

# Overlapping Functions between *SWR1* Deletion and H3K56 Acetylation in *Candida albicans*

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**Nucleosome destabilization by histone variants and modifications has been implicated in the epigenetic regulation of gene expression, with the histone variant H2A.Z and acetylation of H3K56 (H3K56ac) being two examples. Here we find that deletion of *SWR1*, the major subunit of the *SWR1* complex depositing H2A.Z into chromatin in exchange for H2A, promotes epigenetic white-opaque switching in *Candida albicans*. We demonstrate through nucleosome mapping that *SWR1* is required for proper nucleosome positioning on the promoter of *WOR1*, the master regulator of switching, and that its effects differ in white and opaque cells. Furthermore, we find that H2A.Z is enriched adjacent to nucleosome-free regions at the *WOR1* promoter in white cells, suggesting a role in the stabilization of a repressive chromatin state. Deletion of *YNG2*, a subunit of the NuA4 H4 histone acetyltransferase (HAT) that targets *SWR1* activity through histone acetylation, produces a switching phenotype similar to that of *swr1*, and both may act downstream of the GlcNAc signaling pathway. We further uncovered a genetic interaction between *swr1* and elevated H3K56ac with the discovery that the *swr1* deletion mutant is highly sensitive to nicotinamide. Our results suggest that the interaction of H2A.Z and H3K56ac regulates epigenetic switching at the nucleosome level, as well as having global effects.**

Chromatin-level changes, such as histone modifications, variants, and nucleosome remodeling, have been implicated in the epigenetic regulation of gene expression in many systems. In recent years, focus has shifted toward the role of nucleosome dynamics in epigenetic regulation, in particular the importance of nucleosome destabilization in the establishment and inheritance of active chromatin states (1). Increasing evidence has shown that numerous histone modifications affect nucleosome stability and dynamics by altering the rate of nucleosome turnover and DNA access (2).

One such example of a histone modification altering nucleosome dynamics is the acetylation of H3 lysine 56. H3K56 acetylation (H3K56ac) occurs during S phase on newly synthesized histones and facilitates histone deposition onto replicating DNA as well as nucleosome assembly (3). Consequently, H3K56ac is found at high levels on newly synthesized chromatin and is involved in chromatin packing after DNA replication and repair in *Saccharomyces cerevisiae* (3–5). During transcriptional activation, H3K56ac promotes chromatin disassembly (6, 7). Replication-independent deposition of histones is also associated with H3K56ac (8). In addition, H3K56 is located near the DNA entry and exit points on the nucleosome core particle (9), further implicating its role in nucleosome dynamics. In yeast, the acetylation of H3K56 is performed by the histone acetyltransferase (HAT) Rtt109 (10, 11), and deacetylation occurs by the sirtuin class histone deacetylases (HDACs) Hst3 and Hst4 (12, 13).

H3K56 acetylation has been shown to regulate the epigenetic process of white-opaque switching in *Candida albicans*. A major fungal pathogen also found as a commensal with humans, *C. albicans* switches reversibly between two phenotypic states, white and opaque (14, 15). The two states differ in morphologies, transcriptional profiles, and responses to host defenses, and epigenetic switching may enhance the adaption of *C. albicans* to host environments (16–19). As in *S. cerevisiae*, H3K56 acetylation in *C. albicans* is regulated by Rtt109 and the sirtuin Hst3 (20, 21). Deletion of *RTT109* in *C. albicans*, which prevents acetylation of

H3K56, has been shown to impair white-opaque switching, while the *HST3/hst3* mutant promotes switching. Treatment with nicotinamide (NAM), which can inhibit sirtuin activity noncompetitively (22, 23) and increases H3K56ac levels, also promotes switching to the opaque form, and this activity has also been shown to be dependent on *RTT109* (24). Furthermore, hyperacetylation of H3K56 is toxic to *C. albicans* (21, 25).

In *S. cerevisiae*, high H3K56ac levels have been shown to reduce chromatin levels of the histone variant H2A.Z and increase the eviction of H2A.Z-containing nucleosomes (26). This is notable as H2A.Z itself has been implicated in altering nucleosome dynamics and stability. In addition, H3K56ac and H2A.Z have shared roles in genome stability through regulation of chromosome condensation (27, 28). H2A.Z is a highly conserved variant of H2A and is essential in a number of eukaryotic organisms but not in yeast (29–31). H2A.Z is enriched in euchromatin at the promoters of both actively expressed and repressed genes and prevents the spread of heterochromatin (32–34). Nucleosomes containing H2A.Z are enriched at the nucleosome-free regions (NFRs) at promoter regions, in particular those adjacent to the +1/–1 nucleosomes at the transcriptional start site of genes (32, 35). H2A.Z-containing nucleosomes have been noted to have altered stability; however, separate studies have shown H2A.Z nucleo-

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TABLE 1 Strains used in this study

Strain	Description	Genotype	Reference
JYC1	WT	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG</i>	68
HLY4235	<i>SWR1/swr1</i>	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG SWR1/swr1::HIS1</i>	This study
HLY4233	<i>swr1</i>	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG swr1::HIS1/swr1::ARG4</i>	This study
HLY3555	WT + pWOR1-GFP	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG ADH1/adh1::pWOR1-GFP-URA3</i>	47
HLY4234	<i>swr1</i> + pWOR1-GFP	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG swr1::HIS1/swr1::ARG4 ADH1/adh1::pWOR1-GFP-URA3</i>	This study
HLC54	<i>efg1</i>	<i>MTLa/a ura3::1 imm434/ura3::1 imm434 efg1::hisG/efg1::hisG</i>	69
HLY4238	WT + <i>HTA3-HA</i>	<i>MTLa/a ura3::imm434/ura3::imm434 HTA3/3×HA-HTA3-SAT1 ADH1/adh1::pMAL2-WOR1-URA3</i>	This study
HLY4236	<i>swr1</i> + <i>HTA3-HA</i>	<i>MTLa/a ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG swr1::HIS1/swr1::ARG4 ADH1/adh1::pWOR1-GFP-URA3 HTA3/3'HA-HTA3-SAT1 ADH1/adh1::pMAL2-WOR1-URA3</i>	This study
HLY3993	<i>HST3/hst3</i>	<i>MTLa/a ade2/ade2 ura3::ADE2/ura3::ADE2 HST3/hst3::FRT</i>	24
HLY3997	<i>rtt109</i>	<i>MTLa/a rtt109::FRT/rtt109::FRT</i>	24
CLY4	<i>yng2</i>	<i>ura3::1 imm434/ura3::1 imm434 yng2::hisG/yng2::hisG</i>	51
HLY3883	<i>MTLa/a yng2</i>	<i>MTLa/a ura3::1 imm434/ura3::1 imm434 yng2::hisG/yng2::hisG</i>	This study
HLY4035	<i>yng2</i> + <i>YNG2</i>	<i>ura3::1 imm434/ura3::1 imm434 yng2::hisG/YNG2-13MYC-URA3</i>	52
HLY3886	<i>MTLa/a yng2</i> + <i>YNG2</i>	<i>MTLa/a ura3::1 imm434/ura3::1 imm434 yng2::hisG/YNG2-13MYC-URA3</i>	This study

some to be either more or less stable than those with H2A (36, 37). Of particular note is the finding that nucleosomes containing both H3.3, the variant of H3 which is the canonical H3 in yeast, and H2A.Z are more labile (38). In multicellular organisms, H2A.Z has been associated with cellular differentiation and reprogramming (39). H2A.Z is deposited by the Swi/Snf family chromatin remodeling complex SWR1 through the ATP-dependent exchange of H2A.Z-H2B dimers with H2A-H2B (40, 41). *SWR1* encodes the major subunit of the SWR1 complex. Neither the *H2A.Z* nor the *SWR1* gene is essential in yeast; however, *SWR1* activity causes genetic instability in *H2A.Z*-deleted cells (42). The acetylation of canonical histone H2 and H4 tails by the NuA4 HAT complex, in which Yng2 is a subunit of the HAT module, has been shown to recruit the SWR1 complex to deposit H2A.Z (43). In yeast, the NuA4 and SWR1 complexes share four subunits, whereas the two complexes function together as one in higher eukaryotes (44, 45). Recently, NFRs have been shown to be dominant over histone acetylation in targeting SWR1 activity (46). The localization and effect on nucleosomal dynamics of H2A.Z make it a promising subject for the study of epigenetic regulation.

In this study, we find that deposition of H2A.Z by the SWR1 complex is crucial in regulating the white-opaque transition in *C. albicans*. Abrogation of H2A.Z deposition through deletion of *SWR1* results in a phenotype that favors switching to the opaque state and the stability of the opaque phase. We find that nucleosome dynamics on the promoter of *WOR1*, the master regulator of switching (47–49), differ between white and opaque phases and that deletion of *SWR1* changes this dynamic. Furthermore, H2A.Z is present on the *WOR1* promoter at higher levels in the white phase than in the opaque phase, suggesting that its absence destabilizes white cells. We find that deletion of *YNG2* produces a white-opaque phenotype similar to that of the *swr1* deletion mutant. Furthermore, we find that the *swr1* deletion mutant displays a synthetic lethality with nicotinamide but no heightened sensitivity to DNA damage, suggesting a close functional relationship between H2A.Z and H3K56ac in *C. albicans*.

## MATERIALS AND METHODS

**Strains and culturing conditions.** Strains used in this study are listed in Table 1. Strains were maintained on YEP (1% yeast extract, 2% peptone) or SC (synthetic complete) medium with the appropriate carbon sources.

**Strain construction.** *SWR1* was deleted by using the deletion cassettes described previously by Noble and Johnson (50). PCR amplification of *Candida dubliniensis* *HIS1* from plasmid pSN52 or of *Candida dubliniensis* *ARG4* from plasmid pSN69 was performed by using long primers containing sequences flanking *SWR1*, such that the PCR product was flanked by 90-bp sequences homologous to flanking sequences of *SWR1*. The PCR product was then transformed into wild-type (WT) (JYC1) *C. albicans* to replace *SWR1* by homologous recombination. Transformants were selected for survival on SC dextrose histidine or arginine dropout media, and deletion of *SWR1* was verified by PCR.

H2A.Z (*HTA3*) was hemagglutinin (HA) tagged by PCR amplification of the *HTA3* coding sequence with a 3×HA tag at the 5' end of the sequence, with HindIII and PstI sites introduced at the 5' and 3' ends of the amplicon, respectively. The *HTA3* promoter was amplified in a separate reaction, flanked by XhoI and HindIII sites. Subsequently, the HA-*HTA3* and *HTA3* promoter sequences were inserted into the pWOR1-GFP-SAT1 plasmid (24), replacing the *GFP* and *WOR1* promoter sequences, respectively. The plasmid was digested with MscI prior to transformation into the WT or *swr1* strain. All primers used for cloning are listed in Table 2.

**Mating type conversion.** The *MTLa/a yng2* deletion mutant (CLY4) (51) and the *yng2*+*YNG2* strain (HLY4035) (52) were converted to *MTLa/a* by growth on solid SC medium with 2% sorbose at a density of 10<sup>5</sup> cells per plate for 6 days (53). The mating type locus was verified by PCR.

**Switching assays.** Stable white or opaque cells from log-phase cultures were spread onto agarose plates of Lee's medium (modified from reference 54) supplemented with 25 μg/ml uridine and 50 μg/ml arginine, with 1.25% dextrose or *N*-acetyl-D-glucosamine (Spectrum Chemical) as the carbon source. For CO<sub>2</sub> assays, plates were incubated in a Thermo Forma series II water-jacketed CO<sub>2</sub> incubator at 25°C with 5% CO<sub>2</sub>. Colony morphology was scored after 7 days. Both whole-colony and sectored switching events were counted.

**Reverse transcription-quantitative PCR (RT-qPCR).** RNA was isolated from log-phase cultures by using the Qiagen RNeasy kit. cDNA was synthesized from 2 μg total RNA by using the Bio-Rad iScript reverse

TABLE 2 Primers used in this study

Primer	Sequence	Description
SWR1D-F	CGATTTATTTAATTACAATTCGGAAATTAATCTCATCTTGCAGATCCAGC AAACACATCAAGCATATTCTCACAAGAAAACCAGAAA TGCCAGAGAATGGAACCAGTGTGATGGATATCTGC	SWR1 deletion
SWR1D-R	AGACTTATCCTTGATGTCTGTACTAGTACCAACGATACGCAAGAGATTTTT ATAAAAAGAAAGGGCCATTTGCTTAATATGTAATACA AGGAGTGATACCATATAGCTCGGATCCACTAGTAACG	SWR1 deletion
SWR1outsideF	TGTTTACTACTGTAATTTTCTGC	Check integration of deletion cassette at SWR1 locus
SWR1outsideR	ACTATACGATATGGTATCCGT	Check integration of deletion cassette at SWR1 locus
C.dubHIS-F	AACACAACCTGCACAATCTGG	<i>C. dubliniensis</i> HIS1 check
C.dubHIS-R	ATTAGATACGTTGGTGGTTC	<i>C. dubliniensis</i> HIS1 check
C.dubARG-F	ACGGAGTACCACATACGATG	<i>C. dubliniensis</i> ARG4 check
C.dubARG-R	ACACAGAGATACCTTGTACT	<i>C. dubliniensis</i> ARG4 check
SWR1internalF	CTAATTCCAATATCCACCAAGC	SWR1 internal sequence check, 963 bp from 3' end
SWR1internalR	GACCTTGTGATATAACTTTCC	SWR1 internal, 1,134 bp from 5' end of coding sequence
HTA3_F3	CCGCTCGAGTGTGTCAAAGTTCCTACAAAG	Amplification of HTA3 promoter
HTA3_R3	CCCAAGCTTTGCATAGTCCGGGACGTCATAGGGATAGCCCGCATAGTCAGG AACATCGTATGGGTACATTGTAGTGTTTGTTGTGGTTTCCG	Amplification of HTA3 promoter
HTA3_F4	CCCAAGCTTGATCCTATCCATATGACGTTCCAGATTACGCTGCTATGTCTGG GAAGGAAAAGTGC	HA tagging of HTA3
HTA3_R4	GCAACTGCAGACCGCCACTAAATAGAGAGC	HA tagging of HTA3
MTLa-F	TGAAGCGTGAGAGGCAGGAG	MTLa check
MTLa-R	AATTCCTTCTCTTCGATTAGG	MTLa check
MTL $\alpha$ -F	TTTCAGTACATTCTGGTTCGGC	MTL $\alpha$ check
MTL $\alpha$ -R	TGTAACATCCTCAATTGTACCCGA	MTL $\alpha$ check
WOR1-F	AAATAAACCATCGTCATCGGC	WOR1 qPCR
WOR1-R	AGGACCATTACCTAGACTCG	WOR1 qPCR
ACT1-F	TGGAAGCTGCTGGTATTGAC	ACT1 qPCR
ACT1-R	TTCAGCAATACCTGGGAACA	ACT1 qPCR

transcription kit. Quantitative PCR (qPCR) was performed on the Bio-Rad iCycler instrument by using Bio-Rad SYBR green mix. Cycle parameters were 95°C for 1 min and then 39 cycles of 95°C for 10 s, 56°C for 45 s, and 68°C for 20 s. Oligonucleotides used for amplification are listed in Table 2.

**Nucleosome mapping.** Micrococcal nuclease (MNase) digestion was performed with modifications of methods described previously (55, 56). Cells were grown to an optical density at 600 nm ( $OD_{600}$ ) of 2 and then cross-linked with 2% formaldehyde at a growth temperature for 30 min before treatment with zymolyase to spheroplast cells. Spheroplasts were then treated with a titration of 2 to 8 units of MNase, and digested samples were visualized by electrophoresis on a 2% agarose gel. Samples with distinct mononucleosome and dinucleosome band patterns were selected for further analysis. DNA from these samples was isolated by using phenol-chloroform extraction and used for qPCR for nucleosome mapping. *Staphylococcus aureus* MNase was acquired from Sigma-Aldrich. A full list of the primers used for mapping of the *WOR1* promoter is available in Table S1 in the supplemental material. Chromatin immunoprecipitation was performed on mononucleosomal DNA as described previously (57). Anti-HA antibody (Abcam) was used for immunoprecipitation of HA-tagged H2A.Z. Nucleosomal values were normalized to values for the –1 nucleosome peak at the *BUD2* promoter.

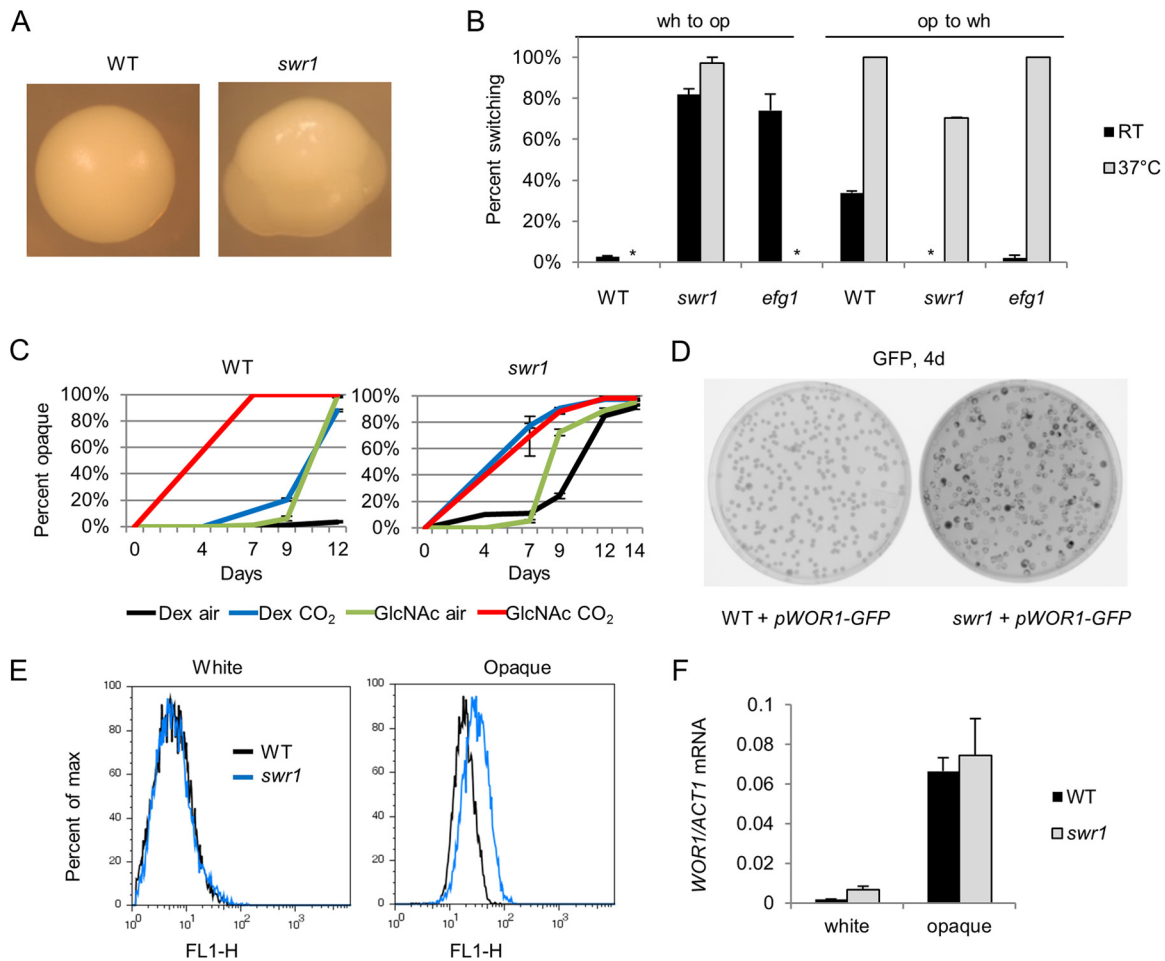
**Spot assays.** *C. albicans* strains were grown in liquid yeast extract-peptone-dextrose (YPD) medium to log phase and diluted to an  $OD_{600}$  of 0.1, after which 5 additional 5-fold serial dilutions were made into double-distilled water ( $ddH_2O$ ) in a 96-well plate. An 8-channel multipipettor was used to transfer 2.5  $\mu$ l of each dilution onto YPD agarose plates. For drug treatment plates, nicotinamide and methyl methanesulfonate

(MMS) were obtained from Sigma-Aldrich. Plates were incubated for 3 days at 30°C, and colony growth was imaged with a Fujifilm LAS-4000 imager.

**Fluorescence-activated cell sorting.** Fluorescence-activated cell sorting (FACS) was conducted by using the BD FACSCalibur system, using cells taken from actively growing liquid cultures. The FL1-H channel was used for green fluorescent protein (GFP) detection, and 10,000 cells in suspension were counted per sample. Results were analyzed with FlowJo cytometry analysis software.

## RESULTS

**The *swr1* deletion mutant increases white-to-opaque switching frequency and stabilizes the opaque phase.** To investigate the role of chromatin remodeling by the SWR1 complex in white-to-opaque switching, we deleted the *SWR1* gene in *C. albicans* using homologous recombination to replace it with auxotrophic markers as described previously by Noble and Johnson (50). In the resulting *swr1* deletion mutant, chromatin immunoprecipitation confirmed that the deposition of H2A.Z into chromatin was abrogated (not shown). High rates of spontaneous switching to the opaque phase in white colonies of the *swr1* mutant were observed at room temperature (RT), often occurring as multiple opaque sectors per white colony (Fig. 1A). The white-to-opaque switching percentage of the *swr1* mutant at room temperature (25°C) was 82%, compared to 2 to 4% for the wild type. Opaque cells of the *swr1* mutant are more stable than those of the wild type, with 0% switching to white after 7



**FIG 1** Deletion of *SWR1* leads to elevated white-opaque switching and opaque stability. (A) Single colonies of white WT (JYC1) and *swr1* (HLY4233) cells after 7 days of growth at room temperature (25°C) on YPD. The *swr1* colony displays multiple opaque sectors typical of the strain. (B) Spontaneous white-to-opaque and opaque-to-white switching rates of the WT, *swr1*, and *efg1* (HLY3600) strains at RT (25°C) and at 37°C after 7 days of growth on SCD medium. The *P* value is <0.01 for *swr1* opaque cells at 37°C compared to the wild type, and the *P* value is <0.005 for all others comparisons to the wild type, based on two-tailed Student's *t* test. \* indicates that 0% switching occurred. (C) Percentage of spontaneous switching to opaque from white cells of the WT and *swr1* strains after 7 days of growth at RT on Lee's medium with 1.25% dextrose (Dex) or 1.25% GlcNac as the carbon source, in air or 5% CO<sub>2</sub>. (D) Fluorescence image of colonies of the WT carrying pWOR1-GFP (HLY3555) and the *swr1* mutant carrying pWOR1-GFP (HLY4234) after 4 days of growth on SCD medium at RT. Opaque colonies and sectors, expressing more GFP, appear darker. (E) FACS profiles of GFP fluorescence in a population of 10,000 white or opaque cells of the WT carrying pWOR1-GFP (HLY3555) and the *swr1* mutant carrying pWOR1-GFP (HLY4234). Cells were from a log-phase liquid culture at RT. (F) Levels of *WOR1* mRNA transcript as assayed by RT-qPCR in white and opaque cells of the WT (JYC1) or the *swr1* (HLY4233) deletion mutant. RNA was collected from log-phase cell cultures in YPD at room temperature. Transcript levels were normalized to *ACT1* levels. All experiments were performed in triplicate.

days of growth at RT, compared to 34% for the WT (Fig. 1B). We next sought to determine the effect of temperature on the *swr1* white-opaque phenotype. At 37°C, white-opaque switching of the *swr1* mutant increased to 97%, while WT cells no longer switched. We also compared the phenotype of the *swr1* mutant to that of the *efg1* mutant, the deletion mutant of the transcriptional regulator *EFG1* which is known to have high rates of white-opaque switching. Unlike the *swr1* mutant, the *efg1* mutant was not able to switch to opaque at high temperatures. At 37°C, opaque stability of the *swr1* mutant was reduced but still greater than that of the WT, with 30% of opaque cells remaining opaque, while all WT opaque cells switched to white. On the other hand, *efg1* opaque cells were less stable than *swr1* cells and switched to white cells readily at 37°C (Fig. 1B). To test if *SWR1* can restore function to the *swr1* mutant, we assessed the white-opaque phenotype of the *swr1*/Tet-*SWR1* mutant, a conditional mutant in which *SWR1* expression is shut down in the presence of doxycycline

(58). When grown on medium containing doxycycline, the strain displayed high levels of white-to-opaque switching and opaque stability, similar to the *swr1* deletion mutant. In the absence of doxycycline, the strain did not readily switch to opaque and also had low opaque stability at 37°C (see Fig. S1 in the supplemental material). This demonstrates that the expression of *SWR1* can restore the wild-type phenotype to an *swr1* mutant. From these results, we find that white-opaque switching and opaque stability are enhanced in the *swr1* mutant even at high temperature.

We proceeded to investigate whether the deletion of *SWR1* promotes switching downstream of known pathways. It has been documented that low concentrations of CO<sub>2</sub> as well as utilization of *N*-acetylglucosamine (GlcNac) as the main carbon source promote white-opaque switching, and these two pathways show a synergistic effect with each other (59, 60). To assay whether Swr1 acts downstream of either of these pathways, we cultured white

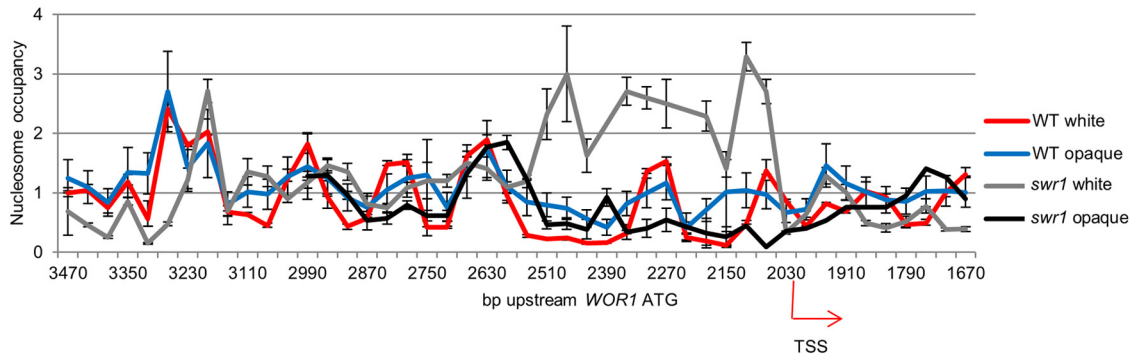
cells of the WT and the *swr1* mutant in air or 5% CO<sub>2</sub> on Lee's medium with dextrose or GlcNAc as the carbon source and tracked switching to opaque over the course of 14 days. In the WT, as previously documented, both CO<sub>2</sub> and GlcNAc promoted white-opaque switching and showed synergy with each other. In the *swr1* mutant, we observed a higher level of spontaneous switching than in the WT, even under control conditions of growth in air on dextrose. Notably, switching of the *swr1* mutant was elevated in the presence of CO<sub>2</sub> regardless of the carbon source, showing a synergy similar to that of GlcNAc and CO<sub>2</sub> (Fig. 1C). Unlike the *swr1* mutant, the *efg1* mutant readily switched to opaque without GlcNAc or CO<sub>2</sub> with a timing similar to that of the wild type in the presence of both GlcNAc and CO<sub>2</sub> (data not shown). Thus, *SWR1* deletion appears to drive white-opaque switching in a pathway synergistic with the Flo8-mediated CO<sub>2</sub>-sensing pathway and may act downstream of the GlcNAc signaling pathway. Although wild-type opaque cells switch to white at 37°C under most conditions, it has been noted that GlcNAc as a carbon source increases switching in a manner synergistic with higher temperatures as well (60). This is consistent with the observation of white-to-opaque switching at high temperature in the *swr1* mutant.

We then explored the mechanisms behind the white-opaque phenotype of the *swr1* mutant through regulation of *WOR1*, the master regulator of switching. First, we investigated whether the *swr1* mutant had a higher level of expression from the *WOR1* promoter by using WT and *swr1* strains transformed with the p*WOR1*-GFP plasmid, which expresses *GFP* from the native *WOR1* promoter. After 4 days of growth on solid SCD medium, starting from white cells, the *swr1* mutant showed numerous instances of darker colonies and sectors than those of the WT, indicative of the expression of p*WOR1*-GFP in cells that had become opaque (Fig. 1D). This further supports elevated white-to-opaque switching in *swr1*. To determine *WOR1* expression levels in white and opaque cells, fluorescence-activated cell sorting (FACS) was performed on WT and *swr1* cells carrying p*WOR1*-GFP. As seen in fluorescence histograms, transcription from the *WOR1* promoter in white cells was not greater for the *swr1* mutant than for the WT. In opaque cells, the level of GFP in the *swr1* mutant was slightly elevated compared to that in the WT (Fig. 1E). Because the basal levels of *WOR1* expression in white cells are extremely low, it could be difficult to detect a slight change in *Wor1* levels by using FACS assays. Given the high white-to-opaque switching frequency in the *swr1* mutant, it may at least be easier to reach the threshold to initiate the positive self-feedback loop of *Wor1* in the mutant than in the WT. Indeed, RT-qPCR analysis of *WOR1* mRNA levels in WT and *swr1* white and opaque cells detected a slightly elevated *WOR1* expression level in white *swr1* cells compared to wild-type cells (Fig. 1F). This difference was not detected by FACS analysis. The *WOR1* level in opaque *swr1* cells was also slightly higher than that in opaque WT cells, consistent with the FACS data. While *WOR1* expression was not greatly elevated in the *swr1* mutant, a moderate increase may make it easier to reach the threshold to initiate the positive self-feedback loop of *Wor1* and promote white-opaque switching or opaque stability through positive feedback.

**Nucleosome positioning on the *WOR1* promoter differs between white and opaque cells and is altered in the *swr1* mutant.** We next sought to investigate whether deletion of *SWR1* altered

the white-opaque phenotype by changing the regulation of *WOR1* through chromatin remodeling at the *WOR1* promoter. As *WOR1* has a long 8-kb promoter as well as a long 5' untranslated region (UTR), nucleosomal dynamics upstream of *WOR1* may be crucial to the regulation of its transcription. Since H2A.Z-containing nucleosomes have been shown to have altered stability and dynamics (36–38, 41), we proceeded with a nucleosome map of the *WOR1* promoter in WT and *swr1* cells, both of which carry HA-tagged *HTA3* (H2A.Z), to facilitate further studies. To obtain nucleosomal DNA, *Staphylococcus aureus* micrococcal nuclease digestion was performed in order to cleave non-nucleosome-bound DNA while leaving nucleosome-bound DNA intact. qPCR was then performed on the nucleosomal DNA, amplifying staggered 100-bp regions along the *WOR1* promoter and providing a map of nucleosomal enrichment in this region. We found that even in the WT, white and opaque cells showed a difference in nucleosome occupancy and positioning along the *WOR1* promoter. While regular nucleosome peaks were observed at similar positions in both cell types, notably, nucleosome peaks were more defined in white cells. In particular, this was true of the +1/–1 nucleosomes adjacent to the transcriptional start site –1,997 bp upstream of the *WOR1* ATG start codon. In white cells, there is a nucleosome-free region (NFR) between the +1 and –1 nucleosomes as well as one ~2,550 to 2,350 bp upstream of the *WOR1* ATG codon, absent in opaque cells. In the *swr1* mutant, we saw a marked change in nucleosome positions on the *WOR1* promoter compared to those in the WT. Nucleosome positioning was altered from that of the wild type, particularly in white cells, and the NFRs were abolished, with nucleosome occupancy being highly elevated in the region spanning bp –2550 to –2350 instead. The highly elevated levels of nucleosome occupancy at NFRs of the *WOR1* promoter in white *swr1* cells likely reflect slower nucleosome turnover, which is consistent with the known function of H2A.Z. In contrast, nucleosome occupancy at the NFRs of the *WOR1* promoter is much lower in opaque *swr1* cells than in wild-type white or opaque cells (Fig. 2). Overall, nucleosome mapping suggests that *Swr1* is required for proper nucleosome positioning and stability at the *WOR1* promoter and for the differential regulation of promoter nucleosomes in white and opaque states. Our data therefore support the importance of H2A.Z deposition in epigenetic regulation of transcription in cell fate determination.

**H2A.Z is enriched at the *WOR1* promoter in white cells.** H2A.Z is known to be enriched at developmentally regulated regions. To further investigate nucleosomal H2A.Z content, we mapped H2A.Z enrichment along the *WOR1* promoter in white and opaque WT cells. In addition, *swr1* cells were used as a control. H2A.Z was HA tagged in all strains used. H2A.Z enrichment was assayed by immunoprecipitation of H2A.Z-containing nucleosomes using anti-HA- on micrococcal nuclease-treated nucleosomes (Fig. 2), followed by qPCR using the same *WOR1* primers as those used for nucleosome mapping to provide coverage along the *WOR1* promoter. As expected, H2A.Z levels on promoter chromatin were very low in the *swr1* mutant. In the wild type, notably, the H2A.Z level was higher in white cells than in opaque cells. Particularly of interest is that H2A.Z was enriched in the –2 and –1 nucleosomes at two NFRs on the *WOR1* promoter (Fig. 3). The location of H2A.Z-enriched nucleosomes at NFRs is consistent with the role of H2A.Z in promoting nucleosome dynamics, as shown in yeast and high eukaryotes (61). The



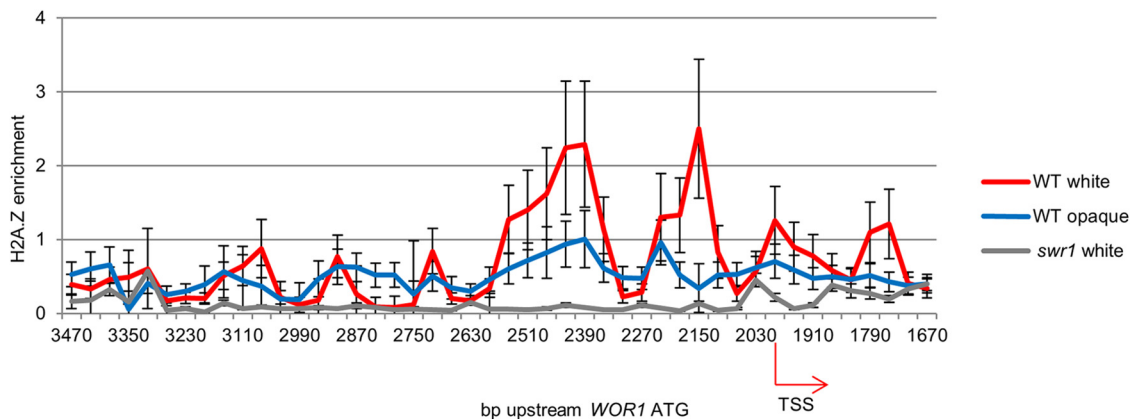
**FIG 2** Nucleosome map of the *WOR1* promoter shows white-opaque differences in the wild type and altered nucleosome positioning and occupancy in the *swr1* mutant. Nucleosomal DNA was isolated from the WT strain carrying *HTA3*-HA (HLY4238) or the *swr1* mutant carrying *HTA3*-HA (HLY4236) after removal of non-nucleosome-bound DNA with micrococcal nuclease digestion. Mapping was performed by qPCR using primers amplifying 100-bp regions along the *WOR1* promoter. The error was calculated from the standard deviation of  $C_T$  values from triplicate experiments. TSS, transcriptional start site.

elevated H2A.Z deposition by the SWR1 complex at the *WOR1* promoter in white cells indicates a role for H2A.Z in stabilizing the repressive chromatin state in white cells, potentially by maintaining NFRs and permitting the binding of repressive factors.

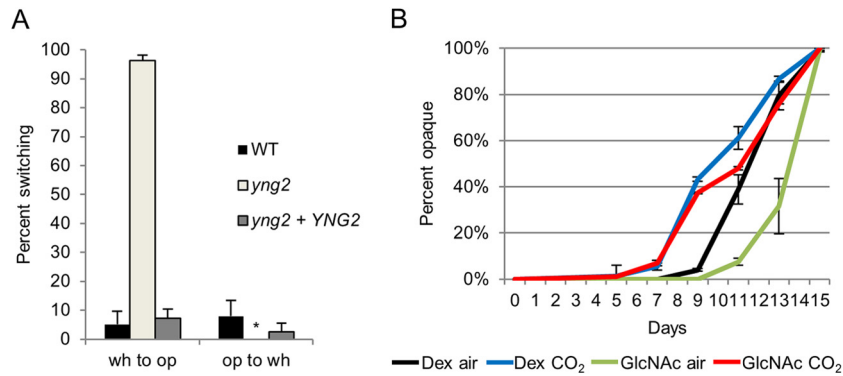
**A deletion mutant of *yng2*, a subunit of the NuA4 complex, produces a white-opaque switching phenotype and synergy with CO<sub>2</sub> similar to those of the *swr1* mutant.** In exploring the regulation of H2A.Z deposition, we considered the recruitment of the SWR1 complex by NuA4 acetylation of H2 and H4 and the substantial overlap in subunits between the SWR1 and NuA4 complexes. Yng2 is a subunit of the NuA4 HAT module, the deletion of which, as we have shown, severely impairs H4 acetylation (51). Thus, we investigated whether deletion of *YNG2* would produce a white-opaque phenotype similar to that of the *swr1* mutant. We found that at room temperature, white cells of the *yng2* mutant switched to opaque at a very high frequency compared to that of the wild type, while opaque cells of the *yng2* mutant were highly stable. A complemented *yng2*+*YNG2* strain showed a phenotype similar to that of the wild type (Fig. 4A). In addition, opaque *yng2* cells were stable at 30°C but not at 37°C (not shown). Because of the relation between NuA4 acetylation of histones and SWR1 de-

position of H2A.Z in the same nucleosome, we speculated that the *yng2* and *swr1* mutants produced their high-switching phenotype through the same pathways and would respond similarly to CO<sub>2</sub> and GlcNAc. To test this, white cells of the *yng2* mutant were grown at room temperature in air or 5% CO<sub>2</sub>, with dextrose or GlcNAc as a carbon source, and switching was assessed. We found that *yng2* cells began switching to opaque sooner when incubated in CO<sub>2</sub>, preceding switching in air by several days. This behavior is similar to that of the *swr1* mutant. Notably, the *yng2* mutant switched more slowly when the carbon source was GlcNAc than when grown on dextrose, showing a more complete lack of synergy with GlcNAc than the *swr1* mutant did (Fig. 4B). Therefore, like SWR1, Yng2/NuA4 also functions downstream of the GlcNAc signaling pathway, separate from the CO<sub>2</sub>-regulated pathway. This is consistent with the known function of NuA4 in histone acetylation, which in turn allows SWR1 recruitment and H2A.Z deposition.

**The *swr1* mutant is highly sensitive to nicotinamide.** In addition to the epigenetic regulation of white-opaque switching, we also sought to investigate the effects of *swr1* deletion on additional cellular functions. H2A.Z is enriched in nucleosomes surrounding transcriptional start sites and is involved in gene expression (62).



**FIG 3** Differential enrichment of H2A.Z at the *WOR1* promoter in white and opaque cells. Chromatin immunoprecipitation of HA-tagged H2A.Z was performed on nucleosomal DNA isolated from white and opaque cells of the WT strain carrying *HTA3*-HA and from white cells of the *swr1* strain carrying *HTA3*-HA. qPCR was performed by using primers used to map the *WOR1* promoter. Results were normalized to the total nucleosome level. TSS, transcriptional start site.

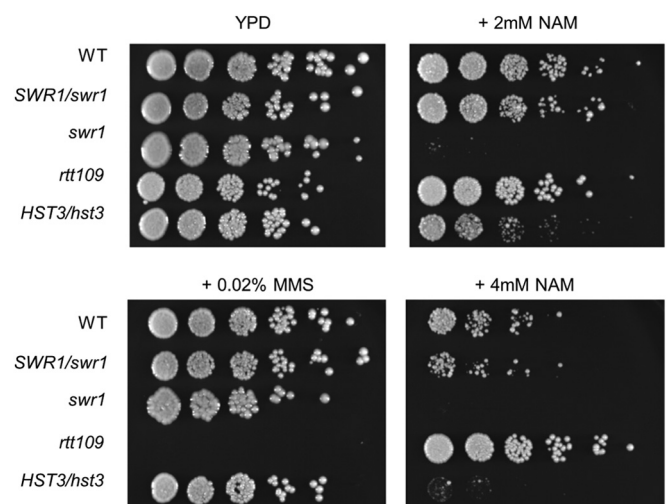


**FIG 4** White-opaque switching of the *yng2* mutant is elevated and shows synergy with CO<sub>2</sub>. (A) Spontaneous white-to-opaque and opaque-to-white switching rates of the *yng2* strain (HLY3883) and the *yng2*+*YNG2* complemented strain (HLY3886) compared to the WT after 14 days at room temperature on SCD medium. The *P* value is <0.005 for the *yng2* and *yng2*+*YNG2* strains compared to the WT. \* indicates that 0% switching occurred. (B) Percentage of spontaneous switching of white *yng2* cells to opaque after 7 days of growth on Lee's medium with 1.25% dextrose or 1.25% GlcNAc as the carbon source, in air or 5% CO<sub>2</sub>, at RT.

Proper deposition of H2A.Z has been linked to numerous global effects, such as euchromatin-heterochromatin boundary establishment, centromeric functions, and DNA repair (33, 63–65). Recent work in *S. cerevisiae* has demonstrated that heightened acetylation of H3K56 is associated with reduced deposition and increased eviction of H2A.Z from chromatin by SWR1, with significant overlap in gene expression profiles between cells with constitutively acetylated H3K56 and *htz1* (H2A.Z) deletion (26). It is known that nicotinamide (NAM) inhibits *C. albicans* growth by an elevation of H3K56ac levels through inhibition of the HDAC Hst3 (21, 24). To further explore the functional relationship between H3K56ac and H2A.Z deposition, we examined the sensitivities of the WT, *SWR1/swr1*, *swr1*, *HST3/hst3*, and *rtt109* strains to NAM by a spot assay for growth on solid medium in the presence of nicotinamide. Cells were grown for 3 days at 30°C. The *HST3/hst3* strain was highly sensitive to NAM and barely grew with 4 mM NAM, while the *rtt109* strain was more resistant to NAM at 4 mM than the WT, consistent with previously reported results (24). Interestingly, the *swr1* strain was even more sensitive to NAM than the *HST3/hst3* strain, barely growing even on 2 mM NAM (Fig. 5). Unlike the *HST3/hst3* strain, NAM sensitivity was not observed for the *SWR1/swr1* strain. The lack of haploid insufficiency indicates that NAM probably does not directly act on Swr1. In contrast to NAM sensitivity, the *swr1* and *HST3/hst3* strains did not show any increased sensitivity to the DNA-damaging agent methyl methanesulfonate (MMS), while the *rtt109* strain was sensitive (Fig. 5). In addition, the *swr1* mutant showed no increased growth sensitivity to hydroxyurea or heat and osmotic stress (not shown). We have previously shown that the *rtt109* mutant has an increased rate of generating aneuploidy (25). However, we did not observe any increase in aneuploidy formation in the *swr1* mutant after growth on L-sorbose medium (not shown). Therefore, *rtt109* (or blocking of H3K56 acetylation) is associated with aneuploidy and sensitivity to DNA damage, while H3K56ac and *swr1* are associated with sensitivity to NAM. From our genetic studies, we suggest that the NAM sensitivity of *swr1* and H3K56ac is associated with their roles in gene expression.

## DISCUSSION

The role of histone variants and modifications on nucleosomal stability is a promising angle for the investigation of epigenetic regulation. In particular, the histone variant H2A.Z as well as the acetylation of H3K56 have been linked to changes in nucleosomal dynamics. Previous studies in our laboratory have demonstrated the effect of H3K56 acetylation on epigenetic regulation of white-opaque switching in *C. albicans* (24, 25). In this study, we show that epigenetic switching is also regulated by chromatin deposition of H2A.Z by SWR1. Both chromatin modifications affect the switching frequency and stability of cell fates but do not dramatically affect transcriptional levels of key cell fate regulators. A small increase in *WOR1* expression was observed in the *swr1* mutant, which may facilitate reaching the threshold for positive self-feedback regulation of Wor1. Additionally, we find synergy between the *swr1* deletion and H3K56ac in *C. albicans*.



**FIG 5** The *swr1* mutant displays high sensitivity to nicotinamide but not to methyl methanesulfonate. Show data from spot assays of the WT, *SWR1/swr1* (HLY4235), *swr1* (HLY4233), *HST3/hst3* (HLY3993), and *rtt109* (HLY3997) strains on YPD, YPD plus 2% MMS, or YPD plus 2 mM or 4 mM NAM. Fivefold serial dilutions from cultures with an OD<sub>600</sub> of 0.1 were made on agar medium plates and incubated for 3 days at 30°C prior to imaging.

H2A.Z enrichment at promoter regions is often associated with both active and inactive transcription (32–34). Many studies have found H2A.Z nucleosomes enriched on the promoters of repressed genes, and H2A.Z deposition is important for proper induction of gene expression. Our study finds an example of these properties in demonstrating that H2A.Z is enriched at the *WOR1* promoter in white cells and that white-to-opaque switching is enhanced in an *swr1* mutant. Thus, H2A.Z is repressive to *WOR1* expression or the opaque state. How does H2A.Z exert both positive and negative effects on transcription? One attractive model is that H2A.Z facilitates the binding of either activating or repressive complexes by keeping regions of promoters accessible (61). We find that nucleosome occupancy at the *WOR1* promoter in white cells is elevated when *SWR1* is deleted. Of particular note, a high H2A.Z level was found at nucleosome-free regions on the *WOR1* promoter in WT cells, and in the *swr1* mutant, these regions were no longer NFRs and instead had very high nucleosome occupancy. The existence of these NFRs in the white but not opaque state in the WT suggests a role for them in the repression of *WOR1* transcription. Chromatin immunoprecipitation with microarray technology (ChIP-chip) studies of *C. albicans* have demonstrated that Efg1, a repressor of *WOR1* and the opaque state, binds at the *WOR1* promoter in white cells at only one major peak in the region between  $-3$  kb and  $-2$  kb, while it binds at multiple regions in opaque cells (66). This finding, in conjunction with our results, suggests that H2A.Z deposition in white cells maintains NFRs that facilitate the binding of repressive factors such as Efg1, favoring a repressive state of *WOR1* transcription. Deletion of *SWR1* may abolish these regions, preventing repressor binding and leading to a more permissive chromatin state for *WOR1* transcription and an increased frequency of white-to-opaque switching. Future work is needed to examine this model of reduced accessibility of the *WOR1* promoter to negative regulators of white-to-opaque switching in the *swr1* mutant.

In exploring the functional relation of Swr1 with known white-to-opaque signaling pathways, we also find that the high-switching phenotype of the *swr1* mutant showed synergy with CO<sub>2</sub> in promoting white-to-opaque switching. This synergy with CO<sub>2</sub> is similar to that of GlcNAc with CO<sub>2</sub>. Furthermore, the *swr1* mutant shows no additional synergy with GlcNAc. Based on these data, we suggest that the SWR1 complex functions in the GlcNAc signaling pathway. A similar high-switching phenotype and synergy with CO<sub>2</sub> are seen in the deletion mutant of *yng2*, a subunit of the NuA4 complex that shares subunits with the SWR1 complex and recruits it through histone H2 and H4 acetylation. Thus, it is likely that reduced histone acetylation in the *yng2* mutant in turn reduces SWR1 deposition of H2A.Z, producing a phenotype similar to that of the *swr1* mutant. Interestingly, *swr1* opaque cells are more stable at high temperature than are those of the *yng2* mutant, showing that there is some difference between the regulatory effects of the two. It has also been noted that the opaque-state-promoting effects of GlcNAc are synergistic with high temperature (56). The chromatin-level changes during temperature-induced opaque-to-white switching are a subject for further exploration. Our genetic data indicate that the GlcNAc signaling in white-to-opaque switching is likely to regulate *WOR1* promoter chromatin in white cells by downregulating histone acetylation and H2A.Z deposition. Notably, the switching phenotype of the *swr1* mutant differs from that of numerous other histone modifier mutants previously described to affect white-to-opaque switching.

The *swr1* mutant produces a higher switching frequency than many such mutants, such as those of the H3K4 methyltransferase *set1*; however, it does not produce complete switching as the *efg1 set1* double mutant does (67). We have also noted that *set1* shows no synergy with GlcNAc or CO<sub>2</sub> (data not shown). Thus, the effects of histone modifications on white-to-opaque switching are manifold, and their interactions with other pathways that regulate switching leave much to be explored.

A recent study has shown that a high level of H3K56ac reduces nucleosome incorporation of H2A.Z (26). Not only is H2A.Z deposition greatly reduced on H3K56ac-containing nucleosomes, the presence of nucleosomal H3K56ac also alters the activity of SWR1, increasing eviction of H2A.Z from nucleosomes. While nucleosomes containing both H3K56ac and H2A.Z are not less stable than those with H2A.Z alone, this increased exchange by H3K56ac can promote nucleosome turnover in H2A.Z-rich regions (26). Whether H2A.Z deposition can in turn promote the turnover of H3K56ac nucleosomes is an attractive prospect. We find in this study that the *swr1* deletion mutant is highly sensitive to NAM. As hyperacetylation of H3K56 in chromatin is toxic to *C. albicans*, the high sensitivity of the *swr1* mutant to NAM could be due to reduced nucleosome turnover and, consequently, the accumulation of H3K56ac on chromatin. In correlation, H2A.Z is enriched at the *WOR1* promoter in white cells, while H3K56 acetylation is enriched in opaque cells (24). Future research is needed to examine if the reduced nucleosome occupancy at the *WOR1* promoter in opaque *swr1* cells is linked to the H3K56ac level. Genome-wide nucleosome mapping should determine whether NAM-treated *swr1* cells show reduced nucleosome occupation at known H2A.Z-enriched regions. The interaction between H2A.Z deposition and H3K56ac may also exist in multicellular organisms, with implications for epigenetic regulation in developmental processes.

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