Hmo1 directs pre-initiation complex assembly to an appropriate site on its target gene promoters by masking a nucleosome-free region

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ABSTRACT

Saccharomyces cerevisiae Hmo1 binds to the promoters of ~70% of ribosomal protein genes (RPGs) at high occupancy, but is observed at lower occupancy on the remaining RPG promoters. In Δhmo1 cells, the transcription start site (TSS) of the Hmo1-enriched RPS5 promoter shifted upstream, while the TSS of the Hmo1-limited RPL10 promoter did not shift. Analyses of chimeric RPSS/RPL10 promoters revealed a region between the RPS5 upstream activating sequence (UAS) and core promoter, termed the intervening region (IVR), responsible for strong Hmo1 binding and an upstream TSS shift in Δhmo1 cells. Chromatin immunoprecipitation analyses showed that the RPSS-IVR resides within a nucleosome-free region and that pre-initiation complex (PIC) assembly occurs at a site between the IVR and a nucleosome overlapping the TSS (+1 nucleosome). The PIC assembly site was shifted upstream in Δhmo1 cells on this promoter, indicating that Hmo1 normally masks the RPS5-IVR to prevent PIC assembly at inappropriate site(s). This novel mechanism ensures accurate transcriptional initiation by delineating the 5′- and 3′-boundaries of the PIC assembly zone.

INTRODUCTION

In Saccharomyces cerevisiae, 138 ribosomal protein genes (RPGs) encode 79 ribosomal proteins (RPs). RPG transcription constitutes ~50% of RNA polymerase II (Pol II)-mediated transcription in rapidly growing cells (1) and consumes an enormous amount of energy and protein resources. RPs are found in equimolar amounts in ribosomes, and their production is coordinately regulated in response to certain environmental conditions, mainly at the transcriptional stage.

During the past 10 years, increasing numbers of factors and/or mechanisms that regulate RPG transcription have been identified. Rap1, the most extensively characterized RPG regulator, binds to most RPG promoters (2,3) and activates transcription by recruiting the NuA4 histone acetyltransferase (HAT) complex and/or TFIID (4–6). Rap1 regulates transcription by forming a nucleosome-free region (NFR) in its target promoters (7–9). Abf1, which binds to fewer RPG promoters, is thought to function similarly to Rap1 in forming NFRs (10), although it is unknown whether Abf1 recruits TFIID and NuA4. Fhl1 also binds to many RPG promoters and recruits the coactivator, Ifh1, or the corepressor, Crf1, in response to environmental stimuli (11–14). Sfp1 regulates RPG transcription and expression of the ribosome biogenesis (Ribi) regulon (15,16) via its translocation between nucleus and cytoplasm in response to certain environmental stresses (17); however, its exact function remains unclear.

Hmo1, a high mobility group B (HMGB) protein, plays roles in Pol I and Pol II transcription, rRNA processing, DNA repair and chromosome/plasmid stability (2,18–25). Previous studies showed that Hmo1 binds to the promoter and coding regions of the 35S rRNA gene in a Pol I-dependent manner (2,20,22,26,27). Hmo1 binds to ~70% of RPG promoters, compared to Rap1 (93%) and Fhl1 (90%), and promotes Fhl1 binding to a subset of RPG promoters. Given that Hmo1 commonly targets both rDNA and RPGs, which are transcribed by two different RNA polymerases (Pol I and Pol II, respectively), one can speculate that it plays a crucial and specialized role in coordinating the transcriptional regulation and synthesis of ribosomes. However, little is known about the molecular function of Hmo1 at either Pol I or Pol II loci. The deletion of HMO1 (Δhmo1) has a milder effect on RPG expression than Δfhl1 (13), or mutating the Rap1

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binding site (28, 29). Furthermore, AhmO1 produces different effects among RPGs, which do not necessarily correlate with the amount of Hmo1 binding, suggesting that the primary role of Hmo1 on RPGs may not be transcriptional activation.

In our previous study, we found that AhmO1 caused an upstream shift in the transcriptional start site (TSS) of Hmo1-enriched RPG promoters and rescued the growth defects of certain sua7 (TFIIB) mutants, which, themselves, caused a downstream TSS shift (23). Such suppression phenotypes for sua7, which probably depend on a TSS shift in the direction opposite to that of sua7, have been found only with mutations in four polypeptides within the pre-initiation complex (PIC): the Tfg1 and Tfg2 subunits of TFIIF (30–33), and the Rpb2 and Rpb9 subunits of Pol II (34–37). Recent studies using a cross-linking technique demonstrated that multiple interactions between TFIIF and Rpb2, which may be reinforced by Rpb9, are critical for Ssu selection (38, 39). Presumably, mutations that affect these interactions may impair the specific function of Pol II that is required for selecting the appropriate TSS, or for stabilizing RNA–DNA hybrids during initiation, leading to an upstream TSS shift (30, 38, 39). In contrast to TFIIF, a direct interaction between Pol II and Hmo1 has not been observed (our unpublished data). Furthermore, in AhmO1 cells, a TSS shift was only observed at Hmo1-enriched RPGs, while in tfg1, tfg2, rpb2 and Δrpb9 cells, a TSS shift was observed for most class II (Pol II-driven) genes, regardless of Hmo1 binding (30, 37). Therefore, we suppose that the upstream TSS shift in AhmO1 is caused by a different mechanism than in other mutants, and reflects a defect in a specialized function(s) of Hmo1 with respect to the regulation of transcriptional initiation at the RPG promoter.

The aim of this study was to unveil such a mechanism by determining how AhmO1 induces an upstream TSS shift in Hmo1-enriched RPG promoters. From the results of extensive chromatin immunoprecipitation (ChIP) and primer extension analyses, we identified the IVR (intervening region) between the upstream activating sequence (UAS) and the core promoter (Core) of RPS5 as the binding site of Hmo1, and found that the IVR is nucleosome depleted. In wild-type (WT) cells, the PIC assembled at a site between the IVR and a nucleosome overlapping the TSS (+1 nucleosome), while it assembled within the IVR in AhmO1 cells. These results strongly suggested that Hmo1 and +1 nucleosome determine the 5′- and 3′-boundaries, respectively, of a zone available for PIC assembly, thereby directing PIC assembly at a biologically relevant site.

MATERIALS AND METHODS

Yeast strains and plasmids

Standard techniques were used for the growth and transformation of yeast (40). The yeast strains used in this study are listed in Supplementary Table S1. Detailed information for each strain is described in the Supplementary Data. The yeast culture conditions for each experiment are described in the figure legends. The detailed protocol used to construct the plasmids in this study is described in Supplementary Data. Oligonucleotides used in this study are listed in Supplementary Table S2.

Primer extension analysis

Transcription start sites were mapped by primer extension analysis as described previously (23). The primers used were TK3212 (RPS5), TK3214 (ADH1), TK9589 (RPL27B), TK9911 (RPS5-mini-CLN2) and TK10595 (ADE2-C reporter). Electrophoretic images were acquired by exposing gels to imaging plates (BAS2500, Fuji Film), and the scanning of each lane was carried out using Multi Gauge version 3.0 software (Fuji Film).

ChIP and sequential ChIP analysis

ChIP analysis was conducted according to the Hahn laboratory protocol (http://labs.fhcrc.org/hahn/Methods/mol_bio_meth/hahnlab_ChIP_method.html) with minor modifications. Briefly, DNA was fragmented by sonication to an average size of 400–500 bp for standard ChIP or 100–200 bp for high-resolution ChIP. Immunoprecipitation was conducted using Dynabeads Protein G (Invitrogen) and monoclonal antibodies against FLAG (Sigma-Aldrich; M2), Pk (AbD Serotec; SV5-Pk1) and Myc (Santa Cruz; 9E10); or polyclonal antibodies against histone H3 (Abcam; ab1791), Rap1 (Santa Cruz; yC-19) and Sua7 (in this study, raised against full-length recombinant Sua7 in rabbit). Real-time quantitative PCR analyses were performed using a KAPA SYBR Fast qPCR kit (KAPA) and Mx3000P (Agilent Technologies). PCR conditions were: 95°C for 40 s; 40 cycles of 95°C for 30 s, 52°C for 30 s and 72°C for 30 s. Each experiment was conducted in triplicate and the average and SD of the ratio of immunoprecipitated DNA versus input DNA (IP/input) was calculated. The positions of amplified regions are depicted in each figure. The primer pairs used for PCR are described in the Supplementary Data.

For sequential ChIP analysis, the first immunoprecipitation was performed as for standard ChIP analysis, except that 5 µg of anti-FLAG antibody and cell extracts containing 5 mg of protein were used. After a final wash with TE, precipitates were eluted by incubating beads with 50 µl of ChIP lysis buffer containing 3xFLAG peptide (200 µg/ml; Sigma-Aldrich; MDYKDDDDKMDYKDDDDK DKDDDDK) at 4°C for 30 min. Elution was performed four times in total, and the combined eluates were diluted with ChIP lysis buffer (to a concentration of 100 µg/ml 3xFLAG peptide), and were subjected to a second immunoprecipitation using an anti-Pk antibody. All steps after the second immunoprecipitation were the same as for standard ChIP analysis.

Northern blot analysis

Northern blot analyses were conducted as described previously (2). For the detection of the TEF2 and ADE2-C reporter genes, DNA fragments were amplified by PCR using the primer pairs TK6965–TK6966 (TEF2) and...
TK10425–TK10426 (ADE2-C) and were then 32P-labelled using random priming.

5’ RLM-RACE

5’ RLM-RACE (RNA Ligase Mediated Rapid Amplification of CDNA Ends) analysis was conducted as previously described (41), using the FirstChoiceTM RLM-RACE Kit (Ambion) with total RNA from H2450 (WT) and YTK8276 (Ahmo1) strains. The experiment was conducted according to the instruction manual of the manufacturer (http://www.ambion.com/jp/techlib/prot/fm_1700.pdf). The nested PCR was conducted using universal primers, which bind the RNA adaptor region, and gene-specific primers TK10942 (outer)/TK11567 (inner) for RPS5 or TK11350 (outer)/TK9589 (inner) for RPL27B.

RESULTS

Ahmo1 complements growth defects and reverses a TSS shift due to an rpb1 mutation in a subset of RPGs

In a previous study, we showed that Ahmo1 caused an upstream TSS shift in Hmo1-enriched RPGs and suppressed the temperature sensitive growth of some sua7 mutants (e.g. sua7-R78C, -E62K), which caused a downstream TSS shift in many class II genes (23). To determine whether the suppressive effect of Ahmo1 is specific to sua7 mutants, we tested for a genetic interaction between HMO1 and RPB1, which encodes the largest Pol II subunit. The mutations, rpb1-N445S (42) and rpb1-R344A (43), are in or near to the active centre of Pol II, and cause a downstream TSS shift. As previously reported (42,43), both rpb1 mutants showed significant growth defects at all temperatures tested, and no growth at 37°C (Figure 1A). In contrast, Ahmo1 cells showed less severe growth defects at high temperature than at low temperature. Importantly, Ahmo1 suppressed the growth defect of the rpb1 mutant at 37°C (Figure 1A), as observed for the sua7 mutants (23).

Next, we tested the effect of these rpb1 mutations on the TSS in the RPS5 and ADH1 promoters by primer extension analysis in the same strains. As previously reported, both rpb1 mutants caused a downstream shift of the TSS in both promoters (Figure 1B and C, lanes 1–3 and 7–9); namely, the ratios of the intensities of the major band and lower bands were altered modestly (RPS5; 36 versus 22) or significantly (ADH1; 38 versus 27). In contrast, Ahmo1 caused an upstream shift of the TSS specifically in the RPS5 promoter (Figure 1B and C), one of the most Hmo1-enriched and transcriptionally Hmo1-dependent RPGs, but not in the ADH1 promoter, which binds Hmo1 weakly and is transcriptionally independent of Hmo1 (Figures 1B and C, compare lanes 7 and 10). Briefly, compared to WT cells, in Ahmo1 cells we observed a decrease in the intensity of two minor bands (26 and 22, Figure 1B, left panel) situated below the most intensely stained band, corresponding to the major TSS (36; marked with an asterisk) in the RPS5 promoter. We also noticed an increase in intensity of two bands (71, 87), and the presence of three new bands (133, 215, 225) for TSSs above the major TSS (Figures 1B and C, compare lanes 1 and 4). Although the upper bands in lane 10 appear stronger than those in lane 7 (Figure 1B), the ratio to the band at 38 (double dagger) was nearly identical in lanes 10 and 7 (Figure 1C).

Note that the upstream TSS shift in Ahmo1 cells can be also detected by another method, 5’ RLM-RACE. The 5’ RLM-RACE is a modified 5’ RACE to amplify selectively the 5’-end of full-length mRNA that contains a base corresponding to TSS. The TAP-dependent bands, which were amplified by PCR, reflect the intact 5’-end of mRNAs of RPS5 (Supplementary Figure S1A, lane 1 and 3). The result shows that TSSs of RPS5 were shifted upstream in Ahmo1 cells (Supplementary Figure S1A, compare lane 1 and 3). A similar result was obtained with the same analysis for RPL27B (Supplementary Figure S1B, compare lane 1 and 3), which showed a more drastic TSS shift than RPS5 in Ahmo1 cells (Supplementary Figure S2A). The results indicate that the upstream TSS shift that was identified by primer extension analysis occurred in Ahmo1 cells.

Importantly, Ahmo1 partly reversed the TSS shift in the RPS5 promoter (Figure 1B and C, compare lanes 1, 4, 5 and 6) but not in the ADH1 promoter (Figure 1B and C, compare lanes 7, 10, 11 and 12) in both rpb1 mutants. This effect was stronger in the Ahmo1 rpb1-R344A mutant (Figure 1B and C, compare lanes 1 and 5, 6), consistent with the observation that Ahmo1 suppressed the growth defect of rpb1-R344A more strongly than for rpb1-N445S (Figure 1A). These results suggested that Ahmo1 suppresses the growth defects of rpb1 mutants by reversing the TSS shift in certain Hmo1-enriched and transcriptionally Hmo1-dependent genes, such as RPS5.

Ahmo1 causes an upstream TSS shift by a different mechanism than tfg1 or Arab9

While all mutant genes that are known to cause the upstream TSS shift encode subunits of Pol II or limited components of PIC (30–37), only Hmo1 is not a component of PIC. This suggests that the upstream TSS shift in Ahmo1 is caused by a different mechanism than in other mutants. Therefore, its mechanism may involve a specialized function(s) of Hmo1 with respect to the regulation of transcriptional initiation at the RPG promoter. To understand the mechanism behind the upstream TSS shift in Ahmo1 and other mutants, transcriptional phenotypes for Ahmo1, Pol II (Arab9) and TFIIF (tfg1-E346A) mutants were compared.

First, we compared the TSS profiles of the RPS5 and ADH1 promoters in three single mutants (Ahmo1, tfg1-E346A and Arab9) and three double mutants (Ahmo1 tfg1-E346A, Ahmo1 Arab9 and tfg1-E346A Arab9) with those in WT cells. As previously reported (31), the tfg1-E346A Arab9 double mutant exhibited more severe growth defects than either of the single mutants (tfg1-E346A, Arab9) at 25°C, 30°C and 35°C (Supplementary Figure 3A). Primer extension analyses revealed that the tfg1-E346A and Arab9 mutations caused upstream TSS shifts in the RPS5 and ADH1 promoters, while the shift...
in each promoter was enhanced in the double mutant (Supplementary Figure S3B and C). Consistent with a previous study (30), although the degree of TSS shift was slightly different between \textit{tfg1-E346A} and \textit{Drpb9}, the positions of the upstream TSS in both promoters were nearly the same in both mutants (Supplementary Figure S3B and C, compare lanes 2, 6 and 3, 7, respectively), suggesting that the mechanism for the TSS shift may be similar in these mutants.

In contrast, the feature of TSS shift was quite different in \textit{Dhmo1} and \textit{tfg1-E346A} mutants (Figure 2). Primarily, \textit{tfg1-E346A} caused an upstream TSS shift in all promoters tested (\textit{RPS5}, \textit{ADH1}, \textit{SPT15}, \textit{HTB1}, \textit{GAL1}, \textit{GAL10}, \textit{HIS3}, \textit{HIS4}, \textit{SNR7}, \textit{SNR14}, \textit{SNR19} and \textit{SNR20}) [in this study, and (30)], while \textit{Dhmo1} shifted the TSS specifically in the Hmo1-enriched RPG promoters, e.g., \textit{RPS5}, \textit{RPL32} (23), and \textit{RPL27B} (Supplementary Figure S2A). Furthermore, it is

Figure 1. Genetic interaction between \textit{HMO1} and \textit{RPB1}. (A) Effect of \textit{Dhmo1} and/or \textit{rpbl} mutations on growth. \textit{Arpbl} and \textit{Arpbl Dhmo1} cells carrying a plasmid encoding the \textit{RPB1} (WT) or \textit{rpbl} mutant (N445S, R344A) were spotted onto YPD (yeast extract, peptone, dextrose) plates at three dilutions and grown for 3 days (30°C, 35°C and 37°C) or 4 days (25°C). (B) Effect of \textit{Dhmo1} and/or \textit{rpbl} mutations on the TSS in \textit{RPS5} and \textit{ADH1} promoters. The strains described in (A) were grown at 25°C. Total RNA (15 \mu g) was prepared and analysed by primer extension. The positions of several TSSs are indicated on the left (\textit{RPS5}) or right (\textit{ADH1}). TSSs are numbered relative to the A (+1) of the start codon, ATG (−22, −26, −36, −71, −87, −133, −215 and −225 for the \textit{RPS5} promoter and −27 and −38 for the \textit{ADH1} promoter). TSSs at −36 and −87 in the \textit{RPS5} promoter and −27 and −38 in the \textit{ADH1} promoter are marked with asterisks (* and **) and daggers († and ‡), respectively. (C) Results shown in (B) were quantified by densitometry. The number in the upper right corner of each panel corresponds to the lane number in (B). Values were normalized to the strongest peak in each panel (i.e. strongest peak set to a value of 1). The horizontal axis represents the position of each band within the region shown in (B). Note that the regions shown in (B) and (C) are identical. Asterisks (* and **) and daggers († and ‡) correspond to the bands at −36 and −87 (\textit{RPS5}), and those at −27 and −38 (\textit{ADH1}), respectively, as shown in (B). Part of the scanned region was enlarged and is shown in the inset to highlight the differences.
noteworthy that transcription from −51A in the RPS5 promoter was markedly enhanced by tfg1-E346A, but not by Δhmo1 (Figure 2B, lanes 1–4). Conversely, TSSs around −220 (−215 and −225) were induced uniquely by Δhmo1 (Figure 2B, lanes 1–4). Similar but weaker effects were observed in Δhmo1, Δrpb9 and Δhmo1 Δrpb9 cells (Figure 3, lanes 1–4). These results suggested that the mechanism(s) underlying the TSS shift in the Δhmo1 cells might be different from that in the TFIIF/Pol II mutants. However, direct evidence will be required to confirm this possibility.

Although the Δhmo1 tfg1-E346A double mutant had more severe synthetic effects on the TSS compared to either of the single mutants, we found no obvious synthetic growth defect for this double mutant when compared to the single mutants. A similar result was obtained for the Δhmo1 Δrpb9 double mutant (data not shown). Therefore, it seems unlikely that the upstream TSS shift itself is the

Figure 2. Genetic interaction between HMO1 and TFG1. (A) Effect of Δhmo1 and/or tfg1-E346A on growth. Growth of Δtfg1 and Δtfg1 Δhmo1 cells carrying the plasmid encoding TFG1 (WT) or tfg1-E346A was analysed as described in Figure 1A. (B) Effect of the Δhmo1 and/or tfg1-E346A mutation on the TSS in RPS5 and ADH1 promoters. Yeast strains were as indicated in (A). Cells were grown at 25°C, and total RNA (15 μg) was prepared and analysed by primer extension as described in Figure 1B. (C) Results shown in (B) were quantified and the data are summarized as described in Figure 1C.
Because Δhmo1 decreased the binding of Fhl1 to the RPG promoter (2,22), it is possible that the upstream TSS shift in Δhmo1 cells is the result of the dissociation of Fhl1 from the RPG promoter. This was tested by analysing the TSS of RPG in Δfhl1 cells by primer extension analysis. Although Δfhl1 shifts the TSS of RPG modestly as compared with WT, the extent of the TSS shift was less severe in Δfhl1 cells than in Δhmo1 cells (Supplementary Figure S2B; TSSs upstream of -87 were not observed in Δfhl1 cells). Given that Δfhl1 decreases the Hmo1 binding to a subset of RPG promoters (22), the TSS shift in Δfhl1 cells may be related to a decrease in Hmo1 binding rather than the direct effect of the loss of Fhl1 function. However, Δhmo1 caused a less pronounced upstream TSS shift in an Hmo1-enriched RPG, RPS18A, whose binding to Fhl1 was not affected in Δhmo1 cells (2), than in the RPG5 promoter (Supplementary Figure S2C). Therefore, it cannot be excluded that Fhl1 has also a direct role in the selection of the correct TSS.

Besides RPG promoters, Hmo1 also binds abundantly to a subset of non-RPG promoters (2,22). Therefore, we tested whether Δhmo1 caused upstream TSS shifts in these promoters. Remarkably, the TSS of FET3, a non-RPG with strong Hmo1 binding activity, was not affected in Δhmo1 cells (Supplementary Figure S2D), suggesting that the function of Hmo1 at non-RPG promoters may be different from that at RPG promoters.

Upstream TSS shift is caused by a pre-PIC assembly defect in Δhmo1 cells and by a post-PIC assembly defect in Arpb9 cells

In S. cerevisiae, a ‘scanning model’ has been proposed to explain TSS selection by Pol II (44–46). In this model, following PIC assembly at the core promoter, Pol II starts scanning downstream and initiates transcription when it encounters an appropriate TSS. According to this model, two different mechanisms for an upstream TSS shift seem possible. The first mechanism involves normal PIC assembly, followed by a defect in the post-PIC assembly function of Pol II, while the second mechanism proposes an upstream shift of the PIC.
assembly site, reflecting a pre-PIC assembly defect but a normal post-PIC assembly function of Pol II.

The rfg1, rfg2, rpb2 and Arpb9 mutants are thought to cause upstream TSS shifts via the post-PIC assembly defect, while Δhmo1 causes the TSS shift by a different mechanism, presumably the pre-PIC assembly defect. If a TSS shift is due to relocation of the PIC assembly site upstream in Δhmo1 cells, we predicted that insertion of a TATA element to induce upstream PIC assembly artificially would not alter the TSS significantly in Δhmo1 cells, but would cause a significant TSS shift in WT cells. As a result, the TSS pattern in WT and Δhmo1 cells would be similar. In contrast, if the TSS shift is caused by a post-PIC assembly defect, such as a change in Pol II activity, we predicted that insertion of an upstream TATA element would generate novel TSS patterns in both WT and Δhmo1 cells, and that these patterns would be different. To test this hypothesis, an ectopic TATA element was engineered at two upstream positions, −125 (TATA1) or −165 (TATA2), in the RPS5 promoter (Figure 3A). The original or modified RPS5 promoters were fused to a mini-CLN2 reporter gene, a non-functional CLN2 lacking part of the ORF (47), and inserted into a low-copy number plasmid. Importantly, the RPS5 promoter on the plasmid had almost identical properties to the chromosomal RPS5 promoter, including abundant Hmo1 binding, Hmo1-dependent Fhl1 binding, and TSS profiles in WT, Δhmo1 and/or Arpb9 cells (Supplementary Figure S4A and B, and data not shown). Primer extension analyses were conducted using WT, Δhmo1, Arpb9 and Δhmo1 Arpb9 cells, each containing a mini-CLN2 reporter plasmid carrying the RPS5, RPS5-TATA1 or RPS5-TATA2 promoters (Figure 3B and C). In all strains tested, RPS5-TATA1 had no significant effect on the TSS, possibly because the endogenous PIC assembly site is close to this site (Figure 3B and C, compare lanes 1–4 with 5–8). On the contrary, RPS5-TATA2 caused a more modest upstream TSS shift in Δhmo1 cells than in WT cells (Figure 3B and C, compare lanes 1, 2 and 9, 10, respectively). As the result, the TSS patterns became similar in WT and Δhmo1 cells, at least in the region downstream of this insertion (Figure 3B, compare lanes 9 and 10). In contrast, RPS5-TATA2 drastically altered the TSS pattern in Arpb9 and Arpb9 Δhmo1 strains (Figure 3B and C, compare lanes 3, 4 and 11, 12). As the result, the TSS patterns of the RPS5-TATA2 promoter in Arpb9 and Arpb9 Δhmo1 became significantly different from those in WT and Δhmo1 (Figure 3B and C, compare lanes 9, 10 and 11, 12). These results suggested that Δhmo1 shifts the TSS by relocating the PIC assembly site upstream, while Arpb9 shifts the TSS by causing a defect(s) in Pol II activity at a post-PIC assembly step.

An intervening region between the UAS and the core promoter is required for Hmo1 binding and the upstream TSS shift caused by Δhmo1 in Hmo1-enriched RPG promoters

Δhmo1 caused an upstream TSS shift in the Hmo1-enriched RPS5 promoter, but not in the Hmo1-limited RPL10 promoter (23). Therefore, to identify the RPS5 promoter region responsible for this shift in Δhmo1 cells, we constructed a series of chimeric promoters in which the UAS, Core or IVR of RPS5 and RPL10 were mutually exchanged (Figure 4A). These modified promoters were integrated into the ADE2 chromosomal locus in WT or Δhmo1 cells. Primer extension analysis for these strains clearly revealed that the IVR of RPS5 was required for an upstream TSS shift in Δhmo1 cells (Figure 4B) while the UAS and Core of RPS5 can be exchanged for those of RPL10 without blocking the TSS shift.

RPL27B showed almost the same properties as RPS5 with respect to Hmo1 binding, Hmo1-dependent transcription and Fhl1 binding. As expected, Δhmo1 also induced an upstream TSS shift in an additional chimeric promoter, constructed from the IVR of RPL27B and the UAS and Core of RPL10 (Figure 4A and B, lanes 17 and 18), implying that the IVR of Hmo1-enriched RPGs is a critical determinant for the TSS shift in Δhmo1 cells.

These results suggested that the IVRs in the RPS5 and RPL27B promoters are also required for abundant Hmo1 binding to these promoters. However, ChIP analyses revealed that Hmo1 binds abundantly to promoters containing the RPS5- or RPL27B-IVR, but not to those containing the RPL10-IVR (Figure 4C, panels 1 and 2). The roles of the IVR, UAS and Core in supporting abundant Hmo1 binding were examined by ChIP analyses of RPS5 promoters lacking each of these segments. The results clearly revealed that the IVR is essential, but the UAS and Core are dispensable, for Hmo1 binding (Figure 4C, panels 3 and 4). Previously, Hall et al. (22) reported that Hmo1 binding to the RPS11B promoter at the HIS3 chromosomal locus was dependent on Rap1 binding sequences. Therefore, we used ChIP analysis to test whether our deleted UAS (ΔUAS) construct contained a cryptic Rap1 binding site. The result confirmed that Rap1 was absent in this construct (Supplementary Figure S5A and B). Furthermore, the ΔUAS construct showed much weaker transcription than WT. These results excluded the possibility that a cryptic Rap1 (or Abf1) binding site would decrease Rap1-dependency of Hmo1 binding in our ΔUAS construct. It is possible that Rap1-dependency of Hmo1 binding differs at each genomic locus.

Further mapping analysis, using a series of promoter constructs, in which 40-bp segments in the RPS5 promoter were deleted systematically, found no segments that were indispensable for Hmo1 binding (data not shown). However, ChIP analysis using promoter constructs, in which the RPS5-IVR between the UAS and Core of RPL10 was deleted serially from upstream or downstream, revealed at least two non-overlapping sequences that supported abundant Hmo1 binding in the RPS5-IVR (~439 to ~260 bp and ~259 to ~127 bp; Supplementary Figure S6A and B). In addition, the ~319 to ~199 bp region (120 bp), which overlaps these two regions, also supported full Hmo1 binding (data not shown). Thus, abundant Hmo1 binding to the RPS5-IVR occurs by the independent or cooperative functioning of multiple specific Hmo1 binding sites.
Figure 4. Mapping of the promoter region required for Hmo1 binding and upstream TSS shift in \textit{AlmB} cells. (A) Schematic diagram depicting chimeric \textit{RPS5}/\textit{RPL10} promoters, \textit{RPS5} promoters lacking one of three segments (UAS, IVR or Core), or a promoter containing the \textit{RPL27B}-IVR. The designation indicated at the left is an abbreviation of each promoter construct. For instance, '5-5-5' at the top denotes a construct that contains \textit{RPS5}-UAS, \textit{RPS5}-IVR, and \textit{RPS5}-Core (+16 bp of 5' region of \textit{RPS5} ORF). All modified promoters were fused to \textit{His3MX6} by PCR and then integrated into the \textit{ADE2} locus with an accompanying deletion of an \~{}1.2-kb DNA region encoding the N-terminal portion of Ade2. The regions amplified by PCR in the ChIP assays are underlined and labelled 'a', 'b' or 'c'. The primer TK10595, used for primer extension, is indicated with an arrow. (B) The promoter region required for the upstream TSS shift in \textit{AlmB} was analysed by primer extension, as described in Figure 1B. The TSSs of the chimeric promoters, described in (A), were examined in \textit{AlmB} (odd-numbered lanes; D) or WT cells (even-numbered lanes; W). TSSs at \~{}36 and \~{}87 in the \textit{RPS5}-Core and \~{}21 in the \textit{RPL10}-Core are marked with asterisks (* and **) and dagger (†), respectively. (C) Hmo1 binding to the test promoters described in (A) was analysed \textit{in vivo} by ChIP assays. The strains carrying modified promoters and expressing Hmo1-FLAG were grown in YPD medium to mid-log phase at 25°C. Cross-linked chromatin was prepared and immunoprecipitated with an anti-FLAG antibody (0.1 μg) and Dynabeads Protein G. Immunoprecipitation was also conducted using an anti-Myc antibody (1 μg) as a negative control (indicated as '−'). Panels 1 and 4 summarize the results for the promoters that contain the \textit{RPS5}-Core (region 'a' is amplified). Panel 2 summarizes the results for (continued)
Alternatively, the length of the IVR might be more critical for Hmo1 binding than a specific DNA sequence because a correlation was observed between the length of the IVR and Hmo1-binding in our deletion analysis of RPS5-IVR (Supplementary Figure S6B) or in endogenous RPG promoters (22) (our unpublished data). To address this possibility, we constructed several promoter constructs that contained different DNA fragments, e.g. a non-promoter sequence from chromosome V (Chr. V) of S. cerevisiae or pBR322 plasmid, or replicate RPL10-IVR between the UAS and Core of RPL10 (Supplementary Figure S6C). Using these modified promoter constructs, the ChIP analysis revealed that while RPS5-IVR (−439 to −260; 180 bp) bound similar levels of Hmo1 as full-length RPS5-IVR, two unrelated sequences of similar lengths showed modest (pBR322) or no (Chr. V) Hmo1 binding (Supplementary Figure S6D). Similarly, replicate RPL10-IVR, which is longer than RPS5-IVR, could not bind Hmo1 (Supplementary Figure S6D). These results suggest that a specific sequence of DNA is more critical than the length of the DNA for Hmo1 binding.

The finding that multiple Hmo1 binding sites exist in RPS5-IVR raised the additional question of whether more than two Hmo1 molecules can bind to an RPS5 promoter simultaneously. To address this question, we conducted sequential ChIP analysis using antibodies against chromatin from yeast cells expressing both of the different Hmo1 species (strain 3). The samples were subjected to sequential immunoprecipitation using an anti-FLAG antibody (first) and an anti-Pk antibody (second). PCR was conducted using both antibodies against chromatin from yeast cells expressing both of the different Hmo1 species (strain 3). This is consistent with our notion that Rap1-dependency of Hmo1 binding may differ depending on the locus. Remarkably, ChIP analyses also showed that the RPS5-IVR is nucleosome-depleted (Figure 5A, panel a) and that the PIC assemblies at a site between the binding peaks of Hmo1 and a nucleosome (3'-side) on the RPS5 promoter (Figure 5A, panel b). Similar binding properties for Hmo1, TFIIB and histone H3 were observed for the other two Hmo1-enriched promoters (RPL27B and HMO1, Figure 5A, panels c-d and g-h, respectively).

Recent ChIP-seq studies revealed that many class II gene promoters have two well-positioned nucleosomes (−1 and +1) (48). The −1 nucleosome, located 150–300 bp upstream of the TSS, regulates the access of transcription factors to this region, while the upstream boundary of the +1 nucleosome lies 10–15 bp upstream of the TSS (49,50). As a result, a relatively wide NFR (~140 bp) is formed between these two nucleosomes. Intriguingly, RPG promoters have a significantly broader NFR than other promoters, possibly due to the lack of the −1 nucleosome (48). Our results showed that, in Hmo1-enriched promoters, Hmo1 apparently binds to the position occupied by the −1 nucleosome in other promoters.

The spatial arrangement of the PIC, +1 nucleosome and Hmo1 within Hmo1-enriched promoters suggests that Hmo1 and the +1 nucleosome direct assembly of the PIC to a specific site. In this regard, Hmo1 is a novel transcription factor involved in determining the 5'-border of a region available for PIC assembly within the core promoter. In contrast to Hmo1-enriched promoters (RPS5, RPL27B and HMO1), the binding peaks of Hmo1 and PIC overlapped in the RPL10 promoter (Figure 5A, panels e and f). Although there was no evidence to exclude the possibility that Hmo1 and the PIC bind together at the same position in the RPL10 promoter, we assume this is due to limitations of the ChIP resolution in the relatively narrow RPL10-IVR. As an alternative possibility, Hmo1 and PIC could bind to the same position, but in different cell populations.

The binding profiles of Hmo1 and nucleosomes raised the possibility that Hmo1 may inhibit nucleosome formation on the IVR. To test this possibility, we used ChIP analysis to compare histone H3 binding profiles in WT and Δhmo1 cells. The results showed that the RPS5 promoter has a similar NFR in both cell types despite the slight increase in histone H3 binding in Δhmo1 cells (Figure 5B), suggesting that Hmo1 does not play a critical role in the formation and/or maintenance of NFRs.

Δhmo1 shifts the PIC assembly site upstream in the RPS5 promoter

The results described above suggested that Hmo1 and the +1 nucleosome determine the 5'- and 3'-border,
respectively, of the PIC assembly zone. If this is so, \( \Delta hmo1 \) should disrupt the 5' border of this zone, allowing ectopic PIC assembly on the IVR. An effect that induces ectopic PIC assembly could account for the upstream TSS shift in \( \Delta hmo1 \) cells. To test this possibility directly, ChIP analysis was conducted to determine the binding positions of several PIC components including TFIIB (Sua7), TFIIF (Tfg1), TFIIE (Tfa2) and TFIIH (Tfb3) on the \( RPS5 \) promoter in WT and \( \Delta hmo1 \) cells. The binding positions of these factors were shifted upstream in \( \Delta hmo1 \) cells (i.e. from position 8 to 7; Figure 6B, compare panels a, c, e, g with panels b, d, f, h, respectively). Our recent mapping analysis for core promoter elements in the \( RPS5 \) promoter revealed that the region corresponding to position 7 cannot bind PIC in the native context in the WT cells, although it can induce PIC assembly when artificially inserted into a different context (51). Therefore, the upstream shift of the PIC assembly site observed in \( \Delta hmo1 \) cells reflects the ectopic PIC assembly in this region (approximately position 7), where PIC intrinsically does not assemble in WT cells. In contrast, a similar upstream shift in the binding position of the PIC component (Sua7) was not observed in \( tfg1-E346A \) and \( Drpb9 \) cells (Figure 6C), indicating that the upstream TSS shift in these mutants was caused by a post-PIC assembly defect. Therefore, we concluded that Hmo1 cooperates with the +1 nucleosome to direct PIC assembly to a site between the IVR and the +1 nucleosome by determining the 5' and 3' boundaries of a zone available for PIC assembly.

**DISCUSSION**

In this study, we aimed to find a function for Hmo1 in the regulation of transcriptional initiation of Hmo1-enriched promoters. (A) Exact binding sites for Hmo1, nucleosome and PIC were identified in several promoters by high-resolution ChIP analyses. The schematic diagrams depicting the endogenous promoter regions tested are indicated above the panels, which summarize the results for each promoter (\( RPS5, \ RPL27B, \ RPL10 \) and \( HMO1 \)). These promoters were divided into UAS, IVR and Core based on information for certain promoter elements (see Supplementary Data). The grey arrows represent the direction of the ORF. The black bars closely aligned to the entire promoter (and part of the ORF) of each gene represent the regions amplified by PCR in ChIP assays. ChIP analyses were conducted as described in Figure 4C, except that chromosomal DNA was fragmented to an average size of 100–200 bp. The ChIP results for Hmo1 (+Pk tag) and histone H3 are indicated in the upper panels (a, c, e, g) with a solid line or broken line, respectively, while results for TFIIB (Sua7-Pk) are indicated in the lower panels (b, d, f, h). The numbers indicated under the panels ('1'–'36') correspond to the regions depicted in the schematic diagram. (B) The binding profiles of histone H3 on the \( RPS5 \) promoter were analysed in WT and \( \Delta hmo1 \) cells by ChIP analysis as described in (A).
Figure 6. $\Delta$hmo1 shifts the PIC assembly site upstream in the RPS5 promoter. (A) Schematic diagram depicting the endogenous RPS5 promoter. (B) Analysis of the effect of $\Delta$hmo1 on the position of PIC assembly on the RPS5 promoter. The exact positions of several PIC components including TFIIB (Sua7-Pk), TFIIF (Tfg1-Pk), TFIIE (Tfa2-Pk) and TFIIH (Tfb3-Pk) on the RPS5 promoter were determined in WT and $\Delta$hmo1 cells by ChIP analysis as described in Figure 5A. The upper and lower panels for each component indicate the results in WT and $\Delta$hmo1 cells, respectively. The numbers under each panel (4'–10') represent the position of regions amplified in the RPS5 promoter as depicted in Figure 5A. (C) Analysis of the effect of $\Delta$rpb9 or $\Delta$tfg1-E346A on the position of PIC assembly on the RPS5 promoter. The TFIIB binding site on the RPS5 promoter was determined in WT, $\Delta$rpb9 and $\Delta$tfg1-E346A cells by ChIP analysis as described in (A), except that immunoprecipitation was conducted by using an anti-Sua7 polyclonal antibody. (D) A model for the proposed role of Hmo1 in Hmo1-enriched RPG promoters (DBD, DNA binding domain; AD: Activation domain). See text for description of the model.
RPGs by determining how $\Delta hmo1$ induces a TSS shift in these genes. The results showed that: (i) the upstream TSS shift in $\Delta hmo1$ cells was due to a pre-PIC assembly defect, while the shifts in $\Delta rpb9$ and $\trianglefigl$-E346A cells were caused by a post-PIC assembly defect; (ii) Hmo1 binds over a broad region corresponding to the RPS5-IVR, which is nucleosome-depleted; (iii) multiple Hmo1 molecules bind to the RPS5-IVR; (iv) PIC assemblies at a site flanked by Hmo1 and the +1 nucleosome; (v) Hmo1 does not play a critical role in the formation and/or maintenance of NFRs, at least in the RPS5 promoter; and (vi) $\Delta hmo1$ causes an upstream shift of the PIC assembly site.

Based on these findings, we proposed a model for a novel function of Hmo1 on its target promoters (Figure 6D). Initially, certain activators on the UAS (e.g. Rap1), remove the nucleosomes around the IVR. In WT cells, Hmo1 then binds to nucleosome-free IVRs to inhibit ectopic PIC assembly in this region, thereby directing PIC assembly to a biologically relevant site in the core promoter (Figure 6D, WT). In contrast, in $\Delta hmo1$ cells, activator(s) facilitate PIC assembly at more proximal site(s) within the IVR, which are devoid of Hmo1 and nucleosomes, leading to an upstream TSS shift (Figure 6D, $\Delta hmo1$).

While $\Delta hmo1$ does not have a severe effect on NFR in the RPS5 promoter, it causes a slight increase in histone H3 binding to the core promoter region of RPS5 (Figure 5B, compare region 8 between WT and $\Delta hmo1$). Because this region is occupied by PIC but not by Hmo1 in WT cells (Figure 5A), we assume that this slight increase in histone H3 binding in $\Delta hmo1$ cells may be caused by a decrease in PIC binding. However, we currently cannot exclude the opposite possibility that the increase in histone H3 binding may cause an upstream shift of the PIC assembly site. In this case, $\Delta hmo1$ would somehow allow invasion of the +1 nucleosome into the core promoter, pushing the PIC towards a more upstream site(s) (Figure 6D, $\Delta hmo1$).

While finding that Hmo1 specifically binds to the NFR on Hmo1 target genes suggests that a nucleosome-free state is a pre-requisite for Hmo1 binding, we have not yet been able to identify specific cis-element(s) for Hmo1 binding in the mapping analysis, possibly because there are multiple binding sites for Hmo1 in RPS5-IVR. Although the IFHL motif and/or GGY(n) repeat were proposed as a binding site for Hmo1 by bioinformatic approaches (22,52), not all Hmo1-enriched RPG promoters contain these element(s) (our unpublished data). Furthermore, deletion of the IFHL motif reduced Hmo1 binding only modestly (22) (our unpublished data). Previously, Hmo1 was isolated in a yeast one-hybrid screen as a CAG repeat binding protein (53). Notably, the CAG repeat and IFHL motifs [or GGY(n) repeat] are GC-rich. In addition, the GC-content in the IVR of Hmo1-enriched RPGs (48.6%) is significantly higher, on average, than in Hmo1-limited RPG (36.1%) (our unpublished data). Therefore, we speculate that Hmo1 may recognize sequences with a relatively high GC-content, as represented by the IFHL motif within the NFR, and conversely that Hmo1 may be excluded from core promoter regions, which are very AT-rich. This speculation seems to be consistent with the results in Supplementary Figure S6C and D.

In previous studies, spt was identified as a suppressor of defects associated with insertions of the Ty1 transposon or $\delta$ sequence into the HIS4 or LYS2 promoters (54–56). Despite the lack of direct evidence, it is likely that relocation of the TSS from the Ty1 or $\delta$ sequence to the original HIS4 or LYS2 promoters is caused by a shift in the PIC assembly site in some spt mutants. It is possible that mutations in TBP (SPT15), SAGA (SPT3, SPT7, SPT8 and SPT20) and Mediator (SPT13) might cause this phenotype through changes in the sequence specificity of PIC components, not due to the unmasking of the NFR. Notably, SPT2 encodes a yeast HMG-like protein and participates in the repression of cryptic transcription in the coding region (57,58), showing similarities to Hmo1. However, it is likely that the spt phenotype is due to destabilization of nucleosomes in spt2 cells (57), and such a defect has also been described in spt6 and spt16 cells (59). Similarly, mutations in histones (SPT11 and SPT12) and in the transcriptional regulators of histone genes (SPT1, SPT10 and SPT21) may produce the spt phenotype by a similar mechanism. In contrast, Hmo1 does not severely affect the position/stability of nucleosomes, but rather functions as if it replaces the function of nucleosomes in the system, ensuring that PIC can be assembled only at a physiologically relevant site by a novel mechanism.

Considering the fundamental importance of nucleosomal structures for cell growth, it is difficult to determine the specific roles of −1/+1 nucleosomes in PIC assembly. Nevertheless, the precise positioning of these nucleosomes suggests that they would determine the 5′- and 3′-boundaries, respectively, of the zone for PIC assembly (i.e. NFR). In a subset of RPG promoters, Hmo1 links to a region that is usually occupied by the −1 nucleosome in other promoters, while $\Delta hmo1$ allows invasion of PIC assembly in this region. These results indicate that Hmo1, instead of the −1 nucleosome, would determine the 5′-boundary of the zone for PIC assembly, at least in Hmo1-enriched RPG promoters, whereas the +1 nucleosome would still determine the 3′-boundary even in these promoters. To our knowledge, this is the first experimental evidence to show that there is indeed a 5′-boundary at the zone for PIC assembly and that the boundary is formed by a protein other than histones, at least in some promoters. Currently, PIC components are thought to bind to core promoters via the recognition of defined (or ill-defined) cis-elements and/or histone modifications of −1/+1 nucleosomes (49). In this context, the mechanism described above, by which certain factor(s) restrict the zone for PIC assembly, should provide an additional layer of specificity to the system, ensuring that PIC can be assembled only at a physiologically relevant site.

Besides a role in directing PIC assembly to an appropriate site, Hmo1 may also promote PIC assembly itself, since the binding of PIC components decreased significantly in $\Delta hmo1$ cells (Figure 6B). In such a role, Hmo1...
could facilitate PIC assembly by recruiting and/or stabilizing TFIIID on its target promoters, as proposed for the −1 and +1 nucleosomes (49), either via direct interaction with TFIIID subunits (23), or through Fhl1/Ifh1 coactivators (2,22). In fact, some yeast (Nhp6α/b) and human (HMGB1/2) HMGB proteins are known to activate transcription by stabilizing the TBP/TFIID–TFIIA promoter complex (60,61).

As another mechanism, Hmo1 might promote transcription by bending or looping a promoter DNA (62). A recent study, using cryo-electron microscopy, revealed that Rap1 on the UAS associates with TFIIA/TFIID at the core promoter, resulting in the looping-out of the region between the UAS and core promoter (i.e. IVR) (63). Intriguingly, the IVRs of Hmo1-enriched RPGs are significantly longer (approximately twice) than those of Hmo1-limited RPGs (22) (our unpublished data). Therefore, Hmo1 might promote and/or stabilize loop-formation, thereby helping Rap1 to affect TFIIA/TFIID efficiently from a distance in RPG promoters containing long IVR. Consistent with this, a TFIIA mutant, toa1–2 (K257A R257A K259A), which has defect in loop-formation (63) or in TFIIA–TFIID interaction (64), showed synthetic growth defects with Δhmo1 (23). At present, although there is no evidence to show that Hmo1-mediated loop-formation occurs in vivo, DNA-looping mediated by Top2 and Hmo1 was proposed to prevent chromosome fragility of variously transcribed intergenic regions in S-phase (21).

In summary, we found that Hmo1 plays a novel role in transcription by forming the 5′-boundary (instead of −1 nucleosome on many other promoters) for the PIC assembly zone on a subset of RPG promoters. Intriguingly, Hmo1 binds to 35S rDNA but only to nucleosome-free (i.e. actively transcribed) repeats in this gene (26). These alternate localizations of Hmo1 and nucleosomes to both RPG and rDNA loci indicate that Hmo1 may have specialized functions that cannot be replaced by nucleosomes. An attractive hypothesis is that the novel function of Hmo1 may play an important role in the coordinated synthesis of RP and rRNA under various environmental conditions. Further studies will be required to define more precisely the roles of Hmo1 in transcription at these two loci.

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

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