Identification of three regions essential for interaction between a σ-like factor and core RNA polymerase

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The cyclic interactions that occur between the subunits of the yeast mitochondrial RNA polymerase can serve as a simple model for the more complex enzymes in prokaryotes and the eukaryotic nucleus. We have used two-hybrid and fusion protein constructs to analyze the requirements for interaction between the single subunit core polymerase (Rpo41p), and the σ-like promoter specificity factor (Mtf1p). We were unable to define any protein truncations that retained the ability to interact, indicating that multiple regions encompassing the entire length of the proteins are involved in interactions. We found that 9 of 15 nonfunctional (petite) point mutations in Mtf1p isolated in a plasmid shuffle strategy had lost the ability to interact. Some of the noninteracting mutations are temperature-sensitive petite (ts petite); this phenotype correlates with a precipitous drop in mitochondrial transcript abundance when cells are shifted to the nonpermissive temperature. One temperature-sensitive mutant demonstrated a striking pH dependence for core binding in vitro, consistent with the physical properties of the amino acid substitution. The noninteracting mutations fall into three widely spaced clusters of amino acids. Two of the clusters are in regions with amino acid sequence similarity to conserved regions 2 and 3 of σ factors and related proteins; these regions have been implicated in core binding by both prokaryotic and eukaryotic σ-like factors. By modeling the location of the mutations using the partial structure of Escherichia coli σ70, we find that two of the clusters are potentially juxtaposed in the three-dimensional structure. Our results demonstrate that interactions between σ-like specificity factors and core RNA polymerases require multiple regions from both components of the holoenzymes.

[Key Words: RNA polymerase; σ factor; transcription initiation; mitochondrial transcription; MTF1; RPO41; two-hybrid]

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The complex RNA polymerases of eukaryotes and prokaryotes require auxiliary factors specific for the initiation phase of transcription. These factors associate with the core polymerases to form holoenzymes competent for promoter recognition, selective DNA binding, and opening of the double-stranded DNA at the start site of transcription. Shortly after initiation, the factors are released as the RNA polymerase makes the transition into its elongating form. The factors associated with eukaryotic nuclear RNA polymerase II (Pol II) before initiation, and released shortly after transcription is initiated include TFIIB, TFIID, TFIIE, TFIIF, and TFIH (Conaway and Conaway 1993; Zawel and Reinberg 1995). In bacterial cells, members of the family of σ factors carry out most of the functions of the many eukaryotic nuclear factors (for review, see Helmann 1994). The interaction of a σ factor with the core polymerase alters the conformation of both the polymerase and the σ factor to expose amino acids critical for promoter recognition, and to allow the loading of the polymerase onto the DNA (Dombroski et al. 1993; Polyakov et al. 1995). Although much has been learned about how σ factors and other sequence-specific binding factors interact with DNA, relatively little is known about the interactions of these factors with the subunits of core polymerases and how these interactions influence conformational changes in both components of the holoenzyme. In this work, we have used the yeast mitochondrial RNA polymerase (mt RNAP) as a simple model to examine the interaction between a promoter specificity factor and a core polymerase.

The core mt RNAP is a single polypeptide encoded by the nuclear gene RPO41. Rpo41p shares nine regions of amino acid sequence similarity with the single subunit RNA polymerases of the T7 and T3 bacteriophage (Masters et al. 1987; Jang and Jaehning 1994). These regions include the amino acids known to be required for struc-
tecture and function of the catalytic domain of the phage polymerases (Delarue et al. 1990; Sousa et al. 1993). However, unlike the phage polymerases that function independently, Rpo41p requires a specificity factor encoded by the nuclear MTF1 gene. Mtf1p has only limited amino acid similarity to σ factors (Jang and Jaehning 1991), but functions in many ways like σ in that it is required for promoter recognition and initiation of transcription. Although Mtf1p does not bind to its simple nine-base promoter (consensus ATATAAGTA; Oshima et al. 1982) on its own, it interacts with the core polymerase in solution to create a holoenzyme capable of promoter recognition (Mangus et al. 1994). Mtf1p is released after a short transcript has been synthesized and is available for interaction with a new core subunit, also reminiscent of σ factors (Mangus et al. 1994). The mitochondrial RNA polymerase therefore undergoes the same cycle of interactions as do the more complicated prokaryotic and eukaryotic nuclear enzymes, but requires only two polypeptides rather than the four to more than 30 used in the more complex systems.

In this work we have investigated the requirements for interaction between Mtf1p and Rpo41p. A comprehensive deletion analysis of both proteins failed to identify a simple interaction region, indicating that several regions of both proteins may be involved in the protein–protein interactions. We have demonstrated that this is the case for Mtf1p by identifying three regions required for interactions with Rpo41p. Two of these regions are similar to regions of σ-like factors shown to have a role in interactions with the prokaryotic and eukaryotic core polymerases. This analysis establishes the yeast mt RNAP as a useful model for the analysis of protein–protein interactions during the transcription cycle, and demonstrates that core polymerase/accessory factor interactions involve complex-binding surfaces on both components of the holoenzyme.

**Results**

The interaction between Mtf1p and Rpo41p can be detected in two-hybrid constructs.

The analysis of interactions between σ-like factors and core polymerases is complicated because of the number of polypeptides in most core RNA polymerases. With the two-component mitochondrial RNA polymerase it is possible to use the powerful technique of two-hybrid analysis (Bartel et al. 1993; Phizicky and Fields 1995) to determine regions and/or specific amino acids of the Rpo41p core and the Mtf1p σ-like specificity factor necessary for protein–protein interactions. Although two-hybrid analyses of proteins that normally interact in the cytoplasm have been successful in the nuclear environment required for the technique, it was critical that we establish that the relatively weak interactions between the mtRNAP subunits (Mangus et al. 1994) could be detected in two-hybrid constructs. Initially, full-length polymerase subunits were tested for interaction in this assay. Mtf1p was fused to the LexA DNA-binding do-

Deletions of Rpo41p and Mtf1p fail to identify a discrete interaction region

Two-hybrid analyses have been used to delineate small regions of proteins necessary and sufficient for protein–protein interactions (Bartel et al. 1993; Phizicky and Fields 1995). Therefore, we asked if deletion constructs could be used to define the region of interaction between the two proteins. As shown in Figure 1, several deletion constructs were made for both Mtf1p (Fig. 1A) and Rpo41p (Fig. 1B) in two-hybrid vector backbones. The six Mtf1p deletions included three carboxy-terminal deletions, one amino-terminal deletion, and two internal deletions. The two internal deletions removed amino acids conserved with σ regions 2.1 and 2.2. All constructs were expressed in yeast cells and produced proteins of the predicted sizes (Fig. 2A; data not shown). Furthermore, the deletion constructs were expressed at levels that were equivalent to the full-length product (Fig. 2A). However, none of the Mtf1p deletion constructs showed any interaction, as measured by β-galactosidase activity, with the core polymerase in this assay (data not shown). These results suggest that it is not a single domain but multiple regions of the folded structure of Mtf1p whose contact with Rpo41p is required to produce a stable interaction.

A similar analysis was conducted with Rpo41p. Rpo41p is larger than its phage relatives and contains an amino-terminal extension of ~300 amino acids that has no similarity to the phage polymerases (Masters et al. 1987). Because the phage polymerases do not require a specificity factor, we have speculated that the amino-terminal extension may have a role in the interaction with the specificity factor (Jaehning 1993). To determine if this sequence alone is capable of interacting with Mtf1p, we fused the amino-terminal sequence (amino acids 1–311) of Rpo41p to the Vp16 activation region. The amino-terminal fragment showed no interaction with Mtf1p by this analysis (data not shown). However, this fusion construct was not detected in vivo by Western blot analysis. Four additional Rpo41p constructs (Fig. 1B) were tested for interaction with Mtf1p. Each of the constructs accumulated in yeast cells (Fig. 2B), but none was capable of interaction with Mtf1p (data not shown). Two constructs (Rpo41p1-597 and Rpo41p1-918) contain the amino-terminal extension, but are unable to mediate interaction with Mtf1p. The carboxy-terminal polymerase region, Rpo41p311–1351, is also incapable of interacting with Mtf1p. Mtf1p may therefore interact with amino acids in both the amino-terminal extension and the polymerase domain.
Isolation of point mutations in Mtf1p that confer a petite phenotype

Because deletions in Mtf1p and Rpo41p failed to identify discrete regions sufficient for interaction, we turned to analysis of point mutations to identify specific residues required for the interaction. Although MTF1 is not essential for yeast cell growth, it is required for the stable replication and transmission of the mitochondrial genome (Lisowsky and Michaelis 1988). Yeast strains lacking functional Mtf1p rapidly lose their full-length mitochondrial DNA, producing a petite phenotype that cannot be complemented by the subsequent expression of MTF1 (Jang and Jaehning 1991). Isolation of nonfunctional MTF1 mutations therefore requires the use of the plasmid shuffle technique where a plasmid bearing a wild-type copy of the gene is used to cover a chromosomal mutation (Sikorski and Boeke 1991). A yeast strain was created for this procedure as outlined in Figure 3.

For mutagenesis, MTF1 was amplified using low fidelity PCR conditions to create random mutations throughout the gene (Materials and Methods). The mutant fragments were transformed into the recipient strain and then grown on 5-fluoro-orotic acid (5FOA) to select for cells that had lost the wild-type MTF1 plasmid with the URA3 marker. The mtf1 mutants were tested subsequently for mitochondrial function by growth on a non-fermentable carbon source. Strains that were unable to grow on glycerol plates (petite), or unable to grow on glycerol plates at 37°C temperature-sensitive petite (ts petite) were collected for further analysis.

From 10,000 primary transformants, 22 petite and ts petite mutants were identified and sequenced to determine the mutation responsible for the petite phenotype. Six of the mutants contained nonsense codons, whereas the other 17 mutants contained either single (6), double (9), or triple (2) missense mutations (Table 1). Missense mutations were identified between codons 40 and 247. To increase the pool of single missense mutations, we reconstructed many of the mutations as single point mutations using site-directed mutagenesis. Oligonucleotides based on the sequenced mutations were designed for mutagenesis of the wild-type sequence (Materials and Methods). The isolated single mutations were retested using the plasmid shuffle to determine the effect of the single mutations in vivo. Nine of the 10 reconstructed mutations were either petite or ts petite when present as single mutations (Table 1). Although not all of the identified mutations were reconstructed, we did isolate a substantial pool of defective Mtf1p mutations. In total, 15 petite or ts petite point mutations were identified.

Figure 1. Two-hybrid deletion constructs. (A) Mtf1p deletions were fused to the LexA DNA-binding region and tested for interaction with a full-length Rpo41p two-hybrid construct. (B) Rpo41p deletions were fused to the Vp16 activator region and tested for interaction with a full-length LexA:Mtf1p construct. Strains, vectors, and protocols are described in Materials and Methods. Numbers refer to amino acid positions retained in the constructs. Shaded boxes in the full-length maps of MTF1 and RPO41 refer to regions conserved between σ factors and Mtf1p (Jang and Jaehning 1991) or between Rpo41p and T7 RNA polymerase (Masters et al. 1987; Jaehning 1993).

Figure 2. Confirmation of the expression of the two-hybrid constructs. Two-hybrid fusion proteins were detected in extracts of whole-cell yeast grown at 30°C. Rpo41p fusion proteins were visualized using an anti-Rpo41p antibody, whereas Mtf1p fusion proteins were detected with an anti-LexA antibody (Materials and Methods). Equal amounts of yeast cell extracts were analyzed for each sample. (A) Mtf1p carboxy-terminal deletion constructs. (B) Rpo41p deletion constructs. (C) Mtf1p point mutations that are defective for interaction with Rpo41p in the two-hybrid system.
Transcription of all classes of mitochondrial genes is reduced in ts mtf1 mutants.

It was possible that some of the defects in Mtf1p function could be attributable to failure to recognize one or more of the promoters in the mitochondrial genome. We therefore analyzed patterns of mitochondrial transcription at the permissive (30°C) and nonpermissive (36°C) temperatures for several ts petite mutants (Fig. 4; data not shown). Strains of yeast bearing the wild-type or temperature-sensitive (L53H and I154T) alleles of MTF1 were grown in minimal glucose media at the permissive temperature and then shifted to the nonpermissive temperature. Samples of yeast were collected at intervals and total RNA was isolated, fractionated, and hybridized with oligonucleotide probes specific for mitochondrial 14S rRNA, mRNAs for the COB and OLI1 genes, and the tRNAs for Glu, Ser1, Trp, ThrACN, f-Met, and Phe, as well as the cytoplasmic 18S rRNA as a normalization control. These genes represent 7 of the 12 known mitochondrial transcription units and include several of the different promoter sequence variations (Dieckmann and Staples 1994). Some of the hybridization analyses are shown in Figure 4A and the quantitation of these data are presented in Figure 4B.

Three conclusions can be drawn from the results in Figure 4. First, mitochondrial gene expression is reduced significantly in the ts mutants relative to wild-type expression, even at the permissive temperature (Fig. 4A). Because the strains bearing the mutant alleles of Mtf1p grow almost as well as wild type on glycerol medium, mitochondrial function can be maintained apparently with only 10%–20% of the wild-type level of mitochondrial transcripts. Second, there is a rapid decrease in mitochondrial RNA abundance in both the wild-type and mutant strains after the shift to the nonpermissive temperature, but the mutants do not recover and RNA levels decrease to almost undetectable levels (Fig. 4B). Finally, the abundance of all mitochondrial transcripts that we analyzed was reduced to a similar extent (Fig. 4A,B; data not shown). These mtf1 mutations therefore affect the transcription of all mitochondrial genes and are not specifically defective for recognition of a subset of mitochondrial promoters. Under the conditions used in these assays, petites do not start to accumulate in the mutant strains until about 24 hr after the shift to the nonpermissive temperature (data not shown). During this period the mutants maintained levels of mitochondrial DNA similar to those found in the wild-type strain (data not shown). Therefore, the petite phenotype caused by the mtf1 mutations is caused by a defect in transcription of mitochondrial genes, not by a rapid loss of mitochondrial DNA. This defect could be caused by failure to interact with the core polymerase, inability to recognize or bind to the mitochondrial promoter, or loss-of-function in other steps in initiation. In the following sections we have tested these mutations for the first of these possible defects.

Some of the mtf1 mutations no longer interact with Rpo41p in the two-hybrid assay.

We introduced each of the point mutations into the two-hybrid LexA fusion vector (Materials and Methods). Unlike the deletion mutations described above, most (10 of 15) of the mutants retained the ability to interact with Rpo41p, as determined by qualitative production of β-galactosidase in filter assays (data not shown). However, five of the 15 mutations (L53H, V135A, I154T, S218R, and D225G; see Fig. 5) were negative for β-galactosidase production, indicating that they had lost the ability to interact with Rpo41p. The inability of these mutants to interact was not attributable to lowered or abolished expression of the fusion constructs, as all of the noninteracting mutants are expressed at levels equivalent to wild-type (Fig. 2C). Because many of the petite MTF1 mutants were isolated as temperature-sensitive mutations, we repeated the filter assays after growing cells at 37°C (data not shown). Two additional mutations (Y42C and K157E) fail to interact at the nonpermissive temperature.

When the qualitative assays were followed by quantitative measurement of β-galactosidase activity, we found that many of the petite mutations associate with Rpo41p at levels that are indistinguishable from the wild-type protein (Fig. 5). However, consistent with the filter assays the noninteracting mutants L53H, V135A,
I154T, S218R, and D225G produce no β-galactosidase activity above background levels. We also identified two mutants that have intermediate levels of interaction. Mutant I221K generates <50% and mutant H44P produces <25% of the β-galactosidase activity of the wild-type construct. The two temperature-sensitive mutations Y42C and K157E produce β-galactosidase at levels that are equivalent to the wild-type protein at their permissive temperatures (30°C and 23°C, respectively), but are reduced severely in interaction at their nonpermissive temperatures.

Table 1. Petite and ts petite mutations identified by PCR mutagenesis of MTF1

| Mutations                  | Phenotype          | Mutations                  | Phenotype          |
|----------------------------|--------------------|----------------------------|--------------------|
| Nonsense                   | single amino acid  |                            |                    |
| K35Stop                    | petite             | Y42C                       | ts petite          |
| Q60Stop                    | petite             | L53H                       | ts petite          |
| K71Stop                    | petite             | I154T                      | ts petite          |
| R189Stop                   | petite             | K157E                      | ts petite          |
| K203Stop                   | petite             | S218R                      | ts petite          |
|                            |                    | I221K                      | petite             |
| Multiple amino acid        | isolated from multiple |                           |                    |
| H44P/E114V                 | petite             | H44P                       | ts petite          |
| R79H/M247T                 | petite             | Y54F                       | petite             |
| I59V/K71E/S81N             | petite             | R79H                       | wild type          |
| C192F/E233D                | ts petite          | S81N                       | petit              |
| Q219R/D225G                | petite             | E114V                      | ts petite          |
| S190P/L228S                | petite             | V135A                      | ts petite          |
| K40R/Q60R                  | ts petite          | C192F                      | ts petite          |
| K67E/Y108N                 | ts petite          | Q219R                      | ts petite          |
| Y68H/D104G/T134S           | ts petite          | D225G                      | petite             |
| L82P/E141V                 | ts petite          | L228S                      | petite             |

Twenty-two petite or ts petite mtf1 mutants were identified by a plasmid shuffle technique (Materials and Methods). Mutants containing nonsense, missense, and multiple missense mutations were identified. Many of the multiple missense mutations were isolated so that single mutations (shown in bold) could be analyzed separately.

Figure 4. Ts mtf1 mutants are defective in all classes of mitochondrial gene transcription. Wild-type (wt) and ts petite strains bearing mtf1 mutations L53H and I154T were grown in minimal glucose medium at 30°C and shifted to 37°C. Total RNA was harvested from cells immediately before and at the indicated time intervals after the shift to the nonpermissive temperature. (A) Blots of the isolated RNAs were hybridized with labeled 14S rRNA, COB, or tRNA<sup>Thr</sup>(ACN) oligonucleotide probes to analyze the abundance of mitochondrial transcripts. (B) Hybridization to the blots was quantitated by PhosphorImager analysis to directly compare levels of transcripts. The blots were hybridized to an 18S rRNA oligonucleotide probe to normalize for loading and transfer. Strains, plasmids, and protocols are described in Materials and Methods.
Sive temperatures (36°C and 30°C, respectively). In all, nine mtf1 mutations are fully, partially, or conditionally defective for interaction with the core polymerase.

Confirmation of the two-hybrid analyses with biochemical association assays

To confirm the two-hybrid data we used a biochemical affinity assay (Mangus et al. 1994) to demonstrate that Rpo41p and Mtf1p interact in solution in the absence of DNA. For this assay the mtf1 mutants were fused to glutathione S-transferase (GST) and mutant and wild-type Mtf1p–GST fusions were bound to a glutathione agarose column (Materials and Methods). A whole-cell yeast extract containing Rpo41p was loaded onto the fusion protein columns that were washed subsequently and step-eluted to release any Rpo41p that had bound to the Mtf1p fusions on the columns (Materials and Methods). Rpo41p that was bound to and eluted from the columns was detected by an anti-Rpo41p antibody (Mangus et al. 1994).

As shown in Figure 6, the column chromatography results confirm the two-hybrid observations. As controls, the wild-type fusion and two interacting mutants, S81N and E114V (Fig. 6; data not shown) were tested for interaction. Rpo41p bound efficiently to all three fusion constructs. Consistent with the two-hybrid results, Rpo41p showed little ability to bind to the noninteracting mutants V135A and S218R (Fig. 6). Noninteracting mutants L53H, I154T, and D225G also show no Rpo41p binding in this assay (data not shown). Mutants H44P (Fig. 6) and I221K (data not shown) show intermediate levels of interaction with Rpo41p in this assay, consistent with the intermediate interaction observed in the two-hybrid assay (note β-galactosidase units in parentheses in Fig. 6). Additionally, mutant K157E, which is temperature-sensitive for interaction in the two-hybrid system, exhibited intermediate binding in this assay (data not shown).

In contrast to the results with the other conditional and partially defective mutants, the Y42C fusion protein did not interact at all with Rpo41p under the conditions of this assay (Fig. 7, top). The buffers used in all of the binding assays shown in Figure 6 contain Tris and are buffered to a pH of 7.9 at room temperature, which is pH 8.3 at the 4°C temperature used for the binding assay. Although the pH of the yeast mitochondrion has not been reported, it seemed likely that the in vivo pH ex-

Figure 5. Identification of point mutations in Mtf1p that fail to interact with Rpo41p. β-Galactosidase activity is shown for the LexA:Mtf1p construct alone and the LexA:Mtf1p construct with the VP16:Rpo41p construct. Interaction between the Mtf1p mutants and Rpo41p was determined for cells grown at 30°C (shaded bar). Activity for ts mutants Y42C and K157E is also shown for cells grown at 37°C (hatched bar) or 23°C (cross-hatched bar), respectively, to demonstrate the temperature-sensitive nature of the interaction. β-Galactosidase activity is expressed in Miller units (Miller 1972).

Figure 6. Biochemical confirmation of the noninteracting Mtf1p mutations. GST fusion constructs of wild-type and the indicated mutant Mtf1ps were isolated by glutathione agarose chromatography. Whole-cell yeast extracts containing Rpo41p were loaded onto the GST–Mtf1p columns. The columns were washed to eliminate nonspecific binding, then step-eluted to release Rpo41p bound to the column. Rpo41p in the input, wash, and elution column fractions was detected by Western blot analysis using an anti-Rpo41p antibody. Protocols are described in Materials and Methods. The percentage of two-hybrid interaction relative to wild-type Mtf1p (from Fig. 5) is shown in parentheses for each mutant.
experienced by the Y42C mutation in the nucleus for the two-hybrid assay or in the mitochondrion for the complementation assay could be significantly lower than pH 8.3. In addition, the change from tyrosine to cysteine alters the residue from hydrophobic to a potentially charged amino acid; the pK of cysteine is in the range of 8–9 depending on context (Walsh 1979). We therefore repeated the binding assay at pH 7.3 to determine if partial deprotonation of the cysteine residue had reduced binding to Rpo41p at the higher pH. As shown in Figure 7 (bottom), at pH 7.3 the Y42C mutant interacts with Rpo41p at levels equivalent to wild type. Furthermore, when this mutant is isolated at the nonbinding pH (pH 8.3) it still retains the ability to interact with Rpo41p when it is shifted to pH 7.3 (data not shown). Binding of the wild type and other mutant Mtf1p constructs is indistinguishable at the two different pHs (data not shown). The simplest explanation of these data is that partial negative charge on the deprotonated cysteine residue directly reduces the affinity of the mutant Mtf1p for Rpo41p. The reversible nature of the interaction is further support of the fact that these point mutations are not unfolded or unstable proteins—they have simply lost the ability to bind to Rpo41p at levels detectable in our assays or sufficient for transcription in vivo.

Discussion

In this work, we have confirmed further the functional homology of the mitochondrial RNA polymerase specificity factor, Mtf1p, with the large family of prokaryotic and eukaryotic nuclear σ-like factors. Despite the limited amino acid sequence similarity between many of the members of this family, there are many shared functions (for review, see Helmann and Chamberlin 1988; Helmann 1994) including roles in suppression of nonspecific interactions with DNA, selective promoter sequence recognition, promoter melting, and, as described in this work, interactions with the core RNA polymerase. The mutational strategy (PCR mutagenesis and plasmid shuffle screening) used in our studies resulted in the identification of a large number of petite and ts petite mutations useful for delineating the functional regions of Mtf1p. As shown in Figure 8, these mutations span much of the length of the MTF1 gene. The mutations that affect interactions with Rpo41p are shown above the linear map of the gene; the nine mutations fall in three discrete clusters that we have designated A, B, and C. Below the map are the six petite mutations that retain the ability to interact with Rpo41p. In several cases, the two classes of mutants are closely apposed.

Additional mutations in Mtf1p have been identified using a site-directed mutagenesis approach (Shadel and Clayton 1995). Although only three of the 14 mutations created in that study resulted in a petite phenotype, the position of the nonfunctional alterations confirmed that the regions of similarity with σ factor were essential for Mtf1p function. Our collection of 15 point mutations serves to establish further the importance of these conserved regions. Ten of the 15 petite mutations lie in the conserved regions identified previously (Fig. 8). The five mutations not localized to the conserved regions lie in the areas between regions 2.1/2.2 and 2.3/2.4, and between regions 2 and 3 (Fig. 8). Although we did not identify point mutations in the region carboxy-terminal to conserved region 3, there are likely to be additional essential residues in this part of the protein based on the deletion analysis reported by Shadel and Clayton (1995). They found that an additional 30 or more amino acids carboxy-terminal of region 3 were required for full function of Mtf1p.

One of our point mutations (Y42C) is in the same position identified as important by Shadel and Clayton (1995); another (L53H) is immediately adjacent to a mutation identified in that study (D52A). The fact that the adjacent ts petite mutant Y54F still interacts with the core (Fig. 5) means that we cannot predict whether the D52A mutation is defective for interaction or another step in the transcription reaction. This region of the pro-
tein therefore appears to have two distinct functions closely interdigitated in the folded structure. It is interesting that both mutations at amino acid 42 (Y42C and Y42R) result in a ts phenotype in vivo (Table 1; Shadel and Clayton 1995). Because we have shown that the Y42C mutation affects core interactions, this is probably also the defect for the Y42R mutation. In an in vitro binding assay we can correct the defect of the Y42C mutation by lowering the pH to create the uncharged form of cysteine (Fig. 7). The fact that both mutations are ts in vivo may therefore reflect alterations in pH or ionic environment in the mitochondrion that occur at elevated temperature. Under these altered conditions, a charged residue (cysteine or arginine) at position 42 does not support interactions with the core.

We found that the temperature shift dramatically decreases mitochondrial transcription even when wild-type Mtf1p is present. When Mtf1p is replaced by a ts noninteracting mutation (I154T or L53H), abundance of all classes of transcripts drops rapidly at the elevated temperature and does not recover to levels that support mitochondrial function. The fact that the reduction was similar for all the promoters we examined is consistent with the idea that the subunits of the holoenzyme could no longer interact in the altered conditions of the mitochondrion at the nonpermissive temperature.

We have not confirmed directly that RNA synthesis shuts off at the nonpermissive temperature. However, our observation that mitochondrial DNA levels do not change during the several cell generations represented by the extended time course of the experiment indicates that the decrease in transcript abundance is not attributable to a loss of template. This observation also calls into question the hypothesized role of the mtRNAP in replication of mitochondrial DNA (Clayton 1991). If, as we have shown, the mutated enzyme is nonfunctional for the synthesis of all classes of transcripts, how can it still be active for primer synthesis? This view is consistent with the work of Fangman et al. (1990), who have shown that mitochondrial DNA can be replicated in the absence of functional Rpo41p.

Comparison to the core interaction regions of σ factors: cluster A mutations/conserved region 2

Previous studies of the core interaction regions of bacterial σ factors have demonstrated clearly that amino acids in conserved regions 2.1 and 2.2 are of critical importance (Shuler et al. 1995; Tintut and Gralla 1995; Joo et al. 1997). The amino acid sequences of these regions of Escherichia coli σ70 and σ32, and Bacillus subtilis σE are shown in Figure 9A aligned with Mtf1p. Also included are a region of the human RAP30 subunit of RNA Pol II factor TFIIF shown to be protected when in a complex with the bacterial core RNA polymerase (McCracken and Greenblatt 1991), and a short region of E. coli σ54 demonstrated to be important for core interactions (Tintut and Gralla 1995).

Although no point mutations of σ70 have been reported that abolish core interactions, Lesley and Burgess (1989) identified several deletions that decreased core binding. A deletion that encompassed region 2.1 had the most deleterious effect, and a synthetic peptide spanning this region (indicated by the broken line in Fig. 9A) was found to bind to core polymerase (Lesley and Burgess 1989). However, this peptide also bound with similar

![Figure 9. Alignment of the cluster A mutations with known core interaction regions of σ and σ-like factors, and location of this region on the structure of σ70.](image-url)
affinities to the holoenzyme form of E. coli RNA polymerase and to $\sigma^70$ (Lesley and Burgess 1989). Severinova et al. (1996) have extended these observations using a tryptic fragment of $\sigma^70$. They found that a fragment encompassing part of region 1 and most of conserved region 2 (amino acids 114–448) bound to the core polymerase. Binding by this fragment is specific, as full-length $\sigma^70$ was able to compete with the fragment for core binding. However, the fragment binds core with an affinity 30-fold lower than full-length $\sigma^70$, suggesting that other regions also make important contributions to core binding.

Recent work has focused on regions 2.1 and 2.2 of $\sigma$ factors to identify individual residues critical for core interactions. Shuler et al. (1995) screened site-directed mutations in regions 2.1 and 2.2 of B. subtilis $\sigma^e$ and identified two point mutations (highlighted in Fig. 9A) that interfered with core interactions. Tintut and Gralla (1995) noted that although $\sigma^{54}$ and $\sigma^{70}$ share very little amino acid sequence similarity, a short motif can be found in both proteins. Mutagenesis of this region of $\sigma^{54}$ resulted in the identification of the residues highlighted in Figure 9A as critical for core interactions (Tintut and Gralla 1995). Joo et al. (1997) found that a mutation in region 2.2 in $\sigma^{32}$ reduces affinity for the core polymerase. Note that the mutations identified in these screens (shown in green) flank the highlighted mutations in cluster A of Mtf1p (shown in red), and in one case, identify the same position in the alignment. Although some of the noninteracting mutations do overlap the original synthetic peptide of Lesley and Burgess (1989), it is clear from this comparison that core interaction requires many amino acids in regions 2.1 and 2.2 in addition to those included in the peptide. This conclusion is consistent with predictions made by Gribskov and Burgess (1986) and Helmann and Chamberlin (1988) based on the high level of amino acid sequence conservation in these regions of the bacterial $\sigma$ factors.

The bacterial $\sigma$ factors and the eukaryotic nuclear factor RAP30 (McCracken and Greenblatt 1991) all form complexes with the bacterial core polymerase. As shown in Figure 9A, the amino acid sequence of this region of the different proteins has not been highly conserved. It is possible, however, that structural elements of the interaction region are similar. We have used the recently reported structure of a portion of $\sigma^{70}$ (Malhotra et al. 1996) to model the position of the cluster A noninteracting mutations from Mtf1p as well as the region 2.1 and 2.2 mutations of $\sigma^{32}$, $\sigma^e$, and $\sigma^{54}$. As shown in Figure 9B, the highlighted positions define a structural domain including the helical region of 2.1 and a turn or bend connecting this element to helical region 2.2.

It is of course difficult to predict the structure of these other $\sigma$-like factors accurately, especially as there are some single amino acid insertions and deletions in the region to be modeled (Fig. 9A). It is however of interest to note the location of the tyrosine residue at an accessible position near the beginning of the turn. We have shown that in Mtf1p, the ionization state of this residue in the Y42C mutation is critical for interaction. Most $\sigma$ factors have a bulky hydrophobic residue in this position (Londo-tto et al. 1992). The fact that $\sigma^{54}$ and RAP30 do not share this residue (Fig. 9A) indicates that the determinants for binding are more complex than simple interactions between single amino acids.

Cluster B mutations

The reported structure of $\sigma^{70}$ only extends to the carboxyl terminus of region 2 (Malhotra et al. 1996), so the additional noninteracting mutations in clusters B and C (Fig. 8) cannot be modeled relative to the cluster A mutations in region 2. However, as indicated by the arrow in Figure 9B, the folded structure does bring the carboxyl terminus of the region 2.4 helix into close juxtaposition with the region 2.1/2.2 interaction domain. This means that the noninteracting mutations that we identified in cluster B are probably very close to those in cluster A in the folded structure. Although there is no obvious amino acid sequence similarity between this region of Mtf1p and the $\sigma$ factors (Jang and Jaehning 1991), common structures may exist. The amino acid differences may be important for the selective interactions with different types of core polymerases. Because none of the original deletion mutations tested by Lesley and Burgess (1989) selectively removed this part of the protein between regions 2 and 3, this question has not yet been addressed.

Cluster C mutations/conserved region 3

The third cluster of mutations identified in our screen lie in conserved region 3. All of these mutations (S218R, I221K, and D225G) are interposed very closely with petite mutations that retain the ability to interact with the core (Q219R and L228S). The spacing between these mutations indicates that they may define two important surfaces (potentially faces of a helical or sheet structure), one of which is critical for core interactions and the other required for another essential function. There is precedent for amino acids in region 3 also having a role in core interactions. Lesley and Burgess (1989) described one deletion that removed region 3 and reduced core interactions by a factor of 5 in vitro. In addition, Zhou et al. (1992) reported that a small deletion in $\sigma^{32}$ reduced significantly the affinity for the core polymerase. Combined with the fact that we were unable to define deletions of either Rpo41p or Mtf1p that retained the ability to interact in vivo, all of these observations strongly support a model of a complex interaction surface created by distant regions in the amino acid sequence brought together in the folded structure. Because our mutagenesis was not exhaustive, it is possible that even more than the three regions that we have identified are important for interactions.

Interaction regions in the core polymerase

There is currently no information on the particular residues or regions of any core polymerase required for factor interactions. $\sigma^{70}$ has been shown to make contacts with all of the subunits of the core [$\beta$, $\beta'$, and $\alpha$ (Coggins et al.

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was cloned as a 1.0-kb Two-hybrid plasmid constructs and assays were carried out by standard methods (Guthrie and Fink 1991). Yeast cells were transformed using the lithium acetate 5-FOA medium was prepared by adding 5-FOA to synthetic media (SC) lacking the appropriate amino acids, and sporulation media (YPD) or glycerol (YPG), synthetic complete media. Standard media such as YP medium containing 2% of either glucose (YPD) or glycerol (YPG), synthetic complete media.

Materials and methods

Media and genetic methods

Standard media such as YP medium containing 2% of either glucose (YPD) or glycerol (YPG), synthetic complete medium (SC) lacking the appropriate amino acids, and sporulation medium were prepared as described by Guthrie and Fink (1991). 5-FOA medium was prepared by adding 5-FOA to synthetic medium at a concentration of 500 mg/liter (Sikorski and Boeke 1991). Yeast cells were transformed using the lithium acetate method (Ito et al. 1983). Mating, sporulation, and dissection were carried out by standard methods (Guthrie and Fink 1991).

Two-hybrid plasmid constructs and assays

MTF1 was cloned as a 1.0-kb EcoRI fragment from pJJ525 (Mangus et al. 1994) into the corresponding site of pBTM116 (Bartel et al. 1993) to create plasmid pJ832. Insert orientation was confirmed by restriction enzyme analysis. For mutant mtf1 constructions, pJ832 was digested with NsI and ligated to produce an internal deletion of 101 bp. The resulting plasmid was digested with either AffII and MscI or AffII and PstI. The fragments were treated with shrimp alkaline phosphatase. mtf1 mutations created by site-directed mutagenesis in pbBlueScript SK(+) were digested with AffII and PstI (vector sequence) and ligated to pJ832. mtf1 mutants isolated as single point mutations (I221K, L53H, I154T, and S218R) were digested with AffII and MscI, and ligated to the corresponding digest of pJ832. Resulting plasmids were tested by restriction analysis to confirm the replacement of the mtf1 deletion with the point mutation constructs. Mutants Y42C and K157E were first cloned into pbBlueScript SK(+), then the AffII to PstI fragments were cloned into pJ832.

Two-hybrid deletion constructs

For carboxy-terminal deletions, plasmid pBTM116 was digested with EcoRI and SmaI. The vector was dephosphorylated with shrimp alkaline phosphatase (U.S. Biochemical), then separated by electrophoresis on an agarose gel and extracted from the gel using Qiagen beads (Qiagen). The vector was ligated to the following MTF1 fragments of pJJ525. For the Mtf1p1-224 construct, the EcoRI to BamHI fragment of MTF1 from pJJ525 was used as the insert. The EcoRI to BgiII fragment of this same vector was used for the Mtf1p1-296 construct. And finally, the EcoRI to XmnI fragment was used to clone the Mtf1p1-315 construct. Inserts were prepared as follows. pJ525 was digested with either BamHI, BgiII, or XmnI. Blunt ends were then created by filling the overhang with Klenow (New England Bio-labs) followed by digestion with EcoRI. Fragments were separated by electrophoresis through 0.8% agarose gel and purified using Qiagen beads (Qiagen). For the amino-terminal deletion (Mtf1p52-341) a Scal to PstI (partial Scal digest) fragment from pJJ525 was inserted into the SmaI and PstI sites of pBTM116. Vector and insert were prepared as described above.

Rpo41p two-hybrid constructs were prepared using Rpo41 fragments generated from the following PCR primers. Restriction sites in the primers are underscored: NOTI-5’, 5’-CAGGCC-TGGCGGCCGCGACATGCTGAGACGCCTATAAAATC3’; NOTI-3’, 5’-CGGATCCGGCCGCGCCGGCAAAAAATTTG-ACTGTTCCTCAATAC-3’; 5’-318, 5’-GCCGGCGCCGGATGACGCCTGCAAGTTTCAAAGCTCT-3’; 3’-318, 5’-CGGCCGCCGGATGACGCCTGCAAGTTTCAAAGCTCT-3’; 3’-318, 5’-CGGCCGCCGGATGACGCCTGCAAGTTTCAAAGCTCT-3’; 3’-597, 5’-GGGCCGATTCCATCTGACGCC-3’.

The PCR fragments were cloned into TA cloning vectors (pGEMT, Promega; pCRII, Invitrogen) and then digested with the appropriate enzymes for cloning. Insert fragments as well as vector fragments were separated by electrophoresis and purified as described above. The Rpo41p1-1351 insert was prepared by the amplification of an Rpo41 plasmid using the NOTI-5’ and NOTI-3’ primers. The insert was created with a NotI digest and cloned into the corresponding site of pVP16 (Hollenberg et al. 1995). The Rpo41p1-311 insert was prepared using the BamiHI-5’ and 3’-311 primers. The insert was created with a BamHI digestion and ligated into the BamHI site of pVP16. Primers 5’-318 and NOTI-3’ were used to amplify the Rpo41p318-1351 insert. The insert was cloned into the BamHI and NotI sites of pVP16 after digestion with the same enzymes. The Rpo41p1-918 insert was created by digesting the Rpo41p1-1351 clone with NotI and MscI. The fragment was ligated to pVP16 that had been digested with EcoRI, filled in with Klenow to make blunt ends, then digested with NotI. The Rpo41p318-597 insert was created with the 5’-318 and 3’-597 primers. The insert was cloned into the BamHI site of pVP16. The Rpo41p1-918 fragment was amplified with the BamHI-5’ and 3’-597 primers. The insert was created by partial digestion with BamHI (there is a BamHI site in the Rpo41p sequence) and ligated to the corresponding site of pVP16. Two-hybrid constructs containing LexA and VP16 fusions were transformed into yeast strains AMR70 and L40, respectively (Hollenberg et al. 1995). To test the constructs for interactions, the haploid strains were mated and diploids were selected on Ura−, Trp−, Leu− medium. The strains were tested initially for β-galactosidase production by filter lift assays (Breen and Nasmyth 1985), and for growth on plates.
lacking histidine and containing 5 mM 3-aminotriazole (Phillizky and Fields 1995). For quantitative assays, strains were grown to mid-log phase in selective medium and β-galactosidase activity was assayed in permeabilized cells (Miller 1972).

Plasmid and strain construction for the MTF1 plasmid shuffle

Plasmids pJH118, pJH121, and pJH124 were constructed by inserting a 1.5-kb EcoRI fragment containing the promoter and entire coding sequence of MTF1 (Mangus et al. 1994) into the EcoRI site of pUC18, pUC7, and pBLUESCRIPT SK(±). Plasmids pJH119 and pJH142 were made by cloning this EcoRI fragment into theURA3* vector YCPlac33, and LEU2* vector YCplac111 (Gietz and Sugino 1988), respectively. To construct plasmid pJH133, a 3.8-kb BglII-BamHI fragment bearing a 3.8-kb hisG-URA3-hisG cassette was isolated from plasmid pNKY51 (Alani et al. 1987), and was inserted into the BglII site of plasmid pJH121, disrupting the MTF1-coding sequence.

The isogenic diploid yH60 was made by crossing yH58 (MATa his4A309 ura3-52 leu2-3,112) and yH59 (MATα his4A309 ura3-52 leu2-3,112) and yH60 was made by one step gene replacement (Rothstein 1983) of the 5.3-kb EcoRI fragment from pJH133. Strain yH60 was made by selecting for loss of the URA3 gene from the heterozygous MTF1/mtf1 yH61 on 5-FOA medium. To construct recipient strain yH71, strain yH64 was transformed with pJH119 containing a functional MTF1 gene and the resulting transformants were sporulated to generate mtf1:hisG (pJH119) haploid progeny whose genotype was tested on both YPG and 5-FOA media. All strain constructions were confirmed using a 1-kb EcoRI fragment of pJH117 (pJ517) as an MTF1 probe.

PCR-based mutagenesis and isolation of mtf1 mutants

Mutagenic PCR was performed essentially as described previously (Leung et al. 1989) to generate in vitro random mutations of MTF1. Plasmid pJH118 containing a 1.5-kb EcoRI fragment of MTF1 was used as a DNA template. A 17-mer of universal sequencing primer and a 23-mer oligonucleotide, 5′-CACAG-GAAACGCTATGACCATG-3′, encompassing the reverse sequencing primer were used as primers. The PCR reactions contained 30 mM Tricine at pH 8.4, 10 ng of pJH118, 0.2 mM dNTPs, 7 mM MgCl2, 0.5 mM MnCl2, 20 pmol of each primer, 5 mM β-mercaptoethanol, 0.01% gelatin, 1 unit of Taq polymerase (the gift of N. Pace, Indiana University). Reactions were done in 100 µl and preheated to 92°C for an initial denaturation step followed by 30 cycles. The PCR-amplified fragment was purified by electrophoresis on a 0.7% agarose gel and used for the introduction of mutations into yeast with the plasmid pH142 gapped with BamHI. The gap-duplex recombinant mutagenesis was performed as described by Muhlrad et al. (1992). Yeast strain yH71 was transformed to Leu+ with 500 ng of PCR-amplified fragment and 100 ng of gapped pH142. Approximately 10,000 individual Leu+ transformants were plated on 5-FOA plates to select those that lost pH119 containing the functional MTF1 gene. 5-FOA resistant Leu+ transformants were screened for loss of mitochondrial function caused by mtf1 mutations in plasmid pH142 by plating them on both YPD and YPG media. The replica plates were subsequently incubated at 30°C and 37°C for 3–6 days. Strains that did not grow on YPG medium at 30°C were isolated as mtf1 mutants, and those that grew on YPG medium at 30°C but not 37°C were also isolated and retested for the conditional phenotype. Mutated plasmids were recovered, passaged through E. coli, and sequenced by the Sanger dideoxy method (Sanger et al. 1977). To ensure that the mtf1 mutant alleles were responsible for the observed phenotypes, the recovered plasmids were transformed into yH71 followed by 5-FOA selection. When the recovered alleles contained multiple point mutations, oligonucleotide-directed mutagenesis reactions were performed to isolate the corresponding single point mutations. The mutagenic oligonucleotides were designed so that each alteration in coding sequence was accompanied by a silent mutation altering a restriction site: H44P, 5′-ATAACATCCGAAGAATTGAAGTACT-3′, delete Scal; V135A, 5′-TTTCTTACAGCTGTAAGT-3′, add PvuII; L228S, 5′-ATGGGAACCCATTTCTATTAG-3′, delete BamHI; S81N, 5′-GAAACGATCAATCTCACA-3′, add PvuII; E114V, 5′-ACTCTGATGATAGGCA-3′, add CiaI; Y54F, 5′-GAAGTGCTTGGACCTTCCATGG-3′, delete Scal; D225G, 5′-AAGATGAGGTTCCATTTTCTTT-3′, delete BamHI; C192F, 5′-CTAGATCTAAATTTCAGTA-3′, add BglII; Q219R, 5′-GATATGTCATGATGAAAG-3′, delete NsiI; R79H, 5′-AGAAAAACACTCGAGCTCT-3′, add Xhol; D2.1, 5′-CA-TAATACTTACATCCAGAAGA-3′, delete 34-42, add BglII; D2.2, 5′-CCAGAAGATTGAGATTTATATATAA-3′, delete 50-62, add BglII. Uracil-containing single-stranded DNA was produced from plasmid pJH124 in E. coli strain CJ236 (dut ung [Kunkel et al. 1987]). Each mutagenic oligonucleotide was phosphorylated with T4 polynucleotide kinase and used separately in vitro mutagenesis reactions. The resulting mutated DNAs were used to transform E. coli strain NM522 (dut ung [Kunkel et al. 1987]) to select for the mutated plasmids. Each mtf1 mutation was confirmed by sequencing and subcloned into the EcoRI site of YCPlac111 to give individual constructs and retested in yH71 as noted above.

RNA isolation and analysis

Yeast were grown to mid-log phase, collected by centrifugation and frozen as pellets in liquid nitrogen. Yeast total RNA was isolated and purified as described by Elder et al. (1983). Yeast total RNA (10–20 µg) was electrophoresed through agarose gels containing formaldehyde and blotted by capillary action to Zetabond nylon membranes (BioRad). Oligonucleotide probes specific for mitochondrial 14S, COB, and trnA1Thr and cytoplasmic 18S RNAs (Ueley et al. 1994) were end-labeled with [γ-32P]ATP (Amershag) and hybridized at 40°C in 0.5 M sodium phosphate buffer (pH 7.5) and 7% SDS. An oligonucleotide specific for 18S rRNA was used to normalize for differences in RNA samples on the blot. Signal intensity was quantitated with a PhosphorImager (Molecular Dynamics).

Western blots

Yeast cell extracts were prepared by growing cells to an OD600 of 0.6. Cells (10 ml) were harvested and resuspended in 1 ml of ice-cold Z buffer (60 mM NaHPO4, 40 mM Na2HPO4, 10 mM KCl, 1 mM MgSO4, and 50 mM β-mercaptoethanol at pH 7.0). Cells were pelleted in a microfuge and resuspended in 200 µl of ice-cold Z buffer. Glass beads (300 µl) were added and the cells were vortexed for 5 min at 4°C. The extract was cleared by a 5-min centrifugation (maximum speed) and the supernatant was collected. Protein concentrations were determined by the Bradford method (Bradford 1976) using BSA as a standard. Total protein (50–100 µg) was separated by SDS-PAGE and blotted to Immobilon-P (Millipore). Rpo41p was detected by polyclonal antibody generously provided by Dr. Roger Brent (Harvard University).
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sity, Cambridge, MA). Hybridization conditions were as described by Harlow and Lane (1988). Detection was by chemiluminescence using ECL kits from Amersham Corp.

GST-fusion plasmid constructs and GST-Mtf1p affinity chromatography

The internal AffI–BglII fragment from the mtf1 mutants was used to replace the wild-type sequence of MTF1 in pGEX-1 (plasmid pJJ526 (Mangus et al. 1994)). pJJ526 was digested with NsiI and religated to produce a deletion of 101 bp. The resulting plasmid was digested with AffI and BglII followed by dephosphorylation of ends by shrimp alkaline phosphatase (U.S. Biochemical). The vector was isolated by gel electrophoresis and ligated to AffI–BglII fragments from the MTF1 mutants. The resulting plasmids were tested by restriction enzyme analysis to confirm insertion of the full-length mutant fragments. Interaction studies with GST–Mtf1p constructs were performed as described previously (Mangus et al. 1994) with the following adjustments. Proteins were eluted with a 5-column volume step of T(500) buffer. The pH of the T(50) and T(500) solutions was 7.9 at 25°C and 8.3 at 4°C. When chromatography was performed at pH 7.3, MOPS buffer was used instead of Tris-HCl in the solutions.

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Identification of three regions essential for interaction between a ς-like factor and core RNA polymerase

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