SUMO functions in constitutive transcription and during activation of inducible genes in yeast

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Transcription factors represent one of the largest groups of proteins regulated by SUMO (small ubiquitin-like modifier) modification, and their sumoylation is usually associated with transcriptional repression. To investigate whether sumoylation plays a general role in regulating transcription in yeast, we determined the occupancy of sumoylated proteins at a variety of genes by chromatin immunoprecipitation (ChIP) using an antibody that recognizes the yeast SUMO peptide. Surprisingly, we detected sumoylated proteins at all constitutively transcribed genes tested but not at repressed genes. Ubc9, the SUMO conjugation enzyme, was not present on these genes, but its inactivation reduced SUMO at the constitutive promoters and modestly decreased RNA polymerase II levels. In contrast, activation of the inducible GAL1, STL1, and ARG1 genes caused not only a striking accumulation of SUMO at all three promoter regions, but also recruitment of Ubc9, indicating that gene activation involves sumoylation of promoter-bound factors. However, Ubc9 inactivation, while reducing sumoylation at the induced promoters, paradoxically resulted in increased transcription. Providing an explanation for this, the reduced sumoylation impaired the cell's ability to appropriately shut off transcription of the induced ARG1 gene, indicating that SUMO can facilitate transcriptional silencing. Our findings thus establish unexpected roles for sumoylation in both constitutive and activated transcription, and provide a novel mechanism for regulating gene expression.

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Protein sumoylation involves the covalent and reversible attachment of the small ubiquitin-like modifier (SUMO) peptide to lysine side chains in acceptor proteins. Mechanistically similar to ubiquitylation, this involves a cascade of enzymatic activities, including activation by an E1 enzyme [Aos1/Uba2] and conjugation to target proteins by a sole E2 enzyme [Ubc9], guided to appropriate substrates by the SUMO E3 ligases (Johnson 2004; Geiss-Friedlander and Melchior 2007). Budding yeast express a single SUMO isoform, encoded by the essential SMT3 gene, whereas three main isoforms are found in mammalian cells: SUMO1, and the highly similar SUMO2 and SUMO3. At the molecular level, attachment of the SUMO peptide to a substrate can promote its association with interacting proteins through recognition of its SUMO-modified form, or the SUMO moiety can interfere with protein–protein interactions by blocking interaction sites. The consequences of altered protein–protein interactions through sumoylation are diverse, and include changes in subcellular localization, protein activity, and protein stability (Geiss-Friedlander and Melchior 2007).

SUMO orthologs have been identified in all eukaryotic species, and sumoylation modifies proteins involved in a wide range of cellular processes, indicating that regulation by sumoylation is widespread (Zhao 2007; Makhnevych et al. 2009). Gene expression, however, appears to be particularly regulated by sumoylation, because a large number of known SUMO conjugates in yeast and mammals are transcription factors (Gill 2005; Zhao 2007; Makhnevych et al. 2009). Blocking sumoylation of gene-specific transcription factors C/EBP [Kim et al. 2002], c-Jun [Muller et al. 2000], ELK-1 [Yang et al. 2003], and many others (Girdwood et al. 2004; Gill 2005) through mutation of SUMO acceptor sites results in increased transcription of target genes. Consequently, sumoylation is generally associated with transcriptional repression. This is supported by other findings, including observations that transcriptional corepressors, including histone deacetylase complexes [HDACs], preferentially associate with sumoylated forms of transcription factors (Garcia-Dominguez and Reyes 2009; Ouyang and Gill
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2009). For example, the coactivator p300 associates with HDAC6 in a SUMO-dependent manner (Girdwood et al. 2003). Additionally, human histone H4 is sumoylated (Shio and Eisenman 2003), as are all four core histones in yeast [Nathan et al. 2006]. Through mutations that attenuate sumoylation, or by the use of histone-SUMO fusion proteins, it was shown that histone sumoylation represses transcription [Nathan et al. 2006]. This is thought to occur through the recruitment of HDACs by SUMO-modified histones, or by interference with transcription-promoting histone modifications such as acetylation or ubiquitylation (Shio and Eisenman 2003; Nathan et al. 2006). However, it is not known whether histone sumoylation is a general mechanism of repression at transcriptionally silent genes. Despite the large number of studies linking SUMO with repression, in a few cases, sumoylation of gene-specific transcription factors is associated with activating transcription [Lyst and Stancheva 2007; Guo and Sharrocks 2009 and references therein], indicating that SUMO does not have a solely repressive role in transcription.

In addition to gene-specific transcription factors, large-scale proteomics screens identified components of the general transcription machinery as SUMO targets in both yeast [Panse et al. 2004; Wohlschlegel et al. 2004; Zhou et al. 2004; Denison et al. 2005; Hannich et al. 2005; Wykoff and O’Shea 2005; Makhnevych et al. 2009] and mammalian [Zhao et al. 2004; Rosas-Acosta et al. 2005] cells. These include multiple subunits of the general transcription factors (GTFs) TFIIA, TFIIF, and TFIIID (TFIIB, as well as several TAFs); Mediator; and subunits of RNA polymerase II (RNAP II) itself. Of the multiple yeast subunits of RNAP II found to be sumoylated, Rpb1 sumoylation was characterized and appears to occur as part of the UV response, but blocking Rpb1 sumoylation does not affect transcription elongation or cell growth [Chen et al. 2009]. Sumoylation of the human TFIIID subunit TAF5 was shown to impair TFIIID binding to promoter DNA [Boyer-Guittaut et al. 2005], suggesting that sumoylation might be a negative regulator of TFIIID as well.

The finding that multiple components of the general transcription machinery are modified by SUMO suggests that, in addition to regulating several gene-specific transcription factors, sumoylation may help control transcription at many genes. Here, we provide evidence that this is indeed the case. We first asked, using chromatin immunoprecipitation (ChIP) assays, whether SUMO is associated with a panel of genes in Saccharomyces cerevisiae, in normally growing cells or under conditions where transcription of certain genes is induced. In apparent contrast to the fact that SUMO is strongly associated with transcriptional repression, we found that SUMO is present at all transcriptionally active genes tested, including both constitutive and inducible promoters, but not at repressed or uninduced genes. We did not detect Ubc9 at constitutive genes, implying that factors are sumoylated prior to association with these genes. In contrast, we show that Ubc9 is recruited to promoters of inducible genes when they are activated. Interestingly, however, impairing Ubc9 function results in increased transcription of these genes. Providing an explanation for this, we found that reducing sumoylation at the induced ARG1 promoter impairs the ability of the cell to shut off its transcription when the activating signal disappears. Our results indicate that sumoylation occurs at promoters of active genes during induced transcription, and functions to regulate transcription levels by facilitating shutting off transcription.

Results

SUMO associates with transcriptionally active genes

Considering the preponderance of studies linking sumoylation of transcription factors with transcriptional repression, we wished to establish whether sumoylation has a general role in repressing transcription at promoters of silenced genes. To this end, we determined whether sumoylated proteins could be detected at a panel of repressed promoters in yeast cells by ChIP using an antibody that recognizes the yeast SUMO peptide Smt3. Chromatin was prepared from cells grown in standard conditions, and samples were immunoprecipitated with the Smt3 antibody [SUMO] or an antibody recognizing the largest subunit of RNAP II as a control [Fig. 1A]. Immunoprecipitated samples were analyzed by PCR for the presence of promoter DNA relative to a nontranscribed region of Chromosome V [Fold Occupancy in Fig. 1B; see the Materials and Methods]. Five inducible genes that are repressed during normal growth but activated under different conditions were analyzed: GAL1, STL1, ARG1, CUP1, and HSP12. Presumably as controls, we also tested three constitutively transcribed genes and an intergenic region of Chromosome VII, representing a transcriptionally silent region. Under normal growth conditions, RNAP II ChIP confirmed that RNAP II was associated with only the transcriptionally active genes [constitutively transcribed PYK1, ADH1, and PMA1], and not the repressed genes or the intergenic region [RNAP II] [Fig. 1A,B]. Surprisingly, however, SUMO ChIP showed that significant levels of SUMO were present on only the three constitutively transcribed gene promoters [Fig. 1B]. We did not detect significant levels of SUMO at the intergenic region, or most notably at any of the five repressed gene promoters [Fig. 1B].

The Smt3 antibody used in our ChIP experiments was described previously [Montpetit et al. 2006], and, when used in a Western blot using yeast whole-cell extract, gave the expected pattern of multiple sumoylated proteins [see Fig. 2A; cf. Wohlschlegel et al. 2004]. However, to confirm that the signal detected by ChIP was indeed due to the presence of sumoylated proteins, we repeated the ChIP experiments using a strain expressing an 8xHIS epitope-tagged version of Smt3 [Wohlschlegel et al. 2004]. As shown in Figure 1C, ChIP using an anti-HIS antibody in the HIS8-SMT3 strain resulted in the same pattern observed with the Smt3 antibody, confirming that the ChIP signal was in fact due to the presence of SUMO at transcriptionally active genes. Low levels of HIS8-Smt3 at the ARG1 promoter likely reflect weak activation of this
gene detected in uninduced cells (Govind et al. 2005). These observations are in contrast to what was expected based on models postulating that SUMO is inherently repressive to transcription (Chupreta et al. 2005), and instead indicate that, in normally growing yeast, SUMO is present at the promoter regions of active genes.

To determine whether SUMO detected at constitutively transcribed genes was restricted to promoter positions, a quantitative ChIP analysis was performed using specific primers corresponding to conserved promoter regions of constitutively active genes. The results showed that SUMO occupancy was significantly higher at these positions compared to silent gene promoters or intergenic regions. This finding suggests that SUMO plays a role in the regulation of transcription in yeast.

Figure 1. SUMO associates with constitutively active genes. [A] Representative ChIP analysis of indicated transcriptionally active and silent gene promoters, and a nontranscribed region of Chromosome VII (Intergenic). Occupancy of SUMO and RNAP II was determined by ChIP with an Smt3 antibody (SUMO) or an antibody recognizing RNAP II (antibody 8WG16; RNAP II). Control primers recognizing sequences in an untranscribed region of Chromosome V were included in the PCR reactions (*), and input chromatin was also analyzed by PCR for comparison and quantification (INPUT). [B] Quantification of ChIP analyses as shown in A. Quantification (described in the Materials and Methods) is fold over background; a value of 1 indicates no signal detected above background, as indicated with a heavy bar. [C] Gene diagrams for PYK1, ADH1, and PMA1 indicating gene length, approximate position of transcriptional start site (bent arrow), and regions amplified by indicated ChIP primers. Lengths of genes and amplification products are to scale with respect to each other. [D] Quantification of detailed ChIP analysis of SUMO occupancy at indicated positions across indicated constitutive genes. [E, left] Cell lysates from UBC9-6HA and a control strain were analyzed by Western blot with anti-HA antibody to confirm tagging of Ubc9. [Right] Occupancy of 6xHA-tagged Ubc9 at promoters of indicated genes in the UBC9-6HA strain, determined by ChIP.

Figure 2. Reduced sumoylation in ubc9-1 cells correlates with reduced RNAP II occupancy at constitutive genes. [A] Whole-cell yeast extracts were prepared from ubc9-1 cells and isogenic wild-type cells, and were analyzed by Western blot with Smt3 antibody (SUMO) and an antibody recognizing the large subunit of RNAP II (Rpb1; antibody 8WG16) as a loading control. Cells were grown at 28°C, or shifted to 37°C for 30 min prior to analysis as indicated. [B] Spot assay comparing growth of ubc9-1 cells and an isogenic wild-type strain on complete minimal medium at the indicated temperature. Approximately 1.5 x 10^4 cells were plated in the first spot on the left, with fivefold dilutions in each subsequent spot toward the right. Growth was for 2 d. All ChIP experiments using ubc9-1 and the isogenic control were performed using cells grown at 28°C. [C] Occupancy of SUMO and RNAP II was determined at the indicated positions (see Fig. 1D) across PYK1, ADH1, and PMA1 in ubc9-1 cells [gray bars] and the isogenic wild-type strain [black bars]. Statistical analysis (Student’s t-test) indicates that SUMO levels and RNAP II levels are significantly different in the wild-type and ubc9-1 data sets (P = 0.004 for SUMO, and P = 0.003 for RNAP II). [D] Semiquantitative RT–PCR analysis of steady-state RNA levels of indicated genes in ubc9-1 cells [gray bars] and in isogenic wild-type cells [black bars]. RNA levels are not significantly different in the wild-type and ubc9-1 data sets (P > 0.1).
regions, we performed more detailed ChIP analyses. Primer pairs representing positions across the PYK1, ADH1, and PMA1 genes were used to determine the level of SUMO cross-linking at promoter, internal, and terminator regions, as indicated in the gene diagrams shown in Figure 1D. Low levels of sumoylated proteins were detected at all positions across the genes, but higher levels were detected around promoters (about twofold to threefold higher than at other regions). This indicates that proteins associated with transcriptionally active genes, particularly those associated with promoters, are sumoylated.

The presence of SUMO at transcriptionally active genes indicates that either "presumoylated" proteins assemble specifically at active genes, or sumoylation of assembled factors takes place at active genes. To distinguish between these possibilities, we determined whether Ubc9 could also be detected at the active genes tested. We constructed a yeast strain expressing 6xHA C-terminal-tagged Ubc9 (Fig. 1F, left) and performed ChIP analysis. As shown in Figure 1F (right), significant levels of Ubc9-6HA were not detected at the PYK1, ADH1, or PMA1 promoter regions, even though Ubc9-6HA was detected at other genes using this strain (see Fig. 3A). Although it is possible that low levels of Ubc9 transiently associate with these genes to catalyze the sumoylation that we detect, our data are consistent with recruitment of proteins that are sumoylated before they associate with the gene.

Lowering SUMO levels at constitutive genes results in reduced RNAP II levels

The presence of sumoylated factors specifically on transcriptionally active genes suggests that SUMO plays a positive role in transcription by RNAP II. To determine whether SUMO affects transcription of these genes, we analyzed the effect of reduced sumoylation on levels of RNAP II on the genes. Both SMT3 and UBC9 are essential, preventing the use of SMT3- or UBC9-null strains. Instead, we used a mutant yeast strain, *ubc9-1*, that expresses a less stable form of the enzyme (Seufert et al. 1995; Betting and Seufert 1996). At normal growth temperatures (28°C–30°C), *ubc9-1* cells grew essentially as well as isogenic wild-type cells, but growth was severely impaired at elevated temperatures (37°C) [Fig. 2B]. However, examination of whole-cell extract by Western blot showed significantly reduced levels of total sumoylation in *ubc9-1* cells compared with wild-type cells at 28°C, while growth at 37°C further exacerbated this effect [Fig. 2A].

To assay the effects of reduced sumoylation on transcription, we compared wild-type and *ubc9-1* cells grown at 28°C, avoiding the possibility of growth-related indirect effects. First, we measured levels of SUMO at positions across the three constitutive genes in wild-type and *ubc9-1* cells. Overall, somewhat less SUMO was detected on the constitutive genes in this strain background (DF5) than in the strain used in our initial analysis (YPH499), but the pattern of SUMO occupancy across the genes was the same (cf. Figs. 1B,E and 2C, top). As expected, the reduction in global sumoylation in *ubc9-1* cells resulted in reduced SUMO levels at the tested genes [Fig. 2C, top]. RNAP II ChIP analysis showed an overall reduction in RNAP II levels across the constitutive genes in *ubc9-1* cells [Fig. 2C, bottom]. Approximately 30% less RNAP II was detected at all positions across the PMA1 gene in *ubc9-1* cells compared with wild-type cells, whereas most of the reduction in the PYK1 and ADH1 genes occurred at promoter-proximal regions, with only a modest reduction detected near the 3’ end. An analysis performed after switching wild-type and *ubc9-1* cells to 37°C for 30 min resulted in similar reductions in RNAP II densities in *ubc9-1* cells [data not shown]. These results indicate that sumoylation is required for efficient recruitment of RNAP II to the promoters of the constitutive genes.
genes tested. However, this did not significantly affect steady-state transcript levels for these genes [Fig. 2D]. This observation is not surprising, considering that ubc9-1 cells grow normally at the permissive temperature, and suggests that the primary function of SUMO at the promoters of constitutive genes likely is not in regulating overall expression levels [see the Discussion].

SUMO and Ubc9 are both recruited to promoters of induced genes during activation

To explore further the function of SUMO in transcription, we next examined whether SUMO associates with activated genes. Three of the inducible genes that did not associate with SUMO when repressed [Fig. 1A,B] were examined in their induced states. GAL1, induced when galactose is the carbon source in growth medium, STL1, an osmoregulated gene induced by the addition of NaCl to the medium, and ARG1, an amino acid biosynthesis gene induced by amino acid starvation conditions, were examined by ChIP. RNAP II ChIP at the GAL1, STL1, and ARG1 promoters showed robust recruitment of RNAP II to the genes under their respective activating conditions [Fig. 3A, left]. Strikingly, analysis of SUMO levels showed the appearance of sumoylated proteins coincident with induction of the genes [Fig. 3A, middle], at levels approximately twice as great as at the constitutive genes tested [cf. Figs. 1B and 3A; data not shown]. The appearance of SUMO at the three inducible genes, under different respective induction conditions, strongly reinforces our assertion that SUMO associates with active sites of transcription in yeast.

We next wished to investigate whether the high SUMO levels detected at the induced genes might reflect active sumoylation at the promoters. To this end, we examined association of Ubc9 with these promoters under induced and uninduced conditions. As the above experiments were conducted in the UBC9-6HA strain, we performed ChIP analysis using an HA antibody. In contrast to the constitutive genes analyzed above [Fig. 1F], significant levels of Ubc9-6HA were present on each of the induced genes during activation [Fig. 3A, right]. Detailed ChIP analysis at positions across GAL1, STL1, and ARG1 demonstrated that the strong SUMO signal detected upon induction was restricted to promoter-proximal regions of the genes [Figs. 3C, 4A,B]. Similar analysis of Ubc9-6HA recruitment along the GAL1 and STL1 genes indicated that Ubc9-6HA was localized primarily to promoter-proximal regions as well [Fig. 3C, right; note that GAL1 promoter elements are shared with the upstream coregulated GAL10 gene]. The concurrent appearance of SUMO and Ubc9 at induced promoters strongly suggests that promoter-bound factors are sumoylated during gene activation. Thus, while SUMO associates with constitutively expressed genes, likely due to recruitment of sumoylated factors, Ubc9-mediated SUMO modification of promoter-bound factors occurs during activation of inducible genes. This represents a previously unappreciated aspect of gene activation in yeast.

Reduced sumoylation at induced promoters correlates with increased transcription

We next wished to determine the effect of impaired sumoylation on activation of inducible genes. To this end, we compared RNAP II recruitment to the STL1 and ARG1 promoters under their respective induction conditions in wild-type and ubc9-1 cells at 28°C. SUMO ChIP indicated that reduced global sumoylation [see Fig. 2A] resulted in less SUMO associated with the promoters of the induced STL1 and ARG1 genes [Fig. 4A,B, left]. Unexpectedly, and in contrast to what we observed at

**Figure 4.** Sumoylation has a negative effect on transcription of inducible genes. (A, B) Occupancy of SUMO [left] and RNAP II [middle] was determined in uninduced and induced conditions [indicated; as in Fig. 3A] for STL1 [A] and ARG1 [B] in ubc9-1 cells [black bars] and isogenic wild-type cells [gray bars]. (Right) Steady-state STL1 [A] and ARG1 [B] mRNA abundance in wild-type and ubc9-1 cells was determined by RT-PCR in untreated or induced cells, as indicated. Values are shown relative to transcript abundance in wild-type cells after induction, which is set to 100. Statistical analysis indicates that RNAP II levels are significantly different in the wild-type and ubc9-1 data sets [P < 0.03 for STL1 data, and P < 0.008 for ARG1 data]. RNA levels were also significantly different in wild-type and ubc9-1 cells, with P-values of 0.02 and 0.03 for STL1 and ARG1, respectively. (C) Growth comparison of wild-type and ubc9-1 cells in 0, 0.4, and 0.6 M NaCl in rich medium [left] or minimal medium lacking Val and Ile, and the same medium supplemented with 0.5 μg/mL SM [right].
constitutive genes, for both STL1 and ARG1, reduced SUMO at promoters correlated with small but consistent increases of RNAP II levels at the promoters and across the genes (Fig. 4A,B, middle). These higher levels of RNAP II resulted in increased steady-state transcript levels [by ~40%] for both genes, as shown by RT–PCR (Figs. 4A,B, right). These results suggest that, although sumoylation is detected at inducible promoters only after activation, it has a negative effect on RNAP II recruitment and, consequently, transcription.

Next, we asked whether growth of ubc9-1 cells (which is normal at 28°C) [see Fig. 2B], is affected under conditions where gene induction is necessary. Induction of osmoreponse genes such as STL1 is essential for growth in the presence of NaCl (Rep et al. 2000). Whereas isogenic wild-type cells showed growth defects at 0.4 and 0.6 M NaCl, ubc9-1 cells grew relatively well across this range of NaCl concentrations (Fig. 4C, left). Under amino acid starvation conditions [for example, in the presence of the drug sulfoemeturon [SM], which inhibits expression of Val and Ile] (Falco and Dumas 1985), induction of amino acid biosynthesis genes such as ARG1 is critical. In this case, reduced sumoylation in the ubc9-1 strain resulted in reduced growth compared with isogenic wild-type cells (Fig. 4C, right). Although reduced sumoylation likely affects multiple systems involved in the response to changes in growth conditions, including exposure to NaCl and amino acid starvation, it is reasonable to speculate that it deregulated, increased transcription of response genes can affect cell growth. For example, increased expression of osmoreponse genes can facilitate sustained growth in the presence of NaCl, whereas extended expression of amino acid biosynthesis genes beyond their need can be energetically costly and result in reduced growth.

**Sumoylation is required for complete deactivation of induced ARG1**

The increased RNAP II densities and elevated transcription of induced genes that we observed in ubc9-1 cells could result in two distinct ways: from increased recruitment of the polymerase to activated promoters, or from prolonged activation of induced genes after the inducing signal is no longer present. In light of our finding that sumoylation has a positive effect on RNAP II recruitment to constitutive genes, we hypothesized that the negative effect of sumoylation on inducible transcription is due to a role in terminating activation of induced promoters, a process that we refer to as deactivation. To investigate whether sumoylation indeed functions in deactivation, we devised a method by which activation of the ARG1 gene could be very rapidly stopped. Upon exposure to SM, depletion of Val and Ile leads to efficient translation and stabilization of the Gcn4 activator protein, which binds the ARG1 promoter and activates its transcription [Qiu et al. 2004, 2005; Govind et al. 2005]. The short half-life of Gcn4 (~5 min under normal growth conditions) [Kornitzer et al. 1994] permits rapid response to changes in amino acid levels. We found that adding a high concentration of Val and Ile [five times their normal concentration, referred to as “stop mix”] to SM-treated ubc9-1 or isogenic wild-type cells resulted in the very rapid disappearance of Gcn4 [no signal in a Western blot 5 min after adding the stop mix] (Fig. 5A), and, as described below, in the complete disappearance of RNAP II from the ARG1 promoter in wild-type cells.

We used the strategy just described to determine whether reduced sumoylation in ubc9-1 cells affects deactivation of ARG1. To this end, wild-type and ubc9-1 cells were treated with SM for 25 min, then treated with the stop mix for 5 min, or simply treated with SM for 25 min with no further treatment. Analysis of RNAP II occupancy at the ARG1 promoter in wild-type chromatin samples showed that the stop mix was highly effective in deactivating ARG1; RNAP II was no longer detectable on the gene 5 min after its addition (Fig. 5B, RNAP II ChIP).
SUMO ChIP with wild-type chromatin indicated that SUMO levels were also greatly reduced at the ARGI promoter upon its deactivation (Fig. 5B, SUMO ChIP). Thus, SUMO associates with ARGI only while it is transcriptionally active. In fact, examination of the ARGI promoter upon activation and deactivation showed that Ubc9, SUMO, and RNAP II show similar patterns of recruitment to the promoter during activation, and all were efficiently cleared from the promoter upon deactivation [Supplemental Fig. S1]. Strikingly, analysis of RNAP II occupancy before and after the stop mix shows that clearance of RNAP II from the ARGI promoter was impaired in ubc9-1 cells compared with wild-type cells. Significant levels of RNAP II remained at the ARGI promoter in ubc9-1 cells 5 min after addition of the stop mix, whereas deactivation was complete at this point in wild-type cells [Fig. 5B, right]. We also examined whether clearance of Gcn4 from the ARGI promoter was affected by reduced global sumoylation in cells expressing Flag-tagged Gcn4. Indeed, as shown in Supplemental Figure S2, significant levels of Gcn4-Flag remained associated with the ARGI promoter in ubc9-1 cells compared with wild-type cells 5 min after addition of the stop mix. This is a strong indication that sumoylation at inducible gene promoters plays an important role in deactivation of transcription by facilitating clearance of promoter-bound factors.

Prolonged activation of genes in ubc9-1 cells after elimination of the activating signal would be expected to result in prolonged transcription and delayed clearance of transcripts. To determine if transcription was prolonged in ubc9-1 cells compared with wild-type cells, we measured steady-state levels of ARGI (which has an approximate half-life of 18–21 min) [Crabbe et al. 1990] by RT–PCR at 20-min intervals post-addition of the stop mix [Fig. 5C, quantification of three experiments is shown at right]. Decay of the ARGI mRNA signal in wild-type cells was as expected, with approximately half as much signal appearing after each 20-min period. In sharp contrast, in ubc9-1 cells, no reduction in ARGI mRNA levels was observed within the first 20 min after adding the stop mix [Fig. 5C]. In fact, a slight increase was detected, possibly reflecting accumulation of the RNA while transcription continued for a short time before deactivation. In the second 20-min interval, after which deactivation was likely completed, decay was normal, with approximately half as much RNA being detected, indicating that the ARGI mRNA half-life was not affected by the ubc9-1 mutation. Taken together with the analysis of RNAP II occupancy, these results indicate that deactivation of ARGI transcription is impaired in ubc9-1 cells, reflecting reduced sumoylation at the promoter.

Discussion

Previous studies have supported the notion that SUMO plays an inherently repressive role in transcription. For example, targeting SUMO isoforms or Ubc9 in human cells to a luciferase reporter gene [through fusions to the Gal4 DNA-binding domain] caused potent reduction of luciferase activity [Holmstrom et al. 2003; Shiio and Eisenman 2003; Chupreta et al. 2005]. In fact, a specific surface on the SUMO moiety, present in all human SUMO isoforms as well as in yeast Smt3, was identified that confers repressive effects in the luciferase reporter assay [Chupreta et al. 2005]. It is surprising, then, that our data shows that SUMO marks transcriptionally active genes in yeast, both constitutive genes and inducible genes, specifically, when they are activated. Furthermore, since reducing levels of SUMO at the constitutive genes resulted in somewhat lower RNAP II density, efficient sumoylation is important for optimal transcription. Thus, while SUMO is able to repress transcription in some contexts, in the natural context of constitutive transcription, other properties of SUMO must come into play, perhaps in the assembly or organization of the transcription machinery [Fig. 6A]. For example, sumoylation plays a role in the appropriate assembly of the human polyadenylation complex on an RNA substrate [Vethantham et al. 2007]. Additionally, several nuclear proteins depend on sumoylation for efficient nuclear localization [Pichler and Melchior 2002]. Detection of sumoylated proteins at constitutively transcribed genes may thus be a consequence of earlier modification.

Our analyses show the appearance of SUMO and recruitment of Ubc9 indicate that promoter-associated factors at inducible genes become sumoylated coincident with activation, and the SUMO signal disappears when the activating signal is no longer present [at least for ARGI]. We did not detect association of either of the two yeast SUMO proteases, Ulp1 and Ulp2, at the ARGI promoter during its activation or deactivation [E Rosonina and JL Manley, unpubl.], indicating that sumoylated promoter-bound factors are cleared from the deactivated promoter, and not simply desumoylated in that process. Activation-coincident sumoylation was observed for genes that are induced under very different conditions: addition of NaCl (STL1), presence of galactose (GAL1), amino acid starvation (ARG1), and addition of Cu (CUP1) [E Rosonina and JL Manley, unpubl.]. Thus, we showed that sumoylation at promoters occurs only while a gene is active, and is likely a general, previously unappreciated aspect of the gene activation process. Because we found that different types of inducible genes all show the same pattern of sumoylation, it is reasonable to assume that components of the general transcription machinery, present at all activated genes, are subject to sumoylation. However, as discussed below, it is also possible that gene-specific transcription factors, a large number of which are known to be sumoylated, are regulated at induced promoters through a common mechanism involving sumoylation.

Our analysis supports a model in which SUMO has multiple roles at actively transcribed genes in yeast. Previously sumoylated proteins assemble at active genes and have a positive effect on RNAP II recruitment [Fig. 6A]. These factors are detected at constitutively transcribed genes, but likely contribute to the SUMO signal detected at activated inducible genes as well [Fig. 6B]. At inducible genes, sumoylation of promoter-bound factors takes place, which has two consequences. First, we found
that sumoylation at induced promoters has a repressive effect on transcription. Recruitment of transcriptional repressors to activated genes has been demonstrated before (e.g., Mot1 and NC2) (Geisberg et al. 2001, 2002; Zanton and Pugh 2004; van Werven et al. 2008), indicating that, in some cases, repressive factors have a role during active transcription. Second, sumoylation functions in shutting off transcription of induced genes at the appropriate time. Both effects of sumoylation on induced transcription can be explained if sumoylation functions in deactivating the promoter, perhaps after every round of transcription initiation. That is, our model posits that sumoylation of some factor(s) is necessary for efficiently clearing the promoter after transcription has initiated, thereby allowing it to respond to another round of activation if a sufficient activation signal [activator] is present, or, if not, to shut down [Fig. 6B]. Failure to clear the promoter efficiently is expected to cause elevated activation and prolonged transcription when the activator is no longer available, as we observed in ubc9-1 cells. Immobilized template experiments, performed in yeast nuclear extract, showed that, once RNAP II has cleared the promoter and initiated transcription, a scaffold of factors remains at the promoter, including a subset of GTFs and Mediator (Yudkovsky et al. 2000). We propose that sumoylation is a general mechanism by which promoters are cleared of these factors and/or the activator, thereby regulating transcription levels and facilitating shut off of inducible genes.

The idea that promoters are cleared after every round of transcription, allowing the cell to respond to activator abundance in the nucleus, has some support, particularly for activators with acidic activation domains (Chi et al. 2001; Tansey 2001; Bhaumik and Malik 2008). Ubiquitin-mediated proteolysis of these activators once they are “spent” is thought to allow activator turnover at induced promoters. This is supported by studies that detect components of the 26S proteasome occupying most genes in yeast [Auld et al. 2006; Sikder et al. 2006]. Interestingly, the strongest evidence for this comes from studies on Gcn4, which is targeted for ubiquitylation by a phosphorylation mark left by a kinase component of the RNAP II Mediator complex Srb10, implying that degradation of Gcn4 is linked to transcriptional activation [Chi et al. 2001]. Promoter-associated factors, such as the repressor Mot1 in yeast, and PARP-1, which associates with the human HSP70.1 promoter [see below], are targets for ubiquitylation via prior sumoylation through the function of SUMO-targeted ubiquitin E3 ligases (STUbLs) [Martin et al. 2009; Wang and Prelich 2009]. Thus, ubiquitin-mediated proteolysis of promoter-associated factors, including activators, is a plausible mechanism by which SUMO facilitates promoter clearance/deactivation at induced promoters [Fig. 6B].

Is the connection between active transcription and the presence of sumoylated proteins that we detected in yeast evolutionarily conserved among eukaryotes? In human cells, heat shock results in recruitment of Ubc9 and the SUMO2 isoform to the heat-shock-induced HSP70.1 promoter, where a number of promoter-associated factors are thought to become sumoylated [Martin et al. 2009], suggesting that our findings may be extrapolated to higher eukaryotes. However, in mouse retina cells, ChIP analysis demonstrated that SUMO is present at both activated and repressed photoreceptor-specific genes [Onishi et al. 2009]. Although this phenomenon might
be specific to this class of genes in the retina, it contrasts with our findings in yeast, where we found no evidence that SUMO is associated with repressed genes. Thus, although the role of sumoylation at activated genes may be evolutionarily conserved, it appears that, in higher eukaryotes, SUMO plays an additional role in maintaining transcriptional repression. Through evolution, higher eukaryotes may have taken advantage of the ability of SUMO to inhibit transcription, which we detect at activated genes, in an additional role in keeping repressed genes silent.

Our findings have several interesting implications. For example, our analysis suggests that one reason so many transcription factors are SUMO substrates is because sumoylation is a general aspect of gene activation that regulates transcription at induced promoters. Furthermore, our results suggest that sumoylation of transcription factors is usually found to impart a negative effect on transcription because sumoylation may be a common mechanism of clearing transcription factors from promoters, such that blocking sumoylation results in apparent increased transcription of target genes. Finally, although the coordinated recruitment of factors to activated promoters has been studied extensively, it is not known how assembled factors are removed from the promoter when the activating signal disappears and the gene product is no longer needed—effectively, how genes are shut off. Our model posits that sumoylation of promoter-bound factors is a general mechanism of deactivating transcription, and suggests that shutting off transcription of many genes, or of individual genes, is a regulated process. In any event, and in contrast to the view that SUMO functions principally at transcriptionally repressed genes, the data presented here indicates that sumoylation is an important and general mechanism by which gene expression is regulated at actively transcribed genes.

Materials and methods

Plasmids and yeast strains

Yeast strains used in this study are listed in Supplemental Table S1. For strain ERYM434B, UBC9 was tagged at its genomic locus by homologous recombination as described previously [Kноп et al. 1999]. To generate plasmid pGCN4-Flag, GCN4, with 1 kb each of flanking upstream and downstream genomic sequence, was cloned into the URA3-marked CEN vector pRS316. Site-directed mutagenesis was used to create a unique EcoRI site at the 3’ end of the GCN4 ORF, upstream of the stop codon, that was used to introduce a Flag epitope tag sequence. Further cloning details for this plasmid and yeast strains used are available on request.

Yeast growth conditions

For galactose induction (induction of GAL1), strains were grown in rich medium containing 2% raffinose until the O.D. of 0.6–0.8, then NaCl was added to a final concentration of 0.4 M [induced sample] or cells were mock-treated [uninduced] and incubated for an additional 10 min. For induction of ARG1, cells were grown in minimal medium lacking Val and Ile to O.D. 0.6–0.8, then SM was added to a final concentration of 0.5 μg/mL [induced] or cells were untreated [uninduced] for 30 min or as indicated. For analysis of constitutive genes, cells were grown in complete minimal medium, then prepared for ChIP or RNA analysis as indicated below.

ChIP

Two-hundred milliliters of cells was grown at 28°C as per the conditions above, and ChIP was performed as described previously [Rosonina et al. 2009]. For SUMO ChIP, 0.8 μL of previously described Smt3 antibody [Montpetit et al. 2006] was used per immunoprecipitation, for RNAP II ChIP, 0.5 μg 8WG16 [Covance] was used, and 1 μg of HA (Sigma) or HIS (Santa Cruz Biotechnology) antibodies was used for HA- and HIS-tagged protein ChIP, respectively. Experiments were performed at least three times for each quantification, which was performed as described previously [Komarnitsky et al. 2000] by determining the ratio of the gene-specific amplification signal to the background amplification signal from the untranscribed region performed in the same reaction, for the immunoprecipitated samples. This ratio was then divided by the same ratio determined for the input material, resulting in a normalized, “fold over background” value and referred to as fold occupancy. Primer sequences used for PCR amplification are listed in Supplemental Table S2. Where statistical analysis was performed to compare data sets, a Student’s t-test was applied, with P-values indicated in the figure legends.

RT–PCR analysis

For preparation of total RNA samples, 50-mL cultures were grown as described above, and RNA was isolated and analyzed by RT–PCR as described previously [Rosonina et al. 2009]. Primer sequences are listed in Supplemental Table S2. Analyses were performed at least three times, and the result of semiquantitative analyses is shown. Where statistical analysis was performed to compare data sets, a Student’s t-test was applied, with P-values indicated in the figure legends.

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SUMO functions in constitutive transcription and during activation of inducible genes in yeast

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