A Nucleosome Surface Formed by Histone H4, H2A, and H3 Residues Is Needed for Proper Histone H3 Lys\(^{36}\) Methylation, Histone Acetylation, and Repression of Cryptic Transcription*

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Histones are subject to an array of post-translational modifications, including methylation, acetylation, phosphorylation, and ubiquitination (1, 2). These covalent histone modifications function to modulate gene expression and other DNA templated processes by altering chromatin structure and/or recruiting additional effector proteins (3–5). In addition, it is believed that aberrant levels or patterns of histone modifications can alter gene expression profiles, leading to improper cell growth and/or differentiation (6).

Set2-mediated H3 Lys\(^{36}\) methylation is a histone modification that has been demonstrated to function in transcriptional elongation by recruiting the Rpd3S histone deacetylase complex to repress intragenic cryptic transcription. Recently, we identified a trans-histone pathway in which the interaction between the N terminus of Set2 and histone H4 Lys\(^{44}\) is needed to mediate trans-histone H3 Lys\(^{36}\) di- and trimethylation. In the current study, we demonstrate that mutation of the lysine 44 residue in histone H4 or the Set2 mutant lacking the histone H4 interaction motif leads to intragenic cryptic transcripts, indicating that the Set2 and histone H4 interaction is important to repress intragenic cryptic transcription. We also determine that histone H2A residues (Leu\(^{116}\) and Leu\(^{117}\)), which are in close proximity to histone H4 Lys\(^{44}\), are needed for proper trans-histone H3 Lys\(^{36}\) methylation. Similar to H4 Lys\(^{44}\) mutants, histone H2A Leu\(^{116}\) and Leu\(^{117}\) mutations exhibited decreased H3 Lys\(^{36}\) di- and trimethylation, increased histone H4 acetylation, increased resistance to 6-azauracil, and cryptic transcription. Interestingly, the combined histone H4 Lys\(^{44}\) and H2A mutations have more severe methylation defects and increased H4 acetylation levels. Furthermore, we identify that additional histone H2A and H3 core residues are also needed for H3 Lys\(^{36}\) di- and trimethylation. Overall, our results show and suggest that multiple histone H4 and Set2 that is essential to mediate trans-histone H3 Lys\(^{36}\) di- and trimethylation is also needed so that intragenic cryptic transcription does not aberrantly occur (12, 13). The mechanism for Set2-mediated H3 Lys\(^{36}\) methylation in inhibiting intragenic cryptic transcription has been determined where H3 Lys\(^{36}\) methylation recruits the Rpd3S histone deacetylase complex by its Eaf3 and Rco1 subunits (14–16). The recruitment of Rpd3S prevents the increased in acetylation of histones H3 and H4 within the coding regions of genes so that transcription does not occur within the body of a gene (13, 16–19).

Recently, a trans-histone pathway was discovered involving Set2-mediated H3 Lys\(^{36}\) methylation (10). This study revealed that interaction between the N terminus of Set2 and histone H4 is needed for histone H3 Lys\(^{36}\) di- and trimethylation. A conserved lysine residue, Lys\(^{44}\), in histone H4 was identified to be important for interacting with the histone H4 interaction motif of Set2 (10). Deletion of this interaction motif resulted in the defects of H3 Lys\(^{36}\) di- and trimethylation. In Saccharomyces cerevisiae, histone H3 Lys\(^{36}\) methylation, catalyzed by the Set2 methyltransferase, is needed for proper transcription elongation. For example, a yeast strain lacking Set2 exhibits increased resistance to 6-azauracil (6-AU),\(^2\) a drug widely used as an indicator for transcription elongation defects (7–9). Furthermore, several studies have indicated that strains carrying histone H3 Lys\(^{36}\) mutations all exhibited increased resistance to 6-AU, suggesting that the 6-AU resistance phenotype is caused by the inability of Set2 to methylate histone H3 Lys\(^{36}\) (7, 10, 11). These findings suggest that Set2-mediated histone H3 Lys\(^{36}\) methylation plays a pivotal role in maintaining transcription elongation.

In this study, we show that the interaction between histone H4 and Set2 that is essential to mediate trans-histone H3 Lys\(^{36}\) di- and trimethylation is also necessary to repress intragenic cryptic transcription in vivo. More importantly, we determine that leucine 116 and leucine 117 residues in the C terminus of H2A, which are in close proximity to histone H4 Lys\(^{44}\), are

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\(^2\) The abbreviations used are: 6-AU, 6-azauracil; WT, wild type; HMT, histone methyltransferase; ChIP, chromatin immunoprecipitation; MOPS, 4-morpholinepropanesulfonic acid; GST, glutathione S-transferase.
required for trans-histone H3 Lys36 di- and trimethylation. Furthermore, both histone H4 and H2A mutants have increases in histone H4 acetylation, resistance to 6-AU, and intragenic cryptic transcription. In addition, the combined histone H4 Lys44 histone H4 acetylation, resistance to 6-AU, and intragenic cryptic transcription is required for H3 Lys36 di- and trimethylation and increases in histone H4 acetylation levels relative to single mutations. Interestingly, mutagenic analysis based on histone residues located within five angstroms of histone H4 Lys44 revealed that additional residues within the histone core of H2A and H3 are also needed for proper H3 Lys36 di- and trimethylation. Overall, our data suggest that histone residues from H4, H2A, and H3 likely contribute to and form a Set2 docking/recognition site on the nucleosome that is needed to maintain proper histone H3 Lys36 di- and trimethylation, histone acetylation levels, and transcription elongation.

EXPERIMENTAL PROCEDURES

Construction of Plasmids and Yeast Strains—Yeast strains used in this work are listed in supplemental Table S1. The set2Δ, ppr2Δ, and wild type (WT) strains of BY4741 were obtained from Open Biosystems. The SDBY1155 strain was generated using PCR amplification of the KanMX cassette from the BY4741 set2Δ strain as described previously (20). Histone H3 and H4 mutants in a CEN, TRP-based plasmid (pH18) were described previously (21). Histone H3 mutant plasmids were generated by site-directed mutagenesis (Stratagene) using pH18 as a template. The pHND20 construct was generated by cloning the wild type HHT2-HHF2 alleles cut from pH18 vector by Sall and Spel restriction sites into a CEN, LEU-based plasmid (pRS415). The pHND21 and pHND22 constructs were generated by site-directed mutagenesis (Stratagene) using the pHND20 as a template. Histone H2A mutant plasmids were generated in a CEN, HIS-based plasmid (pH23) by site-directed mutagenesis. All of these histone constructs were confirmed by sequencing through the coding region of histone H3, H4, or H2A and are listed in supplemental Table S2. The Set2 constructs were generated as described previously (10). All of the Set2 constructs were engineered with a single HA epitope at the C terminus and are listed in supplemental Table S3.

RNA Extraction and Northern Blot Analysis—RNA extraction and Northern blot analyses were performed as described previously (13). Briefly, yeast cells were grown to mid-log phase. Total RNA were prepared by glass bead disruption and phenol extraction. RNA samples were quantified by spectrometer, and equal amounts of RNA samples were resolved on 1% agarose formaldehyde gels running in MOPS buffer and transferred to Immobilon Nylon+ membrane (Millipore). RNA was cross-linked to the membrane by UV irradiation (Stratalinker 2400 UV cross-linker, Stratagene) and dried at 50 °C for 2 h. The membranes were prehybridized in hybridization buffer at 60 °C for 4 h. Probes (STE11 3’ nucleotides 1643–2154; SET11 5’ nucleotides 1–588; FLO8 full-length nucleotides 1–2400; and SCR1 full-length nucleotides 1–517) were PCR-amplified and labeled by random oligonucleotide priming to generate radioactive probes. The labeled probes were purified by G-25 resin column and then added into hybridization buffer to hybridize to the membrane at 60 °C overnight. After washing the membranes with 1× SSC-0.1% SDS buffer at 60 °C twice (20 min each) and at room temperature three times (5 min each), the membranes were exposed to autoradiography x-ray film or by phosphorimaging.

Yeast Extraction and Immunoblot Analysis—Yeast whole cell extracts were prepared as described previously (22). SDS-PAGE and Western blot analyses were also performed as described previously (22). Primary antibodies were used as described previously (23). The α-H3 Lys36 dimethyl-specific antibody (07-274) was obtained from Millipore and was used at a 1:1000 dilution.

GST Binding Assay—GST fusion protein GST-H2A107–131 or GST-H424–50 was expressed in Escherichia coli and purified by glutathione agarose beads. In vitro binding assays were performed as described previously (10, 23).

6-AU Growth Assay—Yeast strains expressing the histone H2A mutants were transformed with CEN, URA3-based plasmid pRS416, and 6-AU growth assays were performed as described previously (10, 23).

Histone Methyltransferase (HMT) Assay—In vitro HMT assays were performed using yeast chromatin substrates isolated from cells expressing WT histones or histone mutants in the absence or the presence of recombinant purified CBP-Set2 (2 μg), along with 2.0 μCi of S-adenosyl-l-[methyl-3H]methionine at 30 °C for 30 min in a total volume of 20 μl. The reaction mixtures were analyzed by liquid scintillation counting. Yeast chromatin substrates were prepared as described previously (10, 23).

Chromatin Immunoprecipitation (ChIP) Analysis—The ChIP assays were performed as described previously using histone H3 (Abcam, ab1791, 1 μl), H3 Lys36 dimethyl (Millipore, 07-274, 2 μl), H3 Lys36 trimethyl (Abcam, ab9050, 1 μl), and H4 acetyl (provided by Dr. David Allis, 1 μl) antibodies (24). Immunoprecipitated DNA was analyzed by quantitative real time PCR (Applied Biosystems) using TaqMan probes (25). Three biological samples with three technical repeats of each were performed. The primers and probes used for ChIP assays are listed in the supplemental information.

RESULTS

The Interaction between Histone H4 and Set2 Is Important to Maintain the Repression of Intragenic Cryptic Transcription—At present, lines of evidence demonstrate that Set2-mediated H3 Lys36 methylation is required for the repression of intragenic transcription in the 3’ ends of transcribed genes, such as STE11 and FLO8 (13, 26). Our recent studies show that the interaction between Set2 and histone H4 mediates the trans-histone H3 Lys36 di- and trimethylation and is needed for 6-AU resistance. However, whether this interaction between Set2 and histone H4 is needed to maintain proper transcriptional elongation is unknown. To determine this, intragenic cryptic transcription was examined at the STE11 or FLO8 gene by Northern blot analysis using SCR1 as a loading control (Fig. 1A and C). As expected, intragenic cryptic transcripts or short transcripts are detected in a set2Δ strain or a strain expressing a histone H3 K36R mutant when using 3’ probes to STE11 or FLO8 (Fig. 1A, lanes 2 and 3). However, short transcripts are not detected when using 5’ probes to STE11 (Fig. 1A). These
results are consistent with the previous reports in which a loss of H3 Lys\textsuperscript{36} methylation results in 3′ STE11 and FLO8 intragenic cryptic transcripts (13, 26). More importantly, the strain expressing a histone H4 K44Q mutant, which disrupts the binding with Set2 and histone H3 Lys\textsuperscript{36} dimethylation, also generates intragenic cryptic transcripts within the 3′ region of FLO8 or STE11, but not at the 5′ region of STE11 (Fig. 1A, lane 4). The strains expressing full-length Set2-HA or Set2-HA\textsubscript{Δ11–15}, which have no defects in H3 Lys\textsuperscript{36} methylation, did not exhibit short transcripts (Fig. 1, B and C, lanes 3 and 7). In contrast, we observe that the yeast strain expressing Set2-HA\textsubscript{Δ31–39}, a mutant lacking the histone H4 interaction motif, or Set2-HA\textsubscript{L117A}, a mutant lacking the Set2-Rpb1 interacting domain, a domain needed to interact with RNA polymerase II, displays defects in histone H3 Lys\textsuperscript{36} di- and trimethylation and intragenic cryptic transcription at STE11 (Fig. 1, B and C, lanes 4 and 5). In addition, a set2Δ strain expressing a Set2 mutant located on the surface of nucleosomes within or near a pocket (Fig. 2, C and D). This observation indicates that these basic and hydrophobic residues may form a binding patch or pocket that allows the histone H4 interaction motif within Set2 to bind so that subsequent trans-histone H3 Lys\textsuperscript{36} di- and trimethylation can occur.

To test these ideas, we first examined whether Leu\textsuperscript{116} and Leu\textsuperscript{117} residues within the C-terminal tail of histone H2A are required for H3 Lys\textsuperscript{36} di- and trimethylation by generating yeast strains that express various H2A mutants. Western blot analyses show that cells expressing H2A single mutation L116A, L116Q, L117A, or L117Q have significantly reduced H3 Lys\textsuperscript{36} trimethylation and intragenic cryptic transcription at STE11 (Fig. 1, B and C, lanes 6). Altogether our results indicate that the interaction between Set2 and histone H4, which is needed for trans-histone H3 Lys\textsuperscript{36} di- and trimethylation, is necessary for suppressing intragenic cryptic transcription.

Histone H2A Residues Leu\textsuperscript{116} and Leu\textsuperscript{117} Are Needed for trans-Histone H3 Lys\textsuperscript{36} Di- and Trimethylation and 6-AU Sensitivity—Our previous study found that H4 Lys\textsuperscript{44} is essential for H3 Lys\textsuperscript{36} methylation in vivo, in which the strains expressing H4 K44Q or K44E severely reduce H3 Lys\textsuperscript{36} di- and trimethylation (10). As shown in both yeast and human nucleosome structures, H4 Lys\textsuperscript{44} is located at the entry and exit point of the nucleosomal DNA, and the side chain of H4 Lys\textsuperscript{44} appears to be surrounded by the C terminus of H2A (Fig. 2A). Upon surveying the nucleosomal region surrounding histone H4 Lys\textsuperscript{44}, we determined that two H2A residues, leucines 116 and 117 of histone H2A, are in close proximity to the histone H4 Lys\textsuperscript{44} residue with a distance of approximately five angstroms (Fig. 2B). Interestingly, as viewed by a surface picture of the nucleosome core particle, H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} are containing the first 261 residues of Set2 (Set2-HA\textsubscript{1–261}), under the control of its endogenous promoter, shows decreased H3 Lys\textsuperscript{36} dimethylation, undetectable H3 Lys\textsuperscript{36} trimethylation, and intragenic cryptic transcription (Fig. 1, A, B, and C, lanes 6). Altogether our results indicate that the interaction between Set2 and histone H4, which is needed for trans-histone H3 Lys\textsuperscript{36} di- and trimethylation, is necessary for suppressing intragenic cryptic transcription.
To determine whether other residues in the C terminus of histone H2A could disrupt H3 Lys\(^{36}\) methylation, a histone H2A K119Q,K120Q double mutant was generated and expressed in yeast. Interestingly, Lys\(^{119}\) has been identified to be an H2A ubiquitination site in higher eukaryotes (27). Up to now, no evidence has been provided to show that H2A is ubiquitinated in \textit{S. cerevisiae}. Nonetheless, we still wanted to test whether these conserved lysine residues could play a role in H3 Lys\(^{36}\) methylation (28). Immunoblots show that this strain has no detectable changes in global histone H3 Lys\(^{3}\), Lys\(^{36}\) or Lys\(^{79}\) methylation, indicating that these conserved lysine residues are not essential for H3 Lys\(^{36}\) histone methylation (Fig. 3A, lane 8). These results further support that H2A Leu\(^{116}\) and Leu\(^{117}\) play an important and specific role in H3 Lys\(^{36}\) di- and trimethylation.

In our previous study, we determine by \textit{in vitro} binding assays that histone H4 Lys\(^{44}\) is the major determinant for Set2 binding. In addition, we also identified a histone H4 interaction motif located within the N terminus of Set2. Given that H2A Leu\(^{116}\) and Leu\(^{117}\) are in close proximity to H4 Lys\(^{44}\) and are required for proper H3 Lys\(^{36}\) di- and trimethylation, we wondered whether H2A Leu\(^{116}\) and Leu\(^{117}\) are also required for interaction with Set2. Therefore, \textit{in vitro} binding assays were performed using purified GST-H2A fusion protein, coding for GST-H2A\(_{107-131}\), incubated with bacterial extracts of recombinant CBP-Set2. Surprisingly, Set2 does not bind to H2A\(_{107-131}\) but binds to H4\(_{24-50}\) efficiently (Fig. 3B). Moreover, incubating GST-H2A\(_{107-131}\) with GST-H4\(_{24-50}\) does not enhance the binding of Set2 (data not shown). Although at this point it is unclear how H2A Leu\(^{116}\) and Leu\(^{117}\) contribute to histone H3 Lys\(^{36}\) di- and trimethylation, it is likely that our \textit{in vitro} binding assay is not sensitive enough to detect interactions, or these residues do not directly contribute to binding but are needed to help correctly position H4 Lys\(^{44}\).

Set2-mediated H3 Lys\(^{36}\) methylation has been demonstrated to play an important role in transcription elongation (13, 17, 18). One of the approaches used to examine this function is the 6-AU sensitivity assay (7, 8). Our previous studies have shown that strains with H3 Lys\(^{36}\) methylation defects exhibit increased resistance to 6-AU when compared with WT cells (10). To examine whether mutations of H2A Leu\(^{116}\) and Leu\(^{117}\) would result in a 6-AU resistance phenotype, cells expressing WT histones or various H2A double deletion or mutations (K119Q,K120Q; ΔL116,L117; L116A,L117A; L116Q,L117Q; and L116E,L117E) were grown on plates with or without 6-AU. As a control for 6-AU sensitivity, a \textit{ppr2Δ} strain was used as a control for 6-AU sensitivity, a \textit{ppr2Δ} strain was used as a control for 6-AU sensitivity.

\textbf{FIGURE 2.} \textit{X-ray crystal structure of the nucleosome core particle indicating the positions of histone H4 Lys\(^{36}\) and H2A Leu\(^{116}\) and Leu\(^{117}\).} A, the structure of the human nucleosome core particle is shown (H3, cyan; H4, green; H2A, gray; H2B, yellow). Histone H3 Lys\(^{36}\) and histone H4 Lys\(^{44}\) are marked in red, and histone H2A Leu\(^{116}\) and Leu\(^{117}\) are marked in blue. B, a zoomed in view of the nucleosome is shown indicating the close proximity of histone H4 Lys\(^{44}\) and histone H2A Leu\(^{116}\) and Leu\(^{117}\) residues. H4 Lys\(^{44}\) is marked in red, and H2A Leu\(^{116}\) and Leu\(^{117}\) are marked in blue. C and D, histone H4 Lys\(^{44}\) and H2A Leu\(^{116}\) and Leu\(^{117}\) form a patch on the surface of the nucleosome within or near a pocket. Histone H3 Lys\(^{36}\) and histone H4 Lys\(^{44}\) are marked in red, and histone H2A Leu\(^{116}\) and Leu\(^{117}\) are marked in blue. All of the figures were generated using Pymol.
negative control. The gene product of PPR2, also called DST1, is known as TFIIS, a known general transcription elongation factor that helps RNA polymerase II to read through blocks that occur during transcriptional elongation (29, 30). As expected, the ppr2/H9004 strain shows a sensitivity to 6-AU, whereas a set2/H9004 strain shows resistance to 6-AU when compared with WT cells (Fig. 3C, rows 1 and 2). Consistent with our previous data, we observe that expression of a H2A K119Q,K120Q double mutant, which has no changes in H3 Lys36 methylation, has WT levels of 6-AU sensitivity (Fig. 3, A, lane 8, and C, row 3). In contrast, cells expressing H2A ΔL116,L117 deletion or various H2A Leu116 and Leu117 mutants (L116A,L117A; L116E,L117E; L116K,L117K; and L116Q,L117Q), which have defects in H3 Lys36 di- and trimethylation, exhibit resistance to 6-AU, similar to the phenotype of a set2/H9004 strain (Fig. 3, A, lanes 11–15, and C, rows 5–8). Taken together, we conclude that histone H2A Leu116 and Leu117 along with H4 Lys44 are needed to mediate trans-histone H3 Lys36 di- and trimethylation and maintain proper transcription elongation.

Both H4 Lys44 and H2A Leu116 and Leu117 Are Needed to Maintain trans-Histone H3 Lys36 Methylation in Vivo and in Vitro—Because H4 Lys44 or H2A Leu116 and Leu117 mutants only affect H3 Lys36 di- and trimethylation, we asked whether combination of these residues would disrupt H3 Lys36 mono-, di-, and trimethylation, and maintain proper transcription elongation.

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ther H4 K44Q single mutant or H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} double mutants (Fig. 4A, lanes 6 and 7 versus lanes 3–5), suggesting that both H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} are needed to mediate H3 Lys\textsuperscript{36} di- and trimethylation. Again, the contribution of H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} to H3 Lys\textsuperscript{36} methylation is specific, because cells expressing the triple histone mutants do not exhibit defects in H3 Lys\textsuperscript{4} and Lys\textsuperscript{79} methylation. In addition, histone H3, H2A, and Set2 proteins levels are similar in WT and all of the histone mutants examined (Fig. 4A).

Next, we examined whether H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} are necessary for \textit{in vitro} Set2 HMT activity on yeast chromatin substrates. Soluble chromatin substrates were isolated from the nuclei of yeast strains that express WT or mutant histones (H3 K36R, H4 K44Q, H2A L116A,L117A, and H4 K44Q with H2A L116A,L117A). \textit{In vitro} HMT assays were performed using these chromatin substrates incubated with or without purified recombinant Set2. After the addition of S-adenosyl-L-[methyl\textsuperscript{3}H]methionine, total [\textsuperscript{3}H]-methyl incorporation is measured. As shown in Fig. 4B, Set2 is not active on H3 K36R substrate, whereas on a H4 K44Q substrate Set2 shows an approximately 3-fold decrease in activity when compared with WT chromatin, which is consistent with our previous report (10). In support of our \textit{in vivo} histone methylation results, when Set2 is incubated with H2A L116A,L117A chromatin substrate, a 4.5-fold decrease in HMT activity is observed (Fig. 4B). In addition, when compared with either a H4 K44Q or a H2A L116A,L117A chromatin substrate, a triple histone mutant chromatin substrate, a H4 K44Q mutation with a H2A L116A,L117A mutation, exhibits a further decrease in Set2 activity (Fig. 4B). Because H4 K44Q, H2A L116A,L117A, and the triple histone mutant show similar levels of H3 Lys\textsuperscript{36} monomethylation \textit{in vitro} when compared with the WT strain, some of the remaining Set2 \textit{in vitro} activity is likely due to H3 Lys\textsuperscript{36} monomethylation on unmodified histone H3 (Fig. 4B). Taken together, our \textit{in vivo} and \textit{in vitro} methylation data are consistent with each other and indicate that both H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} contribute to maintaining the proper H3 Lys\textsuperscript{36} di- and trimethylated states.

Both H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} Are Needed for \textit{trans}-Histone H3 Lys\textsuperscript{36} Di- and Trimethylation at Gene-specific Loci—To determine whether H3 Lys\textsuperscript{36} di- and trimethylation are affected at different coding regions at gene-specific loci, ChIP assays were performed using H3 Lys\textsuperscript{36} di- and trimethyl-specific antibodies. Relative levels of H3 Lys\textsuperscript{36} di- and trimethylation were determined by quantitative real time PCR using TaqMan primer and probe pairs specific to the 5’ and 3’ regions of \textit{STE11} or \textit{FLO8} loci. In our ChIP analysis, we determined that \textit{STE11} and \textit{FLO8} open reading frames in WT cells have H3 Lys\textsuperscript{36} di- and trimethylation present at the 5’ and 3’ regions with an enrichment of trimethylation at the 3’ ends. The detected H3 Lys\textsuperscript{36} methylation is specific because the H3 K36R mutant is used as a negative control (Fig. 5). In addition, the observed changes in histone H3 Lys\textsuperscript{36} methylation are specific and not due to changes in histone levels because all of the ChIP assays are normalized to histone H3 levels. This pattern of H3 Lys\textsuperscript{36} di- and trimethylation is also consistent with results determined by genome-wide high resolution ChIP-ChIP studies (14, 15). Interestingly, when compared with WT cells, there is an approximately 2-fold decrease of H3 Lys\textsuperscript{36} dimethylation in the H4 K44Q mutant or the H2A L116A,L117A mutant, and a 3-fold decrease in the H4 K44Q and H2A L116A,L117A triple mutant (Fig. 5A and supplemental Tables S4 and S5). In a similar manner, H3 Lys\textsuperscript{36} trimethylation is decreased 2-fold in the H4 K44Q mutant, decreased over 4-fold in the H2A L116A,L117A mutant, and decreased ~10-fold in the H4 K44Q and H2A L116A,L117A triple mutant (Fig. 5B and supplemental Tables S6 and S7). Overall, these gene-specific methylation patterns determined by ChIP analysis are in strong agreement with the global methylation patterns that we observed by immunoblot analysis. Therefore, this indicates that both H4 Lys\textsuperscript{44} and H2A Leu\textsuperscript{116} and Leu\textsuperscript{117} residues contribute to maintaining the \textit{trans}-histone H3 Lys\textsuperscript{36} di- and trimethylation pathway at gene-specific loci.

The Histone H4 and H2A Mutants with H3 Lys\textsuperscript{36} Di- and Trimethylation Defects \textit{Exhibit Increases in H4 Acetylation and Intragenic Cryptic Transcription}—Previous studies have correlated Set2-mediated H3 Lys\textsuperscript{36} methylation with Rpd3S-dependent deacetylation of chromatin (13, 17, 18). Disruption of all H3 Lys\textsuperscript{36} mono-, di-, and trimethylation levels, such as set2Δ or H3 K36A mutant, increases histone acetylation levels of
chromatin in the transcribed regions of specific loci (13, 17, 18).

In addition, our previous study has also shown that decreases in H3 Lys36 di- and trimethylation in Set2 mutants lacking the histone H4 interaction motif or the Set2-Rpb1 interacting domain of Set2 increase histone H4 acetylation (10). Therefore, we wanted to determine whether a H4 K44Q mutant or H2A L116A,L117A mutant, and the triple mutants bearing H4 K44Q and H2A L116A,L117A show an increase in histone H4 acetylation predominantly within the 3′ ends of STE11 and FLO8 (Fig. 6, A and B). Particularly, there is a 2–3-fold increase in histone H4 acetylation in the H4 K44Q and H2A L116A,L117A mutants and an over 5-fold increase in histone H4 acetylation the triple histone H4 and H2A mutant (H4 K44Q + L116A,L117A) when compared with WT (Fig. 6, A and B, and Supplemental Tables S8 and S9). Consistent with previous observations, the levels of histone H4 acetylation do not increase significantly at the 5′ ends of STE11 or FLO8 loci in all of the strains examined here, except in the H4 and H2A triple mutant (Fig. 6, A and B). The H4 and H2A triple mutant shows an approximately 2–3-fold increase in histone H4 acetylation as compared with the levels of WT cells, indicating that this mutant might play another role in regulating histone H4 acetylation independent of Rpd3S complex at the 5′ ends of transcribed genes. Intriguingly, abolishing H3 Lys36 methylation in H3 K36R mutant did not substantially increase the levels of histone H4 acetylation at the 3′ region of STE11 or FLO8 loci (Fig. 6, A and B), although this mutant did exhibit short transcripts on STE11 or FLO8 genes (Figs. 1A and 6C, lane 2). Therefore, it appears that small increases in histone H4 acetylation can contribute to cryptic transcription.

We have demonstrated that a H4 K44Q mutant with H3 Lys36 di- and trimethylation defects also generates intragenic cryptic transcripts (Figs. 1A and 4A). Next, we asked whether the strains expressing H2A mutants that have H3 Lys36 di- and trimethylation defects show a cryptic transcription phenotype. To address this issue, Northern blot analyses were performed to examine the occurrence of intragenic initiation within the STE11 gene using the H4 K44Q mutant, H2A double mutants L116A,L117A and L116Q,L117Q, and the histone H4 and H2A triple mutants H4 K44Q with H2A L116A,L117A and H4 K44Q with H2A L116Q,L117Q. We observed that all strains that have defects in H3 Lys36 di- and trimethylation and increases in histone H4 acetylation exhibit short transcripts on the 3′ region of STE11 (Fig. 6C, lanes 3–7), whereas intragenic cryptic transcription is repressed in the WT strain (Fig. 6C, lane 1). Altogether, these observations show that this trans-histone H3 Lys36 methylation pathway mediated by histone H4 Lys44 and H2A Leu116 and Leu117 is essential to maintain proper H3 Lys36 di- and trimethylation, prevent increases in histone H4 acetylation, and repress intragenic cryptic transcription.

Multiple Histone H2A and H3 Residues Located near Histone H4 Lys44 Are Needed for Proper H3 Lys36 Di- and Trimethylation—Because histone H2A residues Leu116 and Leu117 are needed for proper H3 Lys36 di- and trimethylation and are in close proximity to H3 Lys44, we analyzed the nucleosome structure for additional histone residues located within 5 angstroms of histone H4 Lys44. This analysis determined that histone H2A Ile113, His113, Gin114, Asn115, and Pro118 and H3 Leu1, Ile112, Arg2, Ile112, Asn108, and Val117 and H4 Arg39 and Arg45 are all within 5 angstroms of histone H4 Lys44 (Fig. 7A) (31, 32). In addition, many of these residues are exposed at the surface of the nucleosome with the exception of H3 Ile51 and Asn108 and H4 Arg39 (31, 32). To test whether these cis- and trans-histone residues were required for proper Set2-mediated H3 Lys36 methylation, histone H2A and H3 mutants were generated and expressed in yeast. Consistent with what has been reported, cells expressing H3 L48A, H3 I51A, H4 R39A, or H4 R45A are not viable (data not shown) (10, 33, 34). However,
A Nucleosome Surface Is Needed for H3 Lys36 Methylation

In this report, we identify new cis- and trans-histone requirements for Set2-mediated H3 Lys36 methylation. We also show that a H4 K44Q mutant and H2A C-terminal tail Leu116 and Leu117 mutants result in decreased H3 Lys36 di- and trimethylation, increased histone H4 acetylation, and resistance to 6-AU. In addition, this pattern of decreased H3 Lys36 di- and trimethylation and increased H4 acetylation observed in yeast strains expressing H4 Lys44 or H2A Leu116 and Leu117 mutants consistently show aberrant intragenic cryptic transcription. Furthermore, yeast strains expressing triple mutations consisting of histone H4 Lys44 and H2A Leu116 and Leu117 substitution mutations have lower levels of H3 Lys36 di- and trimethylation and increased levels of histone H4 acetylation than either the single H4 Lys44 mutant or the double H2A Leu116 and Leu117 mutants. Finally, we identify several additional histone H2A and H3 core residues that are also required for proper H3 Lys36 di- and trimethylation. Altogether, our results suggest that Set2 likely recognizes and docks on a H4, H2A, and H3 nucleosomal surface so that subsequent histone H3 Lys36 di- and trimethylation allows for proper maintenance of histone acetylation and transcriptional elongation.

Our recent studies have determined that Dot1 and Set2 methylate histone H3 Lys79 and Lys36, respectively, by two different trans-histone methylation pathways (10, 23, 35). In the Dot1 trans-histone methylation pathway, an electrostatic inter-

strains expressing histone H2A I112A, H2A H113A, H2A Q114A, H2A N115A, H2A P1118A, H3 R52A, H3 I112A, H3 N108A, H3 N108D, H3 V117A, or H3 V117Q mutations are viable. Whole cell extracts from viable H2A and H3 mutant strains were analyzed for their histone H3 Lys36 methylation status by immunoblot analysis (Fig. 7, B and C). Interestingly, several of the histone H2A and H3 mutants were defective in H3 Lys36 di- and trimethylation (Fig. 7, B and C). Consistent with the data shown in Fig. 3A, a strain expressing H2A I116A has significantly reduced H3 Lys36 trimethylation and only moderately affects H3 Lys36 dimethylation (Fig. 7B, lane 7). A strain expressing H2A I112A completely abolishes H3 Lys36 di- and trimethylation (Fig. 7B, lane 3), suggesting that H2A Ile112 may play a key role in mediating H3 Lys36 di- and trimethylation. In addition, strains expressing H2A H113A or H2A P1118A moderately reduce H3 Lys36 di- and trimethylation levels (Fig. 7B, lanes 4 and 8), whereas strains expressing H2A Q114A or N115A did not have noticeable changes in H3 Lys36 methylation (Fig. 7B, lanes 5 and 6). Cells expressing histone H3 R52A or H3 N108D mutant almost abolish H3 Lys36 di- and trimethylation (Fig. 7C, lanes 4 and 7). However, cells expressing a H3 I112A mutant significantly reduce H3 Lys36 trimethylation and moderately affect H3 Lys36 dimethylation (Fig. 7C, lane 8). A H3 V117A mutant strain only moderately affects H3 Lys36 trimethylation (Fig. 7C, lane 8), whereas cells expressing H3 N108A or H3 V117Q mutants do not globally change H3 Lys36 methylation (Fig. 7C, lanes 6 and 9). Again, similar to what we have previously published with other histone mutations, H3 Lys36 monomethylation is intact in all of the histone H2A and H3 mutant strains examined (Fig. 7, B and C). In addition, these histone H2A and H3 core residues are also likely to be important for maintaining proper histone acetylation and repression of cryptic transcription similar to histone H4 Lys44 and histone H2A Leu116 and Leu117. Based on our results and the nucleosome structure, histone H4, H2A, and H3 residues appear to form a region on the nucleosomal surface that is optimal for Set2 binding and/or recognition (Fig. 7D). Future studies exploring the co-crystal structure of Set2 and the nucleosome would be interesting and will help determine the precise mechanism of interaction between Set2 and this region of the nucleosome.

DISCUSSION

Our recent studies have determined that Dot1 and Set2 methylate histone H3 Lys79 and Lys36, respectively, by two different trans-histone methylation pathways (10, 23, 35). In the Dot1 trans-histone methylation pathway, an electrostatic inter-

FIGURE 7. Multiple histone H2A and H3 residues located near histone H4 Lys44 are needed for proper H3 Lys36 di- and trimethylation. A, a zoomed in view of the nucleosome is shown indicating multiple residues in the close proximity within 5 angstroms of histone H4 Lys44 residue. Representative residues that are needed for H3 Lys36 di- and trimethylation are labeled. H4 Lys44 is marked in red; H2A Ile112, Leu116, and Leu117 are marked in blue; and H3 Arg52, Ile112, Asn108, and Val117 are marked in yellow. B and C, whole cell extracts prepared from cells expressing WT histones or the indicated histone mutants were immunoblotted with H3 Lys36 methyl-specific antibodies. The asterisks denote nonspecific bands. Immunoblot for histone H3 serve as loading controls. D, the histone residues with H3 Lys36 di- and trimethylation defects have been shown to form a patch on the surface of the nucleosome. H4 Lys44 is marked in red; and H2A Ile112, Leu116, and Leu117 are marked in blue; and H3 Arg52, Ile112, and Val117 are marked in green. H3 Asn108 is not exposed at the surface of the nucleosome. These two structural figures were generated by Pymol.
A Nucleosome Surface Is Needed for H3 Lys36 Methylation

action between the C-terminal tail of Dot1 and histone H4 basic patch is need for proper H3 Lys36 di- and trimethylation (23). In a similar but distinct manner, we have identified that a histone H4 interaction motif found in the N terminus of Set2 interacts with histone H4 Lys44 to mediate H3 Lys36 di- and trimethylation (10). To further understand the Set2-mediated trans-histone pathway, we analyzed the crystal structures of human and yeast nucleosome core particles. Intriguingly, we noticed that the histone H4 Lys44 residue appeared to be surrounded by the C terminus of histone H2A, and we predicted that residues in the H2A C terminus might be needed for H3 Lys36 methylation. We tested this hypothesis and determined that histone H2A residues Leu116 and Leu117 that are in close proximity to the H4 Lys44 residue are critical for Set2-mediated methylation. We also analyzed the nucleosome structure for additional residues located within 5 angstroms of histone H4 Lys44 and found that several H2A and H3 core residues fit this criterion. More importantly, several of these histone core residues were also needed for H3 Lys36 di- and trimethylation. Overall, our studies have identified the first region on the nucleosome surface involving histone H4, H2A, and H3 residues that are required for Set2-mediated H3 Lys36 di- and trimethylation. Given that H4 Lys44 and H2A C-terminal tail mutations increase histone acetylation levels and result in intragenic cryptic transcription, we would expect that H2A and H3 core residues that disrupt H3 Lys36 methylation would also be important in maintaining proper histone acetylation and repression of cryptic transcription. In addition, the histone H3 N-terminal tail was recently determined to help mediate Set2 activity but not binding, suggesting other cis-tail determinants may exist for mediating Set2-mediated H3 Lys36 methylation (36).

Because H4 Lys44 and H2A Leu116 and Leu117 are closely located on the surface of the nucleosome, these residues likely form a docking site for the histone H4 interaction motif of Set2. Although Set2 can bind histone H4 peptides containing H4 Lys44, we were unable to detect Set2 binding to the C terminus of H2A. Therefore, our assay may not be sensitive enough to detect weak Set2 and H2A interactions, or it is quite possible that H2A residues Leu116 and Leu117 are not directly involved in binding to Set2 but are needed to help correctly position H4 Lys44 to form a binding pocket. Further structural studies will be needed to determine precisely how Set2 engages this part of the nucleosome surface and what residues are needed for direct interaction. Because all of the histone H4, H2A, and H3 mutations that disrupt H3 Lys36 di- and trimethylation do not abolish H3 Lys36 monomethylation, additional structural, genetic, and biochemical studies will be needed to understand the determinants for histone H3 Lys36 monomethylation and its function in the cell.

Previous studies have indicated that Set2-mediated H3 Lys36 methylation is needed to recruit the Rpd3S complex and deacetylate histones so that intragenic cryptic transcription within the 3′ region of genes is prevented (13, 17, 18). Consistent with these observations, we show that our previously identified trans-histone methylation pathway involving an interaction between Set2 and histone H4 Lys44 is also needed for repression of intragenic cryptic transcription. In addition, we show that histone H2A C-terminal residues are also required for repression of intragenic cryptic transcription. Recently published reports have indicated that histone H3 Lys36 dimethylation is sufficient for repressing cryptic transcription (37, 38). However, we consistently show that Set2 deletion mutants and histone H4 and H2A mutations still undergo intragenic cryptic transcription, even though these mutants still maintain some level of global and gene-specific H3 Lys36 di- and trimethylation. Therefore, we favor the idea that a combination of H3 Lys36 di- and trimethylation is needed for the proper recruitment or activity of the Rpd3S complex. This idea would also be consistent with the ability of the Rpd3S complex to equally bind H3 Lys36 di- and trimethylated nucleosomes and peptides (17, 38, 39). In addition, it has been reported that other unknown mechanisms besides histone H3 Lys36 methylation may contribute to intragenic cryptic transcription (40). Therefore, additional histone modifications and chromatin factors may help to regulate and control cryptic transcription (40). Although more work will be needed to establish what is required for cryptic transcription, further investigation is also needed to determine why there are so many cryptic promoters and whether they play a functional role in the cell.

Interestingly, during the preparation of this manuscript, a new report showed that a human homologue of yeast Set2, NSD2, exhibits severe reduced activity on H3 Lys36 methylation in vitro on recombinant nucleosome substrates where histone H4 Lys44 was mutated to Gln or Glu (41). These observations reinforce and support our previously published observation regarding Set2-mediated trans-histone pathway in budding yeast (10). Based on our new data and given that histones are highly conserved from yeast to human, it is likely that NSD2 and other Set2-like methyltransferases require this nucleosome surface for proper methyltransferase activity. Therefore, studying how histone H4, H2A, and H3 residues contribute toward maintaining proper H3 Lys36 methylation in yeast may potentially help our understanding of how proper H3 Lys36 methylation and transcription elongation occur in humans. In addition, several Set2 human homologues, NSD1, NSD2, and NSD3, have been found overexpressed or mutated in various human cancers such as acute myeloid leukemia, multiple myeloma, and breast cancer (42–44). Therefore, our studies may also provide insight into how these histone methyltransferases contribute to oncogenesis.

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