

# The box H+ACA snoRNAs carry Cbf5p, the putative rRNA pseudouridine synthase

Denis L.J. Lafontaine, Cécile Bousquet-Antonelli,<sup>1</sup> Yves Henry,<sup>1</sup> Michèle Caizergues-Ferrer,<sup>1</sup> and David Tollervey<sup>2</sup>

Institute of Cell and Molecular Biology, University of Edinburgh, King's Buildings, Edinburgh EH9 3JR, UK; <sup>1</sup>Laboratoire de Biologie Moléculaire Eucaryote, Centre National de Recherche Scientifique (CNRS), 31062 Toulouse Cedex, France

Many or all of the sites of pseudouridine ( $\Psi$ ) formation in eukaryotic rRNA are selected by site-specific base-pairing with members of the box H + ACA class of small nucleolar RNAs (snoRNAs). Database searches previously identified strong homology between the rat nucleolar protein Nap57p, its yeast homolog Cbf5p, and the *Escherichia coli*  $\Psi$  synthase *truB*/P35. We therefore tested whether Cbf5p is required for synthesis of  $\Psi$  in the yeast rRNA. After genetic depletion of Cbf5p, formation of  $\Psi$  in the pre-rRNA is dramatically inhibited, resulting in accumulation of the unmodified rRNA. Protein A-tagged Cbf5p coprecipitates all tested members of the box H + ACA snoRNAs but not box C + D snoRNAs or other RNA species. Genetic depletion of Cbf5p leads to depletion of all box H + ACA snoRNAs. These include snR30, which is required for pre-rRNA processing. Depletion of Cbf5p also results in a pre-rRNA processing defect similar to that seen on depletion of snR30. We conclude that Cbf5p is likely to be the rRNA  $\Psi$  synthase and is an integral component of the box H + ACA class of snoRNPs, which function to target the enzyme to its site of action.

[Key Words: RNA modification; pre-rRNA processing; ribosome synthesis; yeast]

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The rRNAs of all organisms undergo extensive covalent nucleotide modification. In eukaryotes, the rRNAs are generated by post-transcriptional processing of large pre-rRNA species, and these modified nucleotides are formed in the pre-rRNAs, rather than in the mature rRNAs. The most numerous modifications are methylation of the 2'-hydroxyl residue in the ribose moieties (2'-O-methylation) and isomerization of uracil residues to pseudouridine ( $\Psi$ ). Formation of  $\Psi$  residues is thought to occur through base rotation about the C<sub>3</sub>-C<sub>6</sub> axis after cleavage of the glycosyl bond (Goldwasser and Henrikson 1966; for review, see Ofengand et al. 1995). Additionally, a few positions are modified at the base level; the best described example being the universally conserved m<sup>6</sup>Am<sup>6</sup>A doublet at the 3'-end of the 18S rRNA (Lafontaine et al. 1995). Recent data have shown that, in eukaryotes, the sites of both 2'-O-methylation (Cavaillé et al. 1996; Kiss-László et al. 1996; Nicoloso et al. 1996) and  $\Psi$  formation (Ganot et al. 1997a; Ni et al. 1997) in the rRNAs are selected by site-specific base-pairing of small nucleolar RNAs (snoRNAs) to the pre-rRNAs (for review, see Maden 1996; Tollervey 1996; Bachelier and Cavaillé 1997; Smith and Steitz 1997). The snoRNAs involved can be separated into two major groups, which

are designated box C + D snoRNAs and box H + ACA snoRNAs on the basis of conserved sequence elements (Balakin et al. 1996). In the case of 2'-O-methylation, base-pairing of a member of the box C + D class of snoRNAs across the site of methylation positions a conserved sequence element, box D, at a precise distance of 5 bp from the nucleotide to be modified (Cavaillé et al. 1996; Kiss-László et al. 1996; Nicoloso et al. 1996). Presumably, proteins associated with box D use this positional information to select the site of modification. In the case of  $\Psi$  formation, a member of the box H + ACA class of snoRNAs base pairs to nucleotides flanking the substrate uracil, leaving the base of the nucleotide free for interaction with the modifying enzyme (Ganot et al. 1997a; Ni et al. 1997).

Both major groups of snoRNAs are associated with specific proteins in small nucleolar ribonucleoprotein (snoRNP) particles. All box C + D snoRNAs are associated with the protein fibrillarin (Nop1p in yeast) (Lischwe 1985; Schimmang et al. 1989; Ganot et al. 1997b; for review, see Maxwell and Fournier 1995), and mutations in Nop1p can globally block 2'-O-methylation of the yeast rRNAs (Tollervey et al. 1993). Similarly, all box H + ACA snoRNAs are associated with Gar1p (Girard et al. 1992; Balakin et al. 1996; Ganot et al. 1997b), and mutations in Gar1p can globally inhibit  $\Psi$  formation in the rRNA (Bousquet-Antonelli et al. 1997).

<sup>2</sup>Corresponding author.

E-MAIL d.tollervey@ed.ac.uk; FAX 44 131 650 7040.

Nop1p and Gar1p are not required for the synthesis or stability of the snoRNAs with which they are associated. Gar1p is, however, required for the stable association of the box H + ACA snoRNAs with the pre-rRNA (Bousquet-Antonelli et al. 1997); this association presumably underlies the requirement for Gar1p in  $\Psi$  formation.

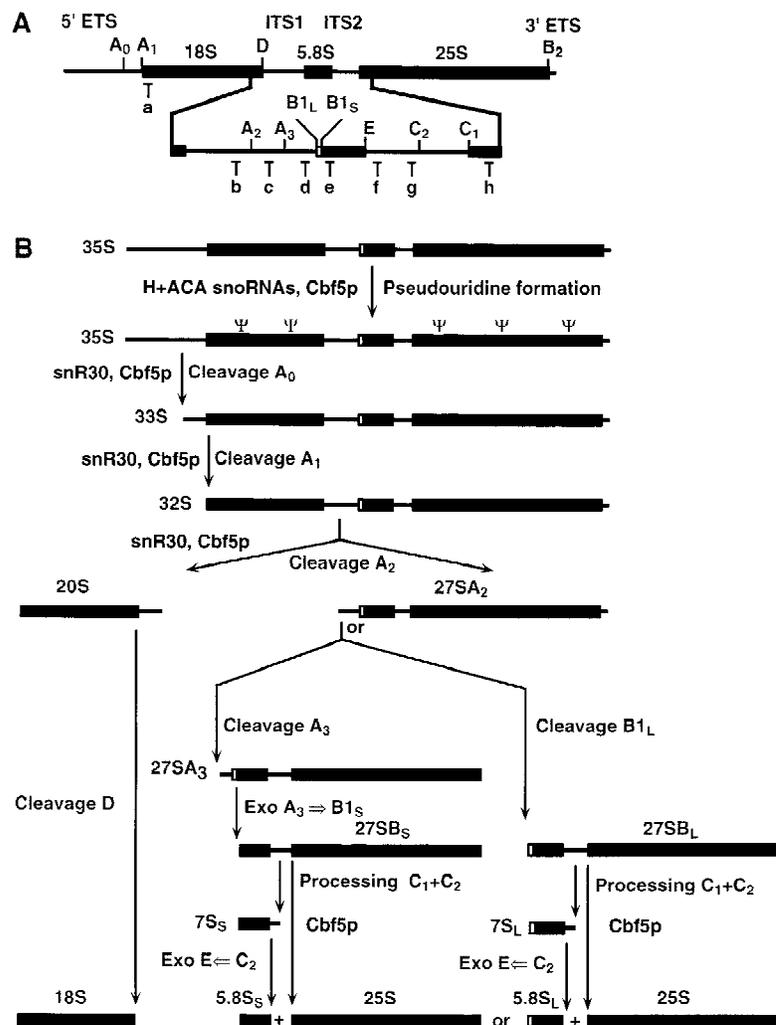
Both classes of snoRNAs include species that probably do not act to select sites of rRNA modification but are required for processing of the pre-rRNA. Genetic depletion of the box C + D snoRNAs U3 (Hughes and Ares 1991) or U14 (Li et al. 1990) or of the box H + ACA snoRNA snR30 (Morrissey and Tollervey 1993), inhibits the early pre-rRNA processing reactions at sites A<sub>0</sub>, A<sub>1</sub>, and A<sub>2</sub>, preventing the synthesis of the 18S rRNA (see Fig. 1). For this reason each of these snoRNAs is essential for cell viability. In contrast, none of the snoRNAs that direct rRNA modification is essential for cell viability, although the absence of the  $\Psi$  guide snoRNA, snR10, leads to some cold sensitivity (Tollervey 1987; Ni et al. 1997).

One of the major unresolved questions concerning these systems of snoRNA-directed modification is the

relationship between the snoRNAs and the modifying enzymes. Specifically, are the enzymes free components that recognize the structure created by the snoRNA-pre-rRNA duplexes in much the same way tRNA-modifying enzymes and bacterial rRNA-modifying enzymes recognize their RNA substrates? Or are the enzymes physically associated with the snoRNAs, which act directly to target the enzymes to their sites of action?

In contrast to  $\Psi$  formation in the eukaryotic pre-rRNA, the formation of  $\Psi$  in tRNAs and bacterial rRNAs is not known to involve RNA cofactors. In these cases, multiple  $\Psi$  synthases exist; four tRNA  $\Psi$  synthases have been characterized in *Escherichia coli* (*truA*, *truB*, *rsuA*, and *rluA*), each of which modifies with high specificity a single site or a number of sites with very similar structures (Kammen et al. 1988; Nurse et al. 1995; Wrzesinski et al. 1995a,b; Simos et al. 1996). A database search (Koonin 1996) revealed that each of these enzymes is a member of a distinct, evolutionarily conserved family of  $\Psi$  synthases. *E. coli* *truB*/P35, which converts U55 to  $\Psi$ 55 in the m<sup>5</sup>U $\Psi$ CG loop in most tRNAs (Nurse et al. 1995), is strongly homologous to two yeast proteins, Cbf5p and

**Figure 1.** Structure of the yeast pre-rRNA and its processing pathway. (A) The 35S pre-rRNA. The sequences encoding the mature 18S, 5.8S, and 25S rRNAs (thick lines) are flanked by the 5' and 3' ETSs and separated by internal transcribed spacers 1 and 2 (ITS1 and ITS2). Sites of pre-rRNA processing are indicated with uppercase letters (A<sub>0</sub> to E) and the positions of hybridization of the oligonucleotides used are indicated with lowercase letters (a-h). (B) The pre-rRNA processing pathway. Processing of the primary 35S precursor starts at site A<sub>0</sub>, yielding 33S pre-rRNA. This molecule is subsequently processed at sites A<sub>1</sub> and A<sub>2</sub>, giving rise successively to the 32S pre-rRNA and to the 20S and 27SA<sub>2</sub> precursors. Cleavage at A<sub>2</sub> separates the pre-rRNAs destined for the small and large ribosomal subunit. The 20S precursor is then endonucleolytically cleaved at site D to yield the mature 18S rRNA. The 27SA<sub>2</sub> precursor is processed by two alternative pathways, forming the mature 5.8S and 25S rRNAs. The major pathway involves cleavage at a second site in ITS1, A<sub>3</sub>, rapidly followed by exonucleolytic digestion to site B1<sub>S</sub>, generating the 27SB<sub>S</sub> precursor. Approximately 15% of the 27SA<sub>2</sub> molecules are processed by the second pathway at site B1<sub>L</sub>, producing the 27SB<sub>L</sub> pre-rRNA. At the same time as processing at B1 is completed, the 3' end of mature 25S rRNA is generated by processing at site B<sub>2</sub>. The subsequent processing of both 27SB species appears to follow a similar pathway. Cleavage at sites C<sub>1</sub> and C<sub>2</sub> releases the mature 25S rRNA and the 7S pre-rRNAs, which undergo rapid 3' → 5' exonuclease digestion to site E generating the mature 3' end of 5.8S rRNA. Cbf5p is required for the early cleavages at sites A<sub>1</sub> and A<sub>2</sub>; loss of these cleavages inhibits formation of the 20S and 27SA<sub>2</sub> pre-rRNA preventing synthesis of 18S rRNA. Cbf5p is also required for efficient processing at site A<sub>0</sub> and efficient processing of the 27SB and 7S pre-rRNAs in ITS2.

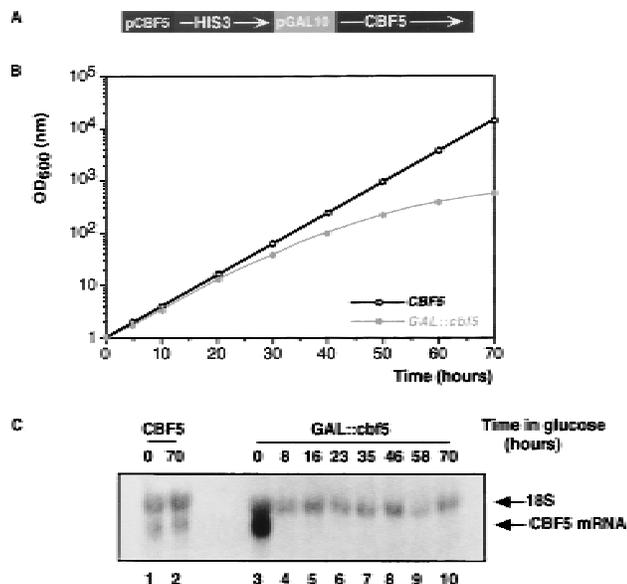


YNL480 (Koonin 1996). Recent data have shown that YNL480 (now designated Pus4p) is the yeast tRNA  $\Psi$ 55 synthase (Becker et al. 1997). Yeast Cbf5p was originally characterized as an essential protein that showed in vitro binding to centromeres and microtubules (Jiang et al. 1993). Subsequently, Cbf5p was found to be highly homologous (64% identity, 79% homology) to the rat nucleolar protein Nap57p and to be localized to the yeast nucleolus (Meier and Blobel 1994). This suggested that Cbf5p and Nap57p might be the rRNA  $\Psi$  synthases in yeast and mammals, respectively. We have therefore investigated this possibility and report here an analysis of the role of Cbf5p in pre-rRNA processing and formation of  $\Psi$  in the rRNA.

## Results

### Construction of a conditional *CBF5* allele

Because *CBF5* is an essential gene (Jiang et al. 1993), we first constructed a conditional allele by placing its expression under the control of an inducible *GAL10* promoter using the one-step PCR method described previously (Lafontaine and Tollervey 1996) (Fig. 2A). On permissive medium [2% raffinose, 2% sucrose, and 2% galactose (rsg)], the growth rate of the *GAL::cbf5* strain is identical to that of the otherwise isogenic parental *CBF5*



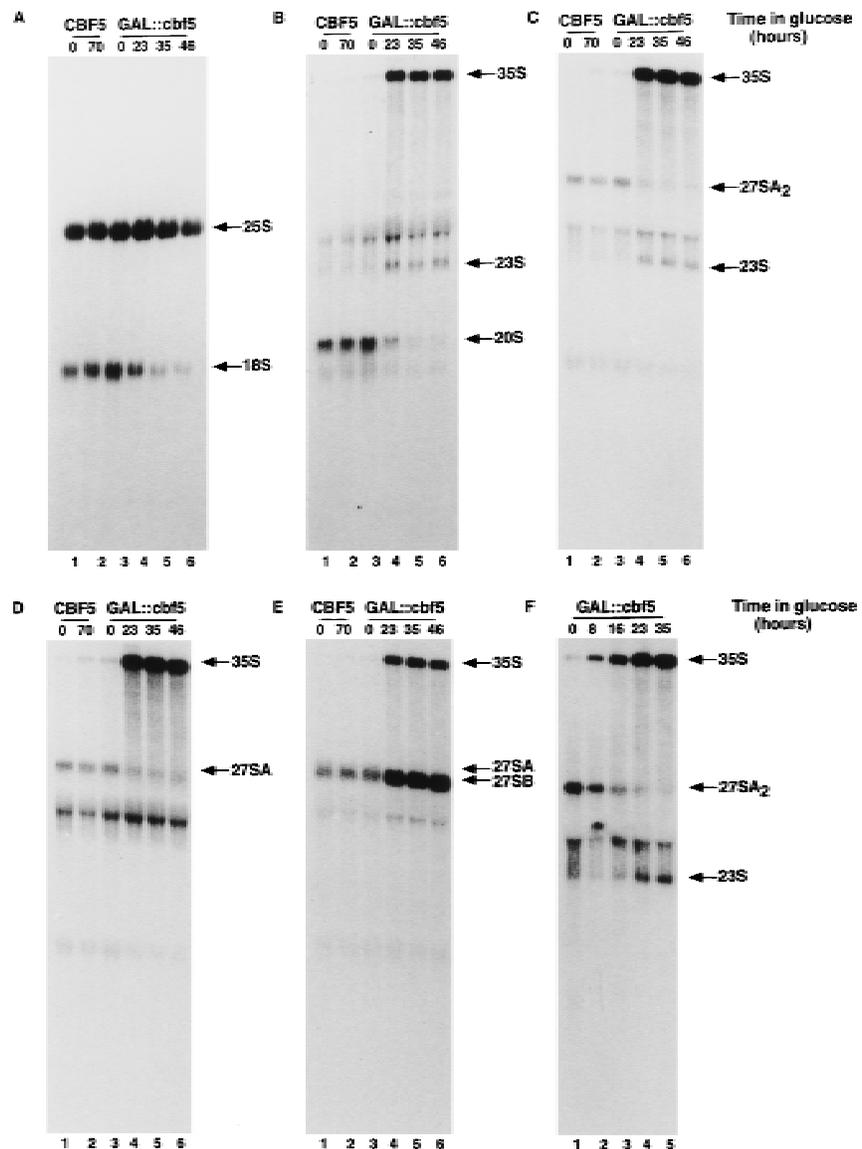
**Figure 2.** Genetic depletion of Cbf5p. (A) Schematic representation of the *GAL::cbf5* allele. (B) Growth of the *GAL::cbf5* (●) and *CBF5* (○) strains following transfer to glucose medium. Cell density was measured at regular intervals, and the cultures were periodically diluted to be continuously kept in exponential growth. (C) Northern hybridization of the *CBF5* mRNA from the *CBF5* strain (lanes 1, 2) and the *GAL::cbf5* strain (lanes 3–10), following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8 to 70-hr lanes). The positions of the *CBF5* mRNA and 18S rRNA are indicated. As expected from the size of its ORF (1452 bp), the *CBF5* mRNA is detected slightly under the 18S rRNA.

strain (doubling every 3 hr). Under these conditions, the level of the *CBF5* mRNA in the *GAL::cbf5* strain is approximately fivefold higher than in the *CBF5* control strain (Fig. 2C, lane 3). This level is expected for a gene whose transcription is driven by the strong *GAL10* promoter. Following transfer of the *GAL::cbf5* strain to glucose (glu) medium the *CBF5* mRNA was rapidly depleted, no mRNA was detected 8 hr after transfer to glu medium (Fig. 2C), and growth slowed progressively, commencing 20 hr after transfer (Fig. 2B). This slow onset in the growth impairment is characteristic of *GAL* depletion of components required for ribosome synthesis.

### *Cbf5p* is required for pre-rRNA processing

Northern hybridization (Fig. 3) shows that the levels of the mature 18S and 25S rRNAs, and all pre-rRNA species are identical in the *CBF5* control strain and in the *GAL::cbf5* strain grown in permissive medium (0 hr lanes in Fig. 3). Following transfer to glu medium (Fig. 3A, 23- to 46-hr lanes) the mature 18S rRNA is progressively depleted in the *GAL::cbf5* strain; at later time points depletion of the 25S rRNA is also observed. Analysis of the pre-rRNAs (Fig. 3B–E) shows that the 35S primary transcript is strongly accumulated, while the 27SA<sub>2</sub> and 20S pre-rRNAs are depleted and an aberrant RNA species, the 23S pre-rRNA, is detected. The 23S RNA extends from the 5' end of the 35S to site A<sub>3</sub> (Fig. 1) and is generated by direct cleavage of the 35S pre-rRNA at site A<sub>3</sub> in the absence of prior cleavage at sites A<sub>0</sub>, A<sub>1</sub>, and A<sub>2</sub>. These effects are characteristic of mutations that inhibit the early pre-rRNA cleavages at sites A<sub>0</sub>, A<sub>1</sub>, and A<sub>2</sub> (see Fig. 1) and were observed following depletion of several different snoRNAs, including the box H + ACA snoRNAs snR10 (Tollervey 1987) and snR30 (Morrissey and Tollervey 1993), and on depletion of Gar1p (Girard et al. 1992), which is associated with the entire class of box H + ACA snoRNAs (Balakin et al. 1996). Depletion of Cbf5p also leads to accumulation of the 27SB pre-rRNAs (Fig. 3E), consistent with the reduction in the levels of the 25S rRNA. A shorter time course following transfer to glucose medium (Fig. 3F) shows that the level of the 35S pre-rRNA is elevated and 27SA<sub>2</sub> is reduced as early as 8 hr (<3 generations) after transfer to glu medium, indicating that these effects rapidly follow the loss of Cbf5p. Analysis of low molecular weight RNA (Fig. 4A) shows that the 7S pre-rRNA is also strongly accumulated following depletion of Cbf5p. This accumulation of the 27SB and 7S pre-rRNAs was not observed on depletion of any characterized snoRNA. In addition, a 5.8S rRNA species intermediate in length between 5.8S<sub>L</sub> and 5.8S<sub>S</sub> was observed on depletion of Cbf5p (\* in Fig. 4B). This has also not been observed previously in snoRNA mutants.

Pre-rRNA processing was further analyzed by primer extension. Analysis of the 5' external transcribed spacer (ETS) shows an increase in the stop corresponding to the 5' end of the 35S pre-rRNA (position +1, Fig. 5A) in the *GAL::cbf5* strain following transfer to glu medium (23-



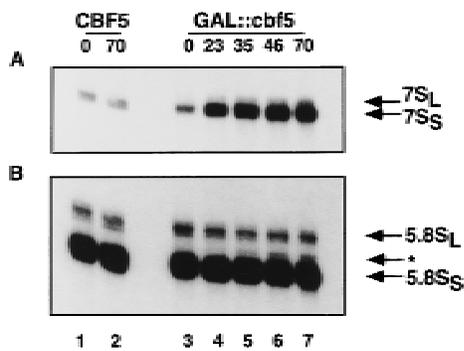
**Figure 3.** Northern analysis of rRNA and pre-rRNA synthesis in a *GAL::cbf5* strain. (A) Probes against mature 25S and 18S rRNA (oligonucleotides a and h); (B) probe against the 5' region of ITS1 (oligonucleotide b), (C); probe against ITS1 between sites A<sub>2</sub> and A<sub>3</sub> (oligonucleotide c); (D) probe against the 3' region of ITS1 (oligonucleotide d); (E) probe against the 5' region of ITS2 (probe f). (F) Shorter time course following transfer to glu medium, hybridized with oligonucleotide c. The oligonucleotides used are depicted in Fig. 1A. RNA was extracted from the *CBF5* and *GAL::cbf5* strains following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8–70-hr lanes) and separated on a 1.2% agarose gel containing formaldehyde.

to 70-hr lanes), consistent with the 35S accumulation detected by Northern hybridization (Fig. 3B–F); however, this was not accompanied by a loss of the pre-rRNA cleaved at site A<sub>0</sub> (Fig. 5B). The primer-extension stop at site A<sub>0</sub> is elevated on depletion of Cbf5p. The accumulation of the 35S pre-rRNA and 23S RNA indicates that cleavage at site A<sub>0</sub> is at least delayed on depletion of Cbf5p. We interpret this observation as showing that cleavages at sites A<sub>1</sub> and A<sub>2</sub> are more sensitive to the depletion of Cbf5p than is cleavage at site A<sub>0</sub>. This phenomenon was observed in strains depleted of snR10, snR30, or Gar1p, but not in strains depleted of U3 (Beltrame et al. 1994).

Primer extension through sites in ITS1 shows the loss of the stop corresponding to cleavage at site A<sub>2</sub>, the 5' end of the 27SA<sub>2</sub> pre-rRNA, and shows the increase of the stops at sites B1<sub>S</sub> and B1<sub>L</sub>, the 5' ends of both the 27SB<sub>S</sub> and 27SB<sub>L</sub> and 7S<sub>S</sub> and 7S<sub>L</sub> pre-rRNAs, respec-

tively (Fig. 6). These observations are in good agreement with the results of Northern hybridization in Figures 3 and 4. The accumulation of the long and short forms of 27SB and 7S, shown by the relative stops at B1<sub>S</sub> and B1<sub>L</sub>, are not obviously different (Fig. 6, cf. A with C, which shows a shorter exposure of the same gel). The level of the 27SA<sub>3</sub> pre-rRNA, shown by the primer-extension stop at site A<sub>3</sub>, is elevated on depletion of Cbf5p (Fig. 6B). This elevation was also observed in strains depleted of snR30, but was not seen on depletion of any other snoRNP component tested (Morrissey and Tollervey 1993).

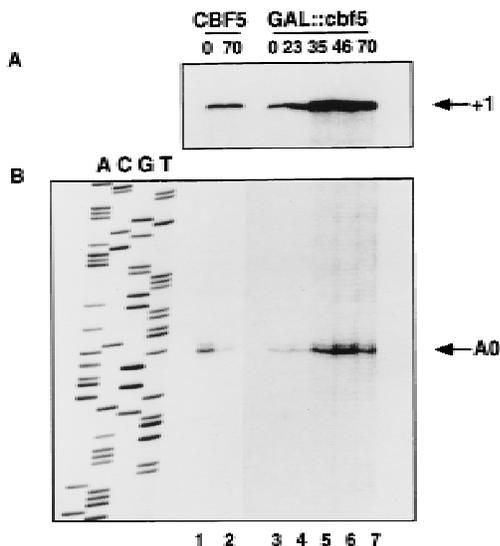
No defects in pre-tRNA processing or accumulation of mature tRNAs were detected by Northern hybridization with probes specific for the intron-containing precursor of tRNA<sup>Trp</sup><sub>CCA</sub>, tRNA<sup>Pro</sup><sub>UGG</sub>, tRNA<sup>Tyr</sup><sub>GVA</sub>, and tRNA<sup>Phe</sup><sub>GAA</sub> and probes specific for mature tRNA<sup>Trp</sup><sub>CCA</sub> and tRNA<sup>Pro</sup><sub>UGG</sub> (data not shown).



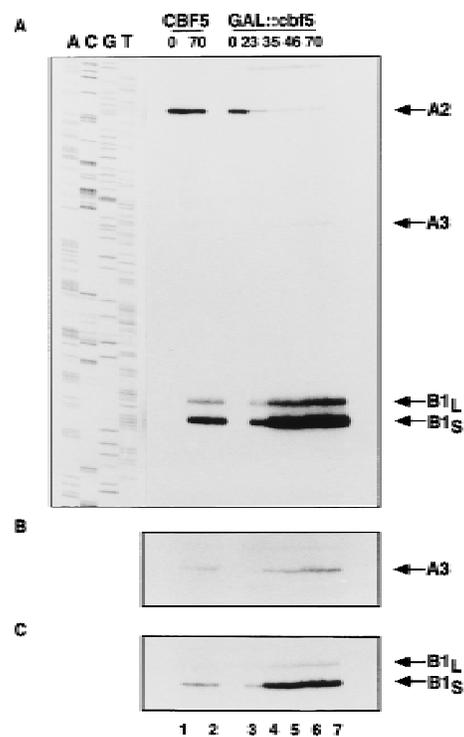
**Figure 4.** Northern analysis of rRNA and pre-rRNA synthesis in a *GAL::cbf5* strain. (A) Probe against ITS2 (probe f); (B) probe against mature 5.8S rRNA (probe e). The oligonucleotides used are depicted in Fig. 1A. RNA was extracted from the *CBF5* and *GAL::cbf5* strains following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8- to 70-hr lanes) and separated on an 8% polyacrylamide gel containing 8 M urea. (\*) 5.8S species with intermediate length.

#### *Cbf5p* is required for formation of $\Psi$ in the pre-rRNA

The effects of depletion of Cbf5p on  $\Psi$  formation were assessed by pulse labeling of the newly synthesized pre-rRNA and rRNA with  $^{32}\text{P}$ . Cells were labeled following growth in permissive rsg medium or following transfer to glu medium for 24 hr. Labeled RNAs, 35S pre-rRNA, mature 25S rRNA, and bulk tRNA, were gel purified, and the nucleotide composition was analyzed by two-dimensional thin layer chromatography (TLC).  $\Psi\text{p}$  content was

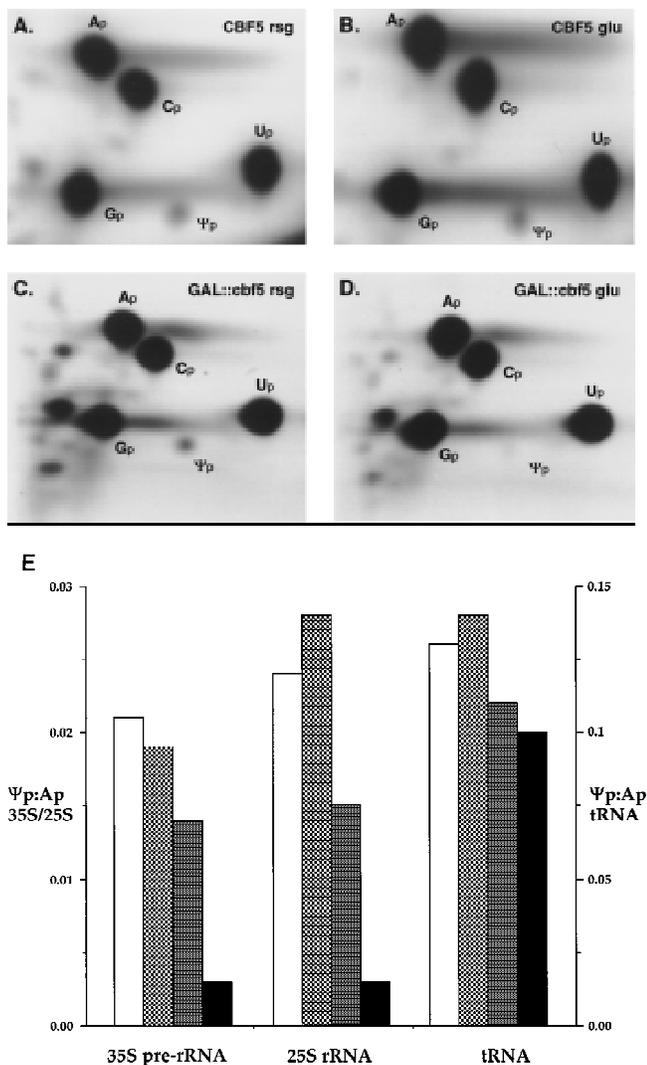


**Figure 5.** Primer-extension analysis of pre-rRNA processing in a *GAL::cbf5* strain. (A) The 5' end of the 35S primary transcript at site +1. (B) Site  $A_0$ . RNA was extracted from the *CBF5* and *GAL::cbf5* strains following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8- to 70-hr lanes) and analyzed by primer extension with oligonucleotide a (Fig. 1A). A DNA sequence made with the same primer is shown as a size marker. Site +1 lies 730 nucleotides from the primer and the sequence is not useful.



**Figure 6.** Primer-extension analysis of pre-rRNA processing in a *GAL::cbf5* strain. (A) Primer extension through sites  $B1_S$ ,  $B1_L$ ,  $A_3$ , and  $A_2$ . (B) Longer exposure of A showing the level of stop at site  $A_3$ . (C) Shorter exposure of A showing the level of stop at site  $B1$ . RNA was extracted from the *CBF5* and *GAL::cbf5* strains following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8- to 70-hr lanes) and analyzed by primer extension using oligonucleotide g (Fig. 1A). A DNA sequence made with the same primer is shown as a size marker.

determined by comparison with other nucleotides present in the same RNA samples. Figure 7E shows the data expressed as the ratio of incorporation into  $\Psi\text{p}$  compared to Ap; comparison of  $\Psi\text{p}$  to Gp or Cp gave similar results (data not shown). The results presented are from one set of experiments; an independent analysis gave similar data (data not shown). The  $\Psi\text{p}$ :Ap ratio in the 25S rRNA from the Cbf5p-depleted strain is ~11% of the value obtained for the wild-type 25S rRNA. This would correspond to a residual level of ~3  $\Psi$  residues per 25S rRNA molecule (the wild-type 25S has 30  $\Psi$  residues) (Ofengand et al. 1995). The  $\Psi\text{p}$ :Ap ratio in the 35S rRNA from the Cbf5p-depleted strain is 16% of the wild-type value, corresponding to ~7  $\Psi$  residues (the wild-type 35S has 43  $\Psi$  residues) (Ofengand et al. 1995). It is not clear whether this corresponds to low residual modification of all sites or whether some sites are preferentially modified. Because the 18S rRNA is not synthesized in the Cbf5p-depleted strain this species cannot be analyzed directly. The reduced modification of the 35S pre-rRNA, however, strongly indicates that formation of  $\Psi$  in the region of the pre-rRNA corresponding to the mature 18S rRNA is also inhibited by depletion of Cbf5p.



**Figure 7.**  $\Psi$  formation in a *GAL::cbf5* strain. Two-dimensional TLC analysis of  $^{32}\text{P}$ -labeled 25S rRNA digested with RNase T2. (A) RNA extracted from the *CBF5* strain following growth in rsg medium. (B) RNA extracted from the *CBF5* strain 24 hr after transfer to glu medium. (C) RNA extracted from the *GAL::cbf5* strain following growth in rsg medium. (D) RNA extracted from the *GAL::cbf5* strain 24 hr after transfer to glu medium. Spots corresponding to Ap, Cp, Gp, Up, and  $\Psi$ p are indicated. (E) Nucleotides separated by two-dimensional TLC were quantitated by PhosphorImager scanning. The ratio between incorporation into  $\Psi$ p and Ap in 35S pre-rRNA, 25S rRNA, and bulk tRNA is shown following growth in rsg medium, and 24 hr after transfer to glu medium. The  $\Psi$ p:Ap ratio in tRNA (right) is shown on a different scale from the 35S and 25S RNA samples (left) because of the greater representation of  $\Psi$ p in tRNA compared to rRNA. (Open bars) *CBF5*<sup>+</sup>/rsg; (light gray bars) *CBF5*<sup>+</sup>/glu; (dark gray bars) *GAL::cbf5* rsg; (solid bars) *GAL::cbf5* glu.

As an example of the data, Figure 7(A–D) presents the analysis of the 25S rRNA from the *CBF5* and *GAL::cbf5* strains. The spot corresponding to  $\Psi$ p can be seen in the *CBF5* samples (Fig. 7A,B) and in the sample from the *GAL::cbf5* strain grown under permissive conditions

(Fig. 7C), but is not readily visible in the sample obtained following depletion of Cbf5p (Fig. 7D).

Surprisingly, the level of  $\Psi$ p in the *GAL::cbf5* strain grown in rsg medium was consistently below the expected level. The  $\Psi$ p:Ap ratios in the 25S and 35S RNA samples was 62% and 67%, respectively, of the values in the corresponding wild-type samples. Northern hybridization indicates that the level of *CBF5* mRNA under permissive conditions is elevated compared to the wild type (Fig. 2C). We speculate that this elevation leads to some excess in Cbf5p synthesis, which leads in turn to a form of squelching, in which excess free protein blocks the interaction of other components with the complex, interfering with its function.

$\Psi$  levels in the tRNA fraction are also mildly reduced by depletion of Cbf5p; the  $\Psi$ p:Ap ratio for the *GAL::cbf5* strain was 85% of the wild-type ratio in rsg medium and 71% of the wild-type ratio in glu medium. *E. coli truB* synthesizes  $\Psi$ 55 in most tRNAs contributing approximately to ~40%–50% of total  $\Psi$  in bulk tRNA (H. Grosjean, pers. comm.), but in yeast, this activity is attributable to Pus4p, another homolog of *truB* (Koonin 1996; Becker et al. 1997). It seems probable that the reduced  $\Psi$ p:Ap ratio is an indirect consequence of the impaired growth of the Cbf5p-depleted strain. It is, however, notable that a number of the box H + ACA snoRNAs can be drawn in the consensus structure to act as  $\Psi$  guides but do not appear to have target sites in the rRNA (Ganot et al. 1997a). The tRNA fraction represents bulk tRNA and we cannot exclude the possibility that some tRNA(s) or other small RNAs present in this fraction are specifically undermodified on depletion of Cbf5p.

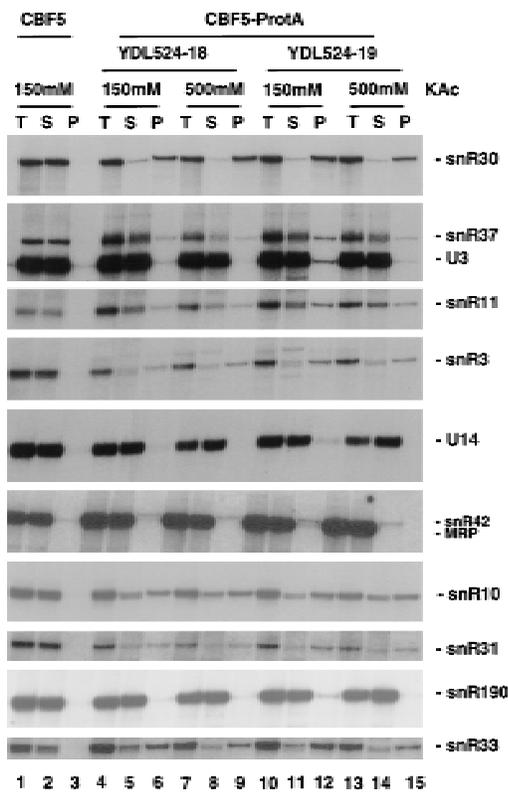
We conclude that, in contrast to *E. coli truB*, Cbf5p is required for  $\Psi$  formation in the pre-rRNA but does not synthesize  $\Psi$ 55 in the tRNAs.

#### *Cbf5p* is a component of the box H + ACA class of snoRNPs

To test for a physical association between Cbf5p and the box H + ACA snoRNAs, a Cbf5p–protein A carboxyl fusion was constructed and integrated at the *CBF5* locus under the control of its own promoter (see Materials and Methods). In the resulting strain, Cbf5p–protein A is the only source of Cbf5p activity in the cell. This strain had a wild-type growth rate (data not shown), showing the Cbf5p–protein A fusion to be fully functional.

Immunoprecipitation of Cbf5p–protein A with IgG–agarose beads resulted in the coprecipitation of all tested box H + ACA snoRNAs (snR3, snR10, snR11, snR30, snR31, snR33, snR37, and snR42) but did not detectably coprecipitate box C + D snoRNAs (U3, U14, and snR190), RNase MRP (Fig. 8) or the U5 snRNA (data not shown). The immunoprecipitations were performed on two independent Cbf5p–protein A strains (YDL524-18 and YDL524-19) at 150 mM and 500 mM Kacetate. Coprecipitation of the box H + ACA snoRNAs with a protein–A fusion to the snoRNP protein Nop1p (fibrillarin) is observed at 150 mM salt but is lost at the higher salt concentration (Ganot et al. 1997b). In contrast, copre-

## A snoRNA-associated pseudouridine synthase



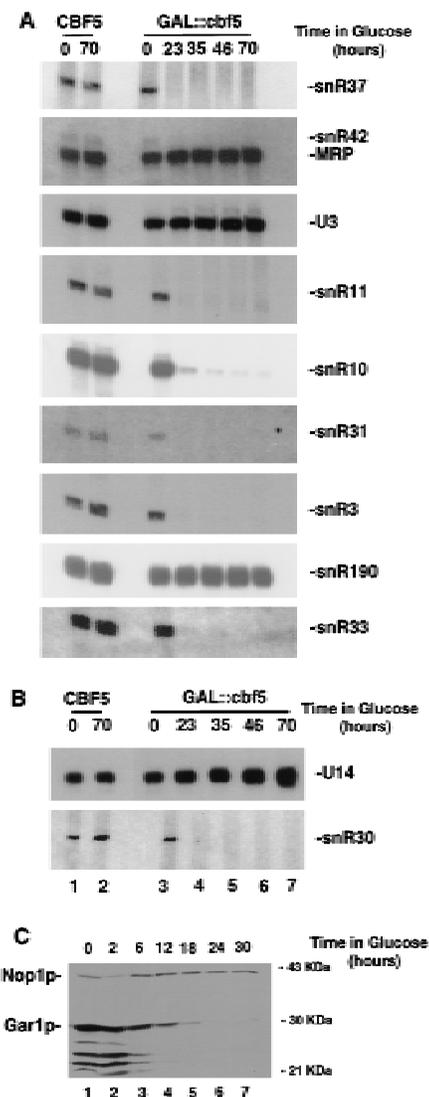
**Figure 8.** The box H + ACA snoRNAs are associated with Cbf5p–protein A (CBFS–Prot.A). Immunoprecipitations were performed at two salts concentration [150 and 500 mM Kacetate (KAc)] on two *CBF5–Prot.A* strains (YDL524-18 and YDL524-19) and at 150 mM Kacetate on the wild-type isogenic control (*CBF5*). RNA was extracted from equivalent amounts of total (T), supernatant (S), and pellet (P) fractions and separated on a 8% polyacrylamide gel containing 8 M urea. Probes used for the hybridizations are described in Materials and Methods.

precipitation of the H + ACA snoRNAs with Cbf5p was observed at both salt concentrations. No precipitation of any RNA was seen with an otherwise isogenic *CBF5* strain expressing only nontagged Cbf5p (Fig. 8, lanes 1–3). The efficiency of coprecipitation of the box H + ACA snoRNAs with Cbf5p–protein A ranges from 30% to 70%, similar to the efficiency with which the box C + D snoRNAs are coprecipitated with a Nop1p–protein A fusion (data not shown).

The levels of the snoRNAs were assessed during depletion of Cbf5p (Fig. 9A,B). Following growth of the *GAL::cbf5* strain on rsg medium, the levels of all tested snoRNAs were the same as in the *CBF5* control strain. In contrast, all tested box H + ACA snoRNAs, snR3, snR10, snR11, snR31, snR33, snR37, snR42 (Fig. 9A), and snR30 (Fig. 9B) were strongly depleted following transfer of the *GAL::cbf5* strain to glu medium. The levels of the box C + D snoRNAs, U3, snR190 (Fig. 9A), and U14 (Fig. 9B) were unaffected, as were the levels of the RNase MRP RNA (Fig. 9A) and the U5 snRNA (data not shown). Analysis of earlier time points during depletion of Cbf5p shows that the major drop in the level of snR30 occurs

between 8 and 16 hr of depletion of Cbf5p (data not shown), in agreement with the onset of the inhibition of processing.

Gar1p, like Cbf5p, is associated with all known members of the family of box H + ACA snoRNAs (Girard et al. 1992). Moreover, a yeast two-hybrid screen has shown that Gar1p interacts physically with Cbf5p (Y. Henry, M. Fromont, P. Legrain, and M. Caizergues-Ferrer, unpubl.). After transfer of the *GAL::cbf5* strain to glu medium, the level of Gar1p also falls dramatically (Fig. 9C).



**Figure 9.** Box H + ACA snoRNP components are codepleted in a *GAL::cbf5* strain. H + ACA snoRNAs (A,B) and Gar1p (C) are codepleted with Cbf5p. Probes used for the hybridizations are described in Materials and Methods. RNA was extracted from the *CBF5* and *GAL::cbf5* strains following growth on rsg medium (0-hr lanes) and at intervals following transfer to glu medium (8- to 70-hr lanes) and separated on an 8% polyacrylamide gel containing 8 M urea. The anti-Gar1p antibody used was described by Girard et al. (1992) and cross-reacts weakly with Nop1p.

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We conclude that Cbf5p is a core component of the box H + ACA snoRNPs that is required for the stability of both the RNA and protein components of the snoRNPs.

## Discussion

We report here a detailed functional analysis of Cbf5p, an essential nucleolar protein (Jiang et al. 1993) and putative  $\Psi$  synthase (Koonin 1996). We found that genetic depletion of Cbf5p inhibits both pre-rRNA processing and formation of  $\Psi$  in the pre-rRNA. Expression of a Cbf5p-protein A fusion protein allowed the coprecipitation of all tested members of the large class of box H + ACA snoRNAs, most of which function as guides for the site-specific formation of  $\Psi$  residues in the pre-rRNA (Ganot et al. 1997a; Ni et al. 1997). Members of the other major class of snoRNAs, the box C + D snoRNAs, were not detectably coprecipitated with Cbf5p-protein A nor were other small RNA species tested (U5 and MRP RNA). Moreover, the depletion of Cbf5p resulted in the codepletion of all tested box H + ACA snoRNAs but did not affect the levels of the box C + D snoRNAs or other small RNA species. We conclude that Cbf5p is an integral component of the box H + ACA class of snoRNPs.

Protein components of other small RNPs are required for the stability of the RNA components of the particles; these include the spliceosomal snRNAs (see Cooper et al. 1995 and references therein) and the RNA components of signal recognition particle (SRP; Brown et al. 1994), RNase P and RNase MRP (Lygerou et al. 1994; Chu et al. 1997). This observation presumably reflects a requirement for the intact structure of the RNP particles to prevent degradation of the RNAs. Unusually, we find that Cbf5p is not only required for the stability of the box H + ACA snoRNAs, but is also required for the stability of another protein component of the snoRNPs, Gar1p. The instability of Gar1p is presumably also attributable to a requirement for the intact RNP structure. In marked contrast to Cbf5p, depletion of Gar1p does not affect the stability of the box H + ACA snoRNAs (Girard et al. 1992). Gar1p depletion, however, does lead to the release of the box H + ACA snoRNA snR36 from the pre-rRNA (Bousquet-Antonelli et al. 1997), indicating that it normally plays a role in establishing the snoRNA-pre-rRNA interaction or in stabilizing the base pairing.

After depletion of Cbf5p, the residual level of  $\Psi$  in the 25S rRNA corresponds to  $\sim 3$   $\Psi$  residues per 25S rRNA molecule, whereas the residual level of  $\Psi$  in the 35S pre-rRNA is estimated at  $\sim 7$  residues per molecule. It is not clear whether the residual  $\Psi$  residues are synthesized by a low residual level of Cbf5p that remains after depletion, or whether these represent a small number of sites at which  $\Psi$  is synthesized by a mechanism that is independent of Cbf5p and the box H + ACA snoRNAs. Surprisingly, the level of  $\Psi$  in the *GAL::cbf5* strain grown under permissive conditions was also consistently below the wild-type values. This result is unlikely to be attributable to a reduced level of Cbf5p; the levels of the box H + ACA snoRNAs is not reduced and pre-rRNA pro-

cessing is unaffected. The *CBF5* mRNA is elevated compared to the wild type under permissive conditions, and the protein is likely to be overexpressed. We speculate that this overexpression leads to a form of squelching, a phenomenon well known from transcription studies (e.g., see Gill and Ptashne 1988; Tasset et al. 1990; Flanagan et al. 1991). Squelching often results from the overexpression of a free form of a component that normally functions in a complex with other proteins. The excess free protein blocks the interaction of other components with the complex, interfering with its function.

In contrast to the  $\Psi$  guide snoRNAs, all of which are nonessential, another box H + ACA snoRNA, snR30, is essential and required for pre-rRNA processing (Bally 1988; Morrissey and Tollervey 1993). Genetic depletion of Cbf5p leads to the codepletion of snR30. The major drop in the level of snR30 occurs between 8 and 16 hr of depletion of Cbf5p, in agreement with the onset of the inhibition of processing. Presumably, as a consequence of this drop, the pre-rRNA processing reactions that require snR30, cleavage at sites  $A_1$  and  $A_2$  and to a lesser extent  $A_0$ , are inhibited by Cbf5p depletion, leading to the loss of 18S rRNA synthesis. This observation explains why Cbf5p is essential for viability. In addition, depletion of Cbf5p leads to a strong delay in the processing of the 27SB and 7S pre-rRNAs, to the accumulation of a 5.8S species of intermediate length, and to a reduced level of mature 25S rRNA. This phenotype has not been observed in any snoRNA mutant and may be a consequence of the absence of  $\Psi$ , possibly because of alterations in the folding of the pre-rRNA or changes in the structure of the preribosomal particle.

Other enzymes involved in RNA metabolism might also be brought to their substrates by carrier RNAs. The box C + D snoRNAs select sites of 2'-*O*-methylation by placing a putative protein-binding site at a precise distance from the nucleotide to be modified (Cavaillé et al. 1996; Kiss-László et al. 1996; Nicoloso et al. 1996). As the rRNA methylase has not yet been identified, it is not clear whether it associates with the box C + D snoRNAs prior to their association with the pre-rRNA. In vivo cleavage of the pre-rRNA at site  $A_0$  in the 5' ETS is absolutely dependent on binding of the U3 snoRNA to an upstream site in the 5' ETS (Beltrame and Tollervey 1995).  $A_0$ , however, can be cleaved in vitro by Rnt1p (an RNase III-like endonuclease) in the absence of cofactors (Abou Elela et al. 1996). It has not yet been established whether U3 carries the enzyme to its site of action in vivo. It seems possible that the role of the box H + ACA snoRNAs in targeting Cbf5p may be a model for the roles of other small RNA species.

Although the data presented here do not formally prove that Cbf5p functions as a  $\Psi$  synthase within the snoRNPs, the homology to *truB* and the strong predictive value of the model make this possibility very likely. The *E. coli* homolog *truB* is the tRNA  $\Psi$ 55 synthase, indicating that the ancestral eukaryotic protein had this function and later acquired the ability to recognize sites made up of a *trans*-acting RNA based-paired to the target site for  $\Psi$  formation. The level of  $\Psi$  in the tRNA popu-

lation is not strongly affected by depletion of Cbf5p, making it unlikely that Cbf5p is the eukaryotic tRNA  $\Psi$ 55 synthase. In fact, Pus4p, another yeast homolog of *truB* (Koonin 1996), is the tRNA  $\Psi$ 55 synthase (Becker et al. 1997). We propose that Pus4p and Cbf5p arose in early eukaryotes by gene duplication followed by divergence of function.

## Materials and methods

### Construction of the GAL::cbf5 and CBF5::Prot.A strains

Standard *Saccharomyces cerevisiae* growth and handling techniques were employed. The transformation procedure was described by Gietz et al. (1992). The wild-type strains used in this study were FY1679-28C and YDL401 (Lafontaine and Tollervey 1996). The GAL::cbf5 strain was created in the YDL401 background by use of a one-step PCR strategy (Lafontaine and Tollervey 1996). The oligonucleotides used for the amplification with plasmid pTL26 were oligonucleotide 1, 5'-TTTCAAATGATGAGATGTTTAGCTTAGGAAAAATATAATGTACTC-TTGGCCTCCTCTAGT-3' and oligonucleotide 2, 5'-CTTAATAACGAAATCCTCCTTTGACATTGTGTATATCGGTCCTCTCGAATTCCTTGAATTTTCAA-3'. Transformants were screened for glucose sensitivity by PCR on yeast colonies and by Southern blot analysis. All the RNA analysis experiments were done in duplicate on two independently isolated GAL::cbf5 strains (YDL521-1 and YDL521-3). The analysis of the  $\Psi$  content of rRNAs was made on strain YDL521-1. The CBF5::Prot.A strains expressing Cbf5-protein A were constructed in strain FY1679-28C by use of the same strategy. The oligonucleotides used for the amplification with pTL54 were oligonucleotide 3, 5'-GAAGACGGTGATTCTGAGGAAAAGAAATCTAAGAAATCTAAGAAAGGCGTGGACAACAAATTC-3' and oligonucleotide 4, 5'-TACAAGTCGTTGATAAGAAATTTTACGTTATTAATATACACTCTGGGTAGAAGATCGGTC-3'. Transformants were screened by PCR on yeast colonies and by Western blot analysis (using PAP antibody, Sigma). Two independently isolated CBF5::Prot.A strains (YDL524-18 and YDL524-19) were used for the immunoprecipitation experiment presented in Figure 8.

### GAL::cbf5 time course, RNA extraction, Northern hybridization, and primer extension

For depletion of Cbf5p, cells growing exponentially in permissive conditions (rsg) at 30°C were harvested by centrifugation, washed, and resuspended in 2% glucose minimal medium. During growth, cells were diluted with prewarmed medium and constantly maintained in exponential phase. RNA extraction, Northern hybridization, and primer extension were as described by Lafontaine et al. (1995). Standard 1.2% agarose/formaldehyde and 8% acrylamide gels were used to analyze the processing of the high and low molecular weight rRNAs species, respectively (Tollervey 1987). Ten percent acrylamide gels were used to analyze the tRNA processing. Nine micrograms of total RNA was used for the Northern and primer-extension experiments presented in Figures 2, 3, 5 and 6, whereas 4.5  $\mu$ g was used for the Northern analysis presented in Figures 4 and 9. Oligonucleotides used for pre-rRNA hybridization were oligonucleotides a, b, c, d, g, h described previously by Lafontaine et al. (1995) as oligonucleotides d, g, h, i, k, l, respectively, and oligonucleotide f described previously as oligonucleotide b by Mitchell et al. (1996). Oligonucleotides anti-U3, U14, MRP, snR10, and snR190 were as described previously (Girard et

al. 1992; Dichtl and Tollervey 1997). Oligonucleotides anti-mature tRNA<sup>Trp</sup><sub>CCA</sub> and tRNA<sup>Pro</sup><sub>UGG</sub> and anti-intronic tRNA<sup>Trp</sup><sub>CCA</sub>, tRNA<sup>Pro</sup><sub>UGG</sub>, tRNA<sup>Tyr</sup><sub>GWA</sub> and tRNA<sup>Phe</sup><sub>GAA</sub> were described by Sharma et al. (1996). Oligonucleotides anti-snR37, snR11, snR3, snR42, snR31, snR33, and U5 were 5'-GATAGTATTAACCACTACTG-3', 5'-GACGAATCGTGACTCTG-3', 5'-TCGATCTTCGTACTGTCT-3', 5'-CTCCCTAAAGCATCACAA-3', 5'-GTAGAACGAATCATGACC-3', 5'-GATTGTCCACACACTTCT-3' and 5'-AAGTTCCAAAAATATGGCAA-3', respectively. Vector pT3/T7 $\alpha$ 18-snR30 (Morrissey and Tollervey 1993) was used to synthesize an snR30 riboprobe. To detect the CBF5 mRNA, a fragment spanning the whole ORF of CBF5 was generated by PCR and labeled by use of the Prime-a-Gene Labeling kit (Promega).

### Analysis of $\Psi$ levels

To determine the  $\Psi$  content in rRNAs and tRNAs, strain YDL521-1 was pregrown at 30°C in rich medium containing 2% raffinose, 2% sucrose, and 2% galactose (YP-RSG). The culture was split in two, washed, and resuspended in either rich glu medium (YPD) or YP-RSG at an OD<sub>600</sub> of 0.03. Both cultures were incubated for 6 hr at 30°C before being washed and transferred to similar medium without PO<sub>4</sub> [following the recipe described by Warner (1991)]. After a further incubation of 18 hr, 50 ml from each culture was labeled for 15 min with 9 mCi of [<sup>32</sup>P] orthophosphate (9000 Ci/mmol). Total RNA was extracted and analyzed by electrophoresis on a 1.2% agarose-formaldehyde gel. 35S pre-rRNA, 25S rRNA, and total tRNAs were purified by electroelution, digested with RNase T2 (in 50 mM Na acetate at pH 5.5 at 37°C), and analyzed by two-dimensional cellulose TLC as described by Filipowicz and Shatkin (1983). The procedure used was essentially as described by Bousquet-Antonelli et al. (1997).

### Immunoprecipitation of Cbf5p-protein A

Yeast whole-cell extracts were prepared according to Séraphin and Rosbash (1989). Lysates were made in buffer A (20 mM Tris HCl at pH 8.0, 5 mM MgCl<sub>2</sub>, 1 mM DTT, 0.2% Triton X-100, 0.5 mM PMSF, and 150 or 500 mM Kacetate), and supernatants were cleared by centrifugation (56,000 rpm at 4°C for 20 min). Immunoprecipitation experiments were performed as described previously (Ganot et al. 1997b). Lysates equivalent to 37.5 OD<sub>600</sub> of cells were incubated on a rotating wheel for 2 hr at 4°C with 100  $\mu$ l of IgG-agarose beads (Sigma A2909), and pre-washed in buffer A in a total volume of 400  $\mu$ l. Pellets were washed four times for 20 min in 1 ml of buffer A. Each gel lane (T, S, and P) was loaded with RNA from a fraction of the preparation equivalent to 10 OD<sub>600</sub> of cells.

### Western blotting

Total protein extracts corresponding to 10<sup>6</sup> cells were loaded in each lane. Affinity-purified anti-Gar1p antibodies were used at 1:200 dilution as described by Girard et al. (1992). The anti-Gar1p antibodies weakly cross-react with Nop1p.

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