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# Nuclear RNA Surveillance in *Saccharomyces cerevisiae*: Trf4p-dependent Polyadenylation of Nascent Hypomethylated tRNA and an Aberrant Form of 5S rRNA

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# Nuclear RNA surveillance in *Saccharomyces cerevisiae*: Trf4p-dependent polyadenylation of nascent hypomethylated tRNA and an aberrant form of 5S rRNA

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## ABSTRACT

1-Methyladenosine modification at position 58 of tRNA is catalyzed by a two-subunit methyltransferase composed of Trm6p and Trm61p in *Saccharomyces cerevisiae*. Initiator tRNA (tRNA<sub>i</sub><sup>Met</sup>) lacking m<sup>1</sup>A58 (hypomethylated) is rendered unstable through the cooperative function of the poly(A) polymerases, Trf4p/Trf5p, and the nuclear exosome. We provide evidence that a catalytically active Trf4p poly(A) polymerase is required for polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> in vivo. DNA sequence analysis of tRNA<sub>i</sub><sup>Met</sup> cDNAs and Northern hybridizations of poly(A)<sup>+</sup> RNA provide evidence that nascent pre-tRNA<sub>i</sub><sup>Met</sup> transcripts are targeted for polyadenylation and degradation. We determined that a mutant U6 snRNA and an aberrant form of 5S rRNA are stabilized in the absence of Trf4p, supporting that Trf4p facilitated RNA surveillance is a global process that stretches beyond hypomethylated tRNA<sub>i</sub><sup>Met</sup>. We conclude that an array of RNA polymerase III transcripts are targeted for Trf4p/Trf5p-dependent polyadenylation and turnover to eliminate mutant and variant forms of normally stable RNAs.

**Keywords:** RNA surveillance; exosome; tRNA turnover; polyadenylation

## INTRODUCTION

Gene transcription by eukaryotic RNA polymerases I, II, or III result in the formation of nascent pre-RNA transcripts that undergo a series of maturation steps involving exo-/endonucleolytic trimming, splicing, post-transcriptional modifications, and addition of 3' polyadenylate tail. The addition of 3' poly(A) tracts to mRNA is important in many aspects of mRNA metabolism. In eukaryotes, the conventional role of poly(A) tails has been in the stabilization, localization, and translation initiation of mRNA transcripts (Wahle and Ruesegger 1999; Proudfoot 2004). In *Xenopus laevis*, polyadenylation of maternally derived cytoplasmic mRNAs stimulates their translation during oocyte maturation (Ballantyne et al. 1997; Dickson et al. 2001). In *Escherichia coli*, poly(A) tails are not normally present in

mRNA, except when mRNA is under the process of turnover (Edmonds 2002).

This traditional view of polyadenylation in eukaryotic RNA metabolism was changed by our previous finding that it plays a role in the degradation of tRNA<sub>i</sub><sup>Met</sup> lacking 1-methyladenosine (m<sup>1</sup>A58) modification in yeast (Kadaba et al. 2004) similar to a mechanism in *E. coli* where a mutant tRNA<sup>Trp</sup> is degraded through polyadenylation (Li et al. 2002). In other instances, polyadenylated forms of other noncoding RNAs like SnRNA, SnoRNAs, and rRNAs have been observed in yeast mutants (Allmang et al. 1999; van Hoof et al. 2000b; Kuai et al. 2004).

The synthesis of m<sup>1</sup>A58 in the TΨC loop of tRNAs is catalyzed by the methyltransferase encoded by *TRM6* and *TRM61* (Anderson et al. 1998). The absence of the m<sup>1</sup>A58 modification from initiator tRNA (tRNA<sub>i</sub><sup>Met</sup>) results in its degradation (Anderson et al. 1998; Kadaba et al. 2004). The selective degradation of tRNA<sub>i</sub><sup>Met</sup> lacking m<sup>1</sup>A58 in *trm6* and *trm61* mutants is presumably due to a weakening of a tertiary interaction between D and T loops that is unique to tRNA<sub>i</sub><sup>Met</sup> (Basavappa and Sigler 1991). Trf4p, a member of nucleotidyltransferase family of proteins (Aravind and Koo-

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nin 1999), and the exosome, a multiprotein complex composed of 3'–5' ribonucleases involved in various RNA degradation and processing (Mitchell et al. 1997; Mitchell 2001; Butler 2002), function cooperatively in a novel surveillance pathway to degrade tRNA<sub>i</sub><sup>Met</sup> lacking m<sup>1</sup>A58 (hypomethylated tRNA<sub>i</sub><sup>Met</sup>) (Kadaba et al. 2004).

Previous studies from our laboratory support that Trf4p is required for the polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Kadaba et al. 2004). In accordance with this observation, recent in vitro studies have shown that Trf4p functions in an oligomeric complex composed of Mtr4p, a putative RNA helicase, and Air2p, a putative RNA binding protein, to polyadenylate in vitro synthesized RNA substrates and stimulate degradation by the exosome (LaCava et al. 2005; Vanacova et al. 2005; Wyers et al. 2005). A limitation of these studies was that the function of Trf4p as a poly(A) polymerase of tRNA substrates was studied in vitro, which may or may not correlate with its actual function in cells. In the present study we show that polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> in vivo is dependent on a functionally active Trf4p.

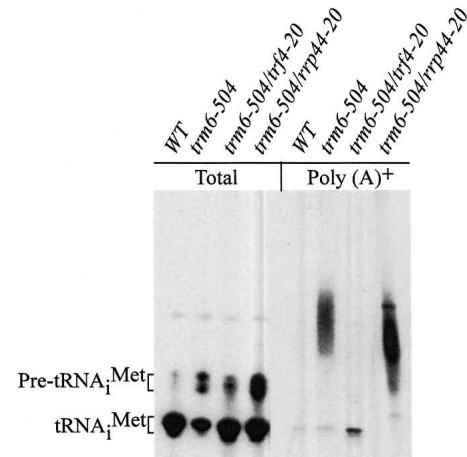
Another interesting aspect of RNA degradation by Trf4p and the exosome that remains to be elucidated is the preferential recognition of mutant forms of RNA by TRAMP and the exosome. In the present study, we show that nascent hypomethylated tRNA<sub>i</sub><sup>Met</sup> with unprocessed 5' leader and 3' extension is the substrate for polyadenylation, thus indicating that polyadenylation by Trf4p and degradation by the exosome occur very early during the biogenesis of hypomethylated tRNA<sub>i</sub><sup>Met</sup>.

Suggesting a more broadly utilized mechanism of RNA turnover, we have identified a mutant U6 SnRNA and a truncated 5S rRNAs that are stabilized by mutations in *TRF4* and the exosome, supporting the idea that adenylation and degradation of aberrant RNAs is an important step in preventing their incorporation into the cellular macromolecular machinery.

## RESULTS

### Trf4p is required for the polyadenylation of hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup>

Previously, we proposed that polyadenylation of pre-tRNA<sub>i</sub><sup>Met</sup> lacking m<sup>1</sup>A58 (hypomethylated) stimulates its degradation by the nuclear exosome based on the observation that tRNA<sub>i</sub><sup>Met</sup> is polyadenylated in a *trm6-504 rrp6Δ* double mutant, where degradation of tRNA<sub>i</sub><sup>Met</sup> is suppressed. To establish that polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> is a consequence of m<sup>1</sup>A58 hypomethylation and not a pleiotropic effect of deleting *RRP6*, we conducted Northern blot analysis of poly(A)<sup>+</sup> RNA from wild-type, *trm6-504* and *trm6-504 rrp44-20* strains. Using equivalent proportions of poly(A)<sup>+</sup> RNA from the above strains we detected poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> in a *trm6-504* strain and to a



**FIGURE 1.** Detection of polyadenylated tRNA<sub>i</sub><sup>Met</sup> in *trm6-504* and *trm6-504* suppressor mutants. Total RNA and poly(A)<sup>+</sup> RNA were isolated from wild-type (Y200), *trm6-504*(Y190), *trm6-504 trf4-20* (sup1), and *trm6-504 rrp44-20*(sup2) strains grown in YPD at 30°C as detailed in Materials and Methods. Total RNA (5 μg) and poly(A)<sup>+</sup> RNA (2–4 μg) were separated on a 6% polyacrylamide (19:1) 8 M urea gel. Membrane-bound tRNA<sub>i</sub><sup>Met</sup> was detected using a radiolabeled oligonucleotide JA11 followed by autoradiography.

greater extent in *trm6-504 rrp44-20* strain, while at the same time there was no poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> detectable in the wild-type strain (Fig. 1), indicating that tRNA<sub>i</sub><sup>Met</sup> is polyadenylated in response to a loss of m<sup>1</sup>A58. Polyadenylation was unique to hypomethylated tRNA<sub>i</sub><sup>Met</sup> and not triggered by a loss of m<sup>1</sup>A58, since we failed to detect poly(A)<sup>+</sup> tRNA<sub>UAU</sub><sup>Ile</sup> or tRNA<sub>e</sub><sup>Met</sup> (data not shown), both tRNAs that normally contain the m<sup>1</sup>A58 modification. Since the *rrp44-20* mutant is defective for degradation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Kadaba et al. 2004), it is not surprising that poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> accumulates to higher levels in the *trm6-504 rrp44-20* than in the *trm6-504* mutant alone. From this we conclude that polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> stimulates degradation by the exosome.

Previously we had observed that overexpression of Trf4p causes an increase in the length distribution of poly(A) tails on hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Kadaba et al. 2004), indicating that Trf4p positively regulates or functions as the poly(A) polymerase that adenylates hypomethylated tRNA<sub>i</sub><sup>Met</sup>. Next we wanted to determine whether Trf4p is required for the polyadenylation of hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup> in vivo. To do this, we tested whether a mutant Trf4p eliminates polyadenylation of tRNA<sub>i</sub><sup>Met</sup> in a *trm6-504* strain. Northern hybridization of poly(A)<sup>+</sup> selected RNA from *trm6-504 trf4-20* detecting tRNA<sub>i</sub><sup>Met</sup> revealed no poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> (Fig. 1), supporting the hypothesis that Trf4p functions as a poly(A) polymerase to polyadenylate only hypomethylated tRNA<sub>i</sub><sup>Met</sup>.

Since Trf4p contains a well-characterized nucleotidyltransferase domain, we wanted to determine whether the *trf4-20* lesion represents a mutation that would predictably

disrupt the poly(A) polymerase function of Trf4p. To this end, the entire *TRF4* open reading frame (ORF) of 1752 nt was amplified from genomic DNA isolated from *trm6-504 trf4-20* using PCR. Plasmid DNA from two independent bacterial clones was analyzed by DNA sequencing, and it was found that each clone contained a single adenosine nucleotide insertion that had occurred at nucleotide 795 of the *TRF4* ORF, which is predicted to cause a +1 translational frame shift during translation of the *TRF4* mRNA to produce a truncated Trf4p (~33 kDa) that is approximately half the size of the full-length Trf4p (~66 kDa). To determine whether the production of a truncated Trf4p was characteristic of other suppressors identified in our original screen and assigned to the *TRF4* complementation group, we randomly chose seven independent isolates that carry mutant *TRF4* alleles for isolation of genomic DNA, amplification and cloning of the *TRF4* ORF, and DNA sequence analysis. From this analysis, we identified three insertion mutants at the same position as *trf4-20*, one single adenosine nucleotide deletion mutant at position 794 of the *TRF4* ORF, which is predicted to cause a -1 frame shift in translational reading frame, and two nonsense mutations: one at position 1375, a T → A mutation that forms an opal codon and one at position 573 a C → A mutation that forms a ochre codon. Each of these changes should result in premature translation termination of *TRF4* mRNA, thus predicted to produce truncated Trf4p. We conclude that lesions arising in *TRF4* that result in *trm6-504* suppression represent a loss of Trf4p function through a failure to make full-length Trf4p.

### Mutation of the polymerase catalytic core in Trf4p eliminates polyadenylation of tRNA<sub>i</sub><sup>Met</sup>

Truncation of Trf4p most likely results in a decreased level or a complete loss of Trf4p, but this does not distinguish between two possible explanations for the role of Trf4p in tRNA<sub>i</sub><sup>Met</sup> degradation: (1) Trf4p functions as a poly(A) polymerase and adenylates tRNA<sub>i</sub><sup>Met</sup>, or (2) Trf4p is required as a positive effector of an unrelated poly(A) polymerase. In order to distinguish between these two possibilities and explore the enzymology of Trf4p, site-directed mutations were made in *TRF4* to change the conserved polymerase core DXD motif to AXA. Mutation of the DXD motif of Trf4p (*trf4-236*) has been shown to lead to diminished DNA and poly(A) polymerase activity in vitro (Wang et al. 2000; Vanacova et al. 2005; Wyers et al. 2005). Since the *trf4-236* mutations do not alter the expression of Trf4p proteins (Wang et al. 2002), we tested if this mutant protein can complement the cold-sensitive phenotype of a *trf4Δ* strain. The *trf4-236* allele in a single-copy number plasmid was used to transform a *trf4Δ* strain, and transformants were tested for growth at 16°C (Fig. 2A). Surprisingly, the *trf4-236* mutant supported growth of *trf4Δ* at 16°C, indicating that the polymerase function of Trf4p is

not required to complement the Cs<sup>-</sup> phenotype of a *trf4Δ* mutant, but stable expression of Trf4p is required for complementation (Fig. 2A) and also supports that the *trf4-236* protein is expressed at normal levels.

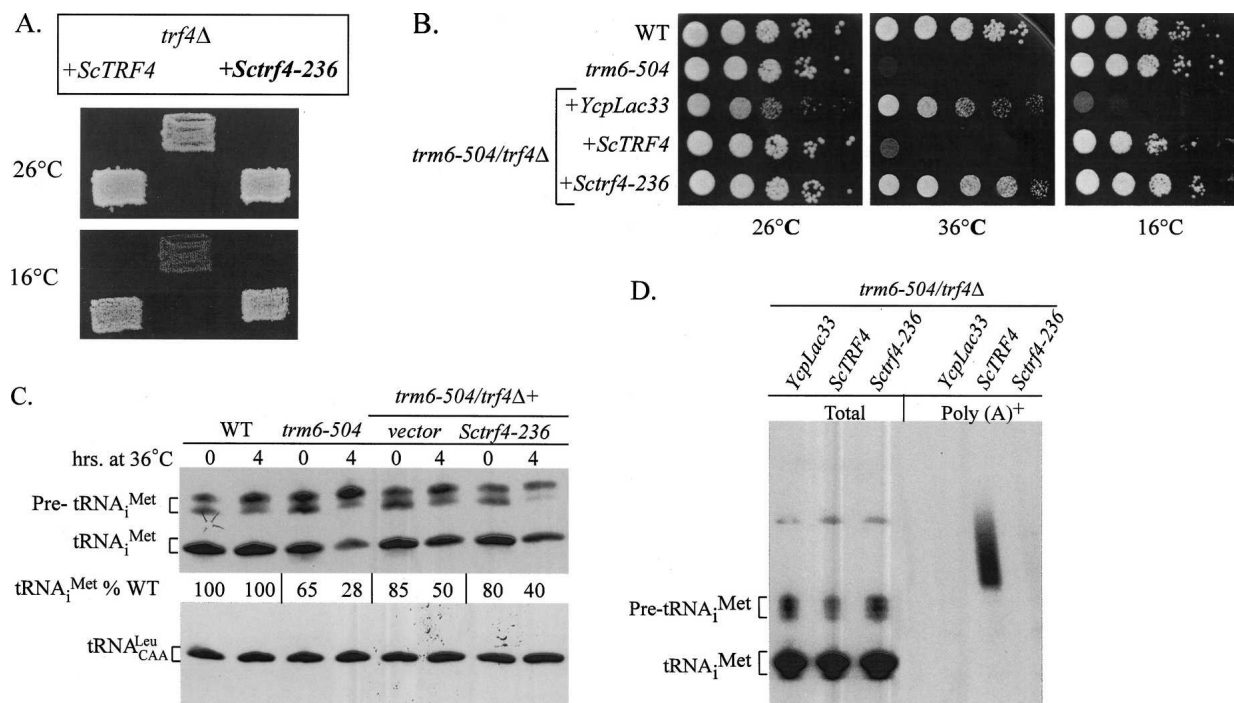
We previously showed that suppression of the temperature-sensitive (Ts<sup>-</sup>) phenotype of a *trm6-504* strain by *trf4-20* is accompanied by an increase in the steady-state levels of hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Anderson et al. 1998; Kadaba et al. 2004). In order to determine if *trf4-236* can function to complement *trf4Δ* and restore the (Ts<sup>-</sup>) phenotype of *trm6-504*, *trf4-236* on a single-copy number plasmid (*Sctrf4-236*) was introduced into a *trm6-504 trf4Δ* strain and individual transformants carrying *ScTRF4*, *Sctrf4-236*, or empty vector (YcpLac33) were plated on selective media and tested for growth at 26°, 36°, and 16°C (Fig. 2B). Comparable growth of *Sctrf4-236 trm6-504 trf4Δ* and YcpLac33 *trm6-504 trf4Δ* transformants at 36°C indicated that *trf4-236* is unable to rescue the *trf4Δ* mutant to restore the *trm6-504* Ts<sup>-</sup> phenotype; thus *trf4-236* functions as a suppressor of *trm6-504*.

To correlate the inability of *trf4-236* to complement *trf4Δ* in a *trm6-504* background with the steady-state level of tRNA<sub>i</sub><sup>Met</sup>, Northern hybridizations were done using equal amounts of total RNA from wild-type, *trm6-504*, *trm6-504 trf4Δ*, and *trm6-504 trf4Δ + Sctrf4-236* at 26° and 36°C. As seen previously, *trm6-504* contains levels of mature tRNA<sub>i</sub><sup>Met</sup> that are 65% and 28% of that seen in a wild-type strain at 26° and 36°C, respectively (Fig. 2C). The level of mature tRNA<sub>i</sub><sup>Met</sup> in *trm6-504 trf4Δ* and *trm6-504 trf4Δ + Sctrf4-236* was increased to 80%–85% of wild type at 26°C and 40%–50% of the wild type at 36°C. Since *trf4-236* resulted in a ~1.5–2-fold stabilization of hypomethylated tRNA<sub>i</sub><sup>Met</sup> at permissive and nonpermissive temperatures, we conclude that mutation of the DXD motif in Trf4p causes it to lose the ability to participate in the degradation of hypomethylated tRNA<sub>i</sub><sup>Met</sup>, presumably due to a lack of its poly(A) polymerase activity.

To determine whether *trf4-236* can function as a poly(A) polymerase to polyadenylate hypomethylated tRNA<sub>i</sub><sup>Met</sup> in vivo, poly(A)<sup>+</sup> RNA was isolated from *trm6-504 trf4Δ*, *trm6-504 trf4Δ + ScTRF4*, and *trm6-504 trf4Δ + Sctrf4-236* and subjected to Northern hybridization. While the strain containing a *ScTRF4* plasmid contained poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup>, we were unable to detect poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> in the strain bearing *Sctrf4-236* (Fig. 2D). Thus, we conclude that Trf4p is the poly(A) polymerase that polyadenylates hypomethylated tRNA<sub>i</sub><sup>Met</sup>.

### Trf4p and Trf5p have redundant functions in tRNA<sub>i</sub><sup>Met</sup> turnover

Trf4p has a structural homolog, Trf5p, that is 55% identical to Trf4p at the amino acid level, and it has been shown genetically that *TRF4* and *TRF5* are functionally redundant (Wang et al. 2000). We wanted to know whether Trf5p

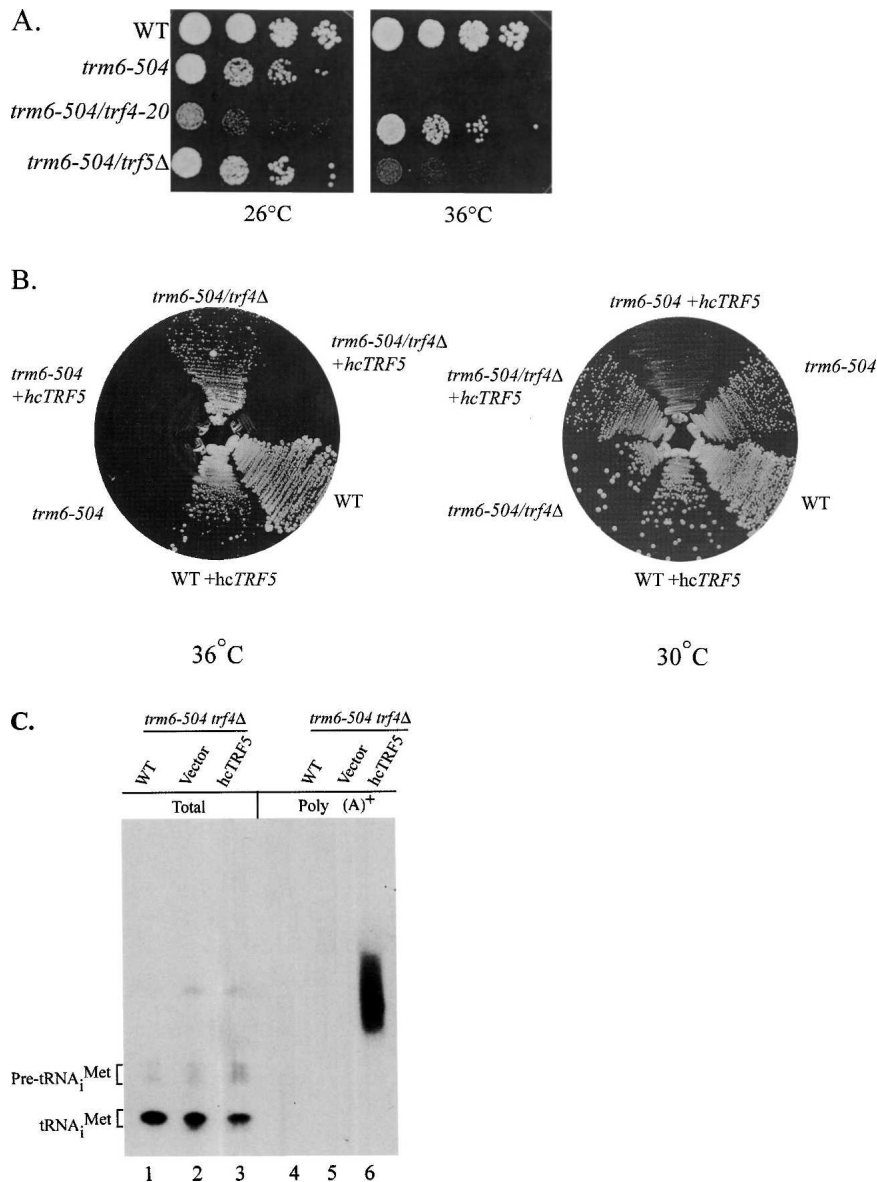


**FIGURE 2.** The DXD motif of Trf4p is crucial for its poly(A) polymerase activity and the degradation of *tRNA*<sub>i</sub><sup>Met</sup>. (A) *Sctrf4-236* complements the cold sensitivity of *trf4Δ*. A strain bearing chromosomal *trf4Δ* was transformed with YCplac33, *ScTRF4* (B254), or *Sctrf4-236* (B316), and the transformants were grown on SC-URA at 30°C, then replica-printed onto SC-URA plates and incubated at 16° and 26°C for 3 d and 1 d, respectively. (B) *trf4-236* suppresses the Ts<sup>-</sup> phenotype of a *trm6-504* mutant. A *trm6-504 trf4Δ* strain (Y301) was transformed with YCpLac33, *ScTRF4* (B254), and *Sctrf4-236* (B316); wild type (Y200) and *trm6-504* (Y190) were transformed just with YCpLac33. Individual transformants were grown to saturation in SC-URA and 10-fold serial dilutions were spotted on SC-URA plates and incubated at 26°, 36°, and 16°C for 2, 3, and 4 d, respectively. (C) Northern hybridization of total RNA (5 μg) from wild type (Y200), *trm6-504* (Y190), *trm6-504 trf4Δ* (sup1) with YCpLac33, or *Sctrf4-236* strains. Total RNA (5 μg) was separated on a 6% polyacrylamide (19:1) 8 M urea gel and transferred to a membrane and probed with JA11 to detect *tRNA*<sub>i</sub><sup>Met</sup> and JA 151 to detect *tRNA*<sub>CAA</sub><sup>Leu</sup>. The tRNAs were visualized by autoradiography and the quantities were determined by phosphorimage analysis using a STORM 640 and ImageQuant software (GE Healthcare). The quantity of *tRNA*<sub>i</sub><sup>Met</sup> in each sample is shown as a percentage of *tRNA*<sub>i</sub><sup>Met</sup> in the wild-type strain after normalization with *tRNA*<sub>CAA</sub><sup>Leu</sup>. This experiment has been repeated and the quantity of *tRNA*<sub>i</sub><sup>Met</sup> was found to be similar each time. (D) The DXD motif is important for its poly(A) polymerase activity in vivo. Total RNA (5 μg) and poly(A)<sup>+</sup> RNA (2–4 μg) isolated from the transformants described in B were separated, and a Northern hybridization was conducted to detect *tRNA*<sub>i</sub><sup>Met</sup> using a radiolabeled oligonucleotide (JA11) and autoradiography.

plays a role in the degradation of hypomethylated *tRNA*<sub>i</sub><sup>Met</sup> along with Trf4p. First we tested whether deleting the *TRF5* gene could suppress the Ts<sup>-</sup> phenotype of *trm6-504* mutant. We found that deletion of *TRF5* in a *trm6-504* strain leads to a slight increase in growth rate at 36°C (Fig. 3A), demonstrating that loss of Trf5p partially suppressed the *trm6-504* mutant phenotype. To determine whether Trf5p and Trf4p have redundant functions in the degradation of hypomethylated *tRNA*<sub>i</sub><sup>Met</sup>, we overexpressed Trf5p from a high-copy number plasmid in a *trm6-504 trf4Δ* mutant and tested growth at 30° and 36°C. If Trf5p is capable of replacing Trf4p function to polyadenylate hypomethylated *tRNA*<sub>i</sub><sup>Met</sup>, then Trf5p overexpression should restore the Ts<sup>-</sup> phenotype of the *trm6-504* mutant. We observed comparable growth levels between a *trm6-504 trf4Δ* strain overexpressing Trf5p and a *trm6-504* mutant grown at 30°C and negligible growth of both strains at 36°C (Fig. 3B), demonstrating that high-level Trf5p expression replaces Trf4p function in the degradation of hypo-

methylated *tRNA*<sub>i</sub><sup>Met</sup>. In addition to complementing *trf4Δ*, *hcTRF5* also leads to an exacerbation of the slow-growth phenotype of a *trm6-504* mutant at 30°C (Fig. 3B *trm6-504* + *hcTRF5*) like what was seen previously when Trf4p was overexpressed in the *trm6-504* mutant (Kadaba et al. 2004).

To determine if there is concomitant polyadenylation of *tRNA*<sub>i</sub><sup>Met</sup> upon restoration of the Ts<sup>-</sup> phenotype of *trm6-504* by overexpression of Trf5p, we isolated poly(A)<sup>+</sup> RNA from wild-type, *trm6-504 trf4Δ*, and *trm6-504 trf4Δ* + *hcTRF5* strains and probed the RNA by Northern hybridization to detect the presence of *tRNA*<sub>i</sub><sup>Met</sup>. We did observe polyadenylated *tRNA*<sub>i</sub><sup>Met</sup> in RNA from the *trm6-504 trf4Δ* *hcTRF5* strain, while it was not detectable in the *trm6-504 trf4Δ* strain bearing an empty plasmid (Fig. 3C). Thus, we can conclude that Trf4p and Trf5p possess a common activity that is central to the degradation of *tRNA*<sub>i</sub><sup>Met</sup>, but possibly differential expression levels or substrate preferences of Trf4p and Trf5p define how they are utilized in tRNA surveillance.



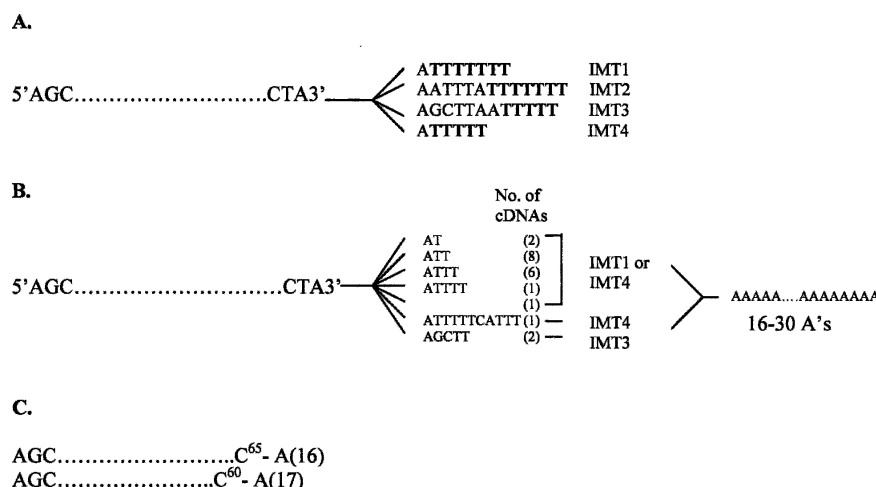
**FIGURE 3.** Trf4p and Trf5p have overlapping functions in the degradation of hypomethylated tRNA<sub>i</sub><sup>Met</sup>. (A) Yeast strains wild type (Y200), *trm6-504* (Y190), *trm6-504 trf4-20* (sup1), and *trm6-504 trf5Δ* were grown to saturation in YPD and serial 10-fold dilutions were plated onto YPD plates and incubated at 26°C or 36°C for 3 d and 2 d, respectively. (B) The same strains from A transformed with empty vector (WT, *trm6-504*, and *trm6-504 trf5Δ*) or a high-copy number plasmid containing *TRF5* (WT, *trm6-504*, and *trm6-504 trf5Δ* + *hcTRF5*) were streaked to synthetic complete media lacking uracil and incubated at 30°C and 36°C for 3 d. (C) Northern hybridization of total RNA (3 μg) and poly(A)<sup>+</sup> RNA (4–5 μg) isolated from wild-type (Y200), *trm6-504 trf4Δ* and *trm6-504 trf4Δ* + *hcTRF5*. The blot was hybridized with a radiolabeled oligonucleotide (JA11) to detect tRNA<sub>i</sub><sup>Met</sup>.

### Polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> occurs before 5' or 3' end processing

The first indication that hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup> is the target for degradation came during a high-copy suppressor analysis of a *trm6-504* mutant (Anderson et al. 1998), and subsequently, support for this hypothesis was derived from an experiment in which it was shown that the treat-

ment of poly(A)<sup>+</sup> RNA from a *trm6-504 rrp6Δ* mutant with oligo-d(T) and RNaseH produced pre-tRNA<sub>i</sub><sup>Met</sup>-sized species (Kadaba et al. 2004), leading to the conclusion that pre-tRNA<sub>i</sub><sup>Met</sup> is polyadenylated in response to a loss of m<sup>1</sup>A58. Transcription termination by RNA polymerase III occurs when a stretch of 4–6 Ts is encountered (Cozzarelli et al. 1983); in tRNA this stretch of Ts usually begins 2–7 nt downstream of the tRNA coding sequence. To determine whether polyadenylated tRNA<sub>i</sub><sup>Met</sup> were processed or unprocessed at their 3' ends, we created and cloned cDNAs of tRNA<sub>i</sub><sup>Met</sup> using oligo-d(T) primed reverse transcription (RT) of total RNA followed by specific amplification of tRNA<sub>i</sub><sup>Met</sup> using PCR (see Materials and Methods). The DNA sequences of 23 cDNAs representing tRNA<sub>i</sub><sup>Met</sup> are summarized in Figure 4. The predicted 3' ends of pre-tRNA<sub>i</sub><sup>Met</sup> derived from IMT1–IMT4 are shown with a partial sequence of tRNA<sub>i</sub><sup>Met</sup> structural RNA (Fig. 4A). The poly(A) tail length of the tRNA<sub>i</sub><sup>Met</sup> cDNAs ranged from 16 to 30 adenylates, with the majority falling between 16 and 20 adenylates (Fig. 4B). Nineteen of the 23 tRNA<sub>i</sub><sup>Met</sup> cDNAs were derived from either *IMT1* or *IMT4* genes, and two were from *IMT3*. The majority of the *IMT1* or *4* derived cDNAs contained 3' ends ending in 2 or 3 Ts (Us in the RNA) just prior to the poly(A) tail. Three *IMT1* or *4* derived cDNAs ended in a single T or no T, and one *IMT4* cDNA extended beyond the expected termination site to a second shorter stretch of Ts (Fig. 4B). Interestingly, none of the cDNAs contained the 3' CCA sequence that is added post-transcriptionally and found only in mature tRNA, suggesting that polyadenylation is limited to 3' end unprocessed hypomethylated tRNA<sub>i</sub><sup>Met</sup>.

The processing of pre-tRNAs includes, in a generally accepted temporal order, 5' end leader removal, intron removal or 3' end trailing sequence processing, and CCA addition (Hopper and Phizicky 2003). Base and nucleotide modifications can occur at any time during the processing steps of tRNA (Grosjean et al. 1997; Jiang et al. 1997), some conducted on nascent transcripts, others after addition of CCA. It is not known at which step in the processing of tRNA that m<sup>1</sup>A58 is added, but we have



**FIGURE 4.** Precursor form of hypomethylated tRNA<sub>i</sub><sup>Met</sup> is being polyadenylated. (A) An abbreviated and stylized representation of the tRNA<sub>i</sub><sup>Met</sup> coding sequence with the predicted 3' ends of each pre-tRNA<sub>i</sub><sup>Met</sup> synthesized from *IMT1-4* shown. The predicted transcription termination sites of pre-tRNA<sub>i</sub><sup>Met</sup> are shown as 5–7 tandem Ts (Us in the transcript) are shown in bold. (B) The abbreviated coding sequence is followed by the exact sequence found at the 3' end of each cDNA, and the poly(A) tail lengths are shown as a range (16–30), but the majority of the poly(A) tails were 16–19 As in length. The number of clones with the identical 3' end sequence are shown in parentheses. (C) The abbreviated coding sequence for the truncated cDNAs found end with the last nucleotide of the tRNA (C<sup>65</sup> or C<sup>60</sup>) followed by the poly(A) tail length for each.

detected m<sup>1</sup>A58 in pre-tRNAs possessing an intron (J. Anderson, unpubl.). While the presence of unprocessed 3' ends in the cDNAs supports the idea that pre-tRNA<sub>i</sub><sup>Met</sup> is targeted for degradation, it does not address how early in a tRNA life the surveillance mechanism is employed.

To better understand when during the processing of tRNA surveillance of hypomethylated tRNA<sub>i</sub><sup>Met</sup> occurs, we tested whether poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> contains the 5' leader that is present in nascent pre-tRNA<sub>i</sub><sup>Met</sup> transcripts by conducting Northern hybridizations on poly(A)<sup>+</sup> RNA from *trm6-504* and wild-type strains using probes that only recognize *IMT1-4* pre-tRNAs containing the 5' end leader sequence (Fig. 5A, 5' *IMT* probes). We detected a significant amount of tRNA<sub>i</sub><sup>Met</sup> using the 5'-specific *IMT* probes in poly(A)<sup>+</sup> RNA from *trm6-504* but not from wild type, supporting the idea that surveillance and polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> occurs very soon after the nascent transcript is formed. Removal of the 3' polyadenylate tail by oligo-d(T) hybridization and RNaseH treatment of poly(A)<sup>+</sup> pre-tRNA<sub>i</sub><sup>Met</sup> had no effect on detection of tRNA<sub>i</sub><sup>Met</sup> by the 5'-end-specific probes (Fig. 5B, lane 3), and the migration distance of 5'-end-containing tRNA<sub>i</sub><sup>Met</sup> was identical to what we had seen previously (Kadaba et al. 2004), confirming that the polyadenylated tRNA<sub>i</sub><sup>Met</sup> species contains unprocessed 5' and 3' ends. The specificity of the 5' *IMT* probes was established by stripping and re-probing the membranes with a probe that hybridizes to an internal portion of tRNA<sub>i</sub><sup>Met</sup> and detects both precursor and mature forms of tRNA<sub>i</sub><sup>Met</sup> (Fig. 5B, lanes 4–6). We

conclude from these data that Trf4p and TRAMP preferentially polyadenylate the precursor form of hypomethylated tRNA<sub>i</sub><sup>Met</sup>.

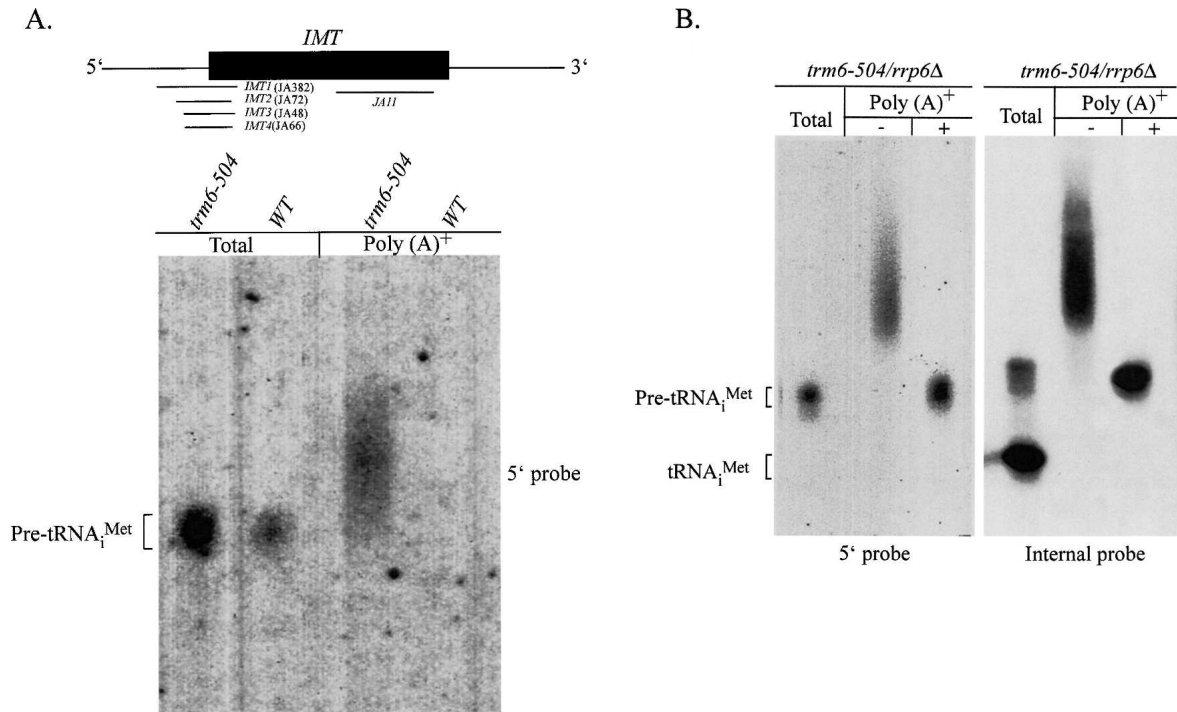
### Mutant U6 snRNA is degraded through the action of Trf4p and the exosome

In order to determine if another non-tRNA mutant RNA might be degraded by polyadenylation and the nuclear exosome, we chose to analyze whether a mutant *SNR6* gene that is rendered unstable by an internal deletion of 14 nt (*snr6Δ59-72*) (Eschenlauer et al. 1993) is targeted for degradation by Trf4p polyadenylation. *SNR6* encodes the U6 SnRNA, which is a component of the spliceosome and required for pre-mRNA splicing. *SNR6* has an internal A block element and a downstream B block element that are important for its transcription. It has been shown that deletion of residues 59–72 of

*SNR6* does not impair transcription in vitro but does result in an unstable U6 snRNA transcript in vivo (Eschenlauer et al. 1993). To test the idea that *snr6Δ59-72* is destabilized by polyadenylation and degradation by the exosome, we isolated total RNA from wild-type, *rrp6Δ*, and *trf4Δ* yeast strains carrying the *snr6Δ59-72* plasmid. Northern hybridization of total RNA from those strains showed that deletion of *RRP6* or *TRF4* resulted in a modest but significant accumulation of the *snr6Δ59-72* compared to that detected in wild type (Fig. 6), demonstrating that Rrp6p and Trf4p are important for degradation of the *snr6Δ59-72* transcript. To confirm that it is the poly(A) polymerase activity of Trf4p that was required to facilitate degradation of the mutant U6 snRNA, we conducted Northern hybridization of RNA from a *trf4Δ* containing the *snr6Δ59-72* plasmid plus either *Sctrf4-236*. The accumulation of *snr6Δ59-72* was similar in the *trf4Δ* and *trf4Δ Sctrf4-236* strains (Fig. 6), supporting that the poly(A) polymerase function of Trf4p is required for degradation of mutant U6 snRNA. The fact that we did not observe levels of mutant U6 snRNA similar to that seen for wild-type U6 snRNA may be due to a lower level of transcription in vivo or another degradation pathway that is still actively degrading *snr6Δ59-72* RNA in *trf4Δ* and *rrp6Δ* mutant strains.

### The exosome and Trf4p facilitate the degradation of 3'-end truncated 5S rRNA transcripts

The nuclear exosome plays a central role in the precise 3' end formation of many RNAs and is involved in the degra-

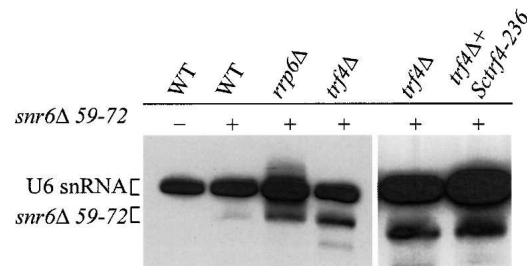


**FIGURE 5.** Polyadenylated pre-tRNA<sup>Met</sup> contains the 5' leader sequence. (A) Northern hybridization analysis of total RNA (5  $\mu$ g) and poly(A)<sup>+</sup> RNA (2–4  $\mu$ g) from *trm6-504* (Y190) and wild type (Y200). (B) Northern hybridization of total RNA (5  $\mu$ g) and poly(A)<sup>+</sup> RNA (2–4  $\mu$ g) from *trm6-504/rrp6Δ* (Y298) with (+) or without (–) pretreatment with oligo d(T) and RNaseH. The blots were hybridized with radiolabeled probes (JA382, JA72, JA66, and JA48) that recognize only pre-tRNA<sup>Met</sup> containing the 5' leader (A,B, lanes 1–3) or a radiolabeled probe (JA11) that is complementary to an internal portion of tRNA<sup>Met</sup> (B, lanes 4–6). The point of migration of pre-tRNA<sup>Met</sup> and mature tRNA<sup>Met</sup> are shown for clarity.

dation of rRNA processing intermediates (Allmang et al. 1999; Raijmakers et al. 2004). Deletion of the gene encoding a nuclear-limited exosome component, Rrp6p, results in the accumulation of polyadenylated species of snRNAs, 5S rRNA, 27S rRNA, and snoRNAs (van Hoof et al. 2000b; Kuai et al. 2004). Whether these polyadenylated RNAs are aberrant transcripts that are being degraded is not known, although it has been proposed that 27S rRNA is polyadenylated and degraded in a 5'–3' direction in an *rrp6Δ* mutant (Kuai et al. 2004). In conducting Northern hybridizations, we routinely use detection of 5S rRNA as an internal loading control, and we noted that *trm6-504* suppressor mutants possessed a small fraction of 5S rRNA that was shorter by 8–20 nt and not detected in wild-type strains (Fig. 7B, JA99). We hypothesized that this short form of 5S rRNA may be the result of aberrant processing and accumulated in *trf4Δ*, *rrp6Δ* and *rrp44-20* mutants because the surveillance mechanism that normally eliminates it is defective. In order to establish the identity of the shorter form of 5S rRNA, we used oligodeoxynucleotide probes that selectively hybridized to 5' or 3' ends of mature 5S rRNA. A probe specific to the first 20 nt of 5S rRNA (Fig. 7B, JA133) detected both the full-length mature and shorter 5S rRNA, whereas a probe that hybridizes to the last 21 nt of 5S rRNA (JA134) detected only the full-length 5S rRNA and not the faster migrating species (Fig. 7B, JA134). This result

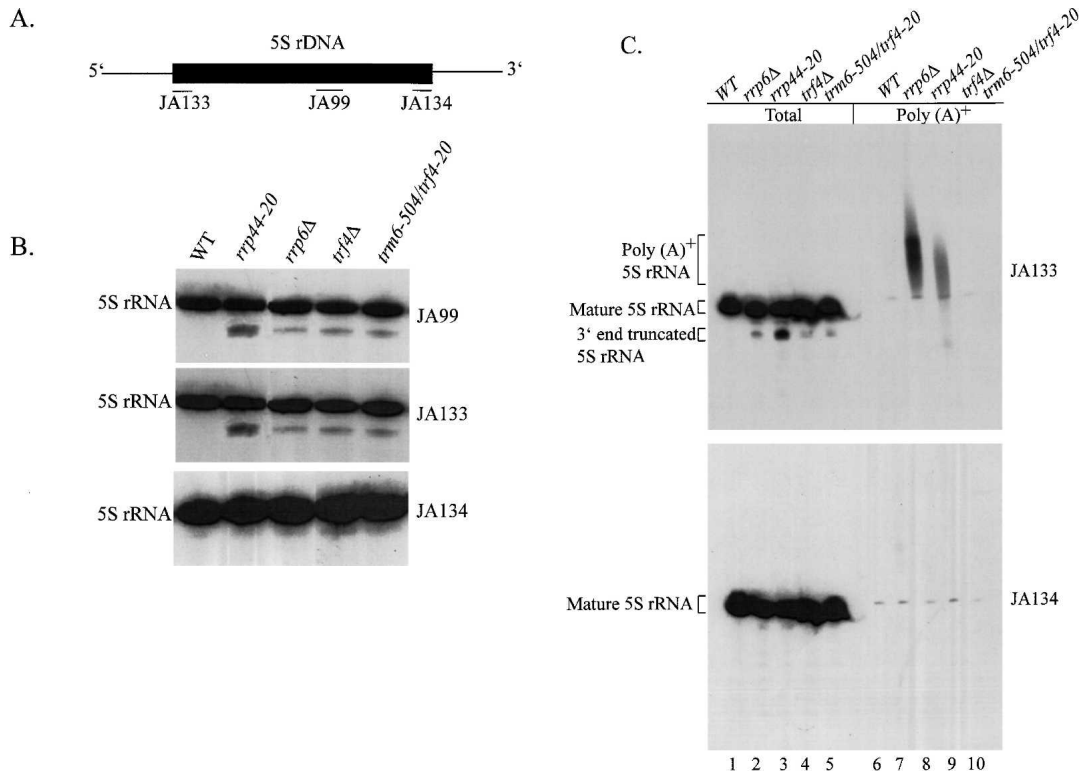
demonstrates that the short 5S rRNA accumulating in exosome and *trf4Δ* mutants is truncated at the 3' end.

To determine whether the 3' truncated 5S rRNA transcript is polyadenylated, and if Trf4p is responsible for its polyadenylation, total RNA and poly(A)<sup>+</sup> RNA from wild-type, *rrp44-20*, *rrp6Δ*, *trf4Δ*, and *trm6-504/rrp6Δ* strains were subjected to Northern hybridization analysis to detect



**FIGURE 6.** Internally truncated U6 snRNA is a substrate for Trf4p polyadenylation and degradation by the nuclear exosome. Wild type (F39), *rrp6Δ* (F23), *trf4Δ* (F22), and *trf4Δ + Sctrf4-236* were transformed with a single-copy number plasmid containing a mutant *SNR6* gene (*snr6Δ59-72*). Total RNA (5  $\mu$ g) from single transformants and a wild-type strain bearing an empty plasmid were analyzed by Northern hybridization using a radiolabeled probe (JA242) that recognizes the endogenous and plasmid-borne U6 snRNAs. The migration positions of the endogenous and *snr6Δ59-72* mutant U6 snRNAs are shown for clarity.





**FIGURE 7.** Trf4p dependent polyadenylation of 3' end truncated 5S rRNA. (A) A schematic diagram of 5S rDNA showing the coding region (filled box) and the 5' and 3' intergenic regions (lines). The approximate positions (not to scale) complementary to the oligonucleotide probes used in this study are shown as lines beneath the coding region. (B) Total RNA (5 μg) isolated from wild type (F39), *rrp6Δ* (F23), *rrp44-20* (Y303), *trf4Δ* (F22), and *trm6-504/trf4Δ* (Y301) strains grown at 30°C was used for Northern hybridization analysis. The membrane was probed with JA99 to detect mature 5S rRNA, JA133 to detect 5S rRNA containing the first 20 nt of 5S rRNA and JA134 to detect 5S rRNA containing the last 20 nt. The 5S rRNA was visualized by autoradiography. (C) Total RNA (5 μg, lanes 1–5) and poly(A)<sup>+</sup> RNA (2–4 μg, lanes 6–10) isolated from the same strains described in B was used for Northern hybridization analysis using probes JA133 (top) and JA134 (bottom). Indicated are the migration positions of the mature 5S rRNA, 3'-end truncated 5S rRNA, and poly(A)<sup>+</sup> 5S rRNA, as well as the probe used for each hybridization.

5S rRNA. The detection of poly(A)<sup>+</sup> 5S rRNA in RNA from *rrp44-20* and *rrp6Δ* using the 5'-specific probe (Fig. 7C, JA133, lanes 7,8) but not the 3'-specific probe (Fig. 7C, JA134, lanes 7,8) indicated that the polyadenylated 5S rRNA lacks the 3' end as we had predicted. Moreover, the presence of 3'-end-shortened poly(A)<sup>+</sup> 5S rRNA was found to be dependent on the expression of a functional Trf4p, since this species was undetectable in poly(A)<sup>+</sup> RNAs isolated from *trf4Δ* or *trm6-504 trf4-20* strains (Fig. 7C, lanes 9,10). We conclude from these studies that mutant and truncated RNAs are recognized, polyadenylated, and degraded by the exosome.

## DISCUSSION

We previously identified a novel tRNA surveillance pathway that utilizes polyadenylation to degrade hypomethylated tRNA<sub>i</sub><sup>Met</sup> through the function of a new poly(A) polymerase, Trf4p, and the nuclear exosome (Kadaba et al. 2004). In the present study, we confirm this by demonstrating that a functional Trf4p is required for polyadenylation and degradation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> in vivo.

We discovered that Trf5p is needed for optimal degradation of hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup>, but not to the same extent that Trf4p is required. Curiously, overexpression of Trf5p completely replaced Trf4p to cause efficient degradation of hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup>. This is in agreement with what was reported by LaCava et al. (2005), that overexpression of Trf5p in a *trf4Δ* mutant blocks the accumulation of U14 pre-snoRNA that is seen in a *trf4Δ* mutant, and Wyers et al. (2005), who showed that Trf5p accounts for a percentage of the degradation of intergenic mRNAs.

We have characterized the tRNA<sub>i</sub><sup>Met</sup> substrate of Trf4p in vivo and found that polyadenylated tRNA<sub>i</sub><sup>Met</sup> is unprocessed at its 5' and 3' ends, demonstrating that the integrity of tRNA is tested very early, possibly soon after transcription termination. We have expanded our understanding of what types of RNA are substrates for the tRNA surveillance mechanism. One such RNA, 5S rRNA, accumulates as a 3'-end-shortened polyadenylated species when the tRNA surveillance pathway is disrupted, suggesting that aberrations in 5S rRNA production or processing are rapidly eliminated in wild-type cells. This novel RNA surveillance pathway must serve a critical role in maintaining the structural or

functional integrity of most or all circulating noncoding RNAs, in addition to its role in controlling the level of intergenic mRNA that was recently reported by Wyers et al. (2005).

### The surveillance of hypomethylated tRNA<sub>i</sub><sup>Met</sup> requires two highly related poly(A) polymerases

We established that Trf4p-directed polyadenylation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> is required for its degradation by introducing mutations in Trf4p that disrupt its polymerase activity (Vanacova et al. 2005; Wyers et al. 2005) and demonstrating that hypomethylated tRNA<sub>i</sub><sup>Met</sup> is no longer polyadenylated, nor is it degraded efficiently. This is strong support for the idea that polyadenylation initiates degradation or stimulates the nuclear exosome to degrade this RNA. Our conclusion is in agreement with that drawn from in vitro experiments showing that purified nuclear exosome alone has very little RNA degradation activity (Mitchell et al. 1997); however, the addition of an oligomeric complex composed of Trf4p, Mtr4p, and Air1p/Air2p (TRAMP) with RNA substrate and exosome led to polyadenylation of the RNA substrate and stimulated RNA degradation by the exosome (LaCava et al. 2005; Vanacova et al. 2005).

The closely related proteins Trf4p and Trf5p are required to achieve a high level of tRNA<sub>i</sub><sup>Met</sup> degradation, but Trf5p is not sufficient to support degradation on its own (Fig. 3). The reason(s) for this is not known, but one explanation could be that Trf5p is expressed at a relatively low level compared to Trf4p, and only when both proteins are expressed optimally do adenylation and degradation occur efficiently. An alternative to this idea is that Trf4p and Trf5p retain the same function to adenylate tRNA<sub>i</sub><sup>Met</sup>, but they work at different stages of degradation. It is possible that Trf4p is required for initial polyadenylation of tRNA<sub>i</sub><sup>Met</sup> substrate and Trf5p plays a more important role during subsequent rounds of adenylation and degradation that has been suggested to occur (LaCava et al. 2005). Another possibility is that Trf4p and Trf5p reside in slightly different TRAMP complexes that show different binding affinities for hypomethylated tRNA<sub>i</sub><sup>Met</sup>; Trf4p TRAMP would exhibit better recognition/binding to hypomethylated tRNA<sub>i</sub><sup>Met</sup> than Trf5p TRAMP. Assuming that Trf4p and Trf5p are interchangeable in these TRAMP complexes, this would help explain why overexpression of Trf5p is required in a *trf4Δ trm6-504* double mutant to restore efficient polyadenylation and degradation of the hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Fig. 3C).

### Degradation of aberrant pre-tRNA<sub>i</sub><sup>Met</sup> and mutant U6 snRNA in vivo

We initially proposed that hypomethylated pre-tRNA<sub>i</sub><sup>Met</sup> is a target for degradation after observing stabilization of

mature tRNA<sub>i</sub><sup>Met</sup> and rapid degradation of tRNA<sub>i</sub><sup>Met</sup> precursor after treating *trm6-504* mutant cells with an RNA polymerase III transcriptional inhibitor (Anderson et al. 1998). Additional support for this original model was obtained when we observed that polyadenylated tRNA<sub>i</sub><sup>Met</sup> isolated from *trm6-504 rrp6Δ* exhibited a gel mobility identical to pre-tRNA<sub>i</sub><sup>Met</sup> after removal of the poly(A) tail by treatment with oligo d(T) and RNaseH (Kadaba et al. 2004). Here we show that poly(A)<sup>+</sup> tRNA<sub>i</sub><sup>Met</sup> isolated from *trm6-504* RNA retains the 5' end leader and 3' end trailer sequences, which strongly supports that the target for polyadenylation is nascent tRNA<sub>i</sub><sup>Met</sup> transcript (Figs. 4, 5). It is generally accepted that tRNA processing begins with removal of the 5'-end leader, followed by either intron splicing or 3'-end processing and 3'-end CCA addition to follow (O'Connor and Peebles 1991; Hopper and Phizicky 2003). The temporal position of base and nucleotide modification during tRNA processing is not well characterized, but it is likely that these modifications occur throughout the production and processing of tRNA (Grosjean et al. 1997; Hopper and Phizicky 2003). Finding that polyadenylated tRNA<sub>i</sub><sup>Met</sup> contains the 5' leader indicates that mutant tRNA<sub>i</sub><sup>Met</sup> has not undergone structural processing before it is polyadenylated and m<sup>1</sup>A58 modification probably occurs during or shortly after tRNA transcription termination.

The mechanism by which hypomethylated initiator tRNA<sup>Met</sup> is recognized for elimination is not yet known, but given that other tRNAs normally containing m<sup>1</sup>A58 are not adversely affected in its absence, it is unlikely that the surveillance mechanism detects the absence of m<sup>1</sup>A58. The dependence of tRNA<sub>i</sub><sup>Met</sup> structure on m<sup>1</sup>A58 is strongly supported by the three-dimensional structure of yeast tRNA<sub>i</sub><sup>Met</sup> (Basavappa and Sigler 1991) showing m<sup>1</sup>A58 forms N6:N7 hydrogen bonds with N1:N6 of adenosine 54; A54 is uniquely found in eukaryotic initiator tRNAs. This is different than the m<sup>1</sup>A58 N6:N7 with rT54 O2:N3, the so-called reverse Hoogsteen base-pairing found in all elongator tRNAs where the three-dimensional structure is known (for review, see Arnez and Moras 1999). The reverse Hoogsteen base pair found in both the yeast tRNA<sup>Phe</sup> and tRNA<sup>Asp</sup> three-dimensional structures is intact regardless of the presence of 1-methylA58 (tRNA<sup>Phe</sup>) (Kim et al. 1974) or its absence (tRNA<sup>Asp</sup>) (Moras et al. 1980), but we believe that the tRNA<sub>i</sub><sup>Met</sup> structure is sensitive to a loss of 1-methylA58. Interestingly, the rT/U replacement of A54 in tRNA<sub>i</sub><sup>Met</sup> in *S. cerevisiae* failed to produce a functional tRNA<sub>i</sub><sup>Met</sup> (von Pawel-Rammingen et al. 1992), supporting that the A54:m<sup>1</sup>A58 interaction in tRNA<sub>i</sub><sup>Met</sup> is contextually different than the rT54:A58/m<sup>1</sup>A58 interaction found in elongator tRNAs. We favor a model in which tRNA that is to be degraded is recognized because of an aberrant structure, rather than one that identifies tRNAs lacking m<sup>1</sup>A58 or because the pre-tRNA is not efficiently processed.

A surveillance mechanism that recognizes and facilitates degradation of aberrant non-protein-encoding RNAs

through polyadenylation might be considered protection against assembly of partial or nonfunctional RNPs. As such, the recognition of aberrant RNAs should occur prior to RNP incorporation, and this may explain why *snr6Δ59–72* RNA is unstable (Fig. 6); mutant U6 snRNA fails to be incorporated into a U4/U6 U5 tri-snRNP. The incorporation of U6 into a U4/U6 snRNP is in part due to base-pairing between U4 and U6 (Hu et al. 1995), which would be severely impaired by deleting 13 of the 24 nt in U6 that form base pairs with U4 snRNA (Hu et al. 1995). The instability of *snr6Δ59–72* RNA is partially suppressed after depletion of Trf4p or Rrp6p (Fig. 6). By partially eliminating mutant U6 degradation, some of the mutant might be incorporated into U4/U6 U5 tri-snRNP, which could cause a measurable defect in pre-mRNA splicing, although we see no evidence to support that stabilization of the mutant U6 causes a change in cell viability that would be expected if it were incorporated into a snRNP and perturbed splicing.

### Global surveillance of nuclear RNAs

Since polyadenylation of noncoding RNAs (Piper et al. 1983; van Hoof et al. 2000b) and hypomethylated tRNA<sub>i</sub><sup>Met</sup> (Kadaba et al. 2004) was reported, others have discovered that several noncoding RNAs (pre-snoRNA, pre-rRNA, and intergenic mRNA) are polyadenylated by the same surveillance machinery (LaCava et al. 2005; Wyers et al. 2005). The reason for polyadenylation of pre-snoRNA and 23S rRNA is unclear, since these precursor RNAs should be amenable to normal processing, ultimately resulting in the formation of functional, mature RNAs. It is formally possible that the polyadenylated forms of pre-snoRNAs and rRNAs observed were aberrantly structured or inaccurately processed, but this went undetected in these studies.

We identified a 3'-end-shortened 5S rRNA that is subjected to RNA surveillance. The shortened 5S rRNA was detected as a faster migrating species representing a small proportion of the total 5S rRNA produced in *trf4Δ*, *rrp44-20*, and *rrp6Δ* mutant strains. The 3'-end-shortened 5S rRNA is polyadenylated in *rrp44-20* and *rrp6Δ* strains (Fig. 7C) but not in a *trf4Δ* mutant, indicating that Trf4p is required for the polyadenylation. By approximating the size of this 3'-end-shortened 5S rRNA, we conclude that it is missing the final 8–20 nt. The processing of 5S pre-rRNA to remove nucleotides of the nascent transcript requires an endo- or a 3'–5' exonuclease (van Hoof et al. 2000a) to trim the 3' end trailing sequence, but because the mature 5' end is formed during transcription initiation (Tekamp et al. 1980; Piper et al. 1983, 1987; Lee and Nazar 1997), no enzymatic alteration of the 5' end is needed. The 5' and 3' termini form stem I, which is required for accurate processing of the 3' end trailer (Tekamp et al. 1980; Piper et al. 1983; Lee and Nazar 1997). The absence of the last 8–20 nt from 5S rRNA would eliminate stem I and could destabilize the 5S rRNA. This 3'-end-shortened 5S rRNA may be a

product of a variant 5S rDNA termed J2\* that is normally turned over very rapidly. The J2\* variant of 5S rDNA resides at the end of the tandem repeated ribosomal RNA genes on chromosome XII (McMahon et al. 1984), and it varies from the wild-type 5S rDNA at three positions near the 3' end of the 5S coding region. These alterations would disrupt two Watson–Crick base pairs (G:C1,120 and U:A3,118) in stem I and replace them with G:U base pairs, which would predictably decrease the stability of the terminal end of stem I. The destabilization could render the 3' end susceptible to a 3'–5' exonuclease activity that could easily remove the 3' end 9 nt that participate in stem I formation and result in a shortened 5S rRNA that is normally rapidly turned over. In support of this, several mutations have been described that, when introduced into stem I of a plasmid-borne copy of 5S rDNA, dramatically affect 5S rRNA stability in vivo (Lee and Nazar 1997) but not 5S rRNA transcription and processing in vitro (Brow 1987). Since the 3'-end-shortened 5S rRNA lacks the final 8–20 nt, we have no way of positively confirming it is a product of the J2\* variant 5S rDNA, but it is either j2\* 5S or incorrectly processed 5S rRNA that is usually rapidly degraded. This stands as a good example of how the surveillance mechanism that eliminates hypomethylated tRNA<sub>i</sub><sup>Met</sup> is also utilized to eliminate aberrantly structured cellular RNAs.

### Substrate recognition and an alternative degradation pathway

Understanding how Trf4p and TRAMP recognize these substrates is a critical question that remains unanswered. It is possible that an RNA chaperone like Lhp1p is involved in distinguishing between normal and abnormal RNA. Some support for this idea comes from a demonstration that the presence of Lhp1p positively influences the stability of a mutant tRNA<sup>Arg</sup> by stabilizing the abnormal anti-codon stem (Chakshusmathi et al. 2003) and hypomethylated tRNA<sub>i</sub><sup>Met</sup> that becomes extremely unstable in the absence of Lhp1p (Calvo et al. 1999; Chakshusmathi et al. 2003). Consistent with these observations, overexpression of Lhp1p partially suppressed degradation of hypomethylated tRNA<sub>i</sub><sup>Met</sup> in *trm6-504* and *trm61-2* mutants (Anderson et al. 1998; Calvo et al. 1999). Whatever the mode of recognition, the elimination of aberrant RNAs is likely to have several faces and require many factors. For instance, Lhp1p is a good candidate for an RNA chaperone of tRNA, but the natural Lhp1p binding site oligo U (Stefano 1984) at the 3' end of most RNA polymerase III transcripts (Campbell and Setzer 1992) is not found at the 3' end of mRNA or rRNA. Thus, the action of Lhp1p may be limited to nascent RNA polymerase III transcripts. Expanding our understanding of what constitutes an aberrant RNA, the mode of recognizing those RNAs, and how the process is regulated will be an important focus of the research on nuclear RNA surveillance and turnover in the next decade.

## MATERIALS AND METHODS

### Yeast strains and media

Standard genetic techniques were employed as described (Sherman et al. 1974). Yeast strains used in this study are described in Table 1. Y303 was created by crossing *trm6-504 rrp44-20* (S94) with F8 strain. The resulting diploid was sporulated in media containing 0.25% yeast extract; 1.5% potassium acetate; 0.05% glucose; 0.004% each of adenine, uracil, and tyrosine; 0.002% each of histidine, leucine, tryptophan, methionine, and arginine; 0.1% phenylalanine; and 0.35% threonine; and subjected to tetrad dissection. Haploids from the resulting tetrads were tested for Ts<sup>-</sup> and Cs<sup>-</sup> phenotypes at 37° and 16°C, respectively. *rrp44-20 TRM6* was identified from a tetrad that showed 2:2 segregation of Ts<sup>-</sup>/Cs<sup>+</sup> (*trm6-504 RRP44*) and Ts<sup>+</sup>/Cs<sup>-</sup> (*TRM6 rrp44-20*), and the presence of *rrp44-20* in the Ts<sup>+</sup>/Cs<sup>-</sup> segregants was confirmed by complementation of the Cs<sup>-</sup> phenotype of *rrp44-20* through introducing a single-copy number plasmid bearing wild-type *RRP44*. A *trm6-504 trf5Δ* (Y374) double mutant was created by gene replacement of *TRF5* with the gene that encodes resistance to G418, *KANMX*, using PCR to create a *TRF5:KANMX:TRF5* product as described (Longtine et al. 1998). The *TRF5:KANMX:TRF5* PCR product was used to transform Y190 (*trm6-504*), and G418 resistant transformants were streaked to YPD plates supplemented with 0.2 mg/mL Gentecin sulphate. The replacement of *TRF5* with *KANMX* was confirmed by PCR using the primer-specific external region of *TRF5* ORF and to the *KANMX* gene.

### Plasmid construction, *trf4* mutant mapping, and site-directed mutagenesis

Plasmids used in this study are shown on Table 2. Plasmid B288 containing a mutant U6 snRNA gene was created by subcloning a 1.4-kb SphI/PstI fragment containing *snr6-59Δ72* from the plas-

mid pΔBE6, a kind gift from Dr. David Brow (University of Wisconsin, Madison) (Eschenlauer et al. 1993) into SphI/PstI digested YcpLac 111. The high-copy number plasmid containing *TRF5* (B232) was created by subcloning a 3.1-kb PstI/BamHI from CB1455 (a generous gift from Dr. Michael Christman, Boston University School of Medicine, Boston, MA) into PstI/BamHI digested YcpLac195.

To map the *trf4* point mutants, genomic DNA was isolated from *trm6-504/trf4-20–27* mutants and the *TRF4* ORF was amplified by PCR in two independent reactions using primers JA194 and JA193 with Pfu Taq polymerase (Stratagene). The PCR product from each reaction was cloned into the pGEMT vector after generating overhanging adenosines using Taq polymerase (Promega) based on the manufacturer's specifications. Single bacterial transformants containing *trf4* mutant ORFs were screened, and at least two individual plasmids for each mutant were purified and subjected to DNA sequence analysis (MWG Biotech). The DNA sequences were aligned to the *TRF4* ORF using the Vector NTI software in order to detect the mutations. The mutation was assigned only if it was found in two or more independently generated clones obtained from the same genomic DNA.

Mutations in *TRF4* ORF were created by a PCR-based Quick Change XL Site-directed mutagenesis kit (Stratagene) in plasmid (B256) as the template using the manufacturer's specifications. Two primers containing the desired mutation were designed so that they hybridized to the same sequence on the opposite strands. The mutations introduced an AfeI restriction site that was used to screen for the desired changes in B315 (*hctrf4-236*). The presence of the desired mutations was confirmed by DNA sequence analysis of the area mutagenized. A single-copy number plasmid carrying *trf4-236* was generated by subcloning a 3.1-kb KpnI/PstI restriction fragment from B315, *trf4-236* into KpnI/PstI digested YcpLac33 to create B316.

### RNA isolation, Northern blotting, purification of poly(A) RNA, and cDNA cloning

Total RNA was isolated as described by Kohrer and Domdey (1990). RNAs were separated using a 10 cm × 14 cm × 0.8 mm 6% polyacrylamide-bis-acrylamide 8 M urea gel in 1× TBE buffer (Maniatis et al. 1982) for 1 h at 450 V. The separated RNA was transferred to a HyBond N+ membrane (Pal Biosciences) in the cold at 14 V for 3 h in 0.5× TBE unless otherwise stated. Blots were probed with radiolabeled deoxyoligonucleotides (0.1 pmol) in 5 mL of hybridization buffer (0.25M Na<sub>2</sub>HPO<sub>4</sub>, 1 mM EDTA, 1% BSA, 7% SDS). Probes were radiolabeled by incubating 10 pmol of deoxyoligonucleotide with γ-<sup>32</sup>P-ATP > 5000 Ci/mmol (Amersham Pharmacia) and T4 polynucleotide kinase (New England Biolabs) at 37°C for 1 h. Unincorporated γ-<sup>32</sup>P-ATP was separated from radiolabeled oligonucleotide by centrifugation through a 0.7-mL G-25 microspin column containing water for 1 min at 2500g. Sequences of all oligodeoxynucleotides used in this study can be found in Table 3.

**TABLE 1.** Yeast strains

Strain	Genotype	Reference
Y190	MATa <i>trm6-504, gcn2-101, his1-29, ura3-52, ino1(HIS4:LacZ, ura3-52)</i>	Kadaba et al. 2004
Y200	MATa <i>TRM6, gcn2-101, his1-29, ura3-52, ino1(HIS4:LacZ, ura3-52)</i>	Kadaba et al. 2004
SUP1	MATa <i>trm6-504, trf4-20, gcn2-101, his1-29, ura3-52, ino1(HIS4:LacZ, ura3-52)</i>	Kadaba et al. 2004
SUP2	MATα <i>trm6-504, rrp44-20, gcn2-101, his1-29, ura3-52, ino1(HIS4:LacZ, ura3-52)</i>	Kadaba et al. 2004
Y301	MATα <i>HIS3 trf4Δ:KANMX4, trm6-504</i>	Kadaba et al. 2004
Y303	<i>rrp44-20, ura3-52</i>	This study
Y374	MATa <i>trm6-504, trf5Δ:KANMX4, gcn2-101, his1-29, ura3-52, ino1(HIS4:LacZ, ura3-52)</i>	This study
F8	MATa <i>ura3-52 ino1 gcn2-1 (HIS4-lacZ, ura3-52)</i>	
F22	MATα <i>his3Δ, leu2Δ, lys2Δ, ura3Δ, trf4Δ: KanMX4</i> Res gen #11777	Invitrogen
F23	MATα <i>his3Δ, leu2Δ, lys2Δ, ura3Δ, rrp6Δ: KanMX4</i> Res gen #16265	Invitrogen
F39	MATα <i>his3Δ, leu2Δ, lys2Δ, ura3Δ</i>	Invitrogen

TABLE 2. Plasmids

Name	Description	Reference
B254	3.1-kb SspI/PmeI fragment of <i>TRF4</i> subcloned from pSup1 into SspI/PmeI YCpLac33	Kadaba et al. 2004
B256	3.1-kb SacI/PstI fragment of <i>TRF4</i> from B254 into SacI/PstI YEpLac195	Kadaba et al. 2004
B315	3.1-kb <i>TRF4</i> gene harboring <i>trf4-236</i> mutation was generated by quick change mutagenesis using B256 as template	This study
B316	KpnI/PstI digested 3.1-kb fragment of <i>trf4-236</i> from B315 subcloned into KpnI/PstI YCpLac33	This study
pΔBE6	<i>snr6</i> Δ59–72	Eschenlauer et al. 1993
B288	1.4-kb SphI/EcoRI fragment of <i>SNR6</i> subcloned from pΔBE6 into SphI/EcoRI YCpLac111	This study
CB1455	3.1-kb PstI/BamHI fragment of <i>TRF5</i> cloned into PstI/BamHI PRS315	Dr. Michael Christman
B232	PstI/BamHI 3.1-kb fragment of <i>TRF5</i> from CB1455 subcloned into PstI/BamHI YEpLac195	This study This study
B180	YCpLac111: <i>CEN4</i> , <i>LEU2</i> cloning vector	Gietz and Sugino 1988
B181	YCpLac33: <i>CEN4</i> , <i>URA3</i> cloning vector	Gietz and Sugino 1988
B184	YEpLac195: <i>URA3</i> high copy vector	Gietz and Sugino 1988

Poly (A)<sup>+</sup> RNA was purified using the same protocol described in Kadaba et al. (2004) except that poly(A)<sup>+</sup> RNA was isolated from 200-mL cultures using 0.05 g Oligo (dT) cellulose.

For the creation of cDNAs of tRNA<sub>i</sub><sup>Met</sup>, 10 μg of total RNA were treated with RNase-free DNase for 20 min at 37°C. The DNase-treated RNA was phenol-extracted and precipitated with ethanol at –20°C for 1 h. The RNA was pelleted by centrifugation, washed with 70% ethanol, and air-dried. One microgram of this RNA was combined with 50 pmol of an oligo d(T) primer (JA311), 1 mM dNTP and incubated at 80°C for 10 min, then immediately placed on ice for 5 min. Reactions were started by the addition of reverse transcriptase buffer (1× final conc.), four units of RNasin (Promega), and 20 units of M-MLV reverse transcriptase (Promega), and incubation at 42°C was carried out for 1.5 h. Two microliters of the completed cDNA reaction were combined with 2.5 μL of each primer (JA312) 20 pmol/μL, and 5′ primer (JA150) 20 pmol/μL, 2 μL dNTP (200 mM) and adjusted to a final volume of 20 μL with dH<sub>2</sub>O, after which it was incubated at 95°C for 5 min. Immediately after preincubation, 5 μL of 10× Taq DNA polymerase buffer A (Promega), 5 μL of MgCl<sub>2</sub> (25 mM), and 0.5 μL of Taq DNA polymerase, 5 U/μL (Promega), and dH<sub>2</sub>O were added to a final volume of 50 μL. The PCR reaction was carried out under the following conditions: 95°C for 30 sec, 55°C for 40 sec, and 72°C for 75

sec for 30 cycles followed by final elongation at 72°C for 7 min. The PCR product was gel-purified (Qiagen) and ligated to pGEMT-Easy vector (Promega) using T<sub>4</sub> DNA li-gase, and the ligation was used to transform DH10B electromax competent cells (Invitrogen). Colony hybridization was performed in order to identify positive colonies carrying tRNA<sub>i</sub><sup>Met</sup> clones with <sup>32</sup>P-labeled JA11 probe specific to tRNA<sub>i</sub><sup>Met</sup> according to the procedure described by Maniatis et al. (1982). Plasmid DNA from the positive colonies was analyzed by DNA sequencing (MWG) using the primer JA294. The DNA sequences were aligned to the DNA sequence of the IMT genes obtained from the *Saccharomyces* genome database using the Vector NTI software program (Invitrogen).

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TABLE 3. Oligonucleotides

Oligo	Target	Sequence
JA11	*tRNA <sub>i</sub> <sup>Met</sup>	TCGGTTTCGATCCGAGGACATCAGGGTTATGA
JA99	*5S rRNA	TCGGTATGGTCACCCACTACA
JA133	*5S rRNA	TGGTAGATATGGCCGCAACC
JA134	*5S rRNA	AGATTGCAGCACCTGAGTTTC
JA151	*tRNA <sub>C<sup>AA</sup></sub> <sup>Leu</sup>	TGGTTGCTAAGAGATTCCGAAC
JA48	*5′ IMT3	ACGGCGCTTAACTTTTATG
JA66	*5′ IMT4	GGCGCTTAGCCAACCTG
JA72	*5′ IMT2	GGCGCTGCTAAATCATGAG
JA382	*5′ IMT1	CACGGCGCTGCTAATCTTGCAATGCCTCT
JA114	*tRNA <sub>U<sup>AU</sup></sub> <sup>Ile</sup>	TGCTCGAGGTGGGGWTTGAACCCACGACGG
JA15	*tRNA <sub>e</sub> <sup>Met</sup>	TGCTCCAGGGGAGGTTCCGAACCTCTCGACC
JA275	<i>trf4-236</i>	CCGATTTGTATTTGCCGTGATCGCCTATTGCTT CGGTGGTAACGAGCG
JA276	<i>trf4-236</i>	CGCTCGTTACCACGCAAGCAATAGCGCTA CCAGGCAAATACAAATCGG
JA311	oligo d(T) adaptor	GGCCACGCGTCGACTACTTTTTTTTTT TTTTTTTTT
JA310	Adaptor	GGCCACGCGTCGACTAC
JA150	tRNA <sub>i</sub> <sup>Met</sup>	TGGCGCAGTGGAAAGCGC
JA193	<i>TRF4</i>	GATGTACAGTTCAGTGCATC
JA194	<i>TRF4</i>	GTGTGAATAAGCAAGGGAAC
JA346	5′ TRF5 deletion	TTATTTTTCAAATAAAACAAACGAGGGCGG AGTTTATTGGGTCGCTGACCGCTGCAGGTCGAC
JA347	3′ TRF5 deletion	AAGAGCCCTGGCCTTTAGAGAGCCAGTA GTCCCTTCTCGTTTGATCATCGATGA ATTCGAGCTC

\*Deoxyoligonucleotides used in Northern hybridization.

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