Roles of Fission Yeast Grc3 Protein in Ribosomal RNA Processing and Heterochromatic Gene Silencing*§

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Grc3 is an evolutionarily conserved protein. Genome-wide budding yeast studies suggest that Grc3 is involved in rRNA processing. In the fission yeast Schizosaccharomyces pombe, Grc3 was identified as a factor exhibiting distinct nuclear dot localization, yet its exact physiological function remains unknown. Here, we show that S. pombe Grc3 is required for both rRNA processing and heterochromatic gene silencing. Cytological analysis revealed that Grc3 nuclear dots correspond to heterochromatic regions and that some Grc3 is also present in the nucleolar peripheral region. Depleting the heterochromatic proteins Swi6 or Clr4 abolished heterochromatic localization of Grc3 and resulted in its preferential accumulation in the perinucleolar region, suggesting its dynamic association with these nuclear compartments. Cells expressing mutant grc3 showed defects in 25 S rRNA maturation and in heterochromatic gene silencing. Protein analysis of Grc3-containing complexes led to the identification of Las1 and components of the IPI complex (Rix1, Ipi1, and Crb3). All of these Grc3-interacting proteins showed a dynamic nuclear localization similar to that observed for Grc3, and those conditional mutants showed defects in both rRNA processing and silencing of centromeric transcripts. Our data suggest that Grc3 functions cooperatively with Las1 and the IPI complex in both ribosome biogenesis and heterochromatin assembly.

Although the eukaryotic nucleus appears relatively unstructured compared with the cytoplasm, it contains a variety of functional compartments (1). The nucleolus, the most striking example of a functional nuclear compartment, is important in ribosome biogenesis (2). High level organization related to nuclear function also exists in the extranucleolar nucleoplasm. For example, heterochromatin, a densely packed nuclear subdomain, is a type of nuclear compartment with essential roles in nuclear organization and chromosome structure (3). Although these two nuclear compartments are distinct, there is increasing evidence that they use similar molecular mechanisms for epigenetic gene regulation (2, 4).

In the fission yeast Schizosaccharomyces pombe, heterochromatin domains are found at telomeres, the silent mating type locus, and on pericentromeric repeats. Heterochromatin establishment correlates with changes in post-translational histone tail modification. Clr4, a mammalian SUV39H homologue, provides histone H3 lysine 9 methylation (H3K9me)3 on heterochromatin, which then serves as a docking site for the HP1 proteins Swi6 and Chp2 (5–8). In most eukaryotes, HP1/Swi6 and H3K9me play a critical role in maintaining higher order chromatin structure and ensuring faithful chromosome segregation (9).

Although heterochromatin has long been thought to be transcriptionally silent, recent studies have shown that it is transcriptionally active and that the processing of transcribed RNAs is tightly coupled with heterochromatin formation. For example, heterochromatin assembly at fission yeast centromeres requires components of the RNAi pathway (10–12). Centromeric dg and dh repeats are transcribed by RNA polymerase II, and the processing of nascent transcripts through the RNAi pathway is coupled with the H3K9me introduction and heterochromatin spreading (13). A second RNA processing pathway, involving Cid14 poly(A) polymerase and the exosome, is required for efficient heterochromatric gene silencing (14, 15). Therefore, it is likely that heterochromatic transcripts are targeted and degraded by the exosome and/or RNAi machinery, with the help of other RNA processing machineries.

Transcribed RNA processing also plays a central role in nucleolar function. In all organisms, the synthesis of ribosomal RNAs (5, 5.8, 18, and 25–28 S rRNA) is not achieved by simply transcribing the individual rRNA species but requires a complex series of post-transcriptional processing steps. Three of the four rRNAs (18, 5.8, and 25 S) are transcribed by RNA polymerase I as a single large precursor, 35 S pre-rRNA, that contains the sequences for mature 18, 5.8, and 25 S rRNAs, along with two external transcribed spacers and two internal transcribed spacers (ITS1 and ITS2). During the maturation process, the pre-rRNA has to undergo a number of modifications and is subjected to cleavages and trimming events. Studies on Saccharomyces cerevisiae have revealed that a remarkable number of trans-acting factors, including small nucleolar...
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RNAs, small nucleolar RNP proteins, putative RNA helicases, endo- and exoribonucleases, and putative ATP/GTPases, are involved in rRNA maturation and assembly into ribosomal sub-units (16, 17). Although recent genome-wide approaches have uncovered a number of additional factors that are potentially involved in rRNA processing (18, 19), their detailed roles in rRNA and ribosome processing remain unresolved.

Grc3 is a widely conserved eukaryotic protein that possesses a conserved ATP/GTPase domain and shows extensive sequence similarity to Cip1, a component of the mRNA cleavage and polyadenylation machinery (20, 21). In S. cerevisiae, Grc3 is essential for growth (22), and genome-wide studies have shown that it is involved in rRNA processing, especially in the ITS2 processing (18). In S. pombe, however, the Grc3 homologue encoded by SPC830.03 was identified in genome-wide localization screenings as a factor displaying “nuclear dot” localization (23–25). Although heterochromatin shares this characteristic nuclear dot localization pattern in fission yeast (6, 26), functional involvement of Grc3 with heterochromatin or rRNA processing remains unknown.

In this study, we show that fission yeast Grc3 dynamically associates with both heterochromatic and perinucleolar regions and is required for both rRNA processing and heterochromatic gene silencing. We further demonstrated that Grc3 associates with both heterochromatic and perinucleolar regions and is required for both rRNA processing and heterochromatin gene silencing. We further demonstrated that Grc3 is functionally linked with the IPI complex and Las1. The protein Grc3 is essential for growth (22), and genome-wide studies have uncovered a number of additional factors that are potentially involved in rRNA processing (18, 27).

EXPERIMENTAL PROCEDURES

Strains and Plasmids—The S. pombe strains used in this study are listed in supplemental Table S1. Tagging to produce GFP and FLAG fusion proteins of endogenous grc3+, crb3+, las1+, rix1+, and ipi1+ was performed using a PCR-based gene targeting protocol (27). Endogenous clr4+ and swi6+ were deleted by a PCR-based method or by standard genetic crosses. To express wild-type or mutant Grc3 from a multicopy plasmid, the wild-type protein was expressed from a multicopy plasmid, and mutation was introduced by in vitro mutagenesis method as described previously (6).

Microscopy Analysis—Immunofluorescence experiments were performed as described previously (6). Wild-type and mutant S. pombe cells expressing each of the GFP fusion proteins were cultured on minimal medium. Single colonies were picked and patched onto new plates. The cells were grown to early log phase in liquid medium and washed twice with deionized H2O, and the DNA was visualized by incubation with 1 µg/ml Hoechst 33342. Microscopic images were obtained on a Zeiss Axioplan2 imaging microscope and an ORCA-ER camera (Hamamatsu) or an Olympus IX71-based three-dimensional microscope system and a CoolSNAP HQ (Photometrics). Three-dimensional optical section images were taken at six focal planes with 0.5-µm focus intervals using MetaMorph (Universal Imaging), and the acquired images were deconvoluted and analyzed by softWorX (Applied Precision).

ChIP—ChIP was performed as described previously (6). Anti-FLAG M2 affinity gel (Sigma) was used to immunoprecipitate FLAG-tagged proteins. The anti-Swi6 polyclonal antibody was described previously (6). PCR products were separated and analyzed by electrophoresis on a 15% polyacrylamide gel. Primers used in this assay are listed in supplemental Table S2.

Protein Affinity Purification—Grc3-TAP, Las1-TAP, and Crb3-TAP purifications were done as described previously (28), with some modifications. The cells were grown in YEA medium (0.5% yeast extract, 3% glucose, 75 µg/ml adenine) to an A595 of 2–3. The cells were spun at 4 °C and washed once with STOP buffer (150 mM NaCl, 50 mM Nait, 10 mM EDTA, 1 mM NaN3, pH 8.0). The cell pellet (1 × 1010 cells in a 50-ml Falcon tube) was resuspended in 0.5 ml of Nonidet P-40 lysis buffer (50 mM HEPES/NaOH, pH 7.9, 150 mM NaCl, 1 mM EDTA, 1% Nonidet P-40, 0.1 mM Na3VO4, 1 mM DTT, 1 mM PMSF), supplemented with protease inhibitor mixture (Complete; Roche Applied Science), and lysed by vortexing with zirconia beads (for 2 min five times at 4 °C, with 2-min intervals on ice). The crude cell lysate was diluted with 10 ml of Nonidet P-40 lysis buffer and incubated with gentle agitation on a disc rotator for 15 min at 4 °C. The extract was spun at 12,000 × g for 10 min at 4 °C. Lysates were then transferred to a 50-ml Falcon tube and incubated with 400 µl of prewashed IgG-Sepharose 6 Fast Flow (GE Healthcare) for 2–3 h at 4 °C. The beads and immobilized proteins were harvested by centrifugation at 400 g, loaded on a Bio-Rad Poly-Prep column, and washed three times with 10 ml of 1% Nonidet P-40 lysis buffer. Bound protein was eluted with 1 ml of tobacco etch virus protease cleavage buffer containing 50–100 units of AcTEV™ Protease (Invitrogen) at 4 °C overnight. The elute was then incubated with 200 µl of calmodulin affinity resin (Stratagene), and the bound protein was eluted as described previously (28). Eluted proteins were precipitated by adding 2 volumes of ethanol, resolved by SDS-PAGE, and subjected to LC-MS/MS analysis.

Mass Spectrometry—Affinity-purified proteins were resolved by 4–20% SDS-PAGE (Daichi). After silver staining, the peptide bands were excised from the gel and subjected to in-gel reduction with 10 mM DTT, alkylation with 55 mM iodoacetamide, and digestion with 10 µg/ml modified Trypsin (Promega) at 37 °C for 16 h. After in-gel digestion, the collected phosphorylated peptides were subjected to mass spectrometry analysis as described previously (29).

Isolation of Temperature-sensitive Mutants—The grc3+ genomic region (from 80 bp upstream of ATG to 650 bp downstream of the stop codon) was first PCR-amplified using KOD polymerase (Toyobo) and cloned into the pREP1 plasmid, and mutation was introduced by KOD mutagenesis method as described previously (6).

PCR products were separated and analyzed by electrophoresis on a 15% polyacrylamide gel. Primers used in this assay are listed in supplemental Table S2. PCR products were separated and analyzed by electrophoresis on a 15% polyacrylamide gel. Primers used in this assay are listed in supplemental Table S2. PCR products were separated and analyzed by electrophoresis on a 15% polyacrylamide gel. Primers used in this assay are listed in supplemental Table S2.
wild-type allele with the mutant allele. Transformed cells were first isolated on minimal medium lacking uracil (for grc3\textsuperscript{ts}) or on YEA medium (0.5% yeast extract, 3% glucose, 75 μg/ml adenine) containing hygromycin (for crb3\textsuperscript{ts}, ipi1\textsuperscript{ts}, las1\textsuperscript{ts}, or rix1\textsuperscript{ts}) at 25 °C, and roughly 2,000 colonies were picked up and tested for temperature-sensitive growth on a nonselective plate at 36 °C. Mutant clones showing temperature-sensitive growth were isolated and used for further analysis. The position of altered amino acids in the isolated temperature-sensitive mutants are listed in supplemental Table S3.

**RNA Extraction and Northern Blot Analysis**—Total RNA was extracted from cells as described previously (6). Purified total RNA samples were separated by electrophoresis on a 1.25% agarose gel containing 6.7% formaldehyde, blotted on nylon membranes, cross-linked, and hybridized with each oligonucleotide probe. The oligonucleotides used are listed in supplemental Table S2. In time course experiments, wild-type and temperature-sensitive mutant \textit{S. pombe} cells were first cultured at the permissive temperature (25 °C) and then shifted to the restrictive temperature (36 °C). Total RNA was prepared from cell samples obtained at each time point after temperature shifting and subjected to Northern analysis.

**Multicopy Suppressor Screening**—The \textit{S. pombe} genomic library (pTN-L1; National Bio Resource Project) was introduced into the grc3 mutant cells (SPM1617, SPM1620, or SPM1623) by electroporation. The transformed cells were incubated on a selected medium (AA-Leu) at 25 °C for 1 day and then at 34 °C for 2–10 days. Approximately 72,000 clones were screened, and 48 clones that grew at 34 °C were isolated. The plasmids were rescued in \textit{Escherichia coli} and reintroduced...
into grc3 cells to confirm their ability to suppress the grc3 mutation. The nucleotide sequences of the inserts were determined by sequencing analysis using the M13-Fw and M13-Rv primers (supplemental Table S2).

**Quantitative RT-PCR Analysis**—To quantify the transcribed RNAs, the total RNA was extracted from yeast cells as described above and then preincubated with RNase-free DNase I (0.4 unit/mg RNA; TaKaRa) to digest the genomic
These results suggested that the budding yeast Grc3 is involved in rRNA processing (18). A homologous protein of Grc3 in S. pombe encoded by SPCC830.03 (hereafter referred to as SpGrc3 or Grc3) (Fig. 1A) was identified as a factor displaying nuclear dot localization (23–25), a pattern reminiscent of heterochromatin proteins (6). To gain insight into the function of S. pombe Grc3, we first examined its subcellular localization by expressing C-terminally GFP-fused Grc3 (Grc3-GFP) from its native promoter. As reported in previous studies (23–25), Grc3-GFP preferentially localized to the nucleus and formed two to three discrete spots in interphase cells (Fig. 1B).

In fission yeast, three heterochromatic regions—centromeres, the mating type region, and telomeres—are visualized by immunostaining for Swi6 or GFP-fused Swi6 (6, 26). Two to five discrete Swi6 foci in interphase cells represent one large cluster entirely composed of centromeres, several clusters of telomeres, and the mating type region (26). To determine whether the dot-like localization of Grc3-GFP corresponds to heterochromatic regions, we constructed strains expressing both Grc3-GFP and N-terminally Kusabira-orange-fused Swi6 (KOR-Swi6) and compared their localization in each cell. Using a fluorescence microscope equipped with a three-dimensional deconvolution system, we confirmed that the Grc3-GFP dots clearly co-localized with the signals for KOR-Swi6 (Fig. 1C). These results suggested that S. pombe Grc3 localizes to the heterochromatin regions.

Heterochromatic Localization of Grc3 Is Dependent on Swi6 and Clr4—Heterochromatin assembly in fission yeast involves H3K9me, mediated by the conserved methyltransferase Clr4(SUV39H). H3K9me creates binding sites for Swi6(HIP1) (7). To analyze the relationship between Grc3 and heterochromatin, we examined the Grc3-GFP localization in swi6-deleted (swi6Δ) or clr4-deleted (clr4Δ) cells. Strikingly, the Grc3-GFP dot-like localization was completely abolished in these mutant cells (Fig. 2A). Furthermore, the delocalized Grc3-GFP in swi6Δ or clr4Δ cells preferentially accumulated in perinuclear regions (Fig. 2, A and B). Careful examination of microscopic images of wild-type cells revealed that, in addition to the dot-like localization, some of the Grc3-GFP was also present in the perinuclear region (Fig. 2C, indicated by white arrowheads). These results confirmed that Grc3 localizes to two nuclear domains, the heterochromatin and perinuclear regions, and further indicated that the Grc3 heterochromatin localization requires an intact heterochromatin structure.

The involvement of Grc3 with rRNA processing was suggested in budding yeast. However, almost nothing is known about the functional involvement of Grc3 with heterochromatin. To examine the association of Grc3 with heterochromatin directly, we performed a ChIP assay. We constructed strains expressing C-terminally 5FLAG-tagged Grc3 (Grc3-FLAG), and its association with the three heterochromatogenic regions, centromeres (dg223), the mating type region (K-R), and telomeres (E12), was determined using the act1 locus as a negative control (Fig. 2D) (6). Consistent with our cytological analyses, Grc3-FLAG associated with the three heterochromatogenic regions, and importantly, this association was eliminated in the Δswi6 or Δclr4 mutant cells (Fig. 2E and data not shown). Together with the cytological analyses, these results indicated that SpGrc3 associates with two distinct nuclear domains, the heterochromatin and perinucleolar regions, and that its heterochromatin association is dependent on Swi6 and Clr4.

Grc3 Is Required for Pre-rRNA Processing—In S. cerevisiae, Grc3 is essential for growth and is involved in ribosomal RNA processing (18, 22). Although a genome-wide deletion analysis revealed that fission yeast Grc3 is essential for cell viability (30), its physiological role and functional involvement in rRNA processing or heterochromatin assembly have not been explored. We replaced an allele for grc3+ in diploid cells with a hygromycin resistance gene (hyg) and confirmed that haploid cells harboring the hyg+ gene were not viable (Fig. 3A).

To examine the cellular function of Grc3, we replaced the original grc3+ allele with a randomly mutagenized gcr3 allele and isolated 12 temperature-sensitive (ts) gcr3 alleles (Fig. 3B and data not shown) (see also “Experimental Procedures” and supplemental Table S3). Using these gcr3-ts mutant strains, we tested whether Grc3 is required for rRNA processing. Wild type and gcr3-7, one of the gcr3-ts mutant strains, were first cultured at the permissive temperature (25 °C) and then shifted to the nonpermissive temperature (36 °C). The total RNAs were extracted from the cells at different time points after the temperature shift and were subjected to Northern hybridization using specific oligonucleotide probes (Fig. 3C). Although no obvious growth defect was observed in the gcr3-7 mutant cells at 25 °C (Fig. 3B, section 7), high levels of rRNA precursors (35 and 27 S) were detected in this mutant even at the permissive temperature (Fig. 3D, gcr3-7, 0 h). After the temperature shift, the high levels of these rRNA precursors became much more pronounced, whereas the mature 25 and 5.8 S rRNAs decreased (Fig. 3D, gcr3-7, 1–6 h). Notably, the level of mature 18 S rRNA was not affected in the gcr3-7 mutant cells, suggesting that the
ITS1 processing was not affected in this mutant. Similar rRNA processing defects were observed in other grc3 mutant cells (data not shown). Together, these results indicated that S. pombe Grc3 is required for pre-rRNA processing, especially ITS2 processing, which is quite consistent with the results for S. cerevisiae Grc3 (18).

Grc3 Regulates Heterochromatin Silencing—Our cytological analyses revealed that the majority of Grc3 localized to hetero-

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**FIGURE 3.** Grc3 is required for proper ribosomal RNA processing. A, Grc3 is essential for cell growth. One grc3<sup>−</sup> allele of a diploid cell was replaced with a hygromycin resistance gene (hyg<sup>r</sup>). After sporulation, each spore was grown on nonselective (N/S) medium, and viable colonies were replicated on YEA medium containing hygromycin (+Hyg). All of the viable cells did not harbor the grc3::hyg<sup>r</sup> allele. A diagram of the tetrad analysis is shown at right. B, temperature-sensitive growth of isolated grc3 mutants. Parental wild-type (SPM1467) and grc3 mutant (grc3-3, grc3-6, grc3-7, grc3-8, or grc3-10) cells were grown on YEA plates at 25 or 36 °C for 3–5 days. C, the ribosomal RNA processing pathways and location of oligonucleotide probes used for Northern hybridization. Section 1, rRNA_18S; section 2, NB629_2; section 3, NB1102_2; section 4, rRNA_5.8S; section 5, rRNA_25S. The processing pathway determined from S. cerevisiae studies is presented; major processing events are thought to be conserved in S. pombe. The transcribed spacer regions are indicated as narrow lines, and mature rRNA is indicated as black rectangles. D, Northern hybridization analyses of rRNAs. Wild-type and grc3-3 cells were cultured at a permissive temperature (25 °C) in YEA medium, and then the culture was shifted to a nonpermissive temperature (36 °C). RNA was isolated from cultures at different time points and subjected to Northern hybridization analysis. rRNAs prepared from exponentially growing wild-type and clr4Δ cells were also analyzed.
chromatic regions, whereas only a minor fraction was associated with perinucleolar regions (Figs. 1 and 2). To examine whether Grc3 is functionally involved in heterochromatic gene silencing or heterochromatin assembly, we first tested the effect of the grc3-ts allele to silence a marker gene inserted in the heterochromatin regions (Fig. 4A). However, the impaired growth of the grc3-ts mutant cells hampered the evaluation by spotting assay (data not shown). Therefore, we assessed the function of Grc3 in heterochromatic silencing by monitoring pericentromeric dh transcripts (Fig. 4B).

In wild-type cells, pericentromeric transcripts are quickly processed by RNAi and/or RNA degradation pathways (13), and low levels of dh transcripts were detected by RT-PCR analysis (31). In grc3 mutant cells, pericentromeric transcript levels were clearly higher than that of wild-type cells even at the permissive temperature (25 °C) (Fig. 4B, left columns). Notably, the transcript levels were elevated at the restrictive temperature (36 °C) (Fig. 4B, right columns). These data suggest that Grc3 is involved in heterochromatin assembly in fission yeast.

Grc3 possesses a conserved ATP/GTPase motif and shows amino acid sequence similarity to Clp1, which is a component of the mRNA cleavage and polyadenylation machinery (Fig. 1A) (20, 21). Human Clp1 has been identified as an RNA kinase that phosphorylates the 5′-end of synthetic siRNAs and tRNA 3′-exons (32). A recent study showed that S. cerevisiae Grc3 has a similar kinase activity (33). Although we have not so far succeeded in detecting a similar kinase activity for S. pombe Grc3 (data not shown), it was conceivable that the ATP/GTPase domain plays a critical role in SpGrc3 function. We introduced an Ala substitution in the ATP/GTPase domain of Grc3 (K344A; Fig. 1A), constructed plasmids to express wild type or mutant Grc3K344A under the nmt1 promoter, and examined its effect on heterochromatin silencing (Fig. 4C).

Wild-type Grc3 expression had only a minor effect on silencing at centromere 1 (otr1R::ura4+) or the mating type region (Kint2::ura4+). However, mutant Grc3K344A expression impaired heterochromatin silencing, which was apparent from its robust growth on a medium lacking uracil and its increased sensitivity to 5-fluoroorotic acid (5FOA) (Fig. 4C). Mutant Grc3K344A may have a dominant-negative role for proper Grc3 function in heterochromatin silencing. Together, these results demonstrated that Grc3 regulates heterochromatic silencing in fission yeast and that its ATP/GTPase domain is critical to its function.

Grc3 Interacts with Las1 and the IPI Complex—To gain further insight into the physiological function of S. pombe Grc3, we purified and analyzed GrC3-interacting proteins. We constructed a strain in which a functional C-terminally TAP-tagged Grc3 protein was expressed from its endogenous promoter and purified this protein using a tandem affinity purification technique (28). Polycrylamide gel electrophoresis and LC-MS/MS analyses revealed that Grc3 interacted with three proteins: Crb3, a WD-repeat protein encoded by SPAC13G7.08c; and two proteins homologous to S. cerevisiae, Las1 and Rix1 (encoded by SPBC16C6.12c and SPCC4G3.18, respectively) (Fig. 5, A and C). Although the functional relationships with the S. cerevisiae counterparts need to be clarified, we hereafter refer to these proteins as SpLas1 and SpRix1, respectively.

In fission yeast, Crb3 has been shown to interact with Cut5 (34) and to assist in maintaining DNA damage checkpoints (35). On the other hand, its S. cerevisiae counterpart, Ipi3, joins with Ipi1 and Rix1/Ipi2 to form the protein complex IPI (involved in processing of ITS2), and all three components are required for proper ITS2 processing in pre-rRNA maturation (18, 19, 36). S. cerevisiae Las1 was originally characterized as an essential protein for cell viability and cell proliferation (37), and a recent report showed that the human nucleolar protein Las1 (Las1L) is essential for cell proliferation and ribosome synthesis (38). Based on sequence similarities, the fission yeast homologues are hypothesized to have a role in rRNA processing, but their physiological function is unknown.

To confirm the physical interaction among these proteins, we constructed strains expressing functional C-terminally
TAP-tagged Crb3 or Las1, performed tandem affinity purification, and analyzed Crb3- and Las1-interacting proteins by LC-MS/MS (Fig. 5, B and C). The results from these independent TAP purifications confirmed the interactions among Grc3, Rix1, Crb3, and Las1. Although we failed to detect a fission yeast homologue of Ipi1 (encoded by SPCC1393.06c) in the initial Grc3-TAP purification (Fig. 5 A), it was detected in the Crb3-TAP purification (Fig. 5 C), thus suggesting that Ipi1, Rix1, and Crb3 form a similar IPI complex in fission yeast and that this IPI complex is functionally linked with Grc3 and Las1.

To rule out the possibility that these proteins interact indirectly via cellular RNAs, we constructed strains expressing Grc3-TAP in conjunction with Crb3-FLAG or Las1-FLAG and conducted pull-down experiments with or without RNase treatment (Fig. 5, D and E, and supplemental Fig. S1). In agreement with the TAP purification, Crb3-FLAG and Las1-FLAG were efficiently co-immunoprecipitated with Grc3-TAP, and these interactions were resistant to RNase treatment (Fig. 5, D and E). We therefore conclude that Grc3 stably interacts with both IPI complex and Las1, in an RNA-independent manner.
Analysis of Grc3 Function in S. pombe

A

Rix1-GFP  Hoechst  Merge

WT

clr4Δ

swi6Δ

B

Crb3-GFP  Hoechst  Merge

B

Crb3-GFP  Hoechst  Merge

C

Ipi1-GFP  Hoechst  Merge

WT

clr4Δ

swi6Δ

D

Las1-GFP  Hoechst  Merge

WT

clr4Δ

swi6Δ

E

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Legend:
- Nuclear dots
- Nuclear dots + perinucleolar
- Nucleoplasm + perinucleolar
- Diffused in nucleolus

Bar scale: 10 μm
Identification of las1 as a Multicopy Suppressor of grc3 Mutation—To search for factors that genetically interact with grc3, we screened for multicopy suppressors of the grc3 mutation. The genomic library of S. pombe was introduced into the grc3-3, grc3-7, or grc3-8 mutant, and viable cells were selected at a nonpermissive temperature (34°C). Among the 1–2 × 10⁵ clones screened, we obtained a total of 48 clones as candidate genes to suppress temperature sensitivity in grc3 mutations. Nucleotide sequencing of the harbored plasmids revealed that 21 clones turned out to contain grc3 genomic DNA (data not shown). Whereas only grc3-containing plasmid was identified for screening using grc3-7 or grc3-8, a number of additional genomic fragments were isolated as suppressors for grc3-3 (Fig. 6A). Notably, the most abundant genomic clones were genomic DNA containing las1. Although several other genomic fragments were isolated as suppressors of the grc3-3 temperature-sensitive phenotype (Fig. 6A), their suppressive effect was much weaker than that of the plasmid containing las1 (Fig. 6B, compare section 2 with other plasmids). It is thus likely that the harbored gene(s) indirectly affected the temperature sensitivity of the grc3-3 mutant cells.

To confirm the suppressive effect of las1, we introduced a las1-containing plasmid (pREP1-las1) into the grc3-3 cells. The temperature sensitivity was indeed relieved at 32°C, although the temperature-sensitive growth was not clearly reversed at 34°C (Fig. 6C). These data indicate that Las1 can suppress the temperature sensitivity of grc3-3 and further strengthened a functional link between Grc3 and Las1. Because the las1-containing plasmid only weakly suppressed the grc3-7 and grc3-8 (data not shown), the grc3-3 allele may have a specific mutation(s) in the region responsible for binding to Las1.

Las1 and IPI Complex Components Localize to Heterochromatic Regions—To examine functional correlations between Grc3-interacting proteins, we examined the subcellular localizations of Crb3, Rix1, Ipi1, and Las1 by expressing C-terminally GFP-fused proteins from their endogenous promoters. Although the subcellular localization of some of these Grc3-interacting proteins has been described previously (24, 25), we confirmed that all of these proteins preferentially localized to the nucleus and formed nuclear spots (Fig. 7). As observed for Grc3-GFP, these nuclear spots were clearly co-localized with the signals for KOR-Swi6 (supplemental Fig. S2). Importantly, deleting swi6 or clr4 caused these nuclear spots to delocalize and led to a characteristic localization pattern surrounding the nucleolus (Fig. 7, A–D, clr4 and swi6Δ). Quantification of the subcellular localization further confirmed a close link between Grc3 and its interacting proteins (compare Fig. 7E with Fig. 2B). These results suggested that Las1 and the IPI complex components also associate with the heterochromatic regions and that they are functionally correlated with Grc3.

Grc3-interacting Proteins Are Required for Both Pre-rRNA Processing and Heterochromatic Gene Silencing—Although functional involvement of IPI complex components in rRNA processing had been described previously in S. cerevisiae (18, 19), the physiological roles of IPI complex components and Las1 in fission yeast have yet to be determined. We first confirmed that these proteins are essential for cell viability (data not shown) (30). To further investigate their cellular function, we isolated temperature-sensitive mutant alleles for crb3+, ipi1+, las1+, or rix1+ by PCR-based mutagenesis method (see “Experimental Procedures” and supplemental Fig. S3 and supplemental Table S3) and tested whether these factors are required for rRNA processing. We found that the temperature-sensitive mutants crb3−3, ipi1−1, las1−1, or rix1−2 accumulated rRNA precursors (35 and 27 S) at the nonpermissive temperature (Fig. 8A). Similar rRNA processing defects were observed in other mutant cells (data not shown).

We next tested whether Grc3-interacting proteins were also required for heterochromatic silencing. We found that higher levels of pericentromeric transcripts were accumulated in crb3−3, las1−1, or rix1−2 mutant cells at the nonpermissive temperature when compared with wild-type cells (Fig. 8B). The effect of ipi1−1 was weaker than that of other conditional mutants. Ipi1 may have distinct functions in heterochromatin silencing. Together, these results suggest that Grc3-interacting proteins are also required for both rRNA processing and heterochromatin assembly in fission yeast.

DISCUSSION

In this report, we demonstrated that the fission yeast Grc3 localizes to both heterochromatin and perinucleolar regions and has a dual role in rRNA processing and heterochromatin silencing. Our biochemical and cytological analyses further revealed that Grc3 is functionally linked with Las1 and the IPI complex. Our findings suggest that rRNA processing and heterochromatic silencing use a shared molecular mechanism that involves Grc3 and its interacting partners.

Previous studies have shown that S. cerevisiae Grc3 is required for ITS2 processing in 27 S pre-rRNA maturation. Using temperature-sensitive mutant cells, we confirmed that S. pombe Grc3 also plays a role in this particular step of rRNA processing. We observed an accumulation of 27 S pre-rRNA and decreased levels of mature 5.8 and 25 S rRNAs in grc3 mutant cells (Fig. 3D). Because the human homologue Nol9 was identified as a nucleolar protein (39), it is conceivable that the Grc3 homologous protein has a conserved role in rRNA processing. Several observations supported the direct involvement of Grc3 in rRNA processing. Grc3 was observed, together with ribosomal proteins, in a protein fraction that was isolated by Ra1 affinity purification (40). In addition, grc3 mutation did not affect the amount of small nuclear RNAs involved in

FIGURE 7. Las1 and IPI complex component localization to heterochromatic regions is dependent on Swi6 and Clr4. A–D, the Grc3-interacting proteins Rix1, Crb3, Ipi1, and Las1 were tagged with GFP at the C terminus and expressed from their native promoter. The in vivo locations of Rix1-GFP (A), Crb3-GFP (B), Ipi1-GFP (C), and Las1-GFP (D) in WT, clr4Δ, or swi6Δ cells were analyzed by fluorescence microscopy. DNA was visualized by Hoechst 33342. Co-localization of each GFP fusion protein with KOR-Swi6 is shown in supplemental Fig. S2. Bar, 5 μm. E, the percentage of cells showing the distinct localization patterns for Rix1-GFP, Crb3-GFP, Ipi1-GFP, or Las1-GFP.
Grc3 has a similar kinase activity and provides enzymatic activity associated with Grc3 may help to explain its role in processing rRNA.

Grc3 has an amino acid sequence similarity with Clp1, a component of the mRNA cleavage and polyadenylation machinery (20, 21). Human Clp1 has been shown to have RNA kinase activity that phosphorylates the 5′-end of synthetic siRNAs and tRNA 3′-exons (32). However, no such activity has been identified for S. cerevisiae Clp1 (41, 42). Because the archaegal Clp1 homologue has polynucleotide kinase activity, it has been suggested that ancestral Clp1 also possesses enzymatic activity (43). A recent study showed that S. cerevisiae Grc3 has a similar kinase activity (33). Considering the high sequence similarity and evolutionary conservation, it is likely that S. pombe Grc3 has a similar kinase activity and provides phosphorylation on the cleaved RNA product for efficient degradation mediated by exonuclease(s). In this study, although we showed that the ATP-GTPase domain is critical for the function of Grc3 in heterochromatic silencing (Fig. 4C), we could not obtain direct evidence of S. pombe Grc3 kinase activity (data not shown). It is possible that Grc3 needs other factors or optimized assay conditions to exert its enzymatic activity or that its ATPase activity is used in other enzymatic processes. Although a recent study showed that S. cerevisiae Grc3 is involved in the transcriptional termination of RNA polymerase I (33), it is not clear how a polymerase I termination defect leads to a ITS2-specific processing defect. We favor the view that Grc3 and other exonuclease activities specifically participate in ITS2 processing. The enzymatic activity of Grc3 may facilitate the efficient degradation of structured RNAs.

In this study, we showed a functional and physical link between Grc3, IPI complex components, and Las1. Although genome-wide studies on S. cerevisiae showed that Grc3 and IPI complex components are required for ITS2 processing, this study provides direct evidence for their functional link. A recent study showing that human Las1L functions in ribosome synthesis (38) further supports our observations. Las1 was originally identified in a genetic screening of S. cerevisiae mutants that could not grow in the absence of an SSD1-1 allele (lethal in the absence of SSD1-1) and was shown to be an essential protein involved in cell morphogenesis and cell surface growth (37). Another recent study revealed that Ssd1 is an RNA-binding protein that preferentially targets transcripts that encode cell wall and bud proteins (44). Although how nuclear-localized Las1 affects Ssd1-bound transcripts remains unclear, Las1 may modulate RNA secondary structure; using such activity, Las1 might alter Ssd1-bound RNAs so they can be efficiently transferred to their particular cellular compartments and similarly prepare ITS2 RNA to be efficiently processed by other factors.

We found that Grc3 and its interacting proteins localize to heterochromatin (Figs. 1 and 7) and that their heterochromatization localization is largely dependent on Swi6 and Clr4. Although heterochromatization localization does not indicate a functional involvement with heterochromatin, it is likely that their activity is linked with heterochromatin assembly. In fission yeast, heterochromatin transcripts are processed by the RNAi pathway and/or by an exosome-mediated degradation process (14, 15). Considering that ITS2 can form complex secondary structures, it is possible that Grc3 and its interacting proteins function to target transcripts with a rigid secondary structure and facilitate their degradation in combination with exonuclease activity such as Rat1. Because heterochromatin localization has not been described for S. cerevisiae counterparts (36), heterochromatin assembly in fission yeast, and probably other higher eukaryotes, may be coupled with active RNA degradation processes involving Grc3 activity.

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Analysis of Grc3 Function in S. pombe