Whether this phenotype is caused by deficient chromatin regulation pathways or is a consequence of altered KT proteins will be further investigated. Finally, we also found genes involved in ubiquitin-dependent proteolysis, including *ubp2* that encodes a deubiquitinating enzyme. ScUbp2 counteracts ubiquitination of the proteasome-associated ubiquitin receptor Rpn10 (Sato et al., 2005). Because ubiquitination of Rpn10 inhibits its capacity to bind to substrates and affects proteasome activity, the lack of Ubp2 might additionally increase cellular Cnp3 levels, providing a likely explanation of the synthetic genetic interaction with the *alm1Δ* mutant.

In fission yeast, the three centromeres cluster together and are anchored at the NE in interphase cells. During mitosis,

they are released from the NE and then attached by the incipient spindle during prophase and metaphase. *S. pombe* has a closed mitosis in which the NE does not disassemble. Thus, except for a brief period during early mitosis, most of the cell cycle centromeres stay in contact with the NE. Intriguingly, the proteasome is also enriched at the nuclear periphery (Wilkinson et al., 1998), and CENP-A^{Cnp1} has been shown to be regulated by proteasomal degradation (Collins et al., 2004; Moreno-Moreno et al., 2006; Ranjitkar et al., 2010; Kitagawa et al., 2014). We have shown that Cnp3 is also regulated by the proteasome. In particular, Cnp3 levels are increased in mutants affected in proteasomal degradation, such as *mts4* or *cut8-563* mutants (Fig. 4, A–E), and ubiquitinated forms of Cnp3 can be detected when proteasomal degradation is compromised (Fig. 4, G and H). Importantly, we find that Alm1 is required to maintain the proteasome and its anchor Cut8 at the NE (Fig. 5, B, D, and F). Thus, Alm1-dependent proteasome localization at the NE is required for Cnp3 proteostasis.

Cut8 has a membrane-binding domain (Takeda et al., 2011), and its enrichment at the nuclear periphery is regulated by the ubiquitin-conjugating enzyme Rhp6 (Rad6), the ubiquitin ligase Ubr1, and the proteasome itself (Takeda and Yanagida, 2005). Thus, although Alm1 may not be directly involved in the recruitment of Cut8 and the proteasome, it may provide a platform for the assembly or tethering of functional proteasomal complexes at the NE. Interestingly, ScMlp1 and ScMlp2 have been isolated in a complex with the NE-associated protein Esc1 (Niepel et al., 2013). Esc1 is involved in a variety of nuclear functions such as telomere NE anchoring and silencing (Taddei et al., 2004), DNA repair (Therizols et al., 2006), or SUMO homeostasis (Lewis et al., 2007), and, recently, it has been shown to bind to the proteasome (Niepel et al., 2013). Therefore, it has been proposed that Mlp1 and Mlp2 are part of a protein–protein interaction network at the NE that includes the SPB, messenger RNPs, chromatin silencing factors, and the proteasome via Esc1 (Niepel et al., 2013). Further studies will contribute to the understanding of the role of proteasomes at the NE.

Our study reveals a novel role of the TPR nucleoporin Alm1 in the homeostasis of KT proteins through the spatial regulation of the proteasome and highlights the emergent role of the NPC in the maintenance of genome integrity.

Materials and methods

Yeast strains, genetic procedures, cell culture, and media

The *S. pombe* strains used in this study are listed in Table S1. Strains expressing GFP-, mCherry-, and tdTomato-tagged (Snaith et al., 2005) proteins and deletion strains were constructed using PCR-based methods and homologous recombination as described previously (Bähler et al., 1998). *ura4+*, *kanMX6* (*Kan*), *hphMX6* (*hph*), and *natMX6* (*nat*) genes were used as selection markers (Bähler et al., 1998; Sato et al., 2005; Snaith et al., 2005). All tagged strains used in this study are expressed from their endogenous loci unless otherwise stated. Yeast transformation was done by the lithium acetate method (Moreno et al., 1991).

Cnp3-GFP from the *ntm81* promoter under repressed conditions (EMM + thiamine). (G) Western blot analysis of total Cnp3-GFP protein from cultures of the indicated genotypes grown in EMM + thiamine, using anti-GFP mAb or antitubulin (TAT1) mAb as a loading control. Molecular mass is given in kilodaltons. (H) Quantification of centromeric Cnp3-GFP intensity levels. *n* = 50 cells. (I) Quantification of mitotic defects (lagging KTs during anaphase B) of the indicated strains. *n* = 50. Graphs represent mean and SD. Bars, 5 μm. A.U., arbitrary units. *, *P* < 0.05; ***, *P* < 0.001.

The *alm1+* gene was deleted with the *ura4+* gene by using the primers Alm1-Ura4.s (5'-TTTCTTCTAACGTGGTATTATTTCTGTACATCGAGGTTGATCACTTACTTCTCGTATACGTACTGGT GCTTCATCGCCAGGGTTTTCCAGTCACGAC-3') and Alm1-Ura4+.a (5'-AAAGTCGTTTTTTTTATTGAAACCAGAAATAA TAATTATGCAGCTAACCTATTTATTATCTGTTTACAAACTCTA GCGGATAACAATTCACACAGGA-3'). The primers used to construct the strain *alm1-tomato-Nat* were Alm1-Tag.s (5'-ACACTA ATTCTCCTCTAAACGGTCCAGTTTCAGACGCTGGTATGGATG TTTCCAATGATGTTAAGAAAGCCAAACCCGGACGGATCCCC GGGTTAATTA-3') and Alm1-Tag.a (5'-AATTAAGTCGTTTT TTTTATTGAAACCAGAAATAATAATTATGCAGCTAACCTATT TTATTATTGTTTACAAACTCTGAATTCGAGCTCGTTTAAAC-3'). Primers to construct the strain *nup211-GFP-kan* were Nup211_Ctag_ FW (5'-CTGCTAAATCCGGCTCCCTTAAAAGACAACGTGACG ATGCGAACAAGGAGGATCCAGTTCGAACCAAAAGAAAGC AAAACGGATCCCCGGGTTAATTA-3') and Nup211_Ctag_RV (5'-TTTACTCATGTCATTATTATAAATCATGTTAACTAAATATG AATAGTCCTAAGAGTGATTTATGAACCATATGAAAACATGAA TTCGAGCTCGTTTAAAC-3').

The primers used to construct the strain *cnp3-GFP-kan* were *cnp3.C-tagFor* (5'-CTGTTTCGACTTTATTATACCCATGCGACCGAC ACCTTGGAAAATAAAGAAGGGGAATAGGGGATTTTCCAAA CGAACGACGGATCCCCGGGTTAATTA-3') and *cnp3.C-tagRev* (5'-AAATTGATAACAGATAAATAATTACATAACGACTAAATGA CTTTTGTACATCGAAATCCATAACACACTATTAGTATTGGAAT TCGAGCTCGTTTAAAC-3').

The primers used to construct the strain *mts4-13Myc-kan* were *mts4-Forward* (5'-CGGAAGCATATACCTTTGACTTCGTTGG AAGGTATTGTTATTTAAAAAAAATACGGAGGACATTGAAA TGACCGCTCGGATCCCCGGGTTAATTA-3') and *mts4-Reverse* (5'-GGGCGCTGAATTTCAACAATCCAGTTTGTTTTATTAATAGC AAAGAAGTGGAAATCAAGCATTAGTACATAAAAATTGTAGAA TTCGAGCTCGTTTAAAC-3').

To construct the strain *pINT-cnp3-GFP-Nat*, *cnp3* was amplified from genomic DNA using the primers BamH1-Cnp3.F (5'-GGA GGATCCATGACGATGAATGAAACGTCT-3') and BamH1-Cnp3.R (5'-GTAGGATCCTCGTTCGTTTGGAAAATCC-3') and cloned into pINTH-41EGFPC plasmid (Fennessy et al., 2014). The strain *nmt81-Cnp3-GFP-Nat alm1::ura4+* was constructed by integrating the *kanMX6-p81nmt* cassette at the *cnp3* promoter locus in the *alm1::ura4+ Cnp3-GFP-Nat* strain. The primers used to amplify the *kanMX6-p81nmt* cassette were *nmt81-Cnp3.F* (5'-AGACAACACTTA CGTCCCCATATATTGTTTTGTTTGGCTATTTAGGCTATATTTTA CTTAATGTTTATTTAATAGAATAGAATTCGAGCTCGTTTAAAC-3') and *nmt81-Cnp3.R* (5'-ACGTACTGCCTACGACACCAATT TCAAAAACACTGATTCTCGTTGTCTTGCCGGAATAGCAGAC GTTTCATTCATCGTCATGATTTAACAAGCGACTATA-3').

Strains harboring Sid2-Tomato, Mis6-GFP, Cut12-GFP, *lys1::LacOp*his7+::LacI-GFP-NLS, *pcs1-GFP*, or *mde4-GFP* were derived from original strains provided by F. Chang (University of California, San Francisco, San Francisco, CA), T. Toda (The Francis Crick Institute, London, England, UK), V. Alvarez (Centro Andaluz de Biología del Desarrollo/Universidad Pablo de Olavide [CABD/UPO], Seville, Spain), P. Nurse (The Francis Crick Institute), and J. Gregan (Research Institute of Molecular Pathology, Vienna, Austria), respectively. Additional strains listed in Table S1 were provided by P. Nurse (The Francis Crick Institute), S. Moreno (Instituto de Biología Funcional y Genómica, Salamanca, Spain), J. Petersen (South Australian Health and Medical Research Institute, Adelaide, Australia), Y. Watanabe (Institute of Molecular and Cellular Biosciences, University of Tokyo, Tokyo, Japan), I. Hagan (Cancer Research UK, University of

Manchester, Manchester, England, UK), Y. Hiraoka (Advanced ICT Research Institute, Kobe, Japan; National Institute of Information and Communications Technology, Kobe, Japan/Graduate School of Frontier Biosciences, Osaka University, Suita, Japan), O. Niwa (Kazusa DNA Research Institute, Chiba, Japan), and the National Bio-Resource Project (Osaka, Japan).

Strain crosses and sporulation were performed in sporulation agar at 25°C, and genetic dissection was performed using a dissection microscope (MSM 400; Singer Instruments). Standard cell culture procedures and media were used (Moreno et al., 1991). Unless otherwise stated, experiments were performed in rich media (YES) at 25°C, and cells were grown until exponential midlog phase. For *mts4-ts* mutant, cells were grown at the permissive temperature of 25°C and shifted to 34°C for 1 h. For *cut8-563-ts* mutant, cells were grown at the permissive temperature of 25°C and shifted to 36°C for 3 h.

For sensitivity drop assays, strains were grown in YES to exponential midlog phase at 25°C. Serial fivefold dilutions were made, starting from an OD of 0.2, and spotted onto YES plates. For TBZ sensitivity assay, cells were spotted onto YES containing DMSO or YES plates containing 15 µg/ml TBZ (T8904; Sigma-Aldrich) and grown at 25°C and 30°C. Pictures were taken after incubation for 3–6 d.

For Cnp3-GFP down-regulation, cells expressing Cnp3-GFP under the control of the weak *nmt81* promoter were grown in Edinburgh minimal media (EMM) at 25°C, and then 15 µM thiamine (T4625; Sigma-Aldrich) was added to the culture medium. Cells were further grown for three generations in the presence of thiamine.

Microscopy and image analysis

Live-cell imaging was performed with a spinning-disk confocal microscope (IX-81; Olympus; Evolve camera, Plan APOchromat 100x, 1.4 NA objective; Roper Scientific). Images were acquired with Metamorph software (Molecular Devices) and analyzed with ImageJ (National Institutes of Health). Unless otherwise stated, time-lapse experiments were performed at 25°C in YES, and images are maximal projections of 18 z sections with a step size of 0.3 µm. In brief, cells were collected by centrifugation and immobilized in soybean lectin (L1395; Sigma-Aldrich)-coated 35-mm glass-bottom culture dishes (P35-1.5-10-C; MatTek) or µ-Slide 8-well dishes (80827; Ibidi). Conditioned media were used to maintain cells during filming. When two strains were filmed in the same microscope field, one of them was stained with either FITC- (F2401; EY Laboratories) or TRI TC-conjugated lectins (T2401; EY laboratories) by incubating 100 µl of cell culture with 1 µl of lectins for 5 min at room temperature. Cells were washed twice with fresh media and immediately mixed and prepared for filming.

For MBC treatment, cells were grown to exponential midlog phase in YES at 25°C and treated with 10 µg/ml MBC (37867-4; Sigma-Aldrich) or DMSO as control 10 min before filming.

Unless otherwise stated, quantification was performed on maximal projections of 18 z sections with a step size of 0.3 µm. Quantification of Nup211-GFP in Fig. 1 B was performed on maximal projections of three z sections of 0.3 µm. Quantification of Cut8-GFP in Fig. 5 B was performed in single-central z sections. For fluorescence signal quantification, background was subtracted after measuring intensity in every region of interest.

Minichromosome loss assay

500–1,000 cells from Ade + colonies of WT and *alm1Δ* backgrounds were plated on YES + 0.15 mg/ml adenine plates and incubated for 3–5 d. The number of colonies with a red sector was counted. The number of chromosome loss events per division is the number of these half-sectored colonies divided by the total number of colonies.

Western blot and kinase assay

Protein extracts for Western blotting were prepared by trichloroacetic acid precipitation. Unless otherwise stated, Western blots were performed from asynchronous cultures. In brief, 45 ml of exponentially growing cultures was quenched by the addition of 5 ml of ice-cold 100% trichloroacetic acid (Sigma-Aldrich). After 30-min incubation on ice, cells were pelleted by centrifugation and washed with 10 ml of ice-cold 100% acetone (Sigma-Aldrich). After removing and drying acetone, cell pellets were resuspended in 500 μ l of beating buffer (8 M urea, 50 mM ammonium bicarbonate, and 5 mM EDTA) with protease inhibitors (cOmplete, Mini, EDTA free; 11836170001; Roche) and lysed in a sample preparation system (FastPrep; MP Biomedicals) with acid-washed glass beads (Sigma-Aldrich). Cell extracts were cleared by centrifugation, the supernatant was recovered, and the protein concentration was measured using an RC DC protein assay kit (5000121; Bio-Rad). Protein samples were loaded on 10% TGX-Fast Cast acrylamide gels (161-0183; Bio-Rad) and blotted on nitrocellulose membranes (Bio-Rad). Blots were probed with primary mouse anti-GFP mAb (11814460001; Roche) and either mouse anti- α -tubulin mAb (TAT1; K. Gull, University of Oxford, Oxford, England, UK) or anti-PSTAIR mAb (P7962; Sigma-Aldrich) at a 1:1,000 dilution, followed by the secondary antibody anti-mouse IgG at a 1:2,000 dilution (A3562; Sigma-Aldrich). ECL (Bio-Rad) or Supersignal West Femto (Thermo Fisher) and Chemidoc XRS+ (Bio-Rad) were used for detection.

For Ark1-kinase assay, *cdc25-22* and *alm1 Δ cdc25-22* cells expressing Ark1-PK were grown to midlog phase in YES at 25°C and then synchronized in G2 by shifting to 36°C for 4 h. Cells were released to 25°C for 45 min, and samples were taken every 15 min. Ark1-kinase assay was performed as previously described (Petersen et al., 2001). In brief, 50 ml of cells was harvested by centrifugation, and the pellets were washed with ice-cold STOP buffer (150 mM NaCl, 50 mM NaF, 10 mM EDTA, and 1 mM Na₃N, pH 8.0) and snap-frozen in liquid nitrogen. The cell pellets were resuspended in 200 μ l HEN buffer (50 mM Hepes, pH 8.0, 150 mM NaCl, 5 mM EGTA, 5 mM EDTA, 1% NP-40, and 50 mM β -glycerophosphate) with protease inhibitors (cOmplete Mini, EDTA free; 11836170001), and extracts were prepared by glass bead lysis at 4°C. Cell extracts were cleared by centrifugation. 30 μ l Sepharose beads (17-0618-01; GE Healthcare) was coupled with 10 μ l of mouse anti-PK mAb (MCA 1360; Serotec) for 1 h at 4°C in an agitation platform. 95 μ l of cleared extract was added to 10 μ l of packed beads and incubated at 4°C for 30 min. The beads with the associated immunoprecipitate were washed three times with 0.5 ml HEN buffer and three times with KAB buffer (50 mM Hepes, pH 7.5, 150 mM NaCl, 1 mM DTT, and 10 mM MgCl₂) and resuspended in 20 μ l KAB buffer. Beads were incubated at 32°C for 5 min, and then 5 μ l of substrate mix (20 μ M ATP and 5 μ g of purified Histone H3; 11034758001; Roche) was added and incubated further for 20 min at 32°C. The reaction was stopped by the addition of 15 μ l 2 \times SDS-PAGE loading buffer (0.25 M Tris-HCl, pH 6.8, 8% [wt/vol] SDS, 0.004% [wt/vol] bromophenol blue, and 20% [vol/vol] 2-mercaptoethanol) and incubated at 98°C for 5 min. 25 μ l of samples was immunoblotted with mouse anti-PK mAb (provided by I. Hagan) to detect Ark1-Pk and rabbit polyclonal anti-phosphoserine 10-histone H3-antibody (provided by I. Hagan). In all experiments, at least three biological repeats were performed.

Cell fixation and immunostaining

For DAPI staining, cells were fixed with 70% cold ethanol, washed with PBS, and resuspended in PBS + 0.2 μ g/ml DAPI (Sigma-Aldrich).

Immunofluorescence microscopy was performed as described previously (Hagan and Hyams, 1988). In brief, 10 ml of exponentially growing culture was fixed with freshly prepared 30% paraformaldehyde

(294474L; VWR) in PEM buffer (0.1 M Pipes, 2 mM EGTA, and 1 mM MgSO₄, pH 6.9) plus glutaraldehyde (G6257; Sigma-Aldrich) at a final concentration of 0.2%, followed by 1-h incubation at room temperature. After washing three times with PEM, the cell wall was digested during 1 h at 37°C with 2.5 mg/ml zymolyase 20T (120491-1; AMSBIO) in PEMS (PEM + 1.2 M sorbitol); cells were permeabilized with 1% Triton X-100 and quenched with 1 mg/ml sodium borohydride. After washing twice with PEM, cell pellet was resuspended in PEMBAL buffer (PEM buffer, 0.1% sodium azide, and 1% BSA) before antibody incubation. Primary antibody was mouse anti-Myc (9E10; Santa Cruz), used at a dilution of 1:100 for mts2-8Myc and 1:200 for mts4-13Myc. The secondary antibody was goat Alexa Fluor 488-tagged anti-mouse (A11029; Invitrogen, Molecular Probes), used at a dilution of 1:1,000.

SGA assay

SGA assay based on TBZ sensitivity was performed as described previously (Verrier et al., 2015), using as query strains WT and *alm1 Δ* mutant. Using the Singer RoToR HDA, WT and *alm1 Δ* cells were crossed with 3,400 gene deletion mutants (*Bioneer* haploid deletion mutant library, v. 3.0) and selected by incubation at 42°C for 4 d (temperature method; Dixon et al., 2008). After haploid selection, two additional steps of double mutant selection were performed, spotting onto EMM-Ura (PMD0410; ForMedium) and YES (Difco) supplemented with 100 μ g/ml G418 (1013027; Invitrogen) for selection of deletion mutants, 100 μ g/ml hygromycin (10687-010; Invitrogen) for selection of marker next to centromere, and 100 μ g/ml ClonNat (Werner Bioreagents) for selection of *alm1 Δ* mutant. Finally, cells were spotted on YES plates containing TBZ at a final concentration of 10 μ g/ml and YES with DMSO as a control. The growth of the single and double mutants was quantified on both media, and the ratio with the median ratio was compared (Barrales et al., 2016). Genetic interactions of double mutants were analyzed based on colony size (area) as a readout of cellular fitness. GO enrichment analysis was performed using the term enrichment service of AmiGO 2 (GO Consortium), searching for biological process without using the Bonferroni correction.

mRNA levels

Cells were grown in liquid YES medium at 25°C to exponential midlog phase, and total RNA was obtained by using a spin column (RNeasy Mini kit; Qiagen) according to the manufacturer's instructions. Transcriptional levels of *act1*, *cnp1*, and *cnp3* were measured by RT-qPCR using iTaq Universal SYBR green one-step kit as indicated (Bio-Rad) in 10 μ l of final volume with 5 μ l iTaq Universal SYBR green reaction mix, 0.125 μ l iScript reverse transcriptase, a mixture of forward and reverse primers (final concentration of 300 nM each), and 500 ng RNA. The primers used were *cnp1.F* (5'-GTTTGGCGTGGCAATCTA CG-3'), *cnp1.R* (5'-CCTGGCTAATGCATGTCTCG-3'), *cnp3.F* (5'-CGTTGAAATGCCAGCAGGAG-3'), *cnp3.R* (5'-ACTGTGACCTCG ATCTTTCCC-3'), *act1.F* (5'-AAGTACCCCATGAGCACGG-3'), and *act1.R* (5'-CAGTCAACAAGCAAGGGTGC-3').

RT-qPCR

Total RNA was isolated from cells grown to exponential midlog phase in YES at 25°C using a TRIzol reagent (Life Technologies). After removing DNA contamination from the total RNA with TURBO DNA-free (Applied Biosystems), samples were subjected to RT-PCR analyses using the SuperScript III First-Strand Synthesis SuperMix (Life Technologies) for cDNA synthesis and treated with RNase. RT-qPCR reactions were carried out in 15 μ l of volume, with 7.5 μ l Light Cyclers 480 SYBR green master mix (Roche), 2.5 μ l of a mixture of forward and reverse primers (1.5 μ M), and 5 μ l cDNA, previously diluted. The primers used were *act1-4.F* (5'-GATTCTCATGGAGCGTGGTT-3'), *act1-4.R*

(5'-CGCTCGTTTCCGATAGTGAT-3'), cen-dh.F (5'-TGAATCGTG TCACTCAACCC-3'), cen-dh.R (5'-TGAATCGTGTCAATCAACCC-3'), cen-dg-1.F (5'-TGCTCTGACTTGGCTTGTCTT-3'), cen-dg-1.R (5'-CCCTAACTTGGAAAGGCACA-3'), SG1953 (5'-TCGCCGTA ACAAAGGATCA-3'), SG1954 (5'-GCATTAGACAACCTCGTTC GATC-3'), imr. MR35.F (5'-GAGCATGGTGGTGTATGGA-3'), and imr. MR36.R (5'-CGACTAAACCGAAAGCCTCGA-3').

ChIP analyses

ChIP was performed as described in Braun et al. (2011) with minor modifications. In brief, Cnp3-GFP cell cultures were incubated overnight in rich medium at 30°C, diluted in fresh rich medium, and incubated until they reached an OD₅₉₅ of ~0.6. Cells were cross-linked with 1% formaldehyde for 10 min at room temperature, and the reaction was quenched by adding glycine to a final concentration of 125 mM. Chromatin was extracted and sheared using a sonicator (QSonica Q800R1; 30 min total, with 30-s on/off cycles). Chromatin from lysates corresponding to 15–30 OD₆₀₀ of cells was immunoprecipitated with 2 µg antibody (anti-GFP, provided by A. Ladurner, Munich University, Munich, Germany) using Dynabeads Protein G (Life Technologies). Immunoprecipitated DNA was quantified by qPCR using Fast SYBR green master mix (Life Technologies) and a 7500 Fast Real-Time PCR system (Applied Biosystems) using a tiling array of primers covering the centromere (Braun et al., 2011).

Measurement of protein stability

For the analysis of the half-life of Cnp3 and Cdc13, WT and *alm1Δ* cells were grown in YES at 25°C to midlog phase, and then 100 µg/ml CHX (01810; Sigma-Aldrich) was added to the cultures. Cells were harvested at the indicated time points, and whole cell extracts were prepared for immunoblotting using mouse anti-GFP mAb (11814460001; Roche) to detect Cnp3-GFP, rabbit anti-Cdc13 pAb (SP4, provided by S. Moreno), and mouse anti-PSTAIR mAb (P7962; Sigma-Aldrich) as a loading control. ImageJ software (version 1.5; National Institutes of Health) was used to quantify Cnp3-GFP and Cdc13 protein levels.

Immunoprecipitation of Cnp3 protein

mts4-ts cells expressing Cnp3-GFP were grown in YES to midlog phase at 25°C and then shifted to 36°C for 3 h. 45 min before harvesting, 5 mM *N*-ethylmaleimide (E3876; Sigma-Aldrich) was added to the culture media to inhibit deubiquitinating enzymes. Cells were harvested by centrifugation, and the pellets were washed with ice-cold STOP buffer and snap-frozen in liquid nitrogen. Native extracts were prepared in PBS with protease inhibitor cocktail (8340; Sigma-Aldrich) and 10 mM PMSF. 6 mg of total protein extracts was incubated with anti-mouse IgG-coated magnetic beads (Dynabeads; 11201D; Thermo Fisher) for 1 h at 4°C. Beads were washed three times with PBS with protease inhibitors, resuspended in 50 µl SDS-PAGE sample loading buffer (0.25 M Tris-HCl, pH 6.8, 8% [wt/vol] SDS, 0.004% [wt/vol] bromophenol blue, and 20% [vol/vol] 2-mercaptoethanol), and boiled for 5 min. Samples were clarified by centrifugation at 13,000 rpm for 10 min, and 20 µl was immunoblotted with mouse anti-GFP mAb (11814460001; Roche) to detect Cnp3-GFP and rabbit antiubiquitin pAb (sc-9133; Santa Cruz; provided by J.A. Sanchez Alcazar, CABD/UPO) to detect ubiquitinated proteins.

Ubiquitin pull-down

Polyubiquitination analysis was performed as previously described (Shiozaki and Russell, 1997). Cells expressing the endogenous *cnp3* tagged with GFP and overexpressing Cnp3-GFP from the medium-strength *nmt41* promoter in a single copy (pINT-Cnp3-GFP) were transformed with the pREP1-His₆-Ubi plasmid (originally provided by H. Seino, National Institute of Genetics, Mishima, Shizuoka, Japan).

Cells were grown in EMM at 25°C in the absence of thiamine for 22 h and then shifted to 36°C for 3 h. 45 min before harvesting, 5 mM *N*-ethylmaleimide (E3876; Sigma-Aldrich) was added to the culture media to inhibit deubiquitinating enzymes. Cells were harvested by centrifugation and washed once with ice-cold STOP buffer, and whole cell extract was prepared in buffer G (6 M guanidine HCl, 0.1 M sodium phosphate, and 50 mM Tris-HCl, pH 8.0). 6 mg of total protein extract (input) was incubated for 1 h with agarose–nitrilotriacetic acid-Ni²⁺ (P6611; Sigma-Aldrich) at room temperature. The beads were then washed three times with buffer U (8 M urea, 0.1 M sodium phosphate, and 50 mM Tris-HCl, pH 8.0) and once with buffer U with 10 mM imidazole. 50 µl SDS-PAGE sample loading buffer was added to the beads and incubated for 5 min at 100°C. 20 µl was immunoblotted with mouse anti-GFP mAb (11814460001; Roche), and 20 µl rabbit antiubiquitin pAb (sc-9133; Santa Cruz) inputs with the same amount of total protein was immunoblotted with anti-GFP mAb (Roche) to detect Cnp3-GFP and mouse antitubulin mAb (Tat1; provided by K. Gull) as a loading control.

Graphs and statistical analyses

Graphs and statistical analyses were performed with Prism 5.0 (GraphPad Software) and Excel (Microsoft). Unless otherwise stated, graphs represent the mean, and error bars represent SD. *n* is the total number of cells scored from at least three independent experiments. Statistical comparison between two groups was performed by unpaired Student's *t* test, considering two-tailed *p*-values exceeding 0.05 to be NS. Asterisks denote *P* < 0.05 (*), *P* < 0.001 (**), *P* < 0.001 (***), and *P* < 0.0001 (****). Data distribution was assumed to be normal, but this was not formally tested.

Online supplemental material

Fig. S1 shows the predicted CC structure of Alm1 (A), its localization and colocalization with Nup107 at the nuclear pore (B and C), and the transient colocalization of Alm1 and Cut12 (D). Fig. S2 shows mitotic chromosome segregation of WT and *alm1Δ* with the histone marker Hht2-GFP (A), Sid2-tomato and lacI-GFP bound to tandem array repeats of lacO at the *lys1* locus at chromosome I (B), and spindle dynamics of WT and *alm1Δ* during mitosis (C–E). Fig. S3 shows localization of Mad2-GFP (A) and Bub1GFP (B) during mitosis in WT and *alm1Δ*, Mad2-GFP localization during interphase (C), total protein levels by Western blot of Mad2 (D), Mad2-GFP localization in MBC metaphase blocked cells (E), and genetic interaction of *alm1Δ* with *mad2Δ* and *bub1Δ* (F). The figure also depicts localization (G) and kinase activity (H) of Ark1 and genetic interaction of *alm1Δ* and *ark1-t7* ts mutant (I and J). Fig. S4 shows four clusters of gene-deletion mutants that have a synthetic effect on TBZ sensitivity with *alm1Δ* (A), images of WT, *alm1Δ*, *mts4*, and *alm1Δ mts4* double mutant expressing Cnp3-GFP (B), images of WT, *alm1Δ*, and *cut8-563* ts mutant expressing Cnp3-Tomato (C), and analysis of Cdc13 degradation in the presence of CHX (D). Table S1 contains the list of strains used in this study.

Acknowledgments

We would like to thank Paul Nurse's lab (The Rockefeller University, New York, NY), our colleagues at the CABD, and members of R.R. Daga's lab and the Cell Cycle group, especially Victor A. Tallada, for helpful discussions, Victor M. Carranco for excellent technical help, and the CABD proteomic and microscopy facility technicians Katherina Garcia and Laura Tomas for their helpful advice.

This work was supported by the Ministerio de Economía y Competitividad of the Spanish government (grant BFU2015-70604-P to R.R. Daga). P. Gallardo was supported by a grant from Universidad Pablo de Olavide (grant Beca Puente PP1-1402).

The authors declare no competing financial interests.

Submitted: 29 December 2016

Revised: 28 May 2017

Accepted: 16 August 2017

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