Shugoshin forms a specialized chromatin domain at subtelomeres that regulates transcription and replication timing

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A chromosome is composed of structurally and functionally distinct domains. However, the molecular mechanisms underlying the formation of chromatin structure and the function of subtelomeres, the telomere-adjacent regions, remain obscure. Here we report the roles of the conserved centromeric protein Shugoshin 2 (Sgo2) in defining chromatin structure and functions of the subtelomeres in the fission yeast Schizosaccharomyces pombe. We show that Sgo2 localizes at the subtelomeres preferentially during G₂ phase and is essential for the formation of a highly condensed subtelomeric chromatin body ‘knob’. Furthermore, the absence of Sgo2 leads to the derepression of the subtelomeric genes and premature DNA replication at the subtelomeric late origins. Thus, the subtelomeric specialized chromatin domain organized by Sgo2 represses both transcription and replication to ensure proper gene expression and replication timing.
Eukaryotic chromosomes consist of various chromatin domains that contribute to genomic integrity and functions. The telomere is a specialized chromatin domain located at chromosome ends that contains repetitive nucleotide sequences called 'telomere repeats' and associated proteins. The telomere protects the chromosome ends and ensures proper chromosome movement during both mitosis and meiosis for genome stability. The subtelomere is the chromosomal region located adjacent to the telomere. Subtelomeric (ST) DNA contains multiple elements that share high sequence similarity among the subtelomeres of an individual organism.

In the budding yeast Saccharomyces cerevisiae, the subtelomeres are highly variable in length and commonly contain an X element and tandem copies of a Y element, which includes the open reading frame (ORF) of an RNA helicase gene. In humans, the subtelomeres comprise a complex mosaic of >40 duplicated segments with a variety of ORFs. Although the sequencing is not yet complete (http://www.pombase.org/status/sequencing-status), the subtelomeres of S. pombe also contain a mosaic region of multiple segments with highly similar sequences that spans ~50 kb of the telomere-proximal site—a site containing multiple ORFs of common genes including the RecQ-type DNA helicase genes (tlh1 and tlh2). In addition, the chromatin structure of the S. pombe subtelomere is composed of two distinct parts: heterochromatin and ST chromatin. The heterochromatin contains histone H3 with methylated Lys9 (H3K9me) and HP1 homologues, and spans ~45–80 kb of the telomere-proximal site. The adjacent telomere-distal ST chromatin spans ~50 kb and shows low sequence similarity among the subtelomeres, but shares a common feature of low levels of methylated and acetylated histones. In contrast to the substantial accumulation of knowledge on telomeres, not much attention has been paid to subtelomeres, and their regulation and physiological function remain largely unknown.

The centromere is a crucial domain for the faithful segregation of chromosomes during cell division. Shugoshin (Sgo) is a conserved centromeric protein that protects centromeric cohesion and promotes loading of the chromosomal passenger complex (CPC) onto centromeres. S. pombe possesses two Sgo paralogues: Sgo1 and Sgo2. Sgo1 is expressed specifically during meiosis I, recruits PP2A phosphatase to the centromeres and thereby counteracts the casein kinase 1-dependent cleavage of cohesion. In contrast, Sgo2 is ubiquitously expressed during the mitotic and meiotic cell cycle, and plays an important role in recruitment of CPC to the centromeres for proper chromosome segregation in M phase. Intriguingly, during interphase, the accumulation of Sgo2 at the centromeres decreases and, instead, a substantial fraction appears at the vicinity of the telomeres. However, details of the regulation and localization of Sgo2 during interphase, in addition to its physiological roles at the telomere vicinity, remain largely unknown.

In this study, we reveal new roles of Sgo2 at the subtelomeres in S. pombe. We show that Sgo2 associates with the subtelomeres preferentially during G2 phase and is essential for the formation of a highly condensed interphase-specific chromatin structure at the subtelomeres. Furthermore, Sgo2 regulates ST gene expression by reducing the occupancy of RNA polymerase II (RNAPII) and maintains normal timing of DNA replication by limiting loading of a replication initiation factor to the late origins within the subtelomeres. Thus, Sgo2 defines the chromatin structure and functions of subtelomeres during interphase in S. pombe.

Results

Sgo2 associates with the subtelomeres of chromosomes 1 and 2. To characterize the precise distribution of Sgo2 across the three chromosomes of S. pombe (Fig. 1a), we performed immunoprecipitation (ChIP)–chip analyses of Flag-tagged Sgo2 in logarithmically proliferating cells, which were primarily in interphase. Sgo2-Flag was enriched not only at the centromeres but also at the subtelomeres of both chromosomes 1 and 2 (subtel1L, 1R, 2L and 2R), extending >100 kb from each telomere (Fig. 1b,c). These Sgo2-enriched regions encompassed the clusters of non-essential genes and partially overlapped with the telomere-adjacent ST heterochromatin, which contains H3K9me and the heterochromatin protein Swi6/HP1 (refs 8,9).

It was not clear whether Sgo2 also localizes near the ends of chromosome 3, because the S. pombe genome database does not cover the regions external to the rDNA repeats that are located near both telomeres of chromosome 3 (tel3) (Fig. 1a). However, Sgo2 was not enriched at the rDNA repeats of chromosome 3 in the ChIP assay (Fig. 1d). Furthermore, Sgo2 was barely detectable by microscopy in the close vicinity of tel3, as indicated by the telomere-binding protein Tax1 (ref. 24) that is associated with the nucleolus protein Gar1 (ref. 25 and Fig. 1e). Thus, it is unlikely to be that Sgo2 accumulates near tel3. The absence of Sgo2 near tel3 is possibly due to the loss of the highly similar ST DNA sequences in chromosome 3 of the wild-type (wt) strain used in this study (Supplementary Fig. 1).

Sgo2 associates with the subtelomeres preferentially in G2. We next examined the cell cycle dependency of the Sgo2 association with the subtelomeres. Cells were arrested in M phase via the nda3-KM311 mutation and the localization of Sgo2 was analysed by ChIP. Sgo2 association with subtel2R was slightly weaker in nda3-KM311 cells even at the permissive temperature (32 °C), as compared with wt cells (Fig. 2a upper); this difference was enhanced when cells were incubated at the restrictive temperature (20 °C), which arrested the nda3 mutant in M phase, while the wt strain remained asynchronously growing (Fig. 2a lower). In contrast, the centromeric localization of Sgo2 was considerably increased in the nda3 mutant (Fig. 2a lower), which is consistent with previous findings that Sgo2 accumulates at centromeres during M phase. Furthermore, time-lapse microscopy revealed that Sgo2 dissociated from the vicinity of the telomeres immediately before chromosome segregation was initiated (Supplementary Fig. 2, 2–4 min.). These data suggest that Sgo2 associates with the subtelomeres preferentially during interphase.

To pinpoint the cell cycle stage during interphase at which Sgo2 accumulates at the subtelomeres, cells were synchronized by means of the cdc25-22 mutation, and the localization of Sgo2 at the centromeres and at the three different loci (20.0, 51.8 and 89.8 kb from tel2R) in subtel2R were analysed by ChIP. Enrichment of Sgo2 at the subtelomeres reached its maximum peak in the middle of G2 phase, which is between the peak of the septation index (S phase) and the peak of centromeric localization of Sgo2 (M phase) (Fig. 2b). These results indicate that the localization of Sgo2 is dynamically regulated during the cell cycle, and that Sgo2 associates with the subtelomeres preferentially in the middle of G2 phase.

H2A phosphorylation is important for the Sgo2 localization. Next, we investigated the requirements for localization of Sgo2 at the subtelomeres. First, we examined whether the telomeric structure is required for Sgo2 to associate with the subtelomeres. Deletion of the telomerase catalytic subunit Trt1 generally leads to cell death due to the loss of telomeric repeat DNA and de-protection of the chromosome ends. However, a fraction of trt1Δ cells survive if all of the chromosomes become self-circularized via chromosomal end fusion at the subtelomeres. A survivor that has lost ~9 kb of subtelomeres from the
chromosome ends is called Type A, whereas one that has lost \( \sim 15 \text{ kb} \) is called Type B\(^3\). We found that Sgo2 remained attached to the subtelomeres in both types of \( \text{trt}1 \Delta \) survivor lacking telomeric DNA (Fig. 3a), indicating that the telomeric structure is dispensable for the association of Sgo2 with the subtelomeres.

We also examined whether well-established chromatin-related factors are required for localization of Sgo2 at the subtelomeres,
including centromeric proteins that play a role in activation of the spindle assembly checkpoint such as Bub1 (ref. 22), Hrk1 (ref. 32) and Mph1 (ref. 33); the core telomere-binding protein Taz1 (ref. 24); and the heterochromatin factors Clr4 and Dcr1 (refs 8,9).

Among the respective deletion mutants, only the bub1\(^+\) deletion abolished the ST localization of Sgo2 almost completely (Fig. 3b and Supplementary Fig. 3). Bub1 is the sole protein kinase that phosphorylates serine 121 of histone H2A (H2A-S121) (Supplementary Fig. 4) and this histone modification is important for the association of Sgo2 with chromatin\(^{1,2}\). Substituting serine 121 of H2A with a non-phosphorylatable alanine (h2a-S121A) resulted in almost the same level of removal of Sgo2 from the subtelomeres as observed in bub1\(^+\) (Fig. 3c), although the amount of Sgo2 protein in bub1\(^+\), h2a-S121A and wt cells was comparable (Supplementary Fig. 5). Moreover, localization of Sgo2 at the subtelomeres was greatly decreased in cells carrying the bub1-KD (kinase-dead) mutation to almost the same level as observed in the h2a-S121A mutant (Supplementary Fig. 6). These data indicate that the phosphorylation of H2A-S121 by Bub1 is important for the association of Sgo2 with subtelomeres, whereas the telomeric structure, the activation of spindle assembly checkpoint and the heterochromatin structure are dispensable.

Sgo2 is crucial for the formation of highly condensed knobs. The primary function of Sgo2 is to recruit CPC to centromeres for the bipolar attachment of kinetochores to spindle microtubules during metaphase in S. pombe\(^{17,18}\). However, our ChIP analyses revealed that any of the CPC components (Ark1, Bir1, Pic1 and Nbl1) is hardly localized at the subtelomeres throughout the cell cycle, in stark contrast to their apparent accumulation at the centromeres during M phase (Supplementary Fig. 7). We therefore explored the idea that Sgo2 has a new function at the subtelomeres in interphase.

‘Knobs’ are highly condensed chromatin bodies in S. pombe that are intensely stained with the DNA-labelling dye 4’,6-diamidino-2-phenylindole (DAPI). They are located in the vicinity of the telomeres, more precisely adjacent to the ST heterochromatin (as indicated by the H3K9me3 signal adjacent to that of Taz1), specifically during interphase\(^{34}\) (see Fig. 4b wt).
Here we found that the subnuclear localization of Sgo2–green fluorescent protein (GFP) always overlapped with the knobs (Fig. 4a), and that the knobs were completely lost in sgo2Δ cells (Fig. 4b,c). Consistent with this, bub1Δ and h2a-S121A mutants, in which Sgo2 was dissociated from the subtelomeres (Fig. 3c), also completely lacked knobs (Fig. 4c). These data demonstrate that the association of Sgo2 with subtelomeres plays a crucial role in the knob formation.

It has been shown that the frequency of the knob formation is substantially decreased by the deletion of Set2 (ref. 34), which is the methyltransferase for H3K36 (refs 35,36 and Fig. 4c). ChIP analyses revealed that localization of Sgo2 at the subtelomeres also decreased markedly in the set2Δ mutant as compared with the wt strain (Fig. 4d). These data further support our conclusion that the knob formation is dependent on localization of Sgo2 at the subtelomeres.

Sgo2 represses transcription of the ST genes. As highly condensed chromatin is generally expected to be inhibitory to transcription37, we investigated the effects of the Sgo2-dependent knob formation on gene expression. Genome-wide expression profiles were obtained by microarray analyses using poly(A)-containing RNAs that were purified from asynchronously growing wt and sgo2Δ cells. We found that the transcript levels of genes at subtel1L, 1R, 2L and 2R were higher in sgo2Δ cells than in wt cells, whereas gene expression was not strongly affected, if at all, by sgo2Δ in other chromosomal regions (Fig. 5a). Notably, the locations of the genes affected by sgo2Δ correlated with the ST distribution of Sgo2, although the expression of some ST genes were barely affected (Fig. 5b and Supplementary Fig. 8), which

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**Figure 3 | H2A-S121 phosphorylation by Bub1 is important for Sgo2 association with the subtelomeres.** (a) ChIP analyses of Sgo2-Flag localization at subtel2R in trt1Δ survivor cells (Types A and B). Relative fold enrichment at subtel2R, normalized to the signal at the his1 locus, is shown. It is noteworthy that the most telomere-proximal data point, which corresponds to the region lost in Type B, is missing for Type B. (b) ChIP analyses of Sgo2-Flag localization in various deletion mutants. Relative fold enrichment at subtel2R and the centromeres (dg), normalized to the signal at the his1 locus, is shown. (c) ChIP analysis of Sgo2-Flag localization at subtel2R in the bub1Δ and h2a-S121ΔA mutants. The relative fold enrichment at subtel2R as normalized by the signals at the his1 locus is plotted. The error bars indicate the s.d. (n = 3).

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**Figure 4 | Sgo2 is required for the formation of knobs.** (a) Co-localization of Sgo2-GFP with knobs during interphase. Cells were grown in EMM at 26 °C and then fixed in formaldehyde (see Methods). 3DSIM images of single optical sections for DAPI and Sgo2-GFP (from immunostaining using anti-GFP antibody) signals in an interphase cell are shown. Arrowheads indicate knobs with intense DAPI signals. Scale bar, 1 μm. (b) Knobs are localized near Taz1-GFP (telomere) and H3K9me3 (subtelomeric heterochromatin) in wt cells, but are absent in sgo2Δ cells. 3DSIM images of single optical sections were taken as in a. Arrowheads indicate knobs. Scale bar, 1 μm. (c) Sgo2 association with subtelomeres is required for knob formation. Frequencies of knob formation in each strain during exponential growth are shown. (d) Sgo2 association with subtelomeres is substantially decreased in the absence of Set2. Sgo2-Flag localization at subtel2R in bub1Δ and set2Δ cells was analysed by ChIP. Relative fold enrichment at subtel2R, normalized to the signal at the his1 locus, is shown. Error bars indicate the s.d. (n = 3).
Figure 5 | Sgo2 represses transcription of the subtelomeric genes. (a) Microarray analysis of genome-wide changes in RNA expression caused by Sgo2 deletion. The signal ratio of each probe on the array is shown. Red dots and grey dots indicate significant ($P<0.002$ for an increase and $P>0.998$ for a decrease) and insignificant changes, respectively. The $P$-value was calculated as change $P$-value (two-sided) by Affymetrix GCOS1.4 with the default settings. (b) Magnified views of the subtelomeric regions in a. Localization of Sgo2 (from the ChIP–chip analyses) is indicated by the pale blue colour. (c) Association of Sgo2 with the subtelomeres is required for subtelomeric gene repression. The transcript levels of the subtelomeric genes and the centromeric repeats (dg) in wt, sgo2Δ, bub1Δ and h2a-S121A cells were analysed by quantitative reverse transcriptase–PCR (RT–PCR). The locations of the genes are shown in the box. Each value was normalized first to that of his1 and then to the wt value. (d) Occupancy of RNAPII at the subtelomeric gene loci is increased in sgo2Δ, bub1Δ and h2a-S121A cells. ChIP analyses of RNAPII at the subtelomeric gene loci and at the centromeric repeats (dg) are shown. The recovery of immunoprecipitated DNA relative to total input DNA was determined by quantitative PCR. (e) ChIP analyses of the levels of H3K4me2 (upper) and H3K36me3 (lower). The recovery of immunoprecipitated DNA relative to total input DNA was measured by quantitative PCR. Error bars indicate the s.d. ($n=3$).
may be due to the unequal ST localization of Sgo2 and/or the difference in the promoter activity of each gene. These data imply that the localization of Sgo2 is important for the observed repressive effect on ST gene expression.

Indeed, reverse transcriptase-PCR (RT-PCR) analyses showed that the transcription of several ST genes was also de-repressed in the bub1Δ, h2a-S121A and set2Δ mutants, as observed in the sgo2A mutant, whereas the expression level of the centromeric repeats (dg) barely changed in these mutants (Fig. 5c and Supplementary Fig. 9a). Furthermore, ChIP analyses of RNAPII showed that RNAPII occupancy at these ST gene loci was considerably higher in the sgo2A, bub1Δ, h2a-S121A and set2Δ mutants than in the wt strain, whereas RNAPII occupancy at the centromeric dg was barely altered (Fig. 5d and Supplementary Fig. 9b). These data indicate that Sgo2 mediates gene repression at the transcriptional level specifically at the subtelomeres. Consistently, the levels of histone modifications associated with transcriptionally active chromatin, such as dimethylation of lysine 4 and trimethylation of lysine 36 in histone H3 (H3K36me3, respectively) were elevated at the ST gene loci (Fig. 9b). These data indicate that Sgo2 mediates gene repression at the transcriptional level specifically at the subtelomeres. Consistently, the levels of histone modifications associated with transcriptionally active chromatin, such as dimethylation of lysine 4 and trimethylation of lysine 36 in histone H3 (H3K36me3, respectively), were elevated at the ST gene loci in sgo2A, bub1Δ, and h2a-S121A cells as compared with wt cells (Fig. 5e and see Discussion).

**Sgo2 forms a new type of repressive chromatin domain.** Swi6/HP1 is known to associate with chromatin in an H3K9me-dependent manner to form transcriptional repressive heterochromatin at the telomere-proximal region of each subtelomere. Therefore, we compared the effects of sgo2Δ and swi6Δ deletion on gene repression by monitoring the expression of an exogenous gene inserted at the subtelomere. As compared with the wt strain, expression of the ura4+ gene placed 7 kb from the telomere (7K:ura4+) increased remarkably in the swi6Δ mutant but not in the sgo2A mutant, indicating that the Swi6 heterochromatin is dominant to repress gene expression at the telomere-proximal region (Fig. 6a middle). In contrast, there was a remarkable increase in the expression of the ura4+ gene when it was placed 52 kb from the telomere (52K:ura4+) in sgo2A cells, whereas only a small increase was observed in swi6Δ cells (Fig. 6a right). bub1Δ and set2Δ cells showed derepression of 52K:ura4+ expression to levels similar to those in sgo2Δ cells (Fig. 6b), indicating that the ST localization of Sgo2 is required for this repressive effect. In contrast, expression of the endogenous ura4+ gene was unaffected in sgo2A cells (Fig. 6a, left), demonstrating that the Sgo2-mediated gene repression was region specific rather than gene specific. These observations were verified by monitoring cell growth on ura4+−selective media (Fig. 6c). Collectively, these results suggest that the association of Sgo2 with subtelomeres forms a new type of repressive chromatin domain that is distinct from the H3K9me-Swi6-mediated heterochromatin.

**Sgo2 limits loading of Sld3 to the replication origins.** Next, we explored which step of replication is regulated by Sgo2. The initiation of DNA replication requires the formation of a pre-replicative complex (pre-RC) at origins during G1 phase, followed by activation of the pre-RC on entry into S phase. We therefore examined the loading of two replication factors, Mcm6 and Sld3, in cells arrested immediately before the stage of replication initiation.

Mcm6, a pre-RC component, was enriched at AT2088 and at the ST late origins (tel-45.7 or tel-21.2) as efficiently as at ars2004 in both wt and sgo2A cells (Fig. 7d left). In contrast, Sld3, which is recruited to origins during the earliest step of pre-RC activation on S phase entry, was enriched at ars2004 but not at the late origins in wt cells, as previously reported (Fig. 7d right). In sgo2A cells, recruitment of Sld3 was accelerated at tel-45.7 and tel-21.2, but not at the internal late origin AT2088 (Fig. 7d right). These data suggest that Sgo2 represses the initiation step of replication by limiting Sld3 loading at the ST late origins, which is required for proper replication timing.

**Discussion**

In this study, we have discovered new regulations and functions of the subtelomere. We have shown that Sgo2, which accumulates at the centromeres and contributes to precise chromosome segregation during M phase, is recruited to the subtelomeres during interphase, especially G2 phase, and plays a role in the organization of the condensed chromatin body called a ‘knob’, in the repression of gene transcription and in the repression of replication, to ensure proper gene expression and replication timing at the subtelomeres (Fig. 8).

It is currently unknown how Sgo2 is targeted to the subtelomeres during interphase. Phosphorylation of H2A-S121 can be observed outside the subtelomeres in interphase (Supplementary Fig. 4), indicating that H2A-S121 phosphorylation alone is not sufficient, and that another factor is required to restrict Sgo2 access to the subtelomeres. One possibility is Tef1—a regulator of histone gene expression that localizes at the...
Set2 is important for Sgo2 localization and knob formation—may support the recruitment of Sgo2. As Sgo2 dynamically changes its location during the cell cycle, some cell cycle regulators may also be involved in Sgo2 recruitment to the subtelomeres.

Notably, knobs are not formed at the centromeres at any stage of the cell cycle, despite the high accumulation of Sgo2 at that location in M phase, suggesting that Sgo2 requires other factors to form knobs. The detailed mechanism underlying the subtelomere-specific formation of knobs remains unclear; however, one possibility is that ST chromatin contributes to this process. ST chromatin is characterized by its unique histone composition with low levels of acetylated and methylated histones, and the histone H2A variant H2A.Z. The low level of H3K9 methylation at the telomere-distal region of the subtelomeres (adjacent to the ST heterochromatin region) is in marked contrast to the high level of H3K9 methylation at the Sgo2-associated outer regions of the centromeres. In fact, knobs are observed preferentially at the telomere-distal regions of the subtelomeres.

Set2 is important for Sgo2 localization and knob formation; however, the level of Set2-dependent H3K36 methylation was found to be increased by Sgo2 deletion, which might seem to be contradictory. H3K36 methylation has been shown to associate with transcription elongation and is observed at actively transcribed regions.

Therefore, it is likely to be that Set2 recruited by the transcription at subtelomeres somehow promotes Sgo2 localization, thereby leading to knob formation. Either the knob or Sgo2 itself may in turn repress the transcription and/or accessibility of histone-modifying enzymes including Set2, thereby forming a negative feedback loop. Removal of Sgo2 from the subtelomeres would resolve the condensed knob structures, leading to the activation of gene expression and the recruitment of Set2, resulting in the observed increase in H3K36 methylation.

Does the knob structure itself regulate gene transcription and replication? The highly condensed nature of a knob may limit access of the transcription and replication machineries to chromatin, which would be consistent with the observation that the association of both RNAPII and the replication regulator Sld3 with the subtelomeres was increased in knob-deficient sgo2Δ cells. Another possibility is that Sgo2 itself recruits some regulatory factors to the subtelomeres. Recent studies in S. cerevisiae have revealed that the telomere-binding protein Rif1 regulates the replication timing of late origins by interacting with PP1 phosphatase to counteract the action of Dbf4-dependent kinase at the initiation step of replication. Interestingly, Sgo1 forms a complex with PP2A phosphatase to protect cohesin in S. pombe, S. cerevisiae and mammals, and human Sgo2 also interacts with PP2A for the protection of cohesion. Thus, S. pombe Sgo2 may also interact with a protein phosphatase to...
counteract with the activity of Dbf4-dependent kinase and thereby regulate Sld3 loading.

It has been predicted that early replication from the subtelomere region will lead to an increase in the length of telomere DNA\(^{33,34}\). We therefore checked whether the early replication caused by Sgo2 deletion leads to abnormal telomere elongation. Telomere Southern analysis showed that the telomere DNA lengths were only slightly shorter in elongation. Telomere Southern analysis showed that the telomere DNA\(^{53,54}\). We therefore checked whether the early subtelomere region will lead to an increase in the length of counteract with the activity of Dbf4-dependent kinase and NATURE COMMUNICATIONS | DOI: 10.1038/ncomms10393 | www.nature.com/naturecommunications

Figure 7 | Sgo2 regulates replication timing at the subtelomeres by limiting Sld3 loading. (a) Replication kinetics of the early and late origins in wt and sgo2Δ cells. The locations of the origins on chromosome 2 are shown schematically (top). \(ars2004\) is an early origin and \(AT2088\) is a late origin. \(tel-21.2\) is located at 21.2 kb from \(tel2R\). The non-origin region (non-ori) is located at 30 kb from \(ars2004\). Temperature-sensitive \(cdc25\) \(22\) mutant cells expressing thymidine kinase and the nucleotide transporter were arrested at the \(G_2/M\) boundary at the restrictive temperature (36°C) and then released from arrest by a temperature shift to 25°C in the presence of BrdU (200 \(\mu M\)). At the indicated time points, the amount of HL DNA and LL DNA, which were separated by caesium chloride (CsCl) density gradient centrifugation, was determined by qPCR (middle and bottom). (b) Replication of the subtelomeric late origins is affected in sgo2Δ cells. The strains used in (a) were released from \(G_2/M\) arrest and their DNA was labelled with BrdU for 90 min in the presence of hydroxyurea (HU: 10 mM). The BrdU-labelled DNA was purified by immunoprecipitation and quantified by quantitative PCR. Each value was normalized to that of wt cells. (Supplementary Fig. 12) Recruitment of Mcm6 to the replication origins. \(cdc25\) \(22\) \(nco4\)-108 mutant cells were released from \(G_2/M\) arrest and then incubated for 180 min at 20°C. ChIP analyses using anti-Mcm6 and anti-Flag (for Sld3-Flag) antibodies were performed. Each value was normalized to that of non-ori. Error bars indicate the s.d. (\(n = 3\)). Asterisks indicate a significant change as compared with the wt strain (\(P < 0.05\), two-tailed t-test).

What are the physiological roles of the ST gene repression caused by Sgo2? Matsuda \textit{et al.}\(^{34}\) reported that the frequency of knob formation substantially decreased when cells were arrested in \(G_1\) phase by nitrogen starvation. In this condition, Sgo2 might dissociate from the subtelomeres, inducing the expression of some ST genes in response to the nutrient stress. Another stress condition—low glucose stress (from 111 to 4.4 \(mM\))—has been shown to induce the expression of a set of hexose transporter genes called \(ght1^+\) and \(ght5^+\) (ref. 35), of which \(ght3^+\) and \(ght7^+\) are located at the subtelomeres. As shown in Fig. 5c, deletion of Sgo2 caused an increase in the expression of \(ght3^+\) and \(ght7^+\).

Figure 8 | Model of the multiple roles of Sgo2 during the cell cycle. During M phase, Sgo2 accumulates at the outer regions of the centromeres, where H3K9me is enriched and recruits CPC for precise chromosome segregation. In interphase, Sgo2 is enriched at the heterochromatin-adjacent region of the subtelomeres, where the levels of the histone modifications such as H3K9me, H3K4me and H3K36me are relatively low. Sgo2 is essential to form the knob structures and to repress both gene transcription and replication, to ensure proper gene expression and replication timing.
We speculate that Sgo2 might dissociate from the subtelomeres under some stress conditions including low glucose, to enable the efficient activation of ST gene expression for cell survival.

In summary, this study has revealed that Sgo defines chromatin functions not only at the centromeres but also at the subtelomeres in a cell cycle-dependent manner. Thus far, ST localization of Sgo in other organisms has not been reported; however, some cases in which Sgo relocates from the centromeres to the chromosome arms have been demonstrated in human cells and in *Xenopus* egg extracts. Thus, one can speculate that the translocation of Sgo to subtelomeres or other chromosomal regions might lead to the local organization of a repressive chromatin domain in other eukaryotes, as it does in *S. pombe*.

### Methods

**S. pombe strains.** All of the experiments were performed using the *S. pombe* strains listed in Supplementary Table 1. Growth media, basic genetics and construction of strains carrying deletion alleles or epitope-tagged proteins were described previously. The primers listed in Supplementary Table 2 were used to generate DNA fragments for homologous recombination to produce the deletion and carboxy-terminal epitope-tagged strains.

**Calculating the distance from a telomere.** We determined the distance from the telomeric repeat DNA on the basis of the following assumptions:

- The sequence between contig P15656 (ref. 7) and tel2R is the same as that in the *pNSU70* vector. This assumption provides the complete subtel2R sequence.
- Other unsequenced regions of the subtelomeres, such as the region between tel2L and contig c212, a gap between contigs c212 and c977, a region between contig c977 and tel2L, and a region between tel2L and contig c1348, have the same sequences as their subtel2R counterparts. This assumption is valid because of the high degree of sequence similarity among the subtelomeres. The sequence of the gap between contigs c1348 and p101D8 has been reported previously.

On the basis of these assumptions, we obtained the following equations for calculating the distances from telomeres:

\[
D_{1L} = 5,977,811 - X_1
\]

\[
D_{2L} = 29,714 + X_1 (1 \leq X_1 \leq 80,201) \text{ or } 47,495 + X_1 (80,302 \leq X_1)
\]

\[
D_{2R} = 4,542,638 - X_2
\]

where **X**1 or **X**2 is the chromosome position number on chromosome 1 or 2, respectively, and **D**1L, **D**2L, or **D**2R is the calculated distance between **X**1 and **X**2, between **X**1 and tel2L, between **X**2 and tel2L, or between **X**1 and tel2R, respectively.

**ChIP assays.** Exponentially growing cells were fixed at 32°C for 30 min in 3% paraformaldehyde (for Sgo2-Flag detection) or 1% formaldehyde (for histone or RNA polymerase II detection). The fixed cells were harvested by centrifugation and washed once with PBS, then with cold 2% paraformaldehyde (for Sgo2-Flag detection) or 1% formaldehyde (for histone or RNA polymerase II detection). The fixed cells were treated with 10 μg ml\(^{-1}\) proteinase K in 0.25% sodium lauryl sulfate (SDS, 37°C overnight), followed by the reverse cross-linking reaction (65°C for 6 h). Associated DNA fragments were purified by phenol chloroform extraction and ethanol precipitation, and then analysed by quantitative PCR (qPCR) using a StepOne real-time PCR system. The sequences of the primer sets used for real-time PCR are listed in Supplementary Table 3.

**ChIP-chip analyses.** DNA labelling, microarray slide hybridization (custom array GPL2777, Agilent Technologies), microarray visualization and data extraction were performed as described previously.

**Microscopy.** For the experiments in Fig. 1e and Supplementary Fig. 2, cells were exponentially grown in the EMM medium at 30°C and 13 optical section images were obtained at focus intervals of 0.3 μm using a DeltaVision microscopy system (Applied Precision) and deconvolved by a three-dimensional deconvolution method using SOFTWORK software.

**Knob observation.** Cells exponentially growing in EMM2-S5 at 26°C were pelleted by gentle centrifugation, chemically fixed by re-suspension in a buffer containing 4% formaldehyde (Polysciences, Inc.), 80 mM HEPES-K, 35 mM HEPES-Na, 2 mM EDTA, 0.5 mM EGTA, 0.5 mM spermidine, 0.2 mM spermine and 15 mM 2-mercaptoethanol pH 7.0, for 10 min at room temperature, and then washed three times in PEMS (100 mM PIPES, 1 mM EGTA, 1 mM MgSO\(_4\), 1.2 M sorbitol pH 6.9) with 2–5 min of rotation. The cell wall was digested by incubation with 0.6 mg ml\(^{-1}\) of zymolyase 100T (Seikagaku Biobusiness) in PEMS at 36°C for 5–30 min. The cells were then treated with 0.1% TritonX-100 in PEMS for 5 min and washed three times with PEMS.

For DAPI staining, the cells were incubated with 0.2 μg ml\(^{-1}\) of DAPI for 10 min and mounted with nPG-Glycerol (100% glycerol for absorptiometric analysis (Wako) with 4% n-propyl gallate, pH 7.0). For immunostaining, the cells were incubated with 1% BSA, 0.1% Na\(_2\)SO\(_4\), 100 mM i-lysine monohydrochloride in PEMS (PEMSBL) for 30 min; the solution was then replaced with antibody in PEMSBL and stained for 1 h. The cells were washed three times with PEMSBL and then secondary antibody was added. Afterwards, the cells were washed, counterstained with DAPI and suspended in mounting medium (nPG-Glycerol). DeltaVision/OMX microscope version 3 (GE Healthcare) was used for three-dimensional structured illumination microscopy (3DSIM) imaging with an objective lens, ×100 UPlanSApo NA1.40 Oil (Olympus). Conspicuously condensed, DAPI-stained bodies were counted as ‘knobs’ by visually inspecting each optical section of the 3D SIM images.

**Antibodies.** For ChIP, immunostaining and western blot analyses, we used the following antibodies: monoclonal anti-Flag M2 (Sigma, F7426; 1:20,000 dilution for ChIP in Figs 1–4 and Supplementary Fig. 3 and 7; 1:8,000 for western blotting), monoclonal anti-Flag M2 (Sigma, F1804; 1:500 for ChIP in Fig. 7 and Supplementary Fig. 6), anti-H2A2S12ph22 (1:500 for ChIP), anti-GFP (Roche, 1181446001; 1:500 for ChIP), anti-H2AS12ph22 (1:500 for ChIP), anti-GFP (Roche, 1181446001; 1:500 for ChIP), anti-RNAPII clone CTD4H8 (Millipore, 05-623; 1:500 for ChIP), anti-H3K9me3 (CMA318 (2F3); 1:500) and anti-GFP (MBL, 598; 1:500) antibodies were used in the observation of knobs.

**RNA quantification.** Exponentially growing cells in YES liquid medium were harvested by centrifugation and washed once with double distilled water. Total RNA was prepared by breaking cells in water-saturated phenol and RNA extraction buffer (20 mM Tris-HCl pH 7.5, 300 mM NaCl, 10 mM EDTA and 1% SDS) using a Multi-beads Shock (Yasui Kika). The total RNA was further purified by 0.2 U μl\(^{-1}\) recombinant DNase I (RNase-free) (Takara Bio) treatment (37°C) followed by phenol chloroform extraction and ethanol precipitation. Complementary DNAs were synthesized from total RNA using the High-Capacity cDNA Reverse Transcription Kit (Applied Biosystems) and then analysed by qPCR using a StepOne real-time PCR system. The sequences of the primer sets corresponding to each gene body used for real-time PCR are listed in Supplementary Table 3.

**Microarray analyses.** Poly(A)-containing RNA was purified from total RNA using a PolyATtract mRNA Isolation System (Promega). The biotin-labelled cRNAs were synthesized from 0.3 μg mRNA using a PolyATtract mRNA Isolation System (Promega). The cRNAs were then hybridized to a GeneChip Yeast Genome 2.0 Array (Affymetrix) in accordance with the manufacturer’s instructions. After staining, the GeneChips were scanned using GeneChip Scanner 3000 7G and GeneChip Operating Software (GCOS 1.4). The scaling and comparative analyses were performed by using GCOS 1.4 software with the default settings and *S. cerevisiae* probe set masking.
Replication timing analyses. \(\text{Chikashige, Y.}\) Chromosome DNA replication was analysed by BrdU incorporation assays as described previously\(^{38}\) with some modifications. Derivatives of \(cdc25-22\) Prmt1-TK PtdOH1-4ENT expressing the herpes simplex virus thymidine kinase gene and the human equilibrative nucleoside transporter gene, respectively, were incubated at 36 °C for 3 h for \(G2/M\) arrest and then released at 25 °C in the presence of 200 \(\mu\)M BrdU (Sigma). Cells collected at the indicated time were fixed with 0.1% sodium azide and total cellular DNA was prepared by the smash and grab method\(^{39}\). For replication kinetics analysis, the cellular DNA was digested with HaeIII, centrifuged in a CsCl density gradient to separate the replicated DNA (heavy–light (HL) density) from unreplicated DNA (light–light (LL) density) and the amount of DNA in the HL and LL densities was determined by qPCR using primers recognizing a real-time PCR system (Applied Biosystems). The primers used for qPCR are listed in Supplementary Table 3. Replication (%) was calculated as the ratio of the amount of heavy–light (HL) density to the sum of the amount of light–light (LL) density plus heavy–light (HL) density times 100%.

For ChIP analyses of Mcm6 and Sld3-Flag, \(\text{wt}\) and \(\text{nda4-108/mcm5 sld3-5flag}\) cross-linking. The immunoprecipitated DNA and the input DNA were analysed by qPCR. For ChIP analyses of Mcm6 and Sld3-Flag, \(\text{wt}\) and \(\text{nda4-108/mcm5 sld3-5flag}\) cross-linking. The immunoprecipitated DNA and the input DNA were analysed by qPCR. Anti-Mouse IgG (Thermo Fisher). The immunoprecipitated DNA was recovered into fragments of around 400 bp, denatured by boiling and used for PCR amplification with anti-FUS antibodies associated with Dynabeads M-280 anti-Mouse IgG (Thermo Fisher). The immunoprecipitated DNA was recovered and purified as described in the Methods for ChIP, except that there was no reverse transcription.

Pulsed-field gel electrophoresis analysis. Intact chromosomes in an agarose plug were electrophoresed using CHEF-DR III Pulsed Field Electrophoresis Systems (Bio-Rad) under the following conditions: 0.8% SeaKem Gold Agarose (Lonza) in 1 x TAE buffer, 14 °C. Block 1: initial switch time 1,200 s, final switch time 1,200 s, run time 24 h, voltage gradient 2 V cm\(^{-1}\) angle 96°; Block 2: initial switch time 1,500 s, final switch time 1,500 s, run time 24 h, voltage gradient 2 V cm\(^{-1}\) angle 100°; Block 3: initial switch time 1,800 s, final switch time 1,800 s, run time 24 h, voltage gradient 2 V cm\(^{-1}\) angle 100°. Chromosomal DNA separated in the gel was transferred to a Hybond-N\(^{\text{\textregistered}}\) (GE Healthcare) membrane according to the manufacturer’s instructions. To generate the probes, the TASS fragments (TAS1, TAS2 and TAS3)\(^{29}\) were excised from pNSU70 (ref. 6) and an rDNA fragment was amplified by PCR from \(S.\) \(\text{pombe}\) expressing the herpes simplex virus thymidine kinase gene and the human equilibrative nucleoside transporter gene, respectively, were incubated at 36 °C for 3 h for \(G2/M\) arrest and then released at 25 °C in the presence of 200 \(\mu\)M BrdU (Sigma). Cells collected at the indicated time were fixed with 0.1% sodium azide and total cellular DNA was prepared by the smash and grab method\(^{39}\). For replication kinetics analysis, the cellular DNA was digested with HaeIII, centrifuged in a CsCl density gradient to separate the replicated DNA (heavy–light (HL) density) from unreplicated DNA (light–light (LL) density) and the amount of DNA in the HL and LL densities was determined by qPCR using primers recognizing a real-time PCR system (Applied Biosystems). The primers used for qPCR are listed in Supplementary Table 3. Replication (%) was calculated as the ratio of the amount of heavy–light (HL) density to the sum of the amount of light–light (LL) density plus heavy–light (HL) density times 100%.

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- \(5’\) CGCTGACATTTGTTAGCGAGGAC3’
- \(3’\) CTAACATGCTCCTAAATTCATC3’

In Southern blot analysis, these DNA fragments were labelled with digoxigenin using the DIG High Prime DNA Labelling and Detection Starter Kit II (Roche).

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