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## RESEARCH COMMUNICATION

# The Spt components of SAGA facilitate TBP binding to a promoter at a post-activator-binding step in vivo

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**The SAGA complex of *Saccharomyces cerevisiae* is required for the transcription of many RNA polymerase II-dependent genes. Previous studies have demonstrated that SAGA possesses histone acetyltransferase activity, catalyzed by the SAGA component Gcn5. However, the transcription of many genes, although SAGA dependent, is Gcn5 independent, suggesting the existence of distinct SAGA activities. We have studied the in vivo role of two other SAGA components, Spt3 and Spt20, at the well-characterized *GAL1* promoter. Our results demonstrate that both Spt3 and Spt20 are required for the binding of TATA-binding protein but not of the activator Gal4 and that this role is Gcn5 independent. These results suggest a coactivator role for Spt3 and Spt20 in the recruitment of TBP.**

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In eukaryotes, transcription initiation by RNA polymerase II (Pol II) requires not only the activities of the general transcription factors and gene-specific activators but also the activities of large coactivator complexes (Hampsey 1998; Hampsey and Reinberg 1999). The SAGA (Spt/Ada/Gcn5/acetyltransferase) complex of *Saccharomyces cerevisiae* is one such coactivator complex. It is required for the full transcriptional activity of a subset of Pol II-dependent genes (Hampsey 1997; Grant et al. 1998). SAGA contains >20 proteins, including Gcn5, a histone acetyltransferase (HAT) whose activity has been shown to be required for the transcription of a subset of genes in vivo (Kuo et al. 1998; Wang et al. 1998). Recent studies have implicated Gcn5 in the recruitment of activators to promoters by SAGA (Cosma et al. 1999).

In addition to the HAT activity of Gcn5, genetic and biochemical studies suggest that SAGA possesses other activities important for transcription (Horiuchi et al. 1997; Roberts and Winston 1997; Dudley et al. 1999; Sterner et al. 1999). In this paper we focus on two other

components of SAGA, Spt20, and Spt3. Spt20 is required for the integrity of SAGA, as the complex cannot be detected in *spt20Δ* mutants (Grant et al. 1997). Spt3 is probably required for a subset of SAGA activities that are independent of Gcn5 and histone acetylation because SAGA purified from an *spt3Δ* mutant still possesses Gcn5-dependent HAT activity, yet *spt3Δ* mutants have multiple transcriptional defects (Roberts and Winston 1997; Dudley et al. 1999; Sterner et al. 1999). Previous genetic and biochemical studies have shown that Spt3 interacts with TATA-binding protein (TBP) (Eisenmann et al. 1992; Lee and Young 1998) and suggest that Spt3 is structurally similar to particular TBP-associated factors (TAFs) (Birck et al. 1998). These findings suggest a model in which Spt3 plays a role in the binding of TBP to particular promoters in vivo.

To better understand the mechanism by which the Spt components of SAGA facilitate the transcription of a subset of RNA Pol II-dependent genes, we chose to study the effects of SAGA mutations on activation by Gal4 at the well-characterized *GAL1* promoter. Studies of activation of the *GAL* promoters by Gal4 have served as a model system for transcriptional regulation for many years (Johnston and Carlson 1992; Ptashne and Gann 1997; Zaman et al. 1998). Furthermore, the *GAL1* promoter is useful for such studies because its regulation is well characterized and it has a relatively simple promoter structure. Under inducing conditions, two proteins, TBP and Gal4, are known to bind the *GAL1* promoter (Johnston and Carlson 1992). Thus, of the genes known to be regulated by the Spt components of SAGA (Roberts and Winston 1997; Sterner et al. 1999), the *GAL* promoters are the only ones for which the Spt dependence of both TBP binding and activator binding can be measured.

In this study we use chromatin immunoprecipitation assays to determine which steps in transcriptional activation require Spt20 and Spt3 at the *GAL1* promoter. Our results demonstrate a severe reduction in TBP binding at the *GAL1* TATA in *spt3Δ* and *spt20Δ* mutants but only a modest decrease in a *gcn5Δ* mutant. These results correlate well with the effects on *GAL1* transcription in these mutants. In contrast, the transcriptional activator Gal4 is bound to the *GAL1* upstream activating sequence (UAS<sub>G</sub>) in all three SAGA mutants. Furthermore, the acetylation of histone H3 at the *GAL1* promoter is only slightly decreased in *gcn5Δ*, *spt3Δ*, and *spt20Δ* mutants. Taken together, our results suggest a Gcn5-independent activity for the Spt20 and Spt3 components of SAGA that is required for the recruitment of TBP, but not of activators, to a subset of Pol II-dependent promoters in vivo.

## Results

### *GAL1* transcription is dependent on Spt3 and Spt20, but not Gcn5

To understand the role of different SAGA functions, we

[Key Words: SAGA complex; *S. cerevisiae*; transcription; Spt3; TBP]

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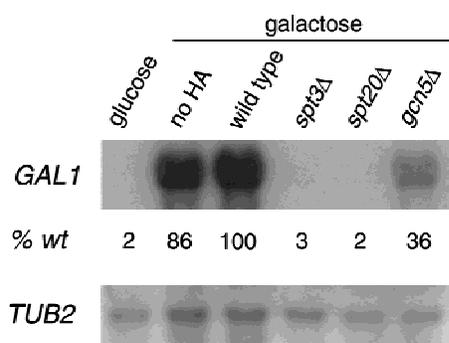
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examined the effects of mutations in three different classes of SAGA genes, *SPT3*, *GCN5*, and *SPT20*, on the well-studied *GAL1* gene of *S. cerevisiae*. Spt3 and Gcn5 are probably required for distinct aspects of SAGA function, whereas Spt20 is believed to be required for all SAGA function (Grant et al. 1997; Sterner et al. 1999). The *GAL1* promoter was particularly useful for analysis of Spt3 and Spt20 because *gcn5Δ* mutants are Gal<sup>+</sup>, whereas *spt3Δ* are moderately Gal<sup>-</sup>, and *spt20Δ* mutants are tight Gal<sup>-</sup> (Roberts and Winston 1997; Sterner et al. 1999). To study primary events of transcriptional activation at the *GAL1* promoter, we analyzed cells at an early point in galactose induction, sufficient for the wild-type strain to have fully induced expression (Materials and Methods). At this point in galactose induction, *GAL1* mRNA levels are decreased in the *spt3Δ* and *spt20Δ* mutants by ~50-fold relative to wild-type levels (Fig. 1). In contrast, a *gcn5Δ* mutant shows only a modest 2.5-fold decrease in *GAL1* mRNA levels. Thus, induction of the *GAL1* promoter is strongly dependent on the Spt3 and Spt20 components of SAGA and only mildly dependent on the Gcn5 HAT.

#### TBP binding at *GAL1* is defective in *spt3Δ* and *spt20Δ* mutants

An important step in transcription initiation is the recruitment of TBP to the TATA region of a promoter. Previous studies of the *GAL1* promoter in vivo demonstrated that TBP is absent from the *GAL1* TATA region under conditions of both glucose repression or noninduction (raffinose or glycerol). However, upon induction with galactose, TBP associates strongly with the *GAL1* TATA region (Selleck and Majors 1987a,b; Kuras and Struhl 1999; Li et al. 1999). To test whether SAGA mutations affect the binding of TBP to the *GAL1* TATA in vivo, we used the chromatin immunoprecipitation assay (Dedon et al. 1991; Orlando and Paro 1993; Strahl-Bolsinger et al. 1997). We observed a strong (>10-fold) defect in the ability of TBP to bind the *GAL1* TATA region in the *spt3Δ* and *spt20Δ* mutants, but only a modest (two-

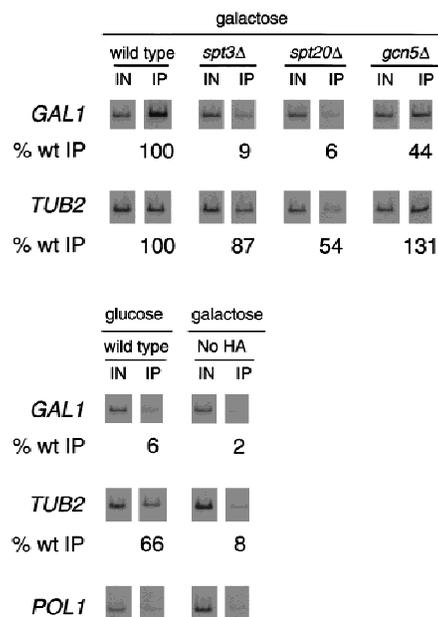


**Figure 1.** *GAL1* mRNA levels in three classes of SAGA mutants. Northern hybridization analysis was performed on total yeast RNA prepared from glucose-repressed wild-type strains and galactose-induced wild-type, *spt3Δ*, *spt20Δ*, and *gcn5Δ* strains. Quantitation was performed by PhosphorImager analysis (Molecular Dynamics).

fold) decrease in TBP binding in the *gcn5Δ* mutant (Fig. 2). These results strongly suggest that the *spt3Δ* and *spt20Δ* defects in *GAL1* activation are caused by an inability to stably bind TBP to the promoter.

#### *Gal4* binding is unaffected in *spt3Δ* and *spt20Δ* mutants

One condition that could account for both the defect in *GAL1* transcription and in TBP binding is the inability of the transcriptional activator Gal4 to bind to UAS<sub>G</sub> in the *spt* mutants. To test this hypothesis, we used chromatin



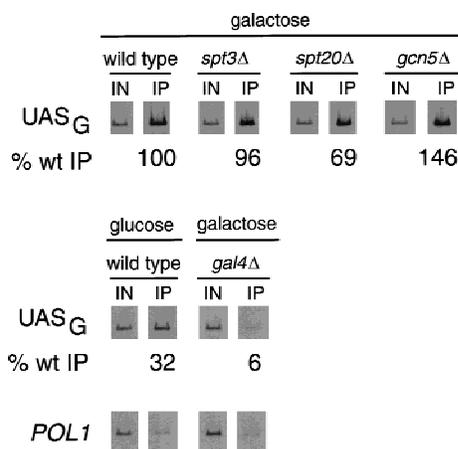
**Figure 2.** TBP is not bound to the *GAL1* TATA in *spt3Δ* and *spt20Δ* mutants. Chromatin immunoprecipitation was performed in parallel with RNA analysis (Fig. 1) from glucose-repressed wild-type strains and galactose-induced wild-type, *spt3Δ*, *spt20Δ*, and *gcn5Δ* strains. A galactose-induced wild-type strain that contained TBP without the triple HA tag (no HA) was included as a negative control for the immunoprecipitation. TBP was immunoprecipitated using the 12CA5 antibody against the HA epitope. PCR products correspond to the *GAL1* TATA region (*GAL1*), the *TUB2* TATA region (*TUB2*), or the *POL1* open reading frame as a negative control. The percentage of DNA immunoprecipitated (% wt IP) in each of the mutants was normalized to the amount immunoprecipitated from the galactose-induced wild-type strain. One set of PCR reactions is shown, and the quantitation represents the average of several experiments. The average values with standard errors for the measurements of TBP binding to the *GAL1* TATA on galactose-grown cells are as follows: *spt3Δ*, 9 ± 2; *spt20Δ*, 6 ± 2; *gcn5Δ*, 44 ± 3; No HA, 2 ± 1. The values for binding to the *TUB2* TATA on galactose-grown cells are: *spt3Δ*, 87 ± 34; *spt20Δ*, 54 ± 10; *gcn5Δ*, 131 ± 22; No HA, 8 ± 2. The values for wild-type glucose-grown cells are as follows: *GAL1*, 6 ± 3; *TUB2*, 66 ± 8. The low level of DNA detected in the No-HA and the *POL1* PCR reactions represents the low amount of TBP-independent DNA precipitated as background in this assay. The *POL1* negative control was performed on all samples, and the results were essentially the same as the example shown.

Dudley et al.

immunoprecipitation assays to analyze the occupancy of the UAS<sub>G</sub> by Gal4 (Fig. 3). Interestingly, in all the SAGA mutants tested we observed in vivo occupancy of the UAS<sub>G</sub> by Gal4 at levels similar to that of wild type. Therefore, Spt3 and Spt20 are not required for Gal4 binding. Together with the defects in TBP binding in *spt3Δ* and *spt20Δ* mutants, these results strongly suggest that Gal4 cannot recruit TBP in the absence of Spt3 or Spt20.

#### Histone H3 acetylation in *gcn5Δ*, *spt3Δ*, and *spt20Δ* mutants

Previous studies have suggested that Spt3 functions in SAGA independently of the HAT activity of Gcn5 (Roberts and Winston 1997; Dudley et al. 1999; Sterner et al. 1999). To test directly whether *spt3Δ*, *gcn5Δ*, or *spt20Δ*

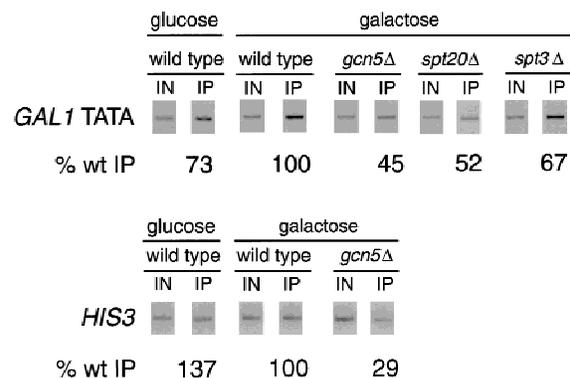


**Figure 3.** Gal4 occupies the UAS<sub>G</sub> in SAGA mutants. Chromatin immunoprecipitation was performed in parallel with RNA analysis (Fig. 1) from glucose-repressed wild-type strains and galactose-induced wild-type, *spt3Δ*, *spt20Δ*, or *gcn5Δ* strains. A galactose-induced *gal4Δ* strain was included as a negative control for the immunoprecipitation. Gal4 was immunoprecipitated using the RK5C1 antibody against Gal4 (Santa Cruz Biotechnology). PCR products correspond to the UAS<sub>G</sub> or the *POL1* ORF as a negative control. The percentage of UAS<sub>G</sub> DNA immunoprecipitated (% wt IP) from each of the mutants was normalized to the amount precipitated from the galactose-induced wild-type strain. One set of PCR reactions is shown, and the quantitation represents the average of several experiments. The average values with standard errors for the measurements of Gal4 binding to the UAS<sub>GAL</sub> on galactose-grown cells are as follows: *spt3Δ*, 96 ± 6; *spt20Δ*, 69 ± 1; *gcn5Δ*, 146 ± 12; *gal4Δ*, 6 ± 1. The measurement for wild type grown on glucose was 32 ± 4. The low level of DNA detected in the *gal4Δ* and the *POL1* PCR reactions represents the low amount of Gal4-independent DNA precipitated as background in this assay. The *POL1* negative control was performed on all samples, and the results for all the other extracts were essentially the same as the example shown. In this experiment we detected a modest, but significant, threefold increase in Gal4 binding to the UAS<sub>G</sub> in galactose-induced cells relative to glucose-repressed cells. These results differ slightly from previous in vivo footprint analyses that detected little or no Gal4 occupancy of the UAS<sub>G</sub> in glucose-repression conditions (Giniger et al. 1985; Selleck and Majors 1987b), a result that highlights the sensitivity of the chromatin immunoprecipitation assay.

mutations alter histone acetylation at the *GAL1* promoter, we used chromatin immunoprecipitation assays to determine the histone H3 acetylation levels of *GAL1* in these SAGA mutants (Fig. 4). As a control, we examined the H3 acetylation levels at *HIS3* and observed a three-fold decrease in a *gcn5Δ* mutant, consistent with previously published results (Kuo et al. 1998). At *GAL1*, our results show that H3 acetylation is only mildly reduced in *spt3Δ*, *gcn5Δ*, and *spt20Δ* mutants. For the *gcn5Δ* mutant, this effect correlates with the weak effect on *GAL1* transcription. However, for the *spt3Δ* and *spt20Δ* mutants, there is not a good correlation between H3 acetylation and the severe *GAL1* transcriptional defects. This result strongly suggests that *GAL1* transcription is not significantly dependent on histone acetylation by SAGA and that the role of Spt3 is unrelated to Gcn5-dependent histone acetylation.

#### Discussion

Coactivators are believed to act as intermediaries between gene-specific activators and TBP or other general transcription factors. However, the molecular mechanisms by which coactivators function are not well understood (Hampsey 1998; Hampsey and Reinberg 1999). Our experiments have defined an in vivo coactivator function for Spt3 and Spt20 of the SAGA complex. In both *spt3Δ* and *spt20Δ* mutants, Gal4 binding is normal, but TBP fails to bind to the *GAL1* promoter, and transcription is reduced >50-fold. Thus, the Spt3 and Spt20



**Figure 4.** Histone H3 acetylation in *spt3Δ*, *gcn5Δ*, and *spt20Δ* mutants. Chromatin immunoprecipitation was performed in parallel with RNA analysis (Fig. 1) from glucose-repressed, wild-type strains and galactose-induced wild-type, *spt3Δ*, *spt20Δ*, or *gcn5Δ* strains. Histone H3 acetylated at Lys-9 and Lys-14 was immunoprecipitated using antisera previously described (Kuo et al. 1998). PCR products correspond to the *GAL1* TATA or the *HIS3* promoter as a control. The percentage of *GAL1* TATA DNA immunoprecipitated (% wt IP) from each of the mutants was normalized to the amount precipitated from the galactose-induced wild-type strain. One set of PCR reactions is shown, and the quantitation represents the average of several experiments. The *spt3Δ* and *spt20Δ* mutants showed some variation, 40%–110% for *spt3Δ* and 36%–70% for *spt20Δ*. At *HIS3*, a threefold decrease in H3 acetylation level was observed in the *gcn5Δ* mutant compared with wild type, similar to previously reported results (Kuo et al. 1998).

proteins are required for Gal4 to recruit TBP to the *GAL1* promoter. These results constitute the first in vivo demonstration of such a coactivator role. Our results fit well with other studies of Gal4 and TBP binding in vivo that demonstrated that TBP binding requires the binding and activation of Gal4 (Selleck and Majors 1987a,b; Kuras and Struhl 1999; Li et al. 1999).

The model most consistent with these results is one in which TBP is physically recruited to the *GAL1* promoter by the Spt components of SAGA. Previous analysis suggests a physical interaction between Spt3 and TBP (Eisenmann et al. 1992; Lee and Young 1998). In addition, the positions of amino acid changes of both TBP and Spt3 mutants that alter their functional interaction suggest specific regions of each protein that may be involved in TBP–Spt3 interactions (Eisenmann et al. 1992). These results are supported by recent studies of TBP–TAF<sub>II</sub>28 interactions (Lavigne et al. 1999), as Spt3 is predicted to have structural similarity to both TAF<sub>II</sub>18 and TAF<sub>II</sub>28 (Birck et al. 1998). Recent results have shown that SAGA can be recruited by several transcriptional activators including VP16 (Ikeda et al. 1999), Gcn4 (Natarajan et al. 1998), and Rtg3 (Massari et al. 1999). Taken together with these results, our data suggest a model for coactivation in which SAGA is targeted to the *GAL1* promoter by interactions with Gal4, followed by recruitment of TBP via interactions with Spt3. The recruitment of TBP by the Spt proteins may be followed by the establishment of previously demonstrated interactions between Gal4 with certain general transcription factors, including TBP, TFIIB, and Srb4 (Melcher and Johnston 1995; Wu et al. 1996; Koh et al. 1998).

Our results, together with previous studies, suggest that other SAGA components may be required for this coactivator function. Spt20, Spt7, and Ada1 are all probably required for the integrity of SAGA (Grant et al. 1997; Sterner et al. 1999). Therefore, the defect observed in *spt20Δ* mutants is probably caused by the loss of all SAGA functions, including Spt3, although we cannot rule out a more direct role. In addition to Spt3, the Spt8 protein has been implicated in SAGA–TBP interactions, and in vitro data suggest a direct role for Spt8 (Sterner et al. 1999). These results suggest that Spt8, along with Spt3, might be required for TBP binding at the *GAL1* promoter. However, an *spt8Δ* mutant is Gal<sup>+</sup>, suggesting that Spt8 plays, at most, a minor role at this promoter. Furthermore, previous studies have demonstrated that a particular mutation in *SPT3* can partially bypass the requirement for Spt8, suggesting that Spt8 plays a more auxiliary role (Eisenmann et al. 1994). Possibly, both Spt3 and Spt8 assist TBP recruitment, but they contribute to different degrees in a promoter-specific fashion. In addition to these Spt proteins, other SAGA components, including the Ada and Taf proteins, may play related roles in the assembly of the preinitiation complex. Finally, recent evidence suggests that the Snf/Swi complex also helps to activate transcription at a step subsequent to activator binding (Ryan et al. 1998).

Our results cannot rule out a model in which the Spt components of SAGA act at another step in transcrip-

tional activation that results in the recruitment of TBP. For example, the Spt proteins could facilitate some aspect of Gal4 activation, subsequent to its DNA binding, that allows Gal4 to recruit TBP to the *GAL1* promoter. Such a role could include helping to determine the correct interactions of Gal4 with either Gal80 or Gal3, both known to interact with Gal4 under inducing conditions (Blank et al. 1997; Yano and Fukasawa 1997). Given that only particular promoters are dependent on the Spt proteins, it is likely that multiple factors determine both their requirement and allowing their function at any Spt-dependent promoter.

Finally, our results demonstrate that the function of SAGA at *GAL1* is largely independent of the Gcn5 HAT. These results provide an interesting contrast to those from a recent study that demonstrated a strong Gcn5 requirement for the binding of the Swi4 activator to the *HO* promoter in vivo (Cosma et al. 1999). The differences between these two sets of results demonstrate that different promoters can require distinct SAGA components for mechanistically distinct functions. The determinants of these requirements at any promoter are an issue for future investigation.

## Materials and methods

### *S. cerevisiae* strains

All *S. cerevisiae* strains are isogenic to a *GAL2*<sup>+</sup> derivative of S288C (Winston et al. 1995). Strains were constructed with the following relevant genotypes: wild type (FY1887 and FY1888), *spt3-202* (FY1889 and FY1890), *spt20Δ100::URA3* (FY1891 and FY1892), and *gcn5Δ::HIS3* (FY1893 and FY1894). Each of these strains also contained an *spt15Δ::LEU2* mutation in the genome and an *HA<sub>3</sub>-SPT15 TRP1 CEN* plasmid (Kuras and Struhl 1999) as the only source of wild-type TBP. The *HA<sub>3</sub>-SPT15* construct fully complemented the *spt15Δ102::LEU2* mutation for all phenotypes tested, including growth on galactose (data not shown). The control strain for the Gal4 chromatin immunoprecipitation experiments, FY760 (*gal4Δ::LEU2*), was *SPT15*<sup>+</sup> in the genome and contained no plasmid. The control strain for the TBP chromatin immunoprecipitation experiments, FY1886 (the “no-HA” control), contained *spt15Δ102::LEU2* in the genome and contained *SPT15*<sup>+</sup> on a plasmid (Kuras and Struhl 1999) as the only source of wild-type TBP. Glucose-repressed strains were grown in YPD (2% glucose). Galactose-induced strains were grown in YPrf (2% raffinose) and induced for 20 min by the addition of galactose to 2%. RNA for Northern analysis and chromatin extracts were prepared from the same cultures grown to cell densities of  $1 \times 10^7$ – $2 \times 10^7$  cells/ml. Northern analysis and chromatin immunoprecipitation assays were performed on both sets of isogenic strains.

### Northern hybridization analysis

Total yeast RNA was prepared as described previously (Swanson et al. 1991). Northern blot analysis was performed on both sets of strains used for chromatin immunoprecipitations. One of the experiments is shown (Fig. 1), and the quantitation represents the average of both sets of strains. The *GAL1* (St. John and Davis 1981) and *TUB2* (Som et al. 1988) probes have been described previously.

### Chromatin immunoprecipitation

Formaldehyde cross-linking extracts were prepared essentially as described previously (Kuras and Struhl 1999) with the following exceptions: First, all centrifugations to pellet the chromatin extract were performed for 1 min at 14,000 rpm in an Eppendorf centrifuge. Second, the separation of soluble chromatin following sonication was accomplished by a 1-hr centrifugation at 14,000 rpm in an Eppendorf centrifuge. Immunoprecipitations of HA<sub>3</sub>-tagged TBP were performed as described previously (Kuras and Struhl 1999). Gal4 immunoprecipitations were performed by the same method used for the HA<sub>3</sub>-tagged TBP with the exception that binding was done in FA lysis buffer containing 150 mM NaCl (Kuras and

Dudley et al.

Struhl 1999) and washes were done three times in the same buffer and once in TE (10 mM Tris-HCl, 1 mM EDTA at pH 8.0). Immunoprecipitation of the hyperacetylated form of histone H3 was performed as described previously (Kuo et al. 1998). PCR reactions were performed essentially as described previously (Kuras and Struhl 1999), with the exception that PCR products were detected by the incorporation of [<sup>32</sup>P]dATP in the reaction. The PCR primers amplify the following regions whose coordinates are given relative to the ATG (+1): *GAL1* TATA primers amplify a 244-bp region from -190 to +54; *GAL1* UAS primers amplify a 260-bp region from -536 to -276; *TUB2* TATA primers amplify a 273-bp region from -186 to +87; *POL1* ORF primers amplify a 219-bp region from +2499 to +2717; and *HIS3* primers amplify a 105-bp region from -28 to +77.

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## The Spt components of SAGA facilitate TBP binding to a promoter at a post-activator-binding step in vivo

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