

A simple and efficient method to search for selected primary transcripts: non-coding and antisense RNAs in the human pathogen *Enterococcus faecalis*

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ABSTRACT

Enterococcus faecalis is a commensal bacterium and a major opportunistic human pathogen. In this study, we combined *in silico* predictions with a novel 5'RACE-derivative method coined '5'tagRACE', to perform the first search for non-coding RNAs (ncRNAs) encoded on the *E. faecalis* chromosome. We used the 5'tagRACE to simultaneously probe and characterize primary transcripts, and demonstrate here the simplicity, the reliability and the sensitivity of the method. The 5'tagRACE is complementary to tiling arrays or RNA-sequencing methods, and is also directly applicable to deep RNA sequencing and should significantly improve functional studies of bacterial RNA landscapes. From 45 selected loci of the *E. faecalis* chromosome, we discovered and mapped 29 novel ncRNAs, 10 putative novel mRNAs and 16 antisense transcriptional organizations. We describe in more detail the oxygen-dependent expression of one ncRNA located in an *E. faecalis* pathogenicity island, the existence of an ncRNA that is antisense to the ncRNA modulator of the RNA polymerase, SsrS and provide evidences for the functional interplay between two distinct toxin-antitoxin modules.

INTRODUCTION

The ubiquitous Gram positive bacterium *Enterococcus faecalis* is a commensal of the intestinal tract and a major opportunistic pathogen responsible for nosocomial

infections (1). *Enterococcus faecalis* has the ability to damage epithelial-cell DNA and may promote colonic inflammation and cancer by producing reactive oxygen species (2). The dual life style of *E. faecalis* hinges on a balance between host physiology, which imposes environmental constraints, and bacterial adaptation processes (1). About a dozen putative virulence genes have been reported in *E. faecalis*. However, the species features genomic diversity, as well as gene expression levels that vary between isolates, indicating that underlying regulatory processes are major determinants of the balance between commensalism and pathogenesis (3–5). Several chromosomal transcription regulators are involved in stress response, commensalism and/or virulence. Those identified to date affect antibiotic resistance, oxidative stress response, lysozyme resistance and tissue invasion, or may have more pleiotropic roles (6–14). In contrast, a sole example of RNA-dependent post-transcriptional control involved in commensalism and virulence has been described thus far, and concerns the utilization of ethanolamine, an abundant product of the gastrointestinal tract. Two RNA-dependent mechanisms participate in the regulation of this metabolic pathway: one involves an attenuation process in which the RNA-binding protein EutV acts in an ethanolamine-dependent manner; the other uses riboswitch elements (see below) that bind adenosylcobalamine, a cofactor required for ethanolamine degradation [(15) and references therein].

Bacteria reprogram their gene expression in response to environmental constraints partly via diverse categories of regulatory RNAs, mainly non-protein-coding RNAs (ncRNAs), antisense RNAs (asRNAs) and 5'-untranslated regions (5'-UTRs). Two classes of ncRNAs are defined according to the nature of their targets: those that bind

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The authors wish it to be known that, in their opinion, the first two authors should be regarded as joint First Authors.

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proteins and change their activities, and those that pair to mRNA targets and affect their encoding capacity (16,17). The latter generally act via a short sequence partially complementary to their mRNA targets, which are transcribed at a different locus. ncRNAs can also act in *cis* when they are present in an antisense configuration with their mRNA targets (18). 5'-UTRs regulate the expression (transcription or translation) of their downstream mRNA sequence by mediating a switch in their folding. These conformational changes are induced by physical parameters (e.g. temperature or pH) or by binding to small ligands, such as metabolites in the case of riboswitches, e.g. (19–22). In addition, several mRNAs and riboswitches are known to exert regulatory functions by pairing to other RNAs, similarly to ncRNAs, e.g. (23–29). The key roles of regulatory RNAs in coordinating adaptation processes in response to environmental cues have been illustrated in many aspects of bacterial physiology, e.g. gene expression reprogramming during the exit from stationary phase, global stress response, virulence, carbon utilization, iron and envelope homeostasis, and antibiotic synthesis, e.g. (17,30–33).

The development of whole-transcriptome studies via tiling arrays or deep RNA-sequencing (RNA-seq) methods in the past two years has transformed our view on bacterial genomes, and RNA-based regulation has been revealed through the discovery of a plethora of unexpected transcripts and potential regulatory RNAs (34). To date, around fifteen 'RNA landscapes' have been inventoried in evolutionary distant species, revealing a general complexity of genomic organization due, in part, to the numerous promoters nested in open reading frames (ORFs), and to the existence of many ncRNAs and asRNAs (10–46% of annotated ORFs). Importantly, they establish that transcriptional organization on bacterial chromosomes cannot be deduced from the current annotation with ORFs alone [see (35) and references therein]. Despite the power of these methods, they remain demanding in terms of cost and expertise, and cannot be routinely performed in most laboratories. In addition, functional insights inferred from the majority of data published thus far are still limited as the full length RNA transcripts cannot be distinguished from the processed transcripts, which is a major issue in understanding gene expression control. Only recently, a differential RNA-seq method (dRNA-seq) has been applied to access primary transcripts by the enzymatic elimination of processed RNA forms (36–39). We present here a simple, reliable and sensitive 5'RACE-derivative method, coined '5'tagRACE', in which 5' RNA extremities are differentially tagged with two different sequences that distinguish full-length from processed RNA species. This method enables via a classical polymerase chain reaction (PCR) to specifically detect and map primary transcripts or processed ends in bacteria whose genome sequences are available. The 5'tagRACE may be used as a stand-alone method as presented here, or it may serve to generate material for high-throughput RNA sequencing to efficiently distinguish primary from processed transcripts. To benchmark the sensitivity and reliability of the 5'tagRACE, we applied the 'classical' approach to

search for *in silico* predicted ncRNAs encoded by the *E. faecalis* chromosome. Out of 45 selected loci in the chromosome of the clinical isolate V583 (40), 28 were active for transcription. Among those, a total of 49 primary transcripts were characterized. Ten novel putative mRNAs and 29 novel ncRNAs were revealed, including 7 ncRNAs expressed from the described pathogenicity island of the *E. faecalis* strain (8). Remarkably, an antisense organization was demonstrated for a third of the 28 loci selected for study, among which we discovered an antisense ncRNA to the housekeeping ncRNA modulator of the vegetative RNA polymerase, SsrS (41). Results of this study present the first characterization of ncRNAs encoded by the chromosome of the human pathogen *E. faecalis*. They also provide the 5'tagRACE, a simple and reliable method which is generally applicable and should be of major interest for routine studies of bacterial gene expression.

MATERIALS AND METHODS

In silico predictions of ncRNAs and selection of chromosomal loci

In silico *E. faecalis* ncRNA predictions were done as we described previously (42), (Supplementary Figure S1). Our search was focused on chromosomal intergenic regions (IGRs) of the clinical *E. faecalis* isolate V583 (40,43). Positions and sequences of IGR (defined as DNA sequences between annotated ORFs) were downloaded from the TIGR website: (http://cmr.jcvi.org/cgi-bin/CMR/shared/MakeFrontPages.cgi?page=intergenic_region&crumbs=genomes). The analysis was restricted to IGRs of at least 150 bp; regions carrying rRNAs, tRNAs, known riboswitches and introns were excluded from further analysis. About 832 such loci were tallied and 236 were picked randomly for *in silico* predictions of ncRNAs, among which 45 IGRs were selected to be probed by 5'tagRACE (see Supplementary Data).

ORF predictions were performed for all the newly discovered transcripts, enabling us to discern non-coding RNAs (ncRNAs) from putative mRNAs (see Supplementary Data).

Construction of a 5'tag-cDNA library

Total RNA was prepared from V583 grown in S and R conditions. Concentration and quality of RNA preparations were evaluated by measurements at 260 and 280 nm, and by denaturing agarose gel electrophoresis. A mixture of both RNA stocks (55 µg each) was used to detect ncRNAs expressed in at least one of the two conditions. DNA contamination of the RNA mixture was eliminated with 22 units of Turbo DNase, following the manufacturer's protocol (AMBION-Applied Biosystems). About 10 µg of the RNA preparation was kept as PCR template to verify the absence of genomic DNA contamination. A 15 nt RNA adaptor (FW.5bt) was added to the remaining 100 µg of RNA at 50 pmol FW.5bt/µg bacterial RNA (Supplementary Table S1). The 5'-monophosphate groups carried by 5'-processed RNA ends or DNA oligonucleotides generated by the DNase treatment were

ligated to FW.5bt using T4 RNA ligase (New England Biolabs) at 4 U/ μ g bacterial RNA; the ligation was performed for 18 h at 16°C in 10% dimethyl sulfoxide (DMSO). The excess of non-ligated adaptor FW.5bt was removed by exclusion chromatography using Micro Bio-Spin® 30 columns (BioRad). In a series of experiments, we previously determined that these columns do not selectively retain folded RNAs, nor affect their integrity, by determining recovery efficiency of synthesized RNA (data not shown). Half of the RNA mixture ($\sim 50 \mu$ g) recovered from the column was then treated with the tobacco alkaline phosphatase (TAP; Epicenter Biotechnologies) at 2 U/ μ g RNA, transforming the 5'-triphosphate groups present at the 5'-ends of primary transcripts ('transcription start site' or 'TSS') into monophosphate groups (44). The TAP-treated RNA stock is referred to as '+T'. Identical treatments were applied to the other half of the RNA mixture, except that TAP was omitted, providing a negative control RNA mixture (-T) for the specific action of TAP on primary transcripts. RNA +T and -T stocks were ligated to a second RNA adaptor (FR.RNA5; Supplementary Table S1) at 75 pmol/ μ g of bacterial RNA. Note that the sequence and length of FR.RNA5 (38 nt long, Supplementary Table S1) may be adapted for specific uses such as high-throughput sequencing. The equivalent of 4 μ g bacterial RNAs +T and -T RNAs were then reverse transcribed using the random primer NONAX and the SuperScript™ III reverse transcriptase (ssIII), (Invitrogen), using 75 pmol NONAX/1 μ g bacterial RNA and 50 U ssIII. Bacterial RNA and the NONAX random primer were mixed, incubated for 5 min at 65°C, and rapidly cooled on ice. The ssIII was then added and the annealing of NONAX to bacterial RNAs was allowed by incubating the mixture 5 min at room temperature. Reverse transcription was performed by four successive steps of 15 min at 42, 50, 55 and 60°C. The reaction was stopped by a 5 min shift to 85°C, and RNA was eliminated by adding RNase H (New England Biolabs) for 20 min at 37°C. Reverse transcribed samples +T and -T both contain single-stranded cDNA synthesized from RNA processed forms tagged at their 5'-ends by the oligonucleotide FW.5bt. The differential use of TAP generates specifically tagged TSSs by the oligonucleotide FR.RNA5 in the +T sample (this constitutes the '5'tag-cDNA library'), in contrast to -T which contains untagged TSSs (Figure 1).

The 5'tag-RACE procedure

Within the 5'tag-cDNA library (+T), a given transcript was detected by PCR using a specific oligonucleotide and the primer FR.DNA5 which specifically hybridizes to the reverse transcribed sequence of FR.RNA5 ligated to TSSs (Supplementary Table S1). Amplifications were performed using 0.5 pmol/ μ l of specific primer, 1.25 pmol/ μ l FR.DNA5, 1.25 U Hot Gold Star DNA polymerase (Eurogentec, S.A.) and 0.5 mM dNTPs in final a 20 μ l reaction. The amount of cDNA template used was arbitrarily defined as the concentration of reverse transcribed bacterial RNA utilized for the PCR

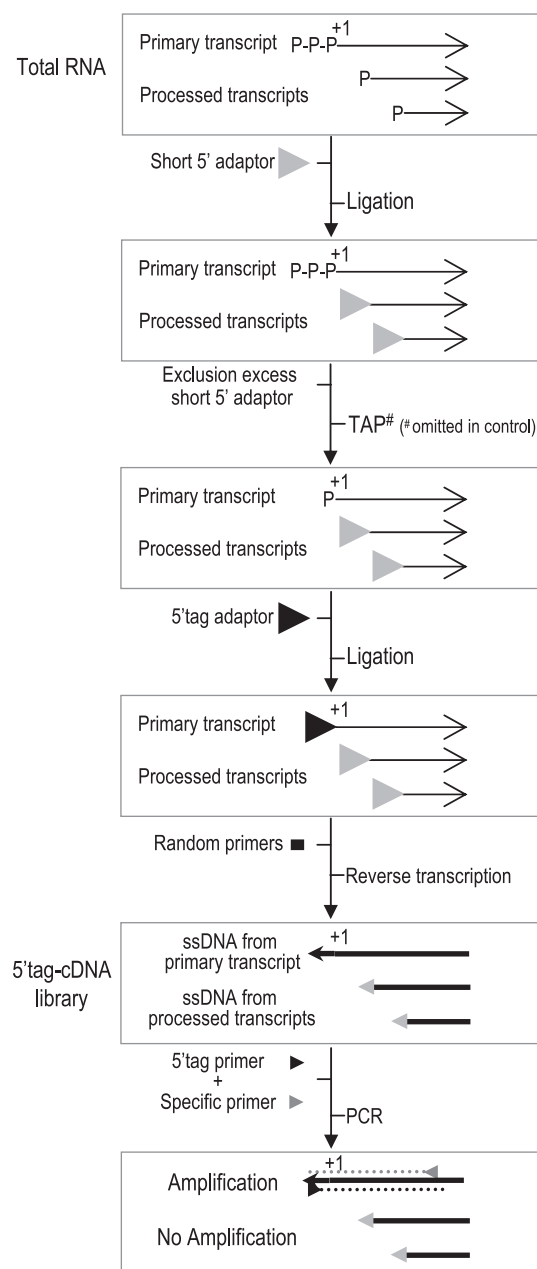


Figure 1. Principle of the 5'tagRACE method. 5' monophosphate RNA extremities generated by processing or degradation are ligated to an excess of a 5' RNA adaptor ('Short 5' adaptor'), resulting in inactive 5'-ends for subsequent ligation steps. The excess of 5' RNA adaptor is eliminated via exclusion chromatography. 5' triphosphate ends of primary transcripts (i.e. '+1' or 'TSS') are then transformed into 5' monophosphate by the TAP enzyme and ligated to a second 5' RNA adaptor ('5'tag adaptor'). Reverse transcription is performed using random primers to generate a cDNA library (5'tag-cDNA library) containing two types of 5'tagged single-stranded-DNA molecules: those ligated to the first 5' adaptor (Short 5' adaptor), i.e. processed ends, and those ligated to the 5' tag adaptor, i.e. TSSs. The use of a specific primer to the 5' tag adaptor and an oligonucleotide specific to the selected RNA (transformed into cDNA) will allow the specific amplification by PCR of the cDNA synthesized only from the primary RNA. An untreated TAP 5'tag-cDNA library is used as negative control (hash sign).

reaction and named hereafter 'Equivalent RNA' (Eq.RNA). We determined a range of Eq.RNA values for which primary transcripts of the predicted ubiquitous ncRNAs (SsrS, SsrA, RnpB and Ffs) were specifically detected (data not shown). For our studies, Eq.RNA values ranged from 1.25 pg/ μ l to 1 ng/ μ l. PCR conditions applied were: 10 min at 94°C, and then 40 cycles (94°C, 40 s; 58°C, 40 s; 4 min at 72°C), and 10 min at 72°C. When using the +T sample as template, the specificity of amplified products was deduced by comparing bands to those obtained with the -T sample. The authenticity of amplicons was then confirmed by sequencing and the TSS was deduced as the first nucleotide of the targeted transcript ligated to the 3'-end of the FR.DNA5 oligonucleotide. Sequencing was performed directly on the amplicon when a single PCR product was obtained or after cloning of the PCR products in the pCRII-Topo vector when several amplicons were obtained.

3'-end mapping

The RNA samples used to generate the 5'-tag-cDNA library were also used for 3' mapping. Eighty micrograms of DNase-treated RNA was dephosphorylated with 40 U of calf intestinal alkaline phosphatase (New England Biolabs) for 1 h at 37°C. The 3'RNA adaptor FR.idT.R1 was then added to the treated RNA to a 75 pmol/ μ g RNA concentration (Supplementary Table S1). The RNA mixture was incubated for 5 min at 90°C and cooled on ice before adding T4 RNA ligase buffer and DMSO to a 10% final concentration. The RNA mixture was then split into two equal volumes each containing ~40 μ g RNA. Four units of T4 RNA ligase per μ g of bacterial RNA was added to one of the samples and replaced by water in the second sample. Samples were then incubated for 18 h at 16°C, and then reverse transcribed using the random NONAX primer and the ssIII system in the same conditions as described for the 5'-tag-cDNA library. The detection of specific 3' extremities was performed by PCR using a specific primer for hybridization with a selected RNA and the oligonucleotide FR.3tail.D1 (Supplementary Table S1), as described for the 5'-end mapping. Specific amplicons obtained with the RNA samples treated here above with T4 RNA ligase were sequenced. The amounts of cDNA template employed for 3'-end mapping were in the same range as those used for the 5'tagRACE (from 1.25 pg/ μ l to 1 ng/ μ l Eq.RNA).

Nomenclature of RNAs characterized

Each primary transcript detected by 5'tagRACE and identified by sequencing was named 'Ref' for RNA in *E. faecalis*. Each *ref* gene was arbitrarily assigned to the number of the original IGR and the name of the specific primer used for the 5'tagRACE, e.g. *ref1C* corresponds to the gene found in the IGR FR1 and whose transcript was detected with the primer FR1C (Supplementary Table S1 and S2). When several TSSs were mapped with a single oligonucleotide, numbers were added with respect to genome coordinates; e.g. *ref8C1* locates upstream *ref8C4*.

Standard procedures

Bacterial strains, growth conditions, plasmids, primers and RNA procedures are provided as supplementary data.

RESULTS AND DISCUSSION

Prediction of non-coding RNAs

The majority of known chromosomal bacterial ncRNAs are expressed from genes nested in sequences flanked by annotated ORFs (IGRs), e.g. (37,39,42,45–50). We therefore searched for ncRNAs in the IGRs of *E. faecalis* V583 chromosome (40). Out of 883 total IGRs, 236 were picked randomly and analyzed *in silico* for putative ORFs, RNA secondary structures and 'orphan' transcriptional terminators as described (42). Among them, 45 IGRs were selected as potentially containing ncRNAs genes and kept for further investigation based on the absence of putative translatable ORF and an empirical favorable defined RNA folding (42), thus excluding 191 IGRs (Supplementary Figure S1 and Table S4). Among the 45 selected IGRs, 4 were predicted in the Rfam database to correspond to the highly conserved bacterial ncRNAs, SsrS (6S), RnpB, Ffs (4.5S) and SsrA (tmRNA), [(42,46) and references therein], and another was flanked by syntenic genes *pnpA* (*ef3064*) and *rpsO* (*ef3065*), and shown in *Escherichia coli* and *Listeria monocytogenes*, two divergent bacterial species, to contain an asRNA to *pnpA*, SraG and RliD, respectively (42,45).

The 5'tagRACE procedure to probe for selected primary transcripts

Thus far, *in silico* predictions of ncRNAs have been experimentally tested by Northern blot [e.g. (42,45,46,48,51)], which confirms the existence, orientation and relative abundance of a given transcript. However, high numbers of predicted candidates render the method tedious, and detection may fail when RNA amounts are below a certain threshold (38). To systematically screen our predictions and as a more information-rich alternative to northern blotting, we developed a simple and sensitive 5'RACE-derivative method that we named 5'tagRACE, which allows the selective amplification of primary RNA 5'-ends (Figure 1). The 5'tagRACE was inspired from the detailed 5'RACE protocol (44), and its original description (52), and consists of: (i) ligation of a first 5'RNA adaptor (FW.5bt) to the 5'-monophosphate group present at the 5' processed ends of RNA species, (ii) enzymatic modification of 5'-triphosphate groups of primary transcript ends (TSSs) into 5'-monophosphate groups, and (iii) ligation of resulting 5'-monophosphate ends to a second 5'-RNA adaptor (FR.RNA5), (Figure 1). The treated RNA is then reverse transcribed to generate a cDNA library ('5'tag-cDNA library') where 5'-RNA ends can be distinguished according to the sequence of the ligated 5'-RNA adaptor. By using primers specific to a selected RNA (in this study, RNAs predicted by bioinformatics) and to the 5'-adaptor ligated to 'TSSs', the PCR amplification step reveals the primary transcripts of interest (Figure 1). In contrast to the

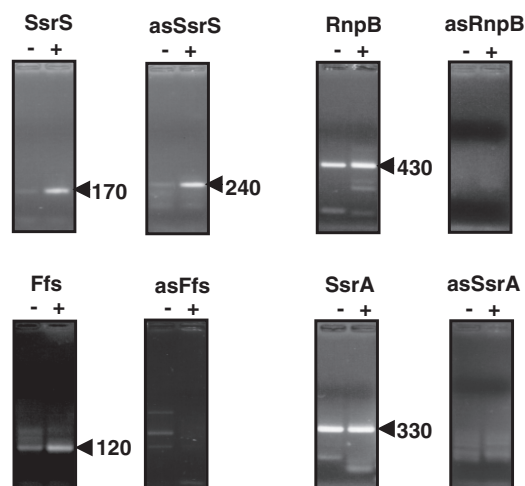


Figure 2. Probing for the housekeeping ncRNAs by 5'tag RACE. Names of RNAs probed are indicated at the top of each panel, antisense RNAs are denoted by 'as'. Amplicons were separated on 2.5% agarose gels and stained with ethidium bromide. At the top of each lane, '-' and '+' refer to untreated and treated TAP RNA samples, respectively. The arrow heads show amplicons that were sequenced and their estimated lengths (nt) on gel. The amount of template used in these set of experiments was 0.05 ng Eq.RNA. With this amount of template, amplicons obtained for RnpB and SsrA with treated and untreated TAP samples were in equivalent amounts.

classical 5'RACE, only DNA strands which are complementary to single strand cDNAs synthesized from primary transcripts by reverse transcription are used as template by the Taq DNA polymerase in the 5'tagRACE procedure (Figure 1).

In a series of pilot experiments we applied the 5'tagRACE method to identify conserved housekeeping ncRNAs SsrS, SsrA, RnpB and Ffs, observed in diverse bacterial species to be highly abundant in their processed and active forms, but whose primary transcripts are present in low amounts [(53) and references therein]. For those ubiquitous ncRNAs, we determined that their detection was specific within Eq.RNA values ranging from 1 ng/ μ l to as low as 1.25 pg/ μ l (see 'Materials and Methods' section; Figure 2). Sequencing of amplicons confirmed the presence of selected primary transcripts in their predicted orientation and allowed us to map TSSs for *ssrS*, *rnpB*, *ffs* and *ssrA* in *E. faecalis* V583. Sequences and orientations obtained for these ncRNAs agreed with sequences predicted in the Rfam database. Note that 5'tagRACE uses at least 10-fold less total RNA than the classical 5'RACE method (44,52). To further confirm the validity of 5'tagRACE, we performed negative controls by searching for *a priori* non-existing RNAs; this was done by probing our 5'tag-cDNA library for RNAs transcribed in the opposite direction (antisense) to *ssrS*, *rnpB*, *ffs* and *ssrA* genes, using specific oligonucleotides nested in the sequences of the four known ncRNAs (Supplementary Table S1). Within the range of cDNA template defined above (from 1.25 pg/ μ l to 1 ng/ μ l of Eq.RNA) and as expected, no asRNAs were detected for RnpB, Ffs and SsrA (Figure 2), attesting to the reliability of the 5'tagRACE method. Unexpectedly, we detected an

antisense transcript to SsrS, which was further confirmed by sequencing and northern blotting (see below).

Taken together data obtained with the housekeeping ncRNAs SsrS, RnpB, Ffs and SsrA validate 5'tagRACE as a robust method to reveal the existence of unknown transcripts. It also lifts uncertainties associated with the classical 5'RACE method and northern blotting, in particular, when the RNA under study is predominant in its processed forms or when it competes with abundant non-specific RNAs, mainly ribosomal RNAs that constitute up to 80–90 % of total RNA (54).

Detection and characterization of predicted ncRNAs within selected IGRs

Primary transcripts encoded by *in silico* selected IGRs (Supplementary Table S2) were probed within the range of Eq.RNA values defined for SsrS, SsrA, RnpB and Ffs ncRNAs (from 1.25 pg/ μ l to 1 ng/ μ l). A single specific primer was used for each candidate with a predicted orientation, as deduced from detection of a transcription terminator (Supplementary Table S2). For candidates whose transcription orientation was uncertain, the two possibilities were probed by using two specific and complementary primers (Supplementary Table S1). In total, 66 transcripts were tested within the 45 IGRs (Supplementary Table S2; the 4 IGRs carrying ubiquitous ncRNAs are considered here). Seventeen of those IGRs either failed to give signals, or gave signals that were complex or non-specific, and were not analyzed further (Supplementary Table S2, Supplementary Figure S2). Among the remaining 28 IGRs, sequencing of resulting products enabled us to map a total of 45 TSSs (Figures 3 and 4 and Supplementary Figure S3). In this set of experiments, the asRNA detected for SsrS (Ref44 or 'asSsrS' in Figure 2), was also sequenced and confirmed (Figure 2 and Supplementary Figure S3). Thus, data obtained with the four housekeeping ncRNAs and the pool of predicted Ref candidates demonstrated the reliability and sensitivity of the 5'tagRACE method (Supplementary Figure S3). For comparison, we independently tested the feasibility of using the classical 5'RACE method to map TSSs for genes *ssrS*, *ref44*, *ref25C* and *ref40* (44). Only Ref40 provided an unambiguous TSS signal identical to that obtained with the 5'tagRACE method, while data from the other samples were inconclusive (Supplementary Figure S4 and data not shown). To further confirm the reliability of 5'tagRACE, the end-mapping data was directly used as a guide for plasmid overexpression of selected Ref RNAs under control of their own promoters (e.g. *ref25C*, *ref40*, *ref44* and *ssrS*). Results obtained confirmed that the characterized TSSs and promoter predictions were real (Supplementary Figure S5, and see below). The above analyses demonstrate the selectivity, sensitivity and reliability of the 5'tagRACE method.

To further illustrate the general interest of the 5'tagRACE method, the 5'tag-cDNA library used in this study was submitted to deep sequencing. From our data set, we recovered identical TSSs as those mapped for several mRNAs by others, e.g. *sodA*, *ace*, *glnQ*, *fsrB* and *gelE* (55–58), attesting to the robustness of the

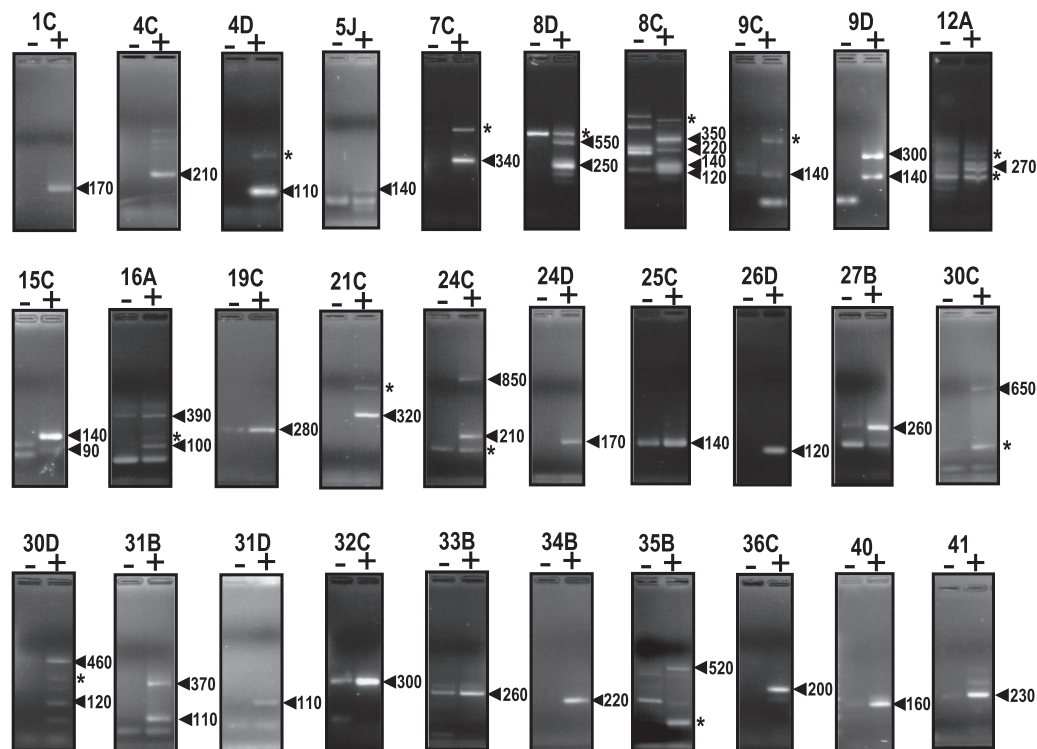


Figure 3. Primary transcripts detected by 5'tagRACE in IGR candidates. Names of oligonucleotides used to probe each RNA candidate are indicated at the top of each panel. Signs '+' and '-' refer to treated or untreated TAP RNA samples, respectively. Numbers on the right indicate lengths estimated for 5'tagRACE PCR products observed on 2.5% agarose gels. Amplicons below ~80 bp were considered as aberrant products due to the average length of the specific primer (~25–30 nt) and the 5'tag RNA adaptor (38 nt), (Supplementary Table S1). Asterisk: signals not sequenced or corresponding to an unrelated primary transcript as determined by sequencing. 5'tagRACE data obtained for rejected IGR candidates are shown in Supplementary Figure S2.

5'tagRACE. In depth analysis of whole genome data by combining 5'tagRACE and RNA-seq technology will be presented elsewhere.

3'-end mapping of unveiled primary transcripts

To get further insight on Ref RNAs, their 3'-ends were mapped using the 3'RACE method and specific primers complementary to those used for 5'tagRACE (Supplementary Table S1). In some cases, 3'-end mapping revealed long amplicons (>500 nt) and complex patterns (Refs 21C, 15C1-2, 26D and 31D), indicating that these transcripts most likely comprise downstream ORFs, and could correspond to 5'-UTRs; these were not analyzed further. In several cases (Refs 1C, 4D, 5J1-2, 19C, 25C, 27B, 32C and 34B, and RnpB), the mapped 3'-ends corresponded to the predicted rho-independent transcription terminators (Figure 4 and Supplementary Figure S3). For Ref44 and Ffs, the 3'-extremities we determined did not correspond to predicted transcription terminators, suggesting rapid processing between the transcription terminator and the specific primer used, and/or steric hindrance during the ligation step due to RNA secondary structures (Supplementary Figure S3). No transcriptional terminators were predicted for Refs 8C1-4, 8D1-2, 24C1-2, 30C and 30D1-2, and SsrS. The 3'-ends of these RNAs overlap with other transcripts expressed from the reverse complementary DNA strand, indicating an antisense

organization (Figure 4 and Supplementary Figure S3). Interestingly, the absence of predictable transcription terminators for these Ref asRNAs evokes recent observations in *Synechocystis* sp., which suggested that asRNA counterpart genes may provoke transcriptional arrest or mutual attenuation without requiring intrinsic terminators (59).

Features of ref genes

Figures 4 and 5 summarize characteristics and positions of the 45 identified *E. faecalis* V583 Ref RNA sequences. Below we classify these RNAs according to their main features.

Non-coding Ref RNAs. Among the characterized transcripts (including SsrS, RnpB, SsrA and Ffs), no translatable ORFs were found in sequences of 31 of them, allowing us to conclude that these transcripts are most certainly ncRNAs. Among those, 27 Ref RNAs were novel and their primary lengths ranged from 128 to 755 nt (Supplementary Table S2 and Figure S3, Figure 4). Two classes of Ref ncRNA genes were distinguished according to their transcriptional organization: (i) ncRNA genes embedded in annotated IGRs (e.g. *ref7C*, *8C1-4*, *19C*, *25C*, *27B*, *rnpB*), resembling the vast majority of ncRNA genes described to act on *trans* encoded mRNAs or proteins; (ii) ncRNA genes spanning protein-encoding genes transcribed in the opposite

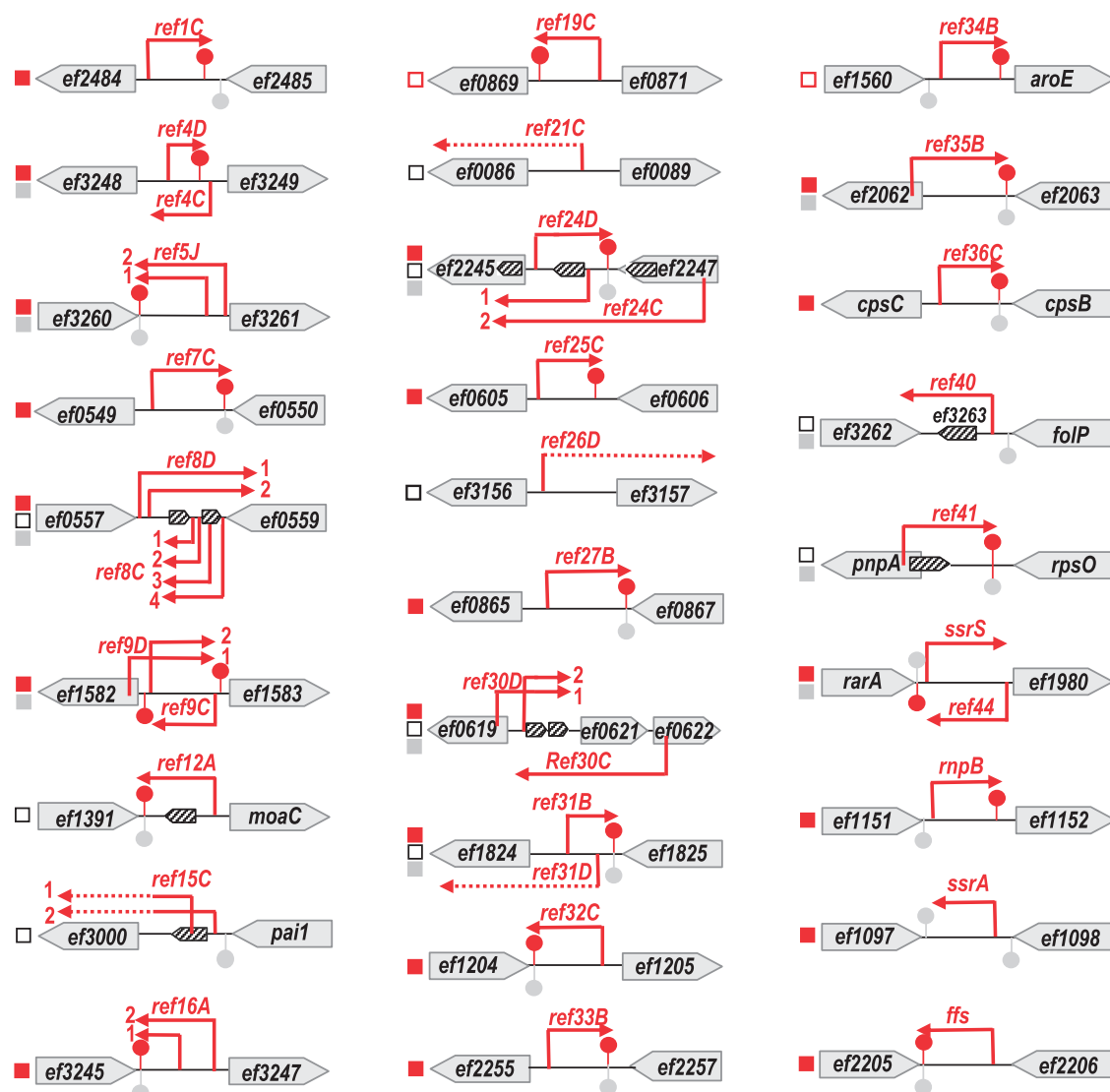


Figure 4. *ref* loci and features of Ref RNAs. Squares are: 'red filled' for ncRNAs; 'red empty' for ncRNAs or riboswitches; 'black empty' for potential 5'-UTRs; and 'gray' for antisense organization. Red arrows symbolize characterized transcripts, dotted lines represent putative 5'-UTRs. Red and gray lollipops represent putative transcription terminators. Annotated ORFs in the V583 genome are shown by gray horizontal arrows, and predicted ORFs in Ref RNA sequences are symbolized by diagonally striped bar arrows. Representation is not at scale; sequences are provided as supplementary data (Supplementary Figure S3).

direction, which may act as antisense. Seven *ref* ncRNA genes (*ref7C*, *8C1-4*, *25C*, *30C*) are located within the V583 pathogenicity island (PAI) and present in closely related *E. faecalis* isolates (Supplementary Table S3). To our knowledge, this is the first description of ncRNAs expressed from the *E. faecalis* chromosome. This collection, and the approach used for its identification, should open the way for characterizing RNA-based processes that govern the physiology of a major bacterial agent of nosocomial infections.

Non-coding RNAs or novel riboswitches? Sequence analysis of Ref19C and Ref34B showed the absence of translatable ORFs, suggesting that these RNAs would also be non-coding transcripts. However, their transcription terminators are spaced by ~100 nt from translation

start codons of the downstream ORFs (Ef0869 and AroE, respectively) transcribed in the same direction, which resembles the organization described for riboswitches or regulatory 5'-UTRs (Figure 4), (21). Although Ref19C and Ref34B RNAs were not predicted within Rfam database as 5'-UTRs regulatory elements, we cannot fully rule out such a possibility (Figure 4 and Supplementary Figure S3), (60).

Ref RNAs encoding peptides. Ten characterized Ref RNAs were predicted to contain small translatable ORFs, ranging in size between 24 and 54 amino acids. None of them was predicted earlier (61). Properties and analysis of these putative ORFs are summarized in Supplementary Figure S3. Among these Ref mRNAs, remarkably Ref8D1 and Ref8D2, located on the PAI,

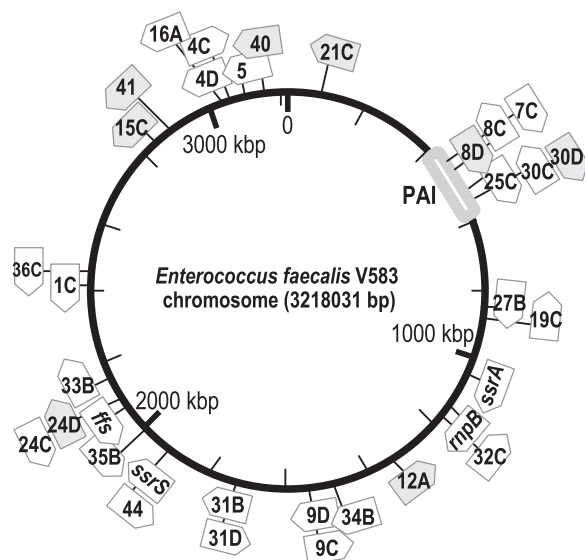


Figure 5. Distribution of Ref RNAs along the chromosome of V583 *E. faecalis* strain. ncRNAs characterized are shown in white; potential mRNA and 5'-UTRs are in gray. Arrows indicate transcription orientation. PAI: pathogenicity island.

are reminiscent of the group I toxin-antitoxin (TA) modules, since they encode a 35 amino acids peptide predicted to contain a trans-membrane domain (the toxin) and are transcribed from the complementary strand to *ref8C1-4* genes expressing antisense ncRNAs (the anti-toxins), (62). Besides putative encoded toxins, peptides are known to participate in regulatory processes related to cell-cell communication in Gram positive bacteria, such as virulence in *Staphylococcus aureus*, competence in *Bacillus subtilis* and *Streptococcus pneumonia* and 'sex' in *E. faecalis* (63–65). Functions of predicted small ORFs in Ref RNAs remain to be established, but as in other species, their occurrence likely reflects a variety of physiological aspects where they participate, e.g. (39,66).

Another peptide-encoding RNA was *ref41*, which bears a potential translatable 43 amino acids ORF transcribed on the opposite strand to the highly conserved *pnpA* gene, encoding the polynucleotide phosphorylase (40). A similar antisense gene was previously reported in two evolutionary distant species, *E. coli* and *L. monocytogenes*, respectively called *sraG* and *rliD* (42,45), indicating a functional relevance and the possible ubiquity of *ref41* in the bacterial kingdom.

Antisense organization of ref loci. Ten of the 28 IGRs displayed transcriptional activity on both DNA strands, thus generating antisense organizations (Figure 4 and Supplementary Figure S3), an occurrence that correlates with the range of frequencies estimated in whole genome analysis using tiling arrays or RNA-seq methods [~30 %; e.g. (39,49,59)]. Three classes of antisense organizations can be defined according to the nature of transcripts involved: (i) ncRNA antisenses to mRNAs, e.g. *ref8C1-4/Ref8D1-2*, *Ref24D/Ref24C1-2*, *Ref30C/ef0621-22* and *ref35B/ef2062*; (ii) ncRNAs antisenses to ncRNAs, e.g. *ref4D/ref4C*, *ref9D1-2/ref9C*, and among

those *ref44* which encodes an asRNA to the RNA polymerase modulator ncRNA, *SsrS* (Figure 4 and Supplementary Figure S3); and (iii) mRNAs antisenses to mRNAs, e.g. *ref40/ef3262* (see below), *ref41/pnpA*, *ref8D1-2/ef0559* and *ref30D1/ef0619*. The diverse antisense transcriptional organizations found in this study for *E. faecalis* IGRs reflect those described in other bacterial species and in the eukaryotic kingdom (35,67). Antisense organization of genes can result in expression regulation at transcriptional and post-transcriptional levels. Generally, the expression of one gene hampers the expression of its counterpart by different, not necessarily exclusive, mechanisms, such as transcription initiation interference, premature transcription elongation arrest, translation repression and degradation of the formed RNA duplex (18,68). Our finding constitutes the first experimental characterization of antisense organizations in the opportunistic human pathogen *E. faecalis* chromosome. The apparent universality of gene expression control by asRNAs and the emerging examples of their contribution to bacterial virulence (69,70) provide to our results a significant importance towards the understanding of the versatile life style of *E. faecalis*.

Conservation of ref genes. *ref* genes can be distinguished according to their conservation in other bacterial genomes and their occurrence within the *E. faecalis* species. Besides the 4 housekeeping ncRNA genes (*ssrS*, *rnpB*, *ssrA* and *ffs*), only a few *ref* genes were found to be significantly conserved in closely related species (*enterococcus* sp.), (*ref30C/D1-2*, *34B*, *35B*, *41*), and in other Firmicutes (*ref7C* in *S. aureus* and *B. anthracis*), (Supplementary Table S3). Among the 25 sequenced *E. faecalis* genomes (40,71,72), *ref* genes were either present in all sequenced isolates (*ref1C*, *9C/D*, *19C*, *27B*, *32C*, *34B*, *35B*, *41*), or limited to a few *E. faecalis* isolates, including those localized in the PAI. Comparative genomic studies revealed that about two-thirds of the *E. faecalis* chromosome corresponds to the core-genome, and as in other bacteria, the pan-genome, defined by genomic islands and mobile elements, harbor virulence and antibiotic resistance traits, as well as accessory factors participating in adaptative functions and niche preferences (5,73). It is thus tempting to speculate that *ref* genes systematically present would ensure a central role in the maintenance of the *E. faecalis* species, while non-conserved *ref* genes would confer specific genetic traits, and each 'ref repertoire' would contribute to individual fitness capacities of isolates.

Expression of Ref RNAs in selected growth conditions

In the presence of oxygen, *E. faecalis* generates massive amounts of superoxide radicals, which can be damaging to the surrounding cellular host environment. However, in the presence of hemin and oxygen, metabolism is rerouted from fermentation to respiration. When this occurs, the production of superoxides is arrested, as reducing equivalents are routed to the respiration chain (74). These differential properties, which depend on hemin availability, are suspected to contribute to

E. faecalis commensalism and pathogenicity (2,75,76). We pursued the characterization of Ref RNAs in northern blot experiments by visualizing their expression in bacterial growth conditions where oxygen and hemin were differentially available. Although northern blots are less sensitive than RT-PCR-based methods, e.g. (38), they provide information on processing and relative expression levels that is unavailable by the latter method. From the 28 IGRs expressing the Ref RNAs discovered via the 5'tagRACE method, 13 were readily visualized (Figure 6): Ref1C, 4D, 16A, 19C, 25C, 27B, 31B, 32C, 40, 44, SsrS, SsrA and Ffs. In most cases, lengths of RNA molecules estimated by northern blot were in good agreement with sizes deduced from the RACE data (Supplementary Table S2). The amounts of Ref RNAs were comparable in both growth conditions used here, except for Ref25C (see below). Complex expression patterns were observed for a few RNAs (Ref27B, 40, SsrS and Ffs), most likely due to specific processing of primary transcripts, e.g. (77–79). The unambiguous mapping of the TSSs via the 5'tagRACE method proved valuable for dissection of complex expression profiles observed on northern blots (e.g. Ref27B), and could be used in tiling array experiments when detecting operonic mRNAs, e.g. (50).

Ref44, an antisense of the RNA polymerase modulator SsrS

SsrS (or 6S) is a ncRNA that accumulates throughout growth and undergoes endoribonucleolytic processing. In *E. coli*, matured SsrS binds to the RNA polymerase holoenzyme $E\sigma^{70}$ and modulates its promoter usage, thereby defining a subset of genes transcriptionally dependent on the ncRNA. When bacteria enter stationary phase, ~80% of $E\sigma^{70}$ is sequestered in a SsrS– $E\sigma^{70}$ complex. Upon outgrowth from stationary phase, SsrS– $E\sigma^{70}$ complex becomes transcriptionally active, and 14–20 nt RNA products (pRNA and pRNA*) are synthesized, leading to complex dissociation. Transcriptional activity of the SsrS– $E\sigma^{70}$ complex responds to nucleotide concentration and SsrS adjusts the bulk of $E\sigma^{70}$ -dependent transcription to metabolic resources (39,41,80).

Unexpectedly, this study revealed the existence of Ref44, an antisense of SsrS. To confirm this finding, a genomic DNA fragment containing both genes was cloned in a multicopy plasmid vector, and introduced into V583. In this construct, Ref44 and SsrS should be expressed from their own promoters as deduced from 5'tagRACE mappings. Total RNA from this strain was separately and specifically probed for SsrS and Ref44. Both ncRNAs were overexpressed, in keeping with the TSS assignments and attesting that *ref44* indeed encodes an antisense RNA to SsrS (Figure 7A). The function of *ref44* is unknown, but as antisense of *ssrS*, these genes could interfere at transcriptional and post-transcriptional levels. The formation of an SsrS/Ref44 RNA duplex could participate in either degradation or maturation of SsrS, as suggested by our data showing that both ncRNAs are mainly present as processed products (Figure 7A and

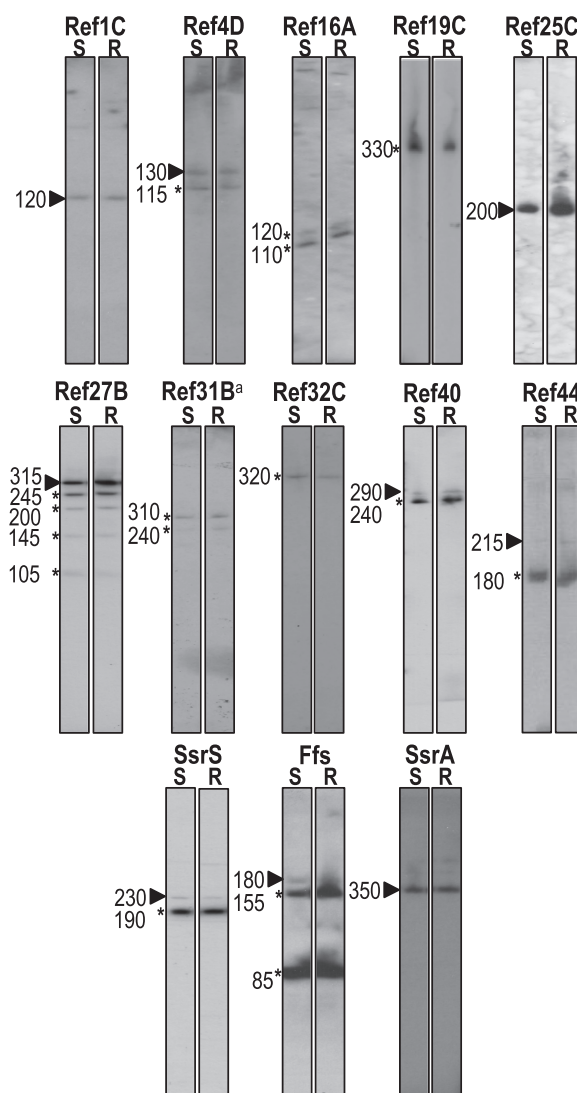


Figure 6. Detection of Ref RNAs by northern blot. Names of detected transcripts are indicated at the top of each panel. Probes used were oligonucleotides designed for 5'tagRACE characterization (Supplementary Table S1). About 10 µg of total RNA was loaded in each lane: on left, static growth conditions (S), and on the right, respiratory (R) growth conditions. Arrow-heads show transcripts with an estimated length agreeing with sizes deduced from end-mapping of primary RNAs, asterisks indicate processed forms. Only 1 µg of total RNA was loaded for SsrS, SsrA and Ffs RNAs. (*): see comments and legend in Supplementary Table S2.

Supplementary Figure S3). In both scenarios, Ref44 may thus modulate the amount of SsrS available to affect $E\sigma^{70}$ activity. To date, we do not know whether such an asRNA is only present in *E. faecalis* species, or beyond, as no antisense to SsrS has been reported in whole-genome transcriptomic data. Our attempts to identify such RNA species by northern experiments in *E. coli*, *B. subtilis* and *S. aureus* did not reveal the existence of an SsrS antisense (our unpublished data).

Ref25C is induced in oxidative stress conditions

We observed that amounts of Ref25C were increased 5- to 10-fold when *E. faecalis* was grown in the presence of

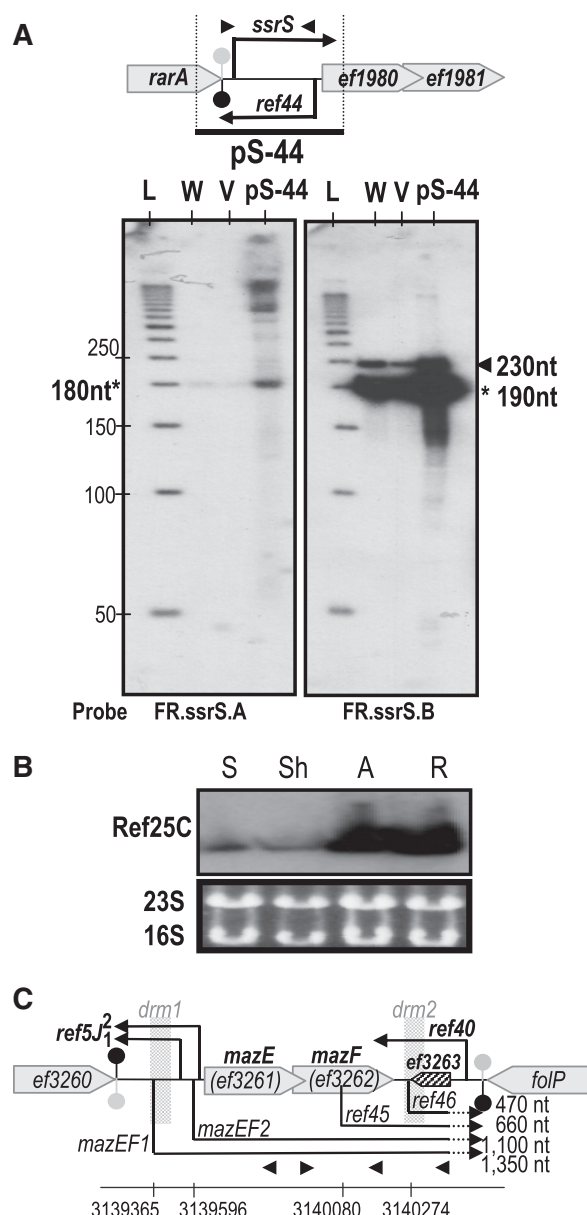


Figure 7. Characterization of three *ref* loci. (A) Expression of SsrS and its antisense Ref44. The genome organization of *ssrS* and *ref44* genes deduced from 5'-tag- and 3'-end mapping is shown in the upper part of the panel. Arrowheads indicate oligonucleotide used as probes for northern blots. 'pS-44': plasmid construct based solely on mapping data and containing the genomic region represented by the black line. The existence of the Ref44 RNA is verified by overexpression-based 5'- and 3'-end mapping (lower part of panel 7A). About 5 µg total RNA was loaded in each lane when probing for Ref44 (right panel), and 1 µg was loaded for SsrS (left panel), respectively. Lanes noted: 'W', wild-type V583 strain; 'V', V583 strain harboring the plasmid vector; 'pS-44', V583 strain transformed by the plasmid pS-44; 'L', 50 base DNA ladder. Lengths indicated for Ref44 and SsrS correspond to the most abundant forms detected. Note that DNA (L) runs slightly faster than RNA. Exposure times were optimized for each panel and signal intensities do not indicate relative abundance between Ref44 and SsrS. Legend is otherwise as in Figure 6. (B) Oxygen-dependent expression of Ref25C ncRNA. Samples were taken at OD₆₀₀ between 0.80 and 0.90. Growth conditions are indicated at the top of each lane: 'S' and 'Sh' correspond to static growth without and with hemin, respectively; 'A' and 'R' correspond to aeration growth without and with hemin, respectively. About 10 µg total RNA was loaded in each lane and the oligonucleotide FR.25C was used as probe for Ref25C.

hemin and oxygen compared to static conditions without hemin ('R' and 'S' conditions, respectively; Figure 6), suggesting a functional relevance of this ncRNA in response to environmental oxygen and/or hemin. To determine whether oxygen, hemin or the combination was responsible for the observed variations, northern blots were performed on total RNAs extracted from bacteria grown in four different environments: static conditions without and with hemin, ('S' and 'Sh, respectively), and aerated conditions without and with hemin ('A' and 'R', respectively), (Figure 7B). Ref25C showed a similar increased abundance in aerated conditions regardless of hemin, compared to static growth conditions, indicating that Ref25C is upregulated in conditions where oxygen is present in the medium (Figure 7B). *ref25C* localizes in the PAI originally described for the V583 isolate and other closely related strains of *E. faecalis* (Figure 5, Supplementary Tables S2 and S3), (8). This gene appears as a single transcriptional unit ending at a rho-independent transcription terminator and is flanked by genes *ef0605* and *ef0606*, encoding a protein of unknown function and one of the two Dps-like proteins contained in the V583 chromosome, respectively, (Figure 4 and Supplementary Figure S3); the other *dps* gene is *ef3233* (40). Dps plays a critical role in oxidative stress by protecting DNA from damage, and contributes to virulence in various bacterial pathogens (81). The location of *ref25C* within the PAI, its vicinity to a *dps* gene and its induction in high oxygen conditions, may suggest its involvement in oxidative stress response. We are currently investigating the functional role of *ref25C* in V583.

Antisense toxin-antitoxin modules I and II

ref5J1-2 and *ref40* genes are transcribed in the same direction and flank the annotated bi-cistronic operon *ef3261-ef3262* expressed from the reverse complementary strand. Remarkably, Ref5J1-2 and Ref40 RNA sequences contain a 167 nt direct repeat motif, called here 'drm1' and 'drm2', respectively (Figure 7C and Supplementary Figure S6). The intervening genes, *ef3261* and *ef3262*, are annotated in the V583 genome as two transcriptional regulators of the broad AbrB- and PemK-protein superfamilies, respectively (40). However, closer inspection by PSI-BLAST revealed Ef3261 (76 amino acids) to be a member of the antitoxin DNA-binding protein MazE

Ribosomal RNAs were used as loading controls (1 µg per lane). (C) Transcriptional organization of the overlapping group I and II TA modules. Location and orientation of specific oligonucleotides used for 5'-tag- and 3'-RACE analysis of *mazEF1-2*, Ref45 and Ref46 transcripts are indicated by arrowheads. Direct repeat motifs *drm1* and *drm2* are shown by gray boxes. A single and common transcription terminator for *mazEF1-2*, Ref45 and Ref46 RNAs was predicted (shown in black) and corresponds to the bidirectional terminator predicted for *folP* (in gray). Putative transcription termination ends for primary transcripts *mazEF1-2*, Ref45 and Ref46 RNAs are shown by dashed lines; numbers at the end of those transcripts are estimated length (see Supplementary Figure S3). Genome coordinates of TSSs mapped for *mazEF1-2*, Ref45 and Ref46 are given at the bottom of the panel.

family, and Ef3262 (121 amino acids) to be a member of the MazE counterpart toxin protein family MazF. Thus, this operon constitutes a type II toxin-antitoxin module, as also suggested in a comparative-genome analysis of TA systems in prokaryotes (82), and will be here called '*mazEF*'. Intriguingly, the small ORF Ef3263 (27 amino acids) contained in Ref40 RNA, lies just downstream MazF and is transcribed from the complementary strand (Figure 7C). Ef3263 was recently described as a member of the TxpA toxin family, a type I TA module in which an asRNA counteracts synthesis of the peptide toxin (83). Thus in V583, the genetic locus flanked by TSSs of *ref5J1-2* and *ref40* genes, comprises type I and type II TA modules transcribed in opposite directions, which suggests that they may impact on the expression of each other. To get a first insight on the possible relationship between these TA systems, we performed in-depth mapping of the transcripts expressed from this locus by 5'tag- and 3'RACE. Data analysis revealed a complex transcriptional organization and rather unusual characteristics for the two distinct groups of TA modules. In addition to Ref5J1-2 and Ref40 RNAs, four other transcripts were found within this region and the complex pattern of overlapping RNA transcripts supports the above hypothesis that expressions of the two TA systems are inter-related (Figure 7C and Supplementary Figure S3).

Interestingly, in the *E. faecalis* OG1RF isolate, the *mazEF* operon is absent, and the duplicated *drm* is resolved into a single 167 bp sequence embedded in a classical *ref40/ref46* group I TA module, making this locus variable among *E. faecalis* species (Supplementary Figure S6 and Supplementary Table S3). TA modules are widespread, including in commensal and pathogenic species where they impact on growth regulation and diverse physiological adjustments such as stress adaptation, antibiotic resistance, biofilm development and persistence in chronic infections (84–86). Biological functions of *mazEF* and *ref40-46* TA modules are unknown in *E. faecalis*, but it may be assumed that these systems also contribute to host-adaptation processes faced by the bacterium when living in the commensal and/or pathogenic state (86). It is also remarkable that in contrast to the *ref40-46* TA system which is conserved among *E. faecalis* strains, the *mazEF* module seems restricted to a limited number of isolates (Supplementary Table S3), and it is thus tempting to speculate that each TA system, separately, might offer a better survival in different niches, and when they cross-regulate, novel fitness capacities could be acquired. Our in-depth RNA mapping provides the basis for molecular and physiological studies of the functional interaction between the two types of TA systems in V583.

CONCLUSION

This study reports a simple, sensitive and general methodological strategy to identify and characterize transcripts, and provides the first data set of ncRNAs in the *E. faecalis* chromosome. Compared to the 5'RACE

method (44,52) from which our method is derived, 5'tagRACE adds a single but crucial RNA ligation step that differentially tags primary and processed RNAs. 5'tagRACE can be utilized for same purposes as the 5'RACE, but in a more reliable manner as illustrated in this work. Most importantly 5'tagRACE can serve a double purpose that is not possible with 5'RACE: (i) simultaneous probing and terminus characterization of unknown transcripts, replacing the two classical successive steps, namely Northern blot and end mapping via 5'RACE; and (ii) a bacterial 5'tag-cDNA library can be used for high-throughput sequencing. By combining 5'tagRACE with RNA-seq, the differential RNA tagging should provide direct access to the ratio between primary and processed forms for a given transcript, thereby offering a dynamic view of bacterial RNA landscapes in terms of regulons through kinetic studies.

Transcriptional analysis of *ref* loci in *E. faecalis* unveiled major features of gene organization, as found with high-throughput methods in other bacterial species, and suggest the importance of RNA-mediated regulatory processes in this species. *E. faecalis* inhabits various ecological niches (soil, water, food, gut) and is able to switch from a normal and harmless commensal bacterium to a life-threatening pathogen (1), attesting to a broad range of regulatory pathways involved in adaptation processes where ncRNAs certainly contribute, as illustrated in other bacterial species, e.g. (17,31). Thus, Ref RNAs being the first ncRNAs described in the *E. faecalis* chromosome, their functional study should have implications in the deciphering of genetic adaptation programs used by the bacterium living in and interacting with the host.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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