

Identifying the methyltransferases for m⁵U747 and m⁵U1939 in 23S rRNA using MALDI mass spectrometry

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ABSTRACT

There are three sites of m⁵U modification in *Escherichia coli* stable RNAs: one at the invariant tRNA position U54 and two in 23S rRNA at the phylogenetically conserved positions U747 and U1939. Each of these sites is modified by its own methyltransferase, and the tRNA methyltransferase, TrmA, is well-characterised. Two open reading frames, YbjF and YgcA, are approximately 30% identical to TrmA, and here we determine the functions of these candidate methyltransferases using MALDI mass spectrometry. A purified recombinant version of YgcA retains its activity and specificity, and methylates U1939 in an RNA transcript *in vitro*. We were unable to generate a recombinant version of YbjF that retained *in vitro* activity, so the function of this enzyme was defined *in vivo* by engineering a *ybjF* knockout strain. Comparison of the methylation patterns in 23S rRNAs from YbjF⁺ and YbjF⁻ strains showed that the latter differed only in the lack of the m⁵U747 modification. With this report, the functions of all the *E.coli* m⁵U RNA methyltransferases are identified, and a more appropriate designation for YbjF would be RumB (RNA uridine methyltransferases B), in line with the recent nomenclature change for YgcA (now RumA).

INTRODUCTION

Transfer and ribosomal RNAs engage in a multitude of precisely coordinated molecular events during protein synthesis, the speed and accuracy of which are governed to a large extent by the ability of RNA to discriminate between functional and non-function interactions. The interactive and discriminatory abilities of RNA are determined by the four nucleotide components that make up its structure. The basic structural repertoire of RNA is expanded in tRNAs and rRNAs by a range of post-transcriptional modifications. These modifications are quite diverse in tRNAs, while rRNA modifications are generally limited to base and sugar

methylations and pseudouridylation (1). Ribosomes of the enterobacterium *Escherichia coli* have 11 modified nucleotides in 16S rRNA and 23 modifications in 23S rRNA (2). Plotting the locations of the modified nucleotides onto the X-ray crystal structures of the ribosome (3–5) shows that they cluster within several discrete regions (6–8) that are concerned with subunit association, mRNA decoding, the binding of auxiliary factors, and peptide bond formation (9–12).

Although it is now generally agreed that all of the modifications within the *E.coli* rRNA have been comprehensively identified and mapped, several of the enzymes responsible for these modifications remain to be accounted for. To date, four of the 16S rRNA modification enzymes have been identified (13–16), while seven of the 23S rRNA enzymes are known (17–23). The incompletely characterised enzymes include those that convert uridine to thymidine (m⁵U), a modification that occurs in both rRNA and tRNAs. In *E.coli*, three RNA uridine nucleotides are converted to m⁵U: one is at the invariant tRNA position U54 (24,25); and the two other sites are at 23S rRNA nucleotides U747 and U1939 (26,27). The methyltransferase enzyme responsible for tRNA U54 methylation, TrmA, has been thoroughly investigated (28–31), whereas at the inception of our studies neither of the two rRNA m⁵U modifications had been linked to its cognate methyltransferase.

In an earlier study, a homology search of the then incomplete *E.coli* database revealed two open reading frames (YgcA and YbjF) with sufficient similarity to TrmA to warrant the authors' conclusion that these represented the U747 and U1939 methyltransferases (32). Repeating the search in the completed *E.coli* genome sequence results in the same conclusion, without revealing additional m⁵U methyltransferase paralogues. Here, we have utilised Matrix Assisted Laser Desorption/Ionisation (MALDI) mass spectrometry to determine whether the putative m⁵U methyltransferases modify RNA transcripts *in vitro*. Using this approach, we have identified the methylation target of recombinant YgcA at U1939, consistent with the findings of Agarwalla *et al.* (21) published during the course of our studies. No recombinant versions of YbjF could be generated that showed *in vitro* activity. In this case a different approach, involving mass spectrometric comparison of the methylation patterns in 23S rRNAs from YbjF⁺ and YbjF⁻ strains, was developed to define the activity of this enzyme *in vivo*.

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Table 1. Strains and plasmids used in the study

Strain/Plasmid	Description	Ref.
<i>E. coli</i> strains		
CP79	<i>thr leu his argA</i> RC ^{rel} F ⁻	(25)
DH1	F ⁻ <i>supE44 recA1 endA1 gyrA96(Nal^r) thi-1 hsdR17 (r_K⁻ m_K⁺) relA1 spoT1</i>	(34)
DH10B	F ⁻ <i>mcrA Δ(mrr-hsdRMS-mcrBC) φ80lacZΔM15 ΔlacX74 deoR recA1 endA1 araD139Δ(ara, leu) 7697 galU galK λ⁻ rpsL nupG</i>	(55)
ER2566	F ⁻ <i>λ⁻ fluA2 [lon] ompT lacZ::T7 gene1 gal sulA11 D(mcrC-mrr)114::IS10 R(mcr-73::miniTn10-TetS)2 R(zgb-210::Tn10) (TetS) endA1 [dcm]</i>	New England Biolabs
IB10	<i>thr leu his argA</i> RC ^{rel} F ⁻ <i>rrmA</i>	(25)
CP79 ΔybjF	<i>thr leu his argA</i> RC ^{rel} F ⁻ <i>ybjF</i>	This study
IB10 ΔybjF	<i>thr leu his argA</i> RC ^{rel} F ⁻ <i>rrmA ybjF</i>	This study
Plasmids		
pTYB11	Cloning/expression vector in T7 polymerase containing host. Amp ^R	New England Biolabs
pMAK705	Ts replication origin. Cm ^R	(41)
pOU12	PQE60 derivative (Qiagen). Amp ^R	Lab. collection
pQErrmA	PQE60 derivative (Qiagen). Amp ^R and Tet ^R	Lab. collection
pSD2KK	pGEM3 derivative (Promega). Amp ^R and Kan ^R	Lab. collection
pMG25	pUHE24-2 derivative (56). Cloning/expression vector. Amp ^R	Lab. collection
pCTM1	<i>ygcA</i> gene cloned into the NcoI/BglII sites in pQErrmA. Amp ^R	This study
pCTM3	<i>ybjF</i> gene cloned into the SphI/HindIII sites of Pou12. Amp ^R	This study
pCTM6	<i>trmA</i> gene cloned into the EcoRI/BglII sites in pQErrmA. Amp ^R	This study
pCTM8	<i>TrmA</i> in the EcoRI/HindIII sites of pMG25. Amp ^R	This study
pCTM9	<i>ygcA</i> in the EcoRI/HindIII sites of pMG25. Amp ^R	This study
pCTM10	<i>ybjF</i> in the EcoRI/HindIII sites of pMG25. Amp ^R	This study
pCTM11	<i>ybjF</i> gene cloned into the SapI/EcoRI sites of pTYB11. Amp ^R	This study
pCTM12	Kanamycin cassette from pSD2KK cloned into pCTM10. Kan ^R and Amp ^R	This study
pCTM13	<i>ybjF</i> gene containing Kan ^R from pCTM12 cloned into pMAK705. Kan ^R and Cm ^R	This study

MATERIALS AND METHODS

Database searches

A BLAST search (33) restricted to the *E. coli* genome revealed two putative gene products, YgcA and YbjF, that have significant similarity to the tRNA m⁵U methyltransferase TrmA. After establishing that YgcA and YbjF are rRNA m⁵U methyltransferases, we used a standard BLASTp to screen for homologous sequences in other organisms. Sequences from the search that fulfilled the following four criteria were considered to be orthologues of the two *E. coli* rRNA m⁵U methyltransferases. First, sequences were required to have an expectation-value of <10⁻¹⁰ as defined by the BLAST program. Second, sequences had to contain all the four conserved m⁵U methyltransferase domains defined by Gustafsson *et al.* (32). Third, they were required to align throughout the entire sequence of YgcA or YbjF. Lastly, sequences exhibiting a higher score in a TrmA BLAST were considered to be tRNA rather than rRNA methyltransferases.

Bacterial strains, plasmids and growth conditions

The *E. coli* strains and plasmid used in this study are listed in Table 1. Strains were grown at 37°C in LB media or on LB-agar plates (34) containing, where appropriate, ampicillin at 100 µg/ml, kanamycin at 25 µg/ml or chloramphenicol at 25 µg/ml.

Cloning of methyltransferase genes

The primers used for PCR amplification of methyltransferase genes (Table 2) were based on gene sequences in the *E. coli* genome database. Template DNA was obtained by suspending a small portion of a single colony of *E. coli* strain DH1 in 50 µl

H₂O and using 1 µl for each amplification reaction with the DNA polymerase Platinum Pfx (Life Technologies). PCR products were purified using High Pure PCR (Boehringer-Mannheim) and QIAEXII (Qiagen) kits. The *ygcA*, *trmA* and *ybjF* genes were inserted into the pQErrmA, pOU12 and pTYB11 cloning vectors to produce proteins with a C-terminal histidine (His-) tag, an N-terminal His-tag, and to form an intein fusion product, respectively (Table 1). Subcloning of tagged genes into plasmid pMG25 and transforming *E. coli* strain DH10B generally produced more efficient protein expression. DNA manipulations and cell transformation with plasmids were carried out using standard techniques (34). Restriction endonucleases and other enzymes for DNA manipulations (New England Biolabs and Roche Molecular Biochemicals) were used in accordance with the suppliers' recommendations. Plasmids were prepared using a mini-prep kit (Bio-Rad), and cloned DNA fragments were sequenced using the CEQ 2000 Dye Terminator procedure (Beckmann).

Purification of methyltransferase enzymes

His-tagged proteins expressed in cells containing plasmids pCTM8, pCTM9 and pCTM10 were purified on Ni-agarose columns (Qiagen). Purified methyltransferases were dialysed against 20 mM Tris-Cl (pH 7.5), 10 mM magnesium acetate, 250 mM NH₄Cl and 10% (w/v) glycerol to remove the imidazol from the elution buffer. The purity of the proteins was monitored by SDS-PAGE, and their sizes were checked against markers of comparable molecular weight (Pharmacia). The identities of the proteins were subsequently confirmed by peptide mapping using mass spectrometry (35).

For expression of the YbjF-intein fusion protein, *E. coli* strain ER2566 containing pCTM11 was grown overnight at

Table 2. Oligodeoxynucleotides used in the study

Name	Sequence (5' to 3')	Purpose
TOFT-1	GCTAAGATCTTTTAAACGCGCGAGAAAAGTA	PCR <i>ycgA</i> 3'-end
TOFT-2	CAGCTTAGCCATGGCGCAATTCTACTCTGC	PCR <i>ycgA</i> 5'-end
TOFT-3	GCCGTAAGCATGCAGTGCCTTTACGA	PCR <i>ybjF</i> 5'-end. Knockout check
TOFT-4	CGTTAGATCTTTGCTTCACCAGCAGCGTCA	PCR <i>ybjF</i> 3'-end. Knockout check
TOFT-5	TATGTAAGCATGCATCACCATCACCAGTGCCTTTACGACGCG	PCR <i>ybjF</i> 5'-end. N-His-tag
TOFT-6	CCACGTTAAGCTTTTGGCTTCACCAGCAGCGTCAGCA	PCR <i>ybjF</i> 3'-end
TOFT-7	CTACAGAATTCATTAAGAGGAGAAATTAATGACCCCGAACACCTTCC	PCR <i>trmA</i> 5'-end
TOFT-8	CACGTTAGATCTTTCGCGGTAGTAATACGC	PCR <i>trmA</i> 3'-end
TOFT-19	TAATACGACTCACTATAGG	Top strand T7 transcription
TOFT-20	GTCGGAACCTTACCCGACAAGGAATTCGCTACCTTAGGACCGTTATAGTTACGG CCCCTATAGTGAGTCGTATTA	T7 RNA transcript with U1939
TOFT-21	TGGAGGGGGCGAAGGGAATCGAACCCCTCGTATAGTGCTTGGGAAGCTCTCGTTC TACCATTGAACTACGCCCCCTATAGTGAGTCGTATTA	T7 tRNA transcript with U54
TOFT-31	GGTGGTTGCTCTTCCAACATGCAGTGCCTTTACGA	PCR <i>ybjF</i> 5'-end. Intein system
TOFT-32	GGTGGTGAATTCCTTGCCTTACCAGCAGCGTCA	PCR <i>ybjF</i> 3'-end. Intein system
SD45	GCCGCATCCGCTAATTTTCAACATTAGTCGTTCCGTCCTCCAGTTAGTGTACC CAACCTTCAACCTGCTCCTATAGTGAGTCGTATTA	T7 RNA transcript with U747

Restriction sites and ATG start codons are in bold in the PCR primer sequences; sequences encoding His-tags are in italics.

15–37°C in medium containing 0.4 mM isopropyl-D-thiogalactopyranoside. Cells were harvested and resuspended in 4 ml of 20 mM Tris–Cl (pH 8.0), 500 mM NaCl, 1 mM EDTA and 0.1% (w/v) Triton X-100 (lysis buffer) prior to sonication. Lysates were loaded onto a chitin column and washed twice with column buffer (lysis buffer without Triton X-100). Incubation in 10 ml column buffer containing 50 mM DTT at 4°C overnight induced cleavage at the intein site. The YbjF protein was eluted with column buffer, and its sequence and purity were checked as described above.

RNA substrates for methylation

RNA transcripts used for *in vitro* methylation assays were synthesised from DNA oligodeoxynucleotide templates encoding the T7 RNA polymerase promoter (36,37). The DNA oligonucleotide TOFT21 encodes a full length tRNA^{pro} transcript containing the TrmA target at U54, and the oligodeoxynucleotides SD45 and TOFT20 encode the regions of *E.coli* 23S rRNA that contain the U747 and U1939 methylation targets, respectively (Table 2; Fig. 1). The RNA transcripts were purified on 10 µl reverse phase Poros 50R2 columns (PerSeptive Biosystems) made in-house (38).

Ribosomal particles from *E.coli* strains CP79, CP79Δ*ybjF*, IB10 and IB10Δ*ybjF* (Table 1) were fractionated on sucrose gradients, and 50S subunits were used to prepare 23S rRNA (39). All RNAs were redissolved and stored at –20°C in H₂O (double distilled in all cases).

In vitro methylation

Purified RNAs (~125 ng) were renatured in 50 µl of 100 mM NH₄Cl, 20 mM Tris–Cl (pH 7.8), 10% (w/v) glycerol, 6 mM β-mercaptoethanol by warming for 5 min at 50°C followed by 10 min at 37°C. The methyl group donor, S-adenosylmethionine (SAM), was added to a final concentration of 1 mM; 1–50 ng of purified methyltransferases were added to start the methylation reactions, which proceeded for 30 min at 37°C. The reactions were stopped by extraction with phenol and chloroform; the RNAs were recovered from the aqueous phase by ethanol precipitation, and were redissolved in 2.5 µl H₂O.

MALDI mass spectrometry analyses of RNAs

All the RNA methylation substrates were digested with ribonucleases to produce a discrete series of oligonucleotide fragments for analysis by mass spectrometry. 2.5 µl RNA (~125 ng) was mixed with 0.5 µl 3-hydroxypicolinic acid (0.5 M in 50% acetonitrile) and 1.5 µl H₂O, and was digested with 10 U RNase T1 (United States Biochemicals) for 3.5 h at 37°C. 3-hydroxypicolinic acid serves as a denaturing agent to ensure complete RNA digestion, in addition to functioning as a matrix for MALDI mass spectrometry (40).

23S rRNA was digested by incubating 2.0 µl RNA (~260 ng) and 0.5 µl 0.5 M 3-hydroxypicolinic acid with 60 U of RNase T1 for 1 h at 37°C. Digestion products were run through Poros 50R2 columns (38), eluting short oligonucleotides with 10 mM triethyl ammonium acetate pH 7.0 (TEAA)/6% acetonitrile, and larger oligonucleotides with 10 mM TEAA/25% acetonitrile. Eluants were dried and redissolved in 3 µl H₂O. The RNA digestion fragments were analysed on MALDI mass spectrometers from Bruker reflex IV (Bruker-Daltonik) or Voyager Elite (PerSeptive Biosystems), recording the spectra in reflector and positive ion mode using delayed ion extraction.

Knockout of the *ybjF* gene

The putative m⁵U methyltransferase gene *ybjF* was inactivated in the chromosome of *E.coli* strains CP79 and IB10 by insertion of a kanamycin resistance cassette. First, the kanamycin resistance cassette from plasmid pSD2KK was inserted in the BglII site of *ybjF* in pCTM10, creating pCTM12 (Table 1). The inactivated *ybjF* gene was excised from pCTM12 with SphI and EcoRV, and was ligated into the SphI/HincII sites of pMAK705, creating pCTM13. Plasmids pMAK705 and pCTM13 possess a temperature-sensitive pSC101 replicon (41); pCTM13 was used to transform strains CP79 and IB10 at the permissive temperature of 30°C. Double crossover recombination events between the genomic- and plasmid-encoded *ybjF* gene copies were facilitated by raising the incubation temperature to 42°C (41), and cells were then screened for kanamycin resistance and chloramphenicol

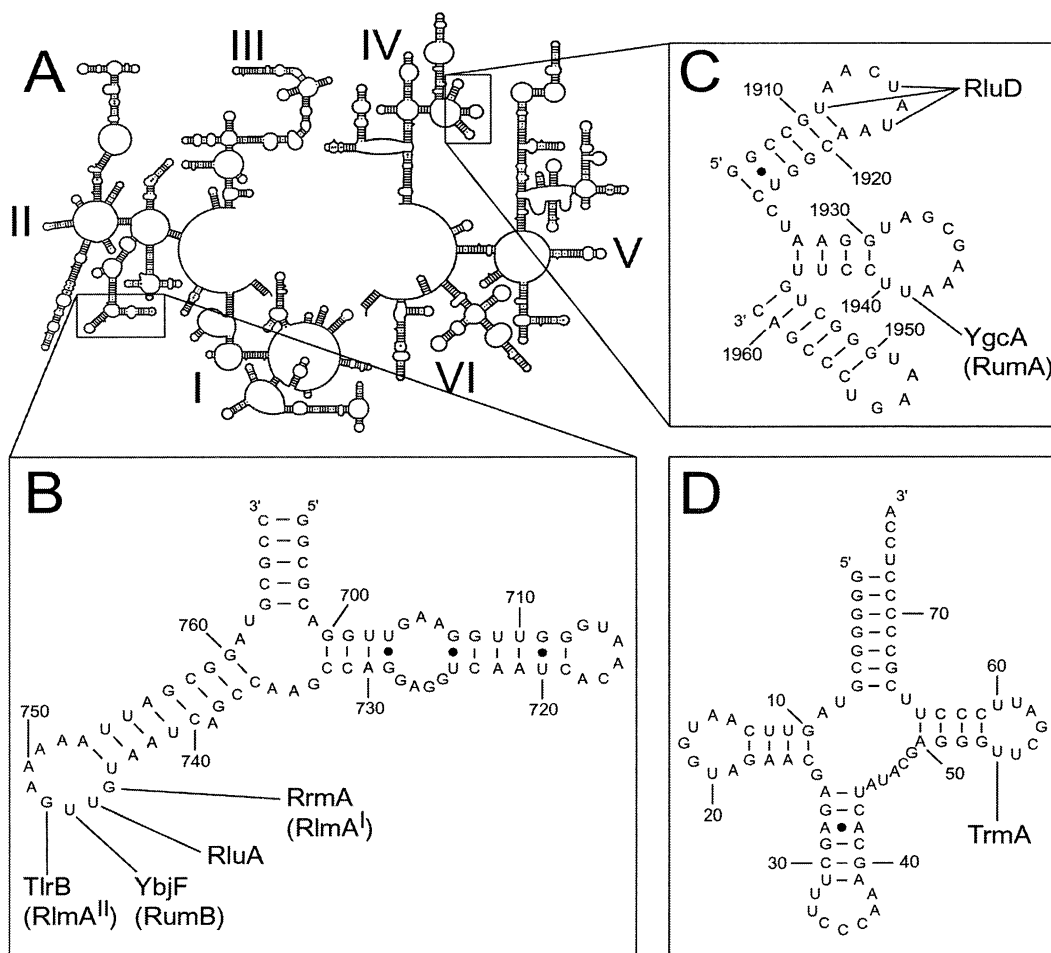


Figure 1. (A) Secondary structure of *E. coli* 23S rRNA (52,53); the domains are labelled with roman numerals. (B) RNA transcript corresponds to domain II nucleotides 694–767, and contains the U747 methylation target. Sites that are post-transcriptionally modified *in vivo* are indicated: RrmA (RlmA^I) (17,37), RluA (18) and YbjF (RumB) (this study) are *E. coli* enzymes; TlrB (RlmA^{II}) is found in Gram-positive bacteria (44). (C) RNA transcript of the three stem-loop structures corresponding to the domain IV sequence 1906–1961 and containing the U1939 target. Sites of modification in *E. coli* rRNA by RluD (54) and YgcA (RumA) (21 and this study), are shown. The terminal base pairs in transcripts (B) and (C) deviate from the *E. coli* 23S rRNA sequence to facilitate *in vitro* transcription and transcript folding. (D) RNA transcript mimicking the structure of *E. coli* tRNA^{Phe} and containing the TrmA target at U54 (28).

sensitivity. The genomic *ybjF* region of potential knockout recombinants was analysed by PCR.

RESULTS

Expression, purification and activity of methyltransferases

The cloned *trmA*, *ycgA* and *ybjF* genes were found to be identical to the sequences in the *E. coli* database (<http://www.ncbi.nlm.nih.gov/Genbank/index.html>, *ycgA* GenBank accession no.: AE000362; *ybjF*, AE000187). Active, recombinant TrmA and YgcA proteins were purified with a C-terminal His-tag. The recombinant YgcA enzyme showed activity specific for the RNA substrate containing U1939 (Fig. 1C), with no cross-reactivity to the other RNA substrates. The RNase T1 oligonucleotide containing U1939 had an expected m/z of 3185.4 (Fig. 2A), and this peak was shifted to an m/z of 3199.5 after treatment of the RNA substrate with YgcA and the methyl group donor SAM (Fig. 2B).

This increase in mass/charge of 14 Da is consistent with the addition of a single methyl group at U1939 by YgcA. Recombinant TrmA efficiently methylated its U54 target in the tRNA transcript (Fig. 1D), shifting the spectrum peak at m/z 1281.2 (UUCG) to 1295.0 (m^5 UUCG), and showed no cross-reactivity with the rRNA transcripts (data not shown).

Recombinant YbjF with a C-terminal His-tag could not be purified efficiently. Reasonable yields of the YbjF protein were obtained both with an N-terminal His-tag and using the intein fusion technique. However, no methylation of any of the transcripts was observed *in vitro*, including the RNA containing U747, which is otherwise an effective substrate for the RrmA and TlrB methyltransferases (37,42) (Fig. 1); recombinant YbjF also failed to transfer a tritiated methyl group from 3 H-SAM to unmethylated 23S rRNA from strains CP79 Δ *ybjF* and IB10 Δ *ybjF* (data not shown). As we were probably inadvertently destroying the activity of YbjF at one of the *in vitro* steps, we elected to study the function of the protein *in vivo*.

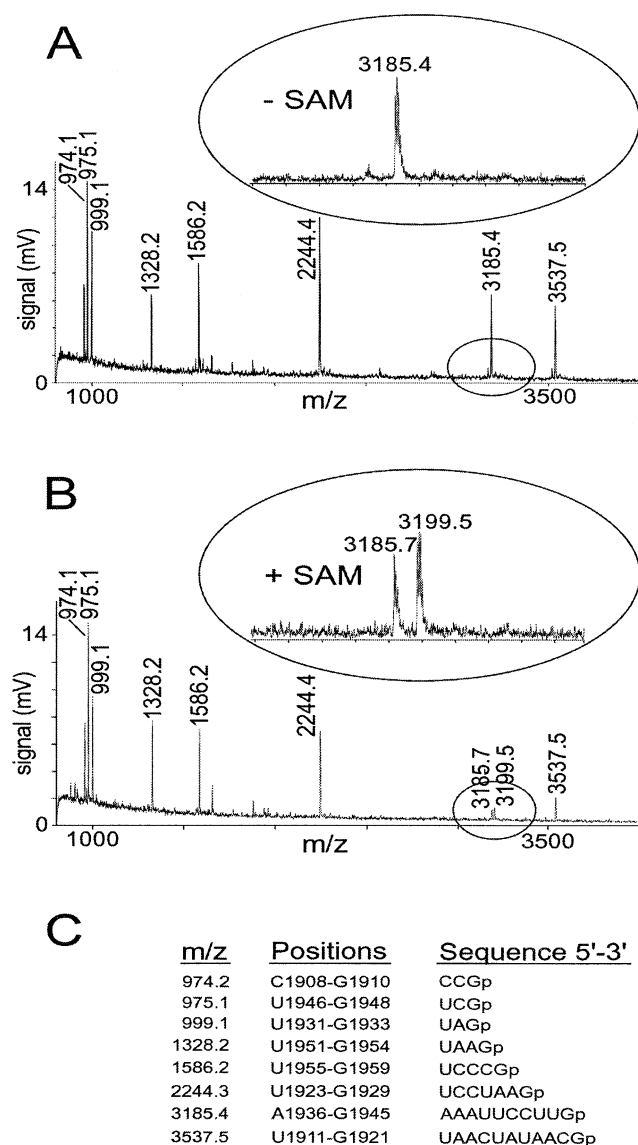


Figure 2. MALDI mass spectra of the U1939 RNA transcript after incubation with recombinant YgcA (RumA) followed by digestion with RNase T1. (A) Control sample without S-adenosylmethionine in the *in vitro* methylation assay (-SAM). (B) Test sample including the methyl group donor (+SAM). The spectral region around AAUCCUUGp is enlarged to show that this peak (theoretical m/z 3185.4) is shifted to m/z 3199.5 when the RNA is methylated. (C) Theoretical singly protonated masses of the RNase T1 digestion products from the U1939 RNA transcript. All measured masses are within 0.3 Da of the theoretical values. The jagged nature of the peaks visible in the enlargements reflects the natural isotopic distribution of ^{12}C and ^{13}C in the RNA.

The YbjF target *in vivo*

The *ybjF* gene was insertionally inactivated on the chromosome of the two *E. coli* strains CP79 and IB10. Potential *ybjF* knockout candidates were screened for kanamycin resistance (indicating the presence of the kanamycin resistance cassette in *ybjF*) and chloramphenicol sensitivity (showing that the plasmid initially harbouring the inactivated *ybjF* had been lost). Approximately 10% of the colonies that were screened displayed this phenotype, and PCR analysis of two candidates

for each strain showed that the chromosomal copy of *ybjF* had been replaced by the inactivated plasmid copy (data not shown).

The 23S rRNAs were isolated from the four strains: CP79, IB10 and their *ybjF* knockouts. Strain IB10 has lost the function of RrmA (RlmA^I) that methylates the N1 position of 23S rRNA nucleotide G745; CP79 is the parent strain with a functional RrmA (25). N1 methylation of G745 renders the nucleotide resistant to ribonuclease T1 digestion (43). Digestion of 23S rRNA with RNase T1 produces a complex array of oligonucleotides, which were purified and recorded in two separate mass spectra (Fig. 3A and B). In the high molecular weight spectrum of the wild-type 23S rRNA, a unique decanucleotide is seen with an m/z of 3253.4 spanning nucleotides 739–748 (Fig. 3C). In corresponding oligonucleotide spectrum from the CP79 $\Delta ybjF$ strain, this peak is shifted to a lower m/z of 3239.6 (Fig. 3D). The loss of ~14 Da indicates that a methyl group is missing from the 739–748 sequence in the $\Delta ybjF$ strain.

The site of YbjF methylation could be pinpointed using IB10 23S rRNA. Here, the unmethylated G745 is cleaved by RNase T1, and the 739–748 decanucleotide is therefore lost from the upper range of the MALDI spectrum (Fig. 3E). The resulting heptanucleotide, 5'-ACUAAUGp (nucleotides 739–745) comes to occupy the spectrum peak at 2268.3 (together with three other 23S rRNA fragments of identical m/z , (Fig. 3A). If the YbjF target lay within the 739–745 sequence, its m/z would be shifted to 2282.3 in the spectrum from the IB10 YbjF⁺ strain. The lack of a spectrum peak at m/z 2282.3 (not shown) makes it evident that the YbjF target resides within the trinucleotide 5'-U746-U747-Gp748. No methylation occurs at nucleotide U746, which is converted to pseudouridine by RluA (18) without changing the mass of the nucleotide (43); and no modification occurs at G748 in *E. coli* 23S rRNA (44). Thus the modification pattern of the trinucleotide is most consistent with the composition 5'- ψ 746- $m^5\text{U}747$ -Gp748.

DISCUSSION

Three $m^5\text{U}$ modifications have previously been identified in *E. coli*: one at tRNA position 54, and two in 23S rRNA at positions 747 and 1939 (2). In this study we have used MALDI mass spectrometry to link the three sites of $m^5\text{U}$ modification with their respective methyltransferase enzymes. The tRNA $m^5\text{U}54$ methyltransferase TrmA is known to retain its activity after purification (30,31), and the enzyme served as a positive control for our *in vitro* methylation assays. A search of the *E. coli* genome at the inception of our studies confirmed that the two most likely $m^5\text{U}$ rRNA methyltransferase candidates were YgcA and YbjF.

Recombinant YgcA proved to be as easy to purify and handle as TrmA, retaining its activity and specificity *in vitro*. YgcA specifically methylated an RNA transcript of 66 nucleotides, which was designed to mimic the structure around 23S rRNA nucleotide U1939 (Fig. 1C). Analysis of the 66mer after RNase digestion showed that YgcA increased the mass of the sequence 1936–1945 by 14 Da in the presence of the methyl group donor, SAM. This is consistent with the addition of a single methyl group at U1939 by YgcA. During the course of our study, Agarwalla and co-workers pinpointed

the YgcA target at U1939 using biochemical approaches; they subsequently renamed this methyltransferase RumA (RNA uridine methyltransferase A) (21).

By a process of elimination, this left YbjF as the most likely candidate to target the U747 site. However, despite the construction of a variety of recombinant versions of YbjF and the use of different enzyme purification methods and RNA substrates, the study of YbjF function *in vitro* proved to be intractable. A second approach was taken, which involved analysis of authentic rRNA. The large subunit rRNAs (5S and 23S rRNAs) give rise to over 900 fragments upon RNase T1 digestion, and analysis of these required stretching the limits of the MALDI technology. Fortunately, the putative target of YbjF occurs within the RNase T1 digestion product 5'-ACUAAUm¹Gψm⁵UGp (nucleotides 739–748), which is a double fragment formed by virtue of the RNase-resistant m¹G745 nucleotide. This fragment forms a distinct peak in the mass spectrum (Fig. 3C), the position of which is shifted 14 Da downstream corresponding to the loss of a single methyl group in the strain without a functional YbjF protein (Fig. 3D). The position of the methyl group added by YbjF could be localised within reasonable doubt to U747 by using strains that lack the m¹G745 modification.

Most rRNA modifications are located within preserved, functionally important regions of the rRNA (6,7). The YgcA (RumA) target U1939 is phylogenetically conserved, with a uridine at this position in >95% of all organisms (Gutell Lab Comparative RNA: <http://www.rna.icmb.utexas.edu>). Similarly, YbjF (now RumB) targets a nucleotide that is conserved as a uridine in >95% of all bacteria, and in >90% of organisms in the other two phylogenetic domains (data from the Gutell web site).

Other enterobacteria possess homologues of *rumA* and *rumB*, and homologues are also evident in less closely related species of archaea (Table 3). The conserved, methylated uridines are situated in regions of the 23S rRNA that have been associated with essential functions in protein synthesis. U1939 is located within the rRNA structure that interacts with the aminoacyl end of A-site bound tRNA (4,45), and could play an active role in sensing uncharged tRNA (21). U747 is situated in a highly conserved and densely modified part of the rRNA that lines a constricted region of the 50S subunit tunnel through which the nascent peptide chain passes during protein

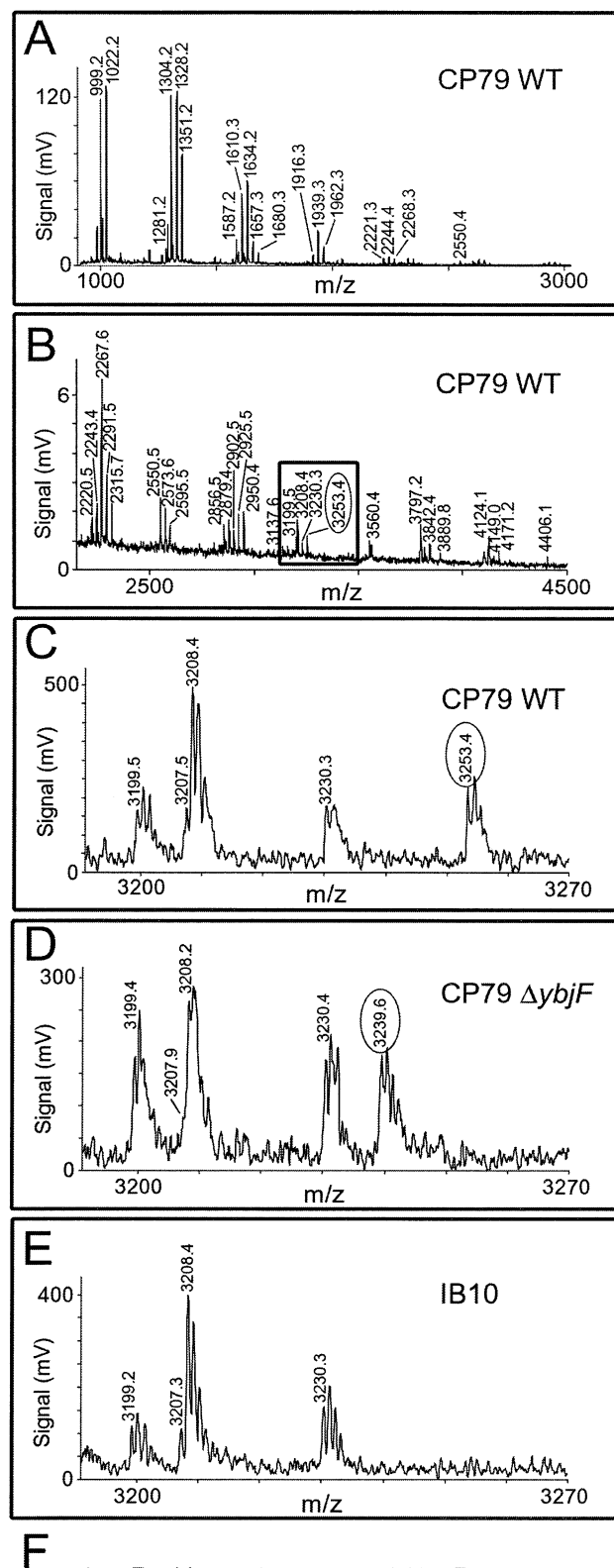


Figure 3. MALDI mass spectra of 23S rRNAs from CP79 wild-type (WT), CP79 $\Delta ybjF$ and IB10 strains after digestion with RNase T1. Fragments arising from 5S rRNA do not appreciably complicate the spectrum and therefore have not been removed. Many oligonucleotides (particularly the smaller ones) have the same m/z values; all fragments of trinucleotides and larger are depicted here, and these have been eluted in two steps. (A) The smaller digestion products eluted with 6% acetonitrile; (B) larger oligonucleotides eluted with 25% acetonitrile. (C) Enlargement of the boxed region shows that the wild-type double fragment 5'-ACUAAUm¹Gψm⁵UGp (m/z 3253.4) is 14 Da heavier than in the $\Delta ybjF$ strain (D). In the IB10 rRNA (E), the double fragment has been lost due to cleavage (see text). (F) The theoretical masses of protonated RNase T1 fragments of 23S rRNA in the region of interest; these match the measured m/z values to within 0.2 Da. The peak at m/z 3199.4 corresponds to a decanucleotide from 23S rRNA positions 1936–1945. This mass includes a methyl group, presumably at the YgcA target m⁵U1939, and is unaffected by inactivation of the *ybjF* gene. The spectra have been electronically smoothed using the Proteometrics Inc 'm/z' software program.

m/z	Positions	Sequence 5'-3'	Remarks
3199.4	A1936-G1945	AAAm ⁵ UUCUUGp	m ⁵ U1939 modification
3207.4	C1140-G1149	CUAACCAUGp	
3208.4	C2558-G2567	CCAUUUAAAGp	
3230.5	A221-G230	AAAUCAACCGp	
3253.4	A739-G748	ACUAAUm ¹ Gψm ⁵ UGp	Wildtype m ⁵ U747 (C)
3239.4	A739-G748	ACUAAUm ¹ GψUGp	Knockout m ⁵ U747 (D)

Table 3. Homologues of YgcA(RumA) and YbjF(RumB)

Bacteria	Gene	GenBank accession no.	Identity/similarity score	
			YgcA(RumA)	YbjF(RumB)
<i>Shigella flexneri</i>	<i>ygca</i>	AE015293	99/99	–
<i>Shigella flexneri</i>	<i>ybjF</i>	AE015110	–	98/99
<i>Salmonella enterica</i>	<i>ygca</i>	AE016843	84/91	–
<i>Salmonella enterica</i>	<i>ybjF</i>	AE016840	–	88/92
<i>Salmonella typhimurium</i>	<i>ygca</i>	AE008835	84/91	–
<i>Salmonella typhimurium</i>	<i>ybjF</i>	AE008737	–	87/92
<i>Shewanella oneidensis</i>	<i>SO3456</i>	AE015782	44/62	–
<i>Shewanella oneidensis</i>	<i>SO0980</i>	AE015542	–	56/73
<i>Haemophilus influenzae</i>	<i>HI0333</i>	U32718	46/65	–
<i>Haemophilus influenzae</i>	<i>HI0958</i>	U32777	–	54/70
<i>Pasteurella multocida</i>	<i>PM1866</i>	AE006224	44/62	–
<i>Pasteurella multocida</i>	<i>PM0070</i>	AE006042	–	56/70
<i>Vibrio cholerae</i>	<i>VC2452</i>	AE004315	48/63	–
<i>Vibrio cholerae</i>	<i>VCA0929</i>	AE004420	–	54/68
<i>Yersinia pestis</i>	<i>YPO1336</i>	AJ414147	–	74/83
<i>Haemophilus somnus</i>	<i>Hsom1363</i>	NZ_AAABO02000010	46/62	–
<i>Pseudomonas fluorescens</i>	<i>Pflu2080</i>	NZ_AAAT02000042	42/57	–
<i>Pseudomonas syringae</i>	<i>Psyr2664</i>	NZ_AAABP02000005	40/56	–
<i>Azotobacter vinelandii</i>	<i>Avin0974</i>	NZ_AAAU02000022	40/57	–
Archaea				
<i>Pyrococcus abyssi</i>	<i>PAB0719</i>	AJ248286	26/44	27/45
<i>Pyrococcus abyssi</i>	<i>PAB0760</i>	AJ248286	26/44	25/41
<i>Pyrococcus horikoshii</i>	<i>PH1137</i>	AP000005	28/46	27/47
<i>Pyrococcus horikoshii</i>	<i>PH1259</i>	AP000005	25/44	24/42
<i>Pyrococcus furiosus</i>	<i>PF1172</i>	AE010226	27/43	27/45
<i>Pyrococcus furiosus</i>	<i>PF0827</i>	AE010198	27/43	26/43

Homologues of *E.coli* YgcA(RumA) and YbjF(RumB) revealed in a BLASTp database search. Regions of similarity were found in 57 bacterial species, but only those with identities of >40% and which fulfilled the four criteria described in Materials and Methods are shown in the table. A dash (–) indicates an identity score of <25% compared to the other methyltransferase. Three species of the archaeon *Pyrococcus* are also listed with sequences similar to both RumA and RumB. These have a lower identity score than many of the bacteria and probably have a comparable, but not identical, function to the *E.coli* methyltransferases. The *Pyrococcus* *PAB0719*, *PH1137* and *PF1172* genes show >95% identity to each other, as do the *PAB0760*, *PH1259* and *PF0827* genes, but there is low similarity between the two groups. No clear homologues were found in other archaea or amongst eukaryotes.

synthesis (3,5). This region of the tunnel additionally forms a binding pocket for macrolides and other antibiotics (39,46,47). Structural changes here confer antibiotic resistance (47,48), and can perturb regulatory interactions with the nascent peptide chain (49–51). Therefore, given its conservation and ribosomal location, the U747 methylation might be expected to be involved in one of more of these processes. However, in competition experiments run in rich and minimal media, we have not been able to detect any growth disadvantage for *ybjF* (*rumB*) knockout strains compared to the wild type (not shown). The presence of the U747 methylation could become important under specific growth or stress conditions, but definition of such a physiological role presently remains elusive.

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