Role of the terminator hairpin in the biogenesis of functional Hfq-binding sRNAs

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ABSTRACT

Rho-independent transcription terminators of the genes encoding bacterial Hfq-binding sRNAs possess a set of seven or more T residues at the 3′ end, as noted in previous studies. Here, we have studied the role of the terminator hairpin in the biogenesis of sRNAs focusing on SgrS and RyhB in Escherichia coli. We constructed variant sRNA genes in which the GC-rich inverted repeat sequences are extended to stabilize the terminator hairpins. We demonstrate that the extension of the hairpin stem leads to generation of heterogeneous transcripts in which the poly(U) tail is shortened. The transcripts with shortened poly(U) tails no longer bind to Hfq and lose the ability to repress the target mRNAs. The shortened transcripts are generated in an in vitro transcription system with purified RNA polymerase, indicating that the generation of shortened transcripts is caused by premature transcription termination. We conclude that the terminator structure of sRNA genes is optimized to generate functional sRNAs. Thus, the Rho-independent terminators of sRNA genes possess two common features: a long T residue stretch that is a prerequisite for generation of functional sRNAs and a moderate strength of hairpin structure that ensures the termination at the seventh or longer position within the consecutive T stretch. The modulation of the termination position at the Rho-independent terminators is critical for biosynthesis of functional sRNAs.

Keywords: Hfq; RNA hairpin; Rho-independent terminator; bacterial sRNA; premature termination

INTRODUCTION

Hfq-binding small RNAs (sRNAs), major regulatory RNAs in bacteria, are induced under specific physiological and/or stress conditions and regulate, along with an RNA chaperone Hfq, the expression of target genes at the post-transcriptional level (Waters and Storz 2009; Gottesman and Storz 2010; Vogel and Luisi 2011; Wagner and Romby 2015). The primary role of Hfq is to accelerate base-pairing between sRNAs and target mRNAs to regulate their translation. The secondary role of Hfq is to recruit RNase E near target mRNAs leading to rapid degradation of sRNA–mRNA hybrids (Massé et al. 2003; Morita et al. 2005). Hfq also plays a role in stabilization of sRNAs by protecting them from the attack of ribonucleases (Massé et al. 2003).

The biosynthesis of sRNAs in cells is regulated primarily at the transcription initiation step. The promoter of the individual sRNA gene is under the control of at least one transcription factor that is modulated by the cognate stress. SgrS and RyhB of Escherichia coli are among well-characterized sRNAs. In response to the glucose-phosphate stress, such as accumulation of glucose-6-phosphate, a transcription factor SgrR is activated to stimulate the transcription of sgrS encoding SgrS (Vanderpool and Gottesman 2007). The transcribed SgrS pairs with target mRNAs, such as the ptsG mRNA encoding the major glucose transporter, to either down- or up-regulate their expression to attenuate the glucose-phosphate stress (Vanderpool and Gottesman 2004; Morita et al. 2005; Papenfort et al. 2013). The promoter of ryhB encoding RyhB is under the control of the Fur repressor. Depletion of Fe^{2+} inactivates Fur, resulting in induction of RyhB, which in turn regulates the translation of several mRNAs encoding Fe-binding proteins through base-pairing (Massé and Gottesman 2002; Massé et al. 2003).

The production of sRNAs is regulated not only at the step of transcription initiation but also at the step of transcription termination (Morita et al. 2015). The sRNA genes possess a typical Rho-independent or factor-independent or intrinsic transcription terminator encoding a GC-rich RNA hairpin followed by a run of T residues (d’Aubenton Carafa et al. 1990; Ray-Soni et al. 2016). We demonstrated previously that transcription termination at the sRNA genes is enhanced under stress conditions and thereby contributes to an efficient production of active sRNAs (Morita et al. 2015). A striking feature of the Rho-independent terminators of sRNA genes is that the length of the T residue stretch is longer
than seven (Otaka et al. 2011; Ishikawa et al. 2012). This feature is a prerequisite for generation of functional sRNAs because a poly(U) tail longer than seven is essential for sRNAs to bind efficiently to Hfq (Otaka et al. 2011). To generate surely the functional sRNAs, however, transcription termination must occur at the seventh or longer position within the T residue stretch by preventing "premature" termination. This implies that the modulation of the termination position at the Rho-independent terminators of sRNA genes is quite important for biosynthesis of functional sRNAs. The stability of the terminator RNA hairpin and the length of the T residue stretch are known to be major determinants for the efficiency of transcription termination, although DNA sequences around terminators also affect the termination efficiency (Lynn et al. 1988; Cheng et al. 1991; Reynolds et al. 1992; Ray-Soni et al. 2016). When the terminator hairpin is too weak, elongating RNA polymerase mostly reads through the terminator, resulting in extended forms of sRNAs, which are nonfunctional (Morita et al. 2015). On the other hand, the termination efficiency is expected to increase when the terminator hairpin is stabilized.

It is an interesting possibility that the stability of the terminator RNA hairpin affects not only the termination efficiency but also the termination position within the polythymidine stretch. The aim of the present work is to examine this possibility by using SgrS and RyhB as model sRNAs. We constructed variant sgrS and rplB genes in which the GC-rich inverted repeat sequences of terminators are extended to stabilize the terminator hairpins. We showed by Northern blotting and 3′ RACE experiments that the extension of the stem leads to production of heterogeneous shorter transcripts possessing shortened poly(U) tails. The shortened transcripts can be generated in an in vitro system with purified RNA polymerase, confirming our working hypothesis that the stabilization of the terminator hairpin causes premature termination. We conclude that the control of the termination position within the T residue stretch is critical for the biogenesis of functional sRNAs, and that the terminator hairpin structure of sRNA genes is optimized to produce efficiently functional sRNAs under stress conditions.

**RESULTS**

**Stabilization of the terminator hairpin enhances termination efficiency**

We demonstrated previously by using a "double terminator system" that RNA polymerase frequently reads through the Rho-independent terminator of sgrS under normal growth conditions (Morita et al. 2015). The moderate termination efficiency of the sgrS terminator could be primarily due to a moderate thermodynamic stability (ΔG) of the terminator hairpin. If so, it is expected that stabilization of the hairpin structure of the sgrS terminator enhances the termination efficiency. To test this possibility, we constructed plasmid pSgrS-S-LS1-rplLT ("LS" stands for "long stem") carrying the sgrS-S-LS1-rplLT by inserting four GC base pairs into the sgrS hairpin stem on plasmid pSgrS-S-rplLT. The full DNA sequence of sgrS-S-rplLT and the sequence around the terminator region of sgrS-S-LS1-rplLT are shown in Figure 1A. The predicted terminator RNA hairpin structures, along with their ΔG (kcal/mol), are shown in Figure 1B. The sgrS-S-rplLT is a hybrid gene in which the second terminator derived from rplLT is placed just downstream from the sgrS-S (Morita et al. 2015). SgrS-S, the 3′ portion of SgrS, represents a minimal functional region of SgrS consisting of the base-pairing region and the Hfq-binding module including the Rho-independent terminator sequence (Otaka et al. 2011; Ishikawa et al. 2012). The insertion of GC pairs increases ΔG of the terminator RNA hairpin of SgrS-S from −10.7 to −24.0. Each plasmid was introduced into TM772 (ΔsgrS Δhfq) cells (Table 1). The sgrS-S gene is under the control of an arabinose-inducible P_{BAD} promoter in these plasmids.

Cells were grown in LB medium and the expression of sRNAs was induced by arabinose. Total RNAs were prepared and analyzed by Northern blotting. We first confirmed the previous observation (Morita et al. 2015) that the

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**FIGURE 1.** Effect of extension of the terminator stem of sgrS on transcription termination. (A) DNA sequences of sgrS-S-rplLT and sgrS-S-LS1-rplLT. The sequence corresponding to the sgrS-S is shown as regular letters, whereas the terminator sequence derived from the rplLT is shown as italic letters. The inverted repeat sequences are indicated by horizontal arrows. Nucleotides are numbered from the site corresponding to the 5′ end of sgrS-S. The inserted sequences to stabilize the terminator hairpin are shown as bold letters. (B) The predicted secondary structures and the thermodynamic stabilities (ΔG, kcal/mol) of terminator RNA hairpins without poly(U) sequence were determined according to the Mfold program (Zuker 2003). (C) Analysis of transcription termination. TM772 (ΔsgrS Δhfq) cells harboring indicated plasmids were grown in LB medium. At A_{600} = 0.6, 0.02% arabinose was added and incubation was continued for 5 min. Total RNAs were prepared, and 10 or 0.25 µg of RNA samples was subjected to Northern blotting using the SgrS-S probe and tmRNA probe, respectively.
readthrough occurs frequently resulting in a significant amount of the readthrough product, SgrS-S-\(rplL\), along with the terminated product, SgrS-S (Fig. 1C, lane 1). When the RNAs generated from the \(sgrS-S-LS1\)-\(rplL\)T were analyzed, the readthrough product markedly decreased, whereas the relative abundance of terminated products increased (Fig. 1C, lane 2). This implies that stabilization of the terminator hairpin enhances the termination efficiency, as expected. Interestingly, the \(sgrS-S-LS1\)-\(rplL\)T generates heterogeneous shorter transcripts (Fig. 1C, lane 2), suggesting that the extension of the terminator stem causes premature termination within or before the polythymidine stretch, resulting in transcripts in which the poly(U) tail is shortened.

**Generation of shorter transcripts is caused by stabilization of the terminator hairpin and enhanced by glucose-phosphate stress**

To focus more on the effect of the strength of the terminator hairpin on the generation of the heterogeneous shorter transcripts, we constructed plasmid pSgrS-S-\(LS1\) carrying the \(sgrS-S-LS1\) by inserting four GC base pairs into the hairpin stem of the \(sgrS-S\) on plasmid pSgrS-S (Fig. 2A). We also constructed plasmid pSgrS-S-\(LS2\) carrying the \(sgrS-S-LS2\) by inserting seven GC base pairs (Fig. 2A). The \(\Delta\)G of the RNA hairpin of SgrS-S-\(LS1\) is \(-24.0\), whereas that of SgrS-S-\(LS2\) is \(-32.2\) (Fig. 2B). Each plasmid was introduced into TM772 (\(\Delta\)sgrS \(\Delta hfq\)) cells. Cells were grown in LB medium and expression of sRNAs was induced by arabinose. Total RNAs were prepared and analyzed by Northern blotting. The \(sgrS-S-LS1\) generated heterogeneous shorter transcripts along with the full-length SgrS-S-\(LS1\) as expected (Fig. 2C, lane 2). The generation of heterogeneous shorter transcripts from the \(sgrS-S-LS2\) (lane 3) was more significant compared with the \(sgrS-S-LS1\), indicating that the extent of the production of the heterogeneous shorter transcripts correlates with the strength of the terminator hairpin.

Because transcription termination is enhanced under stress conditions (Morita et al. 2015), we assume that the glucose-phosphate stress may also enhance the production of shorter transcripts. Then, we tested the effect of glucose-phosphate stress on the generation of shorter transcripts at the \(sgrS\) terminator by using \(\Delta\)sgrS \(\Delta hfq\) cells harboring either pSgrS-S, pSgrS-S-\(LS1\), or pSgrS-S-\(LS2\). Cells were grown in LB medium to exponential phase and exposed to nonmetabolizable glucose analog α-methylglucoside (αMG), and then to arabinose. Total RNAs were prepared and subjected to Northern blotting using the SgrS-S probe (Fig. 2D). Both the \(sgrS-S-LS1\) and \(sgrS-S-LS2\) generated the heterogeneous shorter transcripts along with the full-length transcripts (Fig. 2D, lanes

### TABLE 1. Bacterial strains and plasmids used in this study

| Strain/plasmid | Relevant genotype and property | Source/reference |
|----------------|--------------------------------|------------------|
| Strain          |                               |                  |
| IT1568          |                               | Laboratory stock|
| TM772           |                               | Morita et al. 2015|
| TM542           |                               | Kawamoto et al. 2005|
| TM803           |                               | Morita et al. 2015|
| ECK3152         |                               |                  |
| TM894           |                               |                  |
| ECK1281         |                               |                  |
| TM895           |                               |                  |
| ECK1475         |                               |                  |
| TM896           |                               |                  |
| TM905           |                               |                  |
| TM908           |                               |                  |
| TM635           |                               |                  |
| TM820           |                               |                  |
| Plasmid         |                               |                  |
| pAraX           | Derivative of pMW218 carrying araC-P\(_{BAD}\) | Otaka et al. 2011 |
| pSgrS-S-\(rplL\)T | Derivative of pArX carrying sgrS-S-\(rplL\)T | Morita et al. 2015 |
| pSgrS-S-\(LS1\)-\(rplL\)T | Derivative of pArX carrying sgrS-S-\(LS1\)-\(rplL\)T | This study |
| pSgrS-S         | Derivative of pArX carrying sgrS-S | This study |
| pSgrS-S-\(LS1\) | Derivative of pArX carrying sgrS-S-\(LS1\) | This study |
| pSgrS-S-\(LS2\) | Derivative of pArX carrying sgrS-S-\(LS2\) | This study |
| pSgrS-S-\(LS3\) | Derivative of pArX carrying sgrS-S-\(LS3\) | This study |
| pSgrS-S-\(LS4\) | Derivative of pArX carrying sgrS-S-\(LS4\) | This study |
| pSgrS-S (\(\text{bac}\)) | Derivative of pArX carrying \(P_{\text{bac}}\)-sgrS-S | This study |
| pSgrS-S-\(LS2\) (\(\text{bac}\)) | Derivative of pArX carrying \(P_{\text{bac}}\)-sgrS-S-\(LS2\) | This study |
| pRyhB           | Derivative of pArX carrying ryhB | Otaka et al. 2011 |
| pRyhB-\(LS1\)   | Derivative of pArX carrying ryhB-\(LS1\) | This study |
| pRyhB-\(LS2\)   | Derivative of pArX carrying ryhB-\(LS2\) | This study |
We expect that the poly(U) tail is shortened in the heterogeneous shorter transcripts. If so, the levels of shorter transcripts would not be affected by the hfq backgrounds because SgrS-S possessing a shorter poly(U) tail no longer binds to Hfq (Otaka et al. 2011). Then, we investigated the expression of SgrS-S in TM542 (hfq+) and TM772 (Δhfq) cells (Table 1) under the glucose-phosphate stress condition in which the generation of shorter transcripts is enhanced. The abundance of SgrS-S increased in hfq+ cells compared with in Δhfq cells (Fig. 3A, lanes 1, 2). On the other hand, the levels of shorter transcripts were not affected by the hfq backgrounds, whereas the abundance of full-length SgrS-S-LS1 and SgrS-S-LS2 was significantly elevated in Δhfq+ cells (Fig. 3A, lanes 3–6). This strongly suggests that the shorter transcripts lose the ability to bind to Hfq.

To examine directly the Hfq-binding ability of shorter transcripts, we carried out a pull-down assay using cell extracts of hfq-Flag cells harboring pSgrS-S-LS2. Cell extracts were incubated with anti-Flag M2-agarose beads. Proteins bound to the agarose beads were analyzed by Western blotting using anti-Flag antibodies. The affinity-purified Hfq-Flag was treated with phenol and subjected to Northern blotting. As shown in Figure 3B, the full-length SgrS-S-LS2 but not shorter transcripts copurified with Hfq-Flag. This indicates that the shorter transcripts generated from the sgrS-S-LS2 are not able to bind to Hfq. Thus, it is highly likely that the poly(U) tail is shortened in the heterogeneous shorter transcripts.

The abundance of full-length active SgrS-S variants is markedly decreased with accumulation of inactive shorter transcripts in cells harboring pSgrS-S-LS1 or pSgrS-S-LS2.
in particular under the glucose-phosphate stress. Thus, it is expected that the down-regulation of ptsG mRNA, the target of SgrS, is impaired in these cells. To examine this, we performed Northern blot analysis by using the ptsG probe. First, we analyzed the expression of ptsG mRNA in cells harboring the vector plasmid pAraX, pSgrS-S, pSgrS-S-LS1, or pSgrS-S-LS2 under normal growth conditions. Then, we carried out Northern analysis by using RNA samples prepared from cells exposed to the glucose-phosphate stress (Fig. 3D). The expression of ptsG mRNA was not affected by the stress in cells harboring pAraX (Fig. 3D, lane 1). The dramatic reduction of the ptsG mRNA was observed in cells harboring pSgrS-S as expected (Fig. 3D, lane 2). On the other hand, the abundance of ptsG mRNA clearly increased in cells harboring pSgrS-S-LS1 or pSgrS-S-LS2 under glucose-phosphate stress (Fig. 3D, lanes 3,4), indicating that the down-regulation of ptsG mRNA was partially suppressed in these cells. It is apparent that the marked reduction of the active full-length SgrS-S variants is responsible for this suppression of the down-regulation of ptsG mRNA.

Analysis of 3′ ends of transcripts

To verify that the poly(U) tail is indeed shortened in the heterogeneous shorter transcripts, we determined the 3′ ends of RNAs generated from the sgrS-S-LS2 in hfqΔ cells under the glucose-phosphate stress by the 3′-RACE (rapid amplification of cDNA ends) experiment (Argaman et al. 2001; Kawano et al. 2005). In this experiment, total cellular RNAs and a linker DNA oligo are ligated in vitro and the RNA–DNA junction region is amplified by RT-PCR. The amplified cDNA fragments were cloned into a plasmid and subjected to DNA sequence analysis. The 3′ ends of transcripts can be determined by finding the junction between RNAs and the 5′ end of ligated linker oligo DNA. We first analyzed the 3′ ends of transcripts generated from the sgrS-S. The total RNAs were separated by electrophoresis on a polyacrylamide gel containing 8 M urea. The gel region corresponding to SgrS-S was cut out and RNAs were purified from the gel piece (Fig. 4A). The purified RNAs were subjected to the 3′-RACE experiment. The sequences around the junction region of randomly isolated 15 cDNA clones indicate that 11, three, and one clones correspond to SgrS-S possessing 8U, 7U, and 6U tail, respectively (Fig. 4B). Thus, the sgrS-S is able to produce quite efficiently active SgrS-S possessing a long poly(U) tail. Then, we analyzed the 3′ ends of transcripts generated from the sgrS-S-LS2. The gel region corresponding to heterogeneous RNA bands was divided into three pieces as shown in Fig. 4C. RNAs prepared from each piece were subjected to the 3′-RACE experiment (Fig. 4D). The sequence analysis of randomly isolated cDNA clones indicates that RNAs derived from the upper region exclusively correspond to SgrS-S-LS2 possessing a long poly(U) tail. The length of the poly(U) tail of RNAs derived from the middle region varies from four to eight. The length of the poly(U) tail of RNAs derived from the lower region was <3. The 3′ ends of several RNAs obtained from the lower region were mapped in the sgrS terminator region preceding the poly(U) tail. Thus, we conclude that the poly(U) tail of the heterogeneous shorter transcripts is indeed shortened.
Effect of 3′ exoribonuclease on the generation of shorter transcripts and the stability of transcripts

Our hypothesis is that the shorter transcripts are generated by premature transcription termination within or before the terminator polythymidine stretch. However, it is also possible that the shorter products are generated through trimming of the primary transcripts by 3′ exoribonucleases. To test this possibility, we investigated the expression of sgrS-S and sgrS-S-LS2, TM542 (ΔsgrS) cells harboring pSgrS-S (A) or pSgrS-S-LS2 (C) were grown in LB medium. At A_{600} = 0.6, 0.1% αMG was added and incubation was continued for 10 min, and then 0.02% arabinose was added and incubation was continued for 5 min. Total RNAs were prepared and duplicate RNA samples (10 µg) were resolved side-by-side on a 12% polyacrylamide gel electrophoresis in the presence of 8 M urea. One side of the gel was subjected to Northern blotting (A,C). The region corresponding to transcripts was cut out from the other side of the gel (A,C). A single gel piece containing the SgrS-S band was used for transcripts from sgrS-S (A). The gel was divided into three gel pieces (upper, middle, and lower) for transcripts from sgrS-S-LS2 (C). The upper gel piece corresponds to the full-length transcript, whereas the middle and lower gel pieces correspond to heterogeneous shorter transcripts. RNAs were purified from the gel pieces and subjected to 3′-RACE analysis. DNA sequences corresponding to the 3′ region of transcripts were determined by using randomly picked-up plasmid clones containing amplified cDNAs. The sequences and number of clones analyzed are shown (B,D). Others represent the clones in which the 3′ ends of RNAs were mapped in the region prior to the terminator T residue stretch (D).

pSgrS-S-LS2. Cells were grown to exponential phase, and expression of sRNAs was induced by arabinose. Rifampicin was added to prevent further initiation of transcription. RNAs were isolated at various times after the addition of rifampicin, and subjected to Northern blotting (Fig. 5B). The stability of shorter transcripts was essentially the same as that of the full-length transcripts. As expected, the full-length transcript but not the heterogeneous transcripts were markedly stabilized in the hfq^+ background (data not shown). We also examined the fate of the heterogeneous transcripts under the glucose-phosphate stress. In this case, cells were exposed to αMG prior to the addition of arabinose. Again, the stability of shorter transcripts was the same as that of the full-length transcripts, although both transcripts were significantly stabilized by the stress (Fig. 5C).

These results suggest that the shorter transcripts are generated by transcription termination rather than by degradation of primary transcripts.

### The Shortened Transcripts are Generated in Vitro with Purified RNA Polymerase

To verify directly that the shortened transcripts are generated by premature termination, we tried to perform an in vitro transcription assay. For this, we constructed plasmids, pSgrS-S-(tac) and pSgrS-S-LS2-(tac), in which the sgrS-S and the sgrS-S-LS2 are placed under the constitutive tac promoter (P_{tac}), respectively (Fig. 6A). Each plasmid was introduced into ΔsgrS Δhfq cells. Cells were grown in LB medium, and total RNAs were prepared and analyzed by Northern blotting. As expected, the full-length SgrS-S was expressed in cells carrying pSgrS-S (tac), whereas heterogeneous shorter transcripts along with the full-length SgrS-S-LS2 were generated in cells carrying pSgrS-S-LS2 (tac) (Fig. 6B). We purified Flag-tagged RNA polymerase (RNAP-Flag) from the strain carrying the rpoC-Flag allele (Fig. 6C). Then, we carried out an in vitro transcription experiment using RNAP-Flag, pSgrS-S-(tac), or pSgrS-S-LS2 (tac), and four nucleoside triphosphates (NTPs). If our hypothesis is correct, pSgrS-S-LS2(tac) should produce prematurely terminated transcripts. After the transcription reaction, transcripts were subjected to Northern blotting using the SgrS-S probe (Fig. 6D). When pSgrS-S (tac) was used as a DNA template, a transcript corresponding to the full-length SgrS-S generated in vivo was detected as a

![FIGURE 4. Analysis of the 3′ ends of transcripts derived from sgrS-S and sgrS-S-LS2. TM542 (ΔsgrS) cells harboring pSgrS-S (A) or pSgrS-S-LS2 (C) were grown in LB medium. At A_{600} = 0.6, 0.1% αMG was added and incubation was continued for 10 min, and then 0.02% arabinose was added and incubation was continued for 5 min. Total RNAs were prepared and duplicate RNA samples (10 µg) were resolved side-by-side on a 12% polyacrylamide gel electrophoresis in the presence of 8 M urea. One side of the gel was subjected to Northern blotting (A,C). The region corresponding to transcripts was cut out from the other side of the gel (A,C). A single gel piece containing the SgrS-S band was used for transcripts from sgrS-S (A). The gel was divided into three gel pieces (upper, middle, and lower) for transcripts from sgrS-S-LS2 (C). The upper gel piece corresponds to the full-length transcript, whereas the middle and lower gel pieces correspond to heterogeneous shorter transcripts. RNAs were purified from the gel pieces and subjected to 3′-RACE analysis. DNA sequences corresponding to the 3′ region of transcripts were determined by using randomly picked-up plasmid clones containing amplified cDNAs. The sequences and number of clones analyzed are shown (B,D). Others represent the clones in which the 3′ ends of RNAs were mapped in the region prior to the terminator T residue stretch (D).](image-url)
shown in Figure 7B. Each plasmid was introduced into Δ predicted RNA hairpin structures and the base pairs into the hairpin stem of the TM895 (S-LS2 were grown in LB medium containing 0.2% arabinose. At the indicated time after the addition of rifampicin. Total RNAs were prepared and then rifampicin (250 µg/mL) was added. Total RNAs were prepared and 10 µg of RNA samples was subjected to Northern blotting using the SgrS-S. Arrowheads represent the full-length transcripts.

The poly(U) sequence at the 3′-end of sRNAs is a central element of the Hfq-binding module of sRNAs (Otaka et al. 2011; Ishikawa et al. 2012). Hfq hexamer binds to the poly(U) tail through its proximal face (Sauer and Weichenrieder 2011; Schu et al. 2015). We have previously found that the poly(U) tail must be seven residues or longer to achieve a stable binding to Hfq that is required for the regulatory function of sRNAs (Otaka et al. 2011). Therefore, terminators of sRNA genes are expected to fulfill the structural features that ensure the generation of functional sRNAs possessing a long poly(U) tail. In fact, the polythymidine stretch of Rho-independent terminators is longer than those of many other genes vary from four to eight or more (d’Aubenton Carafa et al. 1990). A recent global study on sRNA-target interactions has also revealed that the poly(U) tails of Rho-independent terminators are longer than those.

Effect of stabilization of the ryhB terminator hairpin on expression of RyhB and sodB mRNA

We also examined the effect of stabilization of the terminator hairpin of the ryhB on expression of RyhB. To do this, we constructed plasmid pRyhB-LS1 carrying the ryhB-LS1 by inserting four GC base pairs into the hairpin stem of the ryhB on plasmid pRyhB (Fig. 7A). We also constructed plasmid pRyhB-LS2 carrying the ryhB-LS2 by inserting seven GC base pairs into the hairpin stem of the ryhB (Fig. 7A). The predicted RNA hairpin structures and the ΔG values are shown in Figure 7B. Each plasmid was introduced into TM635 (ΔryhB hfqΔ) and TM820 (ΔryhBΔhfq) cells. Cells were grown in LB medium to exponential phase and exposed to 2,2′-dipyridyl to deplete Fe²⁺, and then arabinose was added to induce the transcription of ryhB. Expression of RyhB and its variants was analyzed by Northern blotting. As shown in Figure 7C, shorter transcripts, along with a decreasing amount of the full-length transcripts, were generated from the ryhB-LS1 and the ryhB-LS2 but not from the wild-type ryhB genes. The generation of shorter transcripts was more significant in the case of ryhB-LS2 compared to ryhB-LS1. Consistent with the increase in shorter transcripts, the amount of the full-length transcripts decreases more significantly in the ryhB-LS2. The abundance of the full-length RyhB, RyhB-LS1, and RyhB-LS2 in the hfqΔ cells was significantly higher than that in the ΔhfqΔ cells, reflecting Hfq binding of the full-length RyhB variants. On the other hand, the hfqΔ backgrounds did not affect the level of the shorter transcripts, suggesting that the poly(U) tails of these RyhB variants are shortened. Then, we examined the expression of sodB mRNA in the hfqΔ cells harboring the vector plasmid pAraX, RyhB, RyhB-LS1, or RyhB-LS2 under the Fe²⁺ depletion. The sodB mRNA is well expressed in cells harboring pAraX (Fig. 7D, lane 1). A marked reduction of the sodB mRNA was observed in cells harboring pRyhB (Fig. 7D, lane 2). The reduction of sodB mRNA expression was moderate in cells harboring pRyhB-LS1 or pRyhB-LS2 (Fig. 7D, lanes 3,4), indicating that the down-regulation of sodB mRNA was partially suppressed in these cells. The suppression of the sodB mRNA down-regulation was stronger in cells harboring pRyhB-LS2 in which the reduction of levels of the full-length active RyhB variant was more significant. Thus, we conclude that the stabilization of the terminator hairpin of the ryhB also leads to generation of the shortened transcript with concomitant reduction of the active full-length transcript.

DISCUSSION

The poly(U) sequence at the 3′ end of sRNAs is a central element of the Hfq-binding module of sRNAs (Otaka et al. 2011; Ishikawa et al. 2012). Hfq hexamer binds to the poly(U) tail through its proximal face (Sauer and Weichenrieder 2011; Schu et al. 2015). We have previously found that the poly(U) tail must be seven residues or longer to achieve a stable binding to Hfq that is required for the regulatory function of sRNAs (Otaka et al. 2011). Therefore, terminators of sRNA genes are expected to fulfill the structural features that ensure the generation of functional sRNAs possessing a long poly(U) tail. In fact, the polythymidine stretch of Rho-independent terminators of sRNA genes are longer than seven (Otaka et al. 2011; Ishikawa et al. 2012), whereas those of many other genes vary from four to eight or more (d’Aubenton Carafa et al. 1990). A recent global study on sRNA–target interactions has also revealed that the poly(U) tails of Rho-independent terminators are longer than those.
Concerning the roles of the terminator hairpin and the T residue stretch in transcription termination at Rho-independent terminators, although the DNA sequences within and near the hairpin also affect the termination efficiency (Lynn et al. 1988; Cheng et al. 1991; Reynolds et al. 1992; Ray-Soni et al. 2016). Indeed, we showed that the extension of the terminator stem markedly enhances termination at the sgrS terminator (Fig. 1). The central finding in the present study is that the extension of the stem leads to generation of significant amounts of heterogeneous transcripts in which the poly(U) tail is shortened (Figs. 1–4, 6, 7). The transcripts with shortened poly(U) tails lose the ability to bind to Hfq and therefore are no longer able to repress target mRNAs (Figs. 3, 7). Importantly, we found that heterogeneous transcripts can be generated in an in vitro transcription system with purified RNA polymerase (Fig. 6), confirming that premature transcription termination within and/or before the terminator polythymidine stretch is at least partly responsible for the generation of shortened transcripts. Thus, the stability of the terminator RNA hairpins of sgrS and ryhB genes is optimized (not too strong and not too weak) to ensure an effective termination at the seventh or longer position of the consecutive T stretch. The moderate thermodynamic stability of the terminator hairpin is associated with most well-characterized sRNAs (Table 2), whereas the hairpin stability fluctuates more widely in many other terminators (d’Aubenton Carafa et al. 1990). The present study, along with previous studies (Otaka et al. 2011; Ishikawa et al. 2012; Morita et al. 2015), led us to conclude that the Rho-independent terminators of sRNA genes possess two common features to generate functional sRNAs: a long T residue stretch (longer than seven) and a moderate strength of hairpin structure that allows the termination at the seventh or longer position of the consecutive T stretch.

In the present work, we studied the role of the terminator hairpin in the biogenesis of sRNAs, focusing on how the hairpin stability affects the termination position at Rho-independent terminators. We constructed variant sgrS and ryhB genes in which the inverted repeat sequences of terminators are extended to stabilize the terminator hairpin. The stabilization of a hairpin at a given terminator is expected to enhance the termination efficiency because the strength of the terminator hairpin, along with the length of the T residue stretch, is an important element to determine the efficiency of termination at Rho-independent terminators, although the DNA sequences within and near the hairpin also affect the termination efficiency.
sRNAs. How the stability of the terminator hairpin controls the termination position is certainly an important question to be addressed for understanding the molecular mechanism of transcription termination at Rho-independent terminators. In this connection, it is interesting to note that shortened transcripts were generated at a Rho-independent terminator in an in vitro transcription system with a low concentration of NTPs (McDowell et al. 1994).

Another interesting question regarding the mechanism of transcription termination is how the stress enhances the termination. We demonstrated previously that stress conditions lead to the enhancement of termination at not only sRNA genes but also a gene encoding an mRNA (Morita et al. 2015). It is interesting that low NTP concentrations are reported to enhance transcription termination at Rho-independent terminators in vitro (Reynolds et al. 1992). In addition, it is reported that shortened transcripts were generated at a Rho-independent terminator in vitro when a low concentration of NTPs was used (McDowell et al. 1994). Therefore, one plausible effect of stress is to reduce the level of NTPs in cells.

The simplest type of Hfq binding module consists of a long poly(U) tail, terminator hairpin, and an internal U-rich sequence just before the hairpin (Ishikawa et al. 2012). The poly(U) tail binds to the proximal face of the Hfq hexamer,
whereas the internal U-rich sequence preferentially binds to the rim of the Hfq hexamer (Sauer and Weichenrieder 2011; Sauer et al. 2012; Schu et al. 2015). Thus, the roles of the poly(U) tail and internal U-rich sequence in Hfq binding are clear. On the other hand, it remains to be studied whether the terminator RNA hairpin plays no role in Hfq binding itself. Further studies are needed to know whether the hairpin structure of the terminator RNA hairpins without the poly(U) sequence were determined according to the Mfold program (Zuker 2003).

### MATERIALS AND METHODS

**Bacterial strains and plasmids**

The *E. coli* K12 strains and plasmids used in this study are listed in Table 1. IT1568 (W3110 *Δ*arcZ) was used as a parent strain. To construct TM894, TM895, and TM896, the alleles Δ*pnp*::*cat* and Δ*rnb*::*cat* of ECK3152, ECK1281 and ECK4175, respectively, were moved to TM542 (Kawamoto et al. 2005) by P1 transduction. ECK3152, ECK1281, and ECK4175 were donated from H Mori, NAIST (pers. comm.). The *rpoC*-Flag-*cat* allele was constructed according to the modified Datsenko–Wanner protocol using pSU313 harboring the *Flag*-cat sequence (Uzzau et al. 2001). To construct TM905 in which the chromosomal *rpoC* gene was replaced with the *rpoC*-Flag encoding carboxy terminally Flag-tagged β’, the *rpoC*-Flag-*cat* allele was moved to IT1568 by P1 transduction. TM908 was constructed by removing the *cat* gene flanked by two FRT sequences from TM905.

The DNA primers used are listed in Table 3. Plasmid pSgrS-S-LS1-rpILT was constructed as follows: pSgrS-S-rpILT was used to amplify the DNA fragment containing the sgrS-S-LS1 sequence and rpILT sequence with primers 1585 and 1544. The amplified DNA fragment was digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S, pSgrS-S-LS1, pSgrS-S-LS2, pSgrS-S-LS3, and pSgrS-S-LS4 were constructed as follows: pSgrS was used to amplify the DNA fragment containing the sgrS-S, sgrS-S-LS1, sgrS-S-LS2, sgrS-S-LS3, or sgrS-S-LS4 sequence with primers 1127 and 1839, 1840, 1923 or 1924, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (Otaka et al. 2011) was used to amplify the DNA fragment containing the sgrS-S-LS2, sgrS-S-LS4 sequence with primers 1127 and 1839, 1840, 1923 or 1924, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac*~-sgrS-S and *P*~*tac*~-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac*~-sgrS-S or *P*~*tac*~-sgrS-S-LS2 sequence with primers 1919 and 1839 or 1841, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac*~-sgrS-S and *P*~*tac*~-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac*~-sgrS-S or *P*~*tac*~-sgrS-S-LS2 sequence with primers 1919 and 1839 or 1841, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac*~-sgrS-S and *P*~*tac*~-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac*~-sgrS-S or *P*~*tac*~-sgrS-S-LS2 sequence with primers 1919 and 1839 or 1841, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac*~-sgrS-S and *P*~*tac*~-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac*~-sgrS-S or *P*~*tac*~-sgrS-S-LS2 sequence with primers 1919 and 1839 or 1841, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac**-sgrS-S and *P*~*tac**-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac**-sgrS-S or *P*~*tac**-sgrS-S-LS2 sequence with primers 1919 and 1839 or 1841, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX. Plasmids pSgrS-S (tac) and pSgrs-S-LS2 (tac) carrying the *P*~*tac**-sgrS-S and *P*~*tac**-sgrS-S-LS2, respectively, were constructed as follows: pSgrS or pSgrs-S-LS2 was used to amplify the DNA fragment containing the *P*~*tac**-sgrS-S or *P*~*tac**-sgrS-S-LS2 sequence with pri
and HindIII, and cloned into pAraX. Plasmids pRyhB-LS1 and pRyhB-LS2 were constructed as follows: pRyhB (Otaka et al. 2011) was used to amplify the DNA fragment containing the rhyB-LS1 or rhyB-LS2 sequence with primers 1144 and 1845 or 1844, respectively. The amplified DNA fragments were digested with XbaI and HindIII, and cloned into pAraX.

**Northern blotting**

Cells carrying the indicated plasmids were grown at 37°C to mid-log phase in LB medium supplemented with kanamycin (15 µg/mL) and indicated amounts of arabinose and αMG when needed. Total RNAs were isolated as previously described (Aiba et al. 1981). To detect SgrS-S and RyhB RNAs, RNA samples were resolved by 12% and 10% polyacrylamide gel electrophoresis, respectively, in the presence of 8 M urea and blotted onto a Hybond-N+ membrane (GE Healthcare). To detect tmRNA, ptsG mRNA, and sodB mRNA, RNA samples were resolved by 1.5% (tmRNA) and 1.2% (mRNAs) agarose gel electrophoresis in the presence of formaldehyde and blotted onto a Hybond-N+ membrane (GE Healthcare). The RNAs were visualized using digoxigenin (DIG) reagents and kits for nonradioactive nucleic acid labeling and a detection system (Roche Applied Science) according to the procedure specified by the manufacturer. The RNA probes SgrS-S and RyhB corresponding to the antisense of portion (+168 to +198) of sgrS and portion (+1 to +55) of ryhB, respectively, were prepared by the DIG RNA Labeling Kit (Roche Applied Science). The following DIG-labeled DNA probes were prepared by PCR using DIG-dUTP: a 363-bp fragment corresponding to the tmRNA (tmRNA probe); a 305-bp fragment corresponding to the 5′ region of ptsG (ptsG probe); a 210-bp fragment corresponding to the 5′ region of sodB (sodB probe). Dyna Marker, RNA Low II (BioDynamics Laboratory Inc.) was used as RNA size markers. The positions of RNA markers were shown on the left of the figures of Northern analysis.

**3′-RACE**

Total cellular RNAs prepared from cells harboring pSgrS-S or pSgrS-S-LS2 were resolved by electrophoresis on a 12% polyacrylamide gel in the presence of 8 M urea. After electrophoresis, the gel pieces corresponding to the full-length and shorter transcripts were cut out from the gel. Eluted samples were treated with phenol, precipitated, and washed with ethanol. The precipitants were dissolved in 10 µL of H2O containing RNase inhibitor (TOYOBO). cDNA was synthesized by the 3′ Full RACE Core Set (TAKARA) according to manufacturer’s instructions with RT primer 1786. The

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**TABLE 3. DNA primers used for construction of plasmids**

| Primer | Sequence | Plasmid |
|--------|----------|---------|
| 1585   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS1- rplL (F) |
| 1544   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS1- rplL (R) |
| 1127   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS2 (F) |
| 1839   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS2 (R) |
| 1840   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS3 (R) |
| 1841   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS4 (R) |
| 1144   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS1 (F) |
| 1839   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS1 (R) |
| 1840   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS2 (R) |
| 1841   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS3 (R) |
| 1844   | CCCAAGCTTAGAATGAAAAACGAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | pSgrS-S-LS4 (R) |
| 1786   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | pRyhB-LS2 (R) |
| 1787   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | RT primer |
| 1787   | GCGCTCTAGATAGTGTGACTGAGTATTGGTGTAAAATCACCCGCCAGCAGCCCCATT | RT PCR primer |
| 1901   | TGCTGAACGCAGGCTTGGCGGTCGACTAATACGAAAGGCTGCTGACTTATCGCAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | rpoC-Flag-cat P1 |
| 1902   | CAGAAATTCCTTTAAATATTTCTGACAGAATATCGCAGGCTCCTTGTGACCTGCTTGGTGTAAATCACCCGCCAGCAGCCCCATT | rpoC-Flag-cat P2 |

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**www.rnajournal.org 1429**
cDNA was amplified with primers 1127 and 1787 by using PCR SuperMix (Life Technologies). Cycling conditions were followed as: 98°C/30 sec; 20 cycles of 95°C/10 sec, 53°C/30 sec, 72°C/60 sec; 72°C/5 min. The amplified DNA fragments were purified by 6% native PAGE, followed by digestion with XbaI and EcoRI, and cloning into pTWV228 (TAKARA). Positive colonies were randomly picked up and plasmid DNAs were extracted using the QIAprep Spin Miniprep Kit (QIAGEN). Inserted cDNA sequences were analyzed with −47 sequence primer using CEQ8000 sequencer (SCIEX).

**Pull-down assay**

Cells were grown in 200 mL of LB medium at 37°C. At $A_{660} = 0.6$, 0.1% dMG was added and incubation was continued for 10 min, and then 0.02% arabinose was added and incubation was continued for 5 min. Cells were harvested, and washed by 15 mL STE buffer (100 mM NaCl, 10 mM Tris-HCl at pH 8.0, and 1 mM EDTA). The cells pellet was suspended in ice-cold 1 mL IP buffer 1 (20 mM Tris-HCl at pH 8.0, 0.1 M KCl, 5 mM MgCl₂, 10% glycerol and 0.1% Tween20). The cell suspension was crushed by µT-01 Beads Crusher (TITEC) with 0.500 mm of glass beads, followed by centrifugation at 10,000g for 10 min at 4°C. The supernatant (crude extract [CE]) was incubated with 50 µL of anti-Flag M2-agarose suspension (Sigma-Aldrich) in 10 mL of IP buffer 1 for 20 min at 4°C. The mixture was filtered by using a mini chromatography column (Bio-Rad). The agarose beads were washed twice by 10 mL of IP buffer 1. The proteins bound to the beads were eluted with 50 µL of IP buffer 1 containing 0.4 mg/mL Flag peptide (Sigma-Aldrich) and used as bound fraction (B). To analyze proteins, CE (10 µL) and B (10 µL) were treated with phenol, precipitated with ethanol. The pellets were washed with ethanol, and dissolved in RNA buffer. The samples were subjected to Northern blotting using the SgrS-S RNA probe.

**In vitro transcription assay**

Flag-tagged RNA polymerase (RNAP-Flag) in which the carboxy-terminus of the β′ subunit is tagged with the Flag sequence was purified as follows. TM908 carrying the rpoC-Flag allele was grown in 400 mL of LB medium at 37°C. At $A_{660} = 0.8$, cells were harvested and washed with 40 mL of STE, and suspended in ice-cold 4 mL of IP buffer 2 (20 mM Tris-HCl at pH 8.0, 0.2 M KCl, 5 mM MgCl₂, 10% glycerol, and 0.1% Tween20). The cell suspension was crushed by the µT-01 Beads Crusher (TITTEC) with 0.350–0.500 mm of glass beads, followed by centrifugation at 10,000g for 10 min at 4°C. The supernatant was incubated with 200 µL of anti-Flag M2-agarose suspension (Sigma-Aldrich) in 10 mL of IP buffer 2 for 30 min at 4°C. The proteins bound to the beads were eluted with 200 µL of IP buffer 2 containing 0.4 mg/mL Flag peptide (Sigma-Aldrich) and concentrated by Amicon Ultra (0.5 mL 30K Centrifugal Filters (Millipore). The purity and the concentration of RNAP-Flag were estimated by electrophoresis using Blot 4–12% Bis-Tris Plus gel and SimplyBlue SafeStain (Invitrogen) (Fig. 6C). An equal amount of glycerol was added to the concentrated RNAP-Flag solution and used for an in vitro transcription assay. Plasmids pSgrS-S (tac) and pSgrS-S-LS2 (tac) were purified from cells harboring the respective plasmid by using QIAprep Spin Miniprep Kit (QIAGEN); then they were used as DNA templates for in vitro transcription assay.

Plasmid DNA (1.6 µg) was incubated in 10 µL of 0.1× TE containing 40 mM NaCl for 2 min at 37°C, and then cooled down to 30°C. The DNA was mixed with 4 nM RNAP-Flag and 25 µM NTPs in 20 µL (final volume) of transcription buffer (50 mM Tris- HCl at pH 8.0, 50 mM NaCl, 3 mM MgCl₂, 0.1 mM EDTA, 0.1 mM DTT, 0.01% BSA) containing RNase inhibitor (TOYOBO). The reaction mixture was incubated for 30 min at 37°C. The reaction was terminated by adding phenol. Nucleic acids in the aqueous phase were precipitated with ethanol. The pellets were washed with ethanol, and dissolved in RNA buffer. The samples were subjected to Northern blotting using the SgrS-S RNA probe.

**ACKNOWLEDGMENTS**

We thank Hirotada Mori (Nara Institute of Science and Technology) for the E. coli strains ECK3152, ECK1281, and ECK4175, and Mitsuoki Kawano (Kawasaki Medical University) for the information about the 3′ RACE experiment. This work was supported by Grants-in-Aid from the Ministry of Education, Culture, Sports, Science and Technology of Japan (to H.A. and T. M.) and the Takeda Science Foundation (to T.M.).

Received January 17, 2017; accepted May 30, 2017.

**REFERENCES**

Aiba H, Adhya S, de Crombrugghe B. 1981. Evidence for two functional gal promoters in intact Escherichia coli cells. *J Biol Chem* **256**: 11905–11910.

Argaman L, Hershberg R, Vogel J, Bejerano G, Wagner EG, Margalit H, Altvia S. 2001. Novel small RNA-encoding genes in the intergenic regions of *Escherichia coli*. *Curr Biol* **11**: 941–950.

Cheng SW, Lynch EC, Leason KR, Court DL, Shapiro BA, Friedman DI. 1991. Functional importance of sequence in the stem-loop of a transcription terminator. *Science* **254**: 1205–1207.

Churchman LS, Weissman JS. 2011. Nascent transcript sequencing visualizes transcription at nucleotide resolution. *Nature* **469**: 368–373.

d’Aubenton Carafa Y, Brody E, Thermes C. 1990. Prediction of rho-independent *Escherichia coli* transcription terminators. A statistical analysis of their RNA stem-loop structures. *J Mol Biol* **216**: 835–858.

Dimastrogiovanni D, Fröhlich KS, Bandyra KJ, Bruce HA, Hohensee S, Vogel J, Luisi BF. 2014. Recognition of the small regulatory RNA product Hfq. *eLife* **3**: e05375.

Gottesman S, Storz G. 2010. Bacterial small RNA regulators: versatile roles and rapidly evolving variations. *Cold Spring Harb Perspect Biol* **1**: a003798.

Gusarov I, Nudler E. 1999. The mechanism of intrinsic transcription termination. *Mol Cell* **3**: 495–504.

Ishikawa H, Otaka H, Maki K, Morita T, Aiba H. 2012. The functional Hfq-binding module of bacterial sRNAs consists of a double or single hairpin preceded by a U-rich sequence and followed by a 3′ poly (U) tail. *RNA* **18**: 1062–1074.

Kawamoto H, Morita T, Shimizu A, Inada T, Aiba H. 2005. Implication of membrane localization of target mRNA in the action of a small
RNA: mechanism of post-transcriptional regulation of glucose transporter in Escherichia coli. Genes Dev 19: 328–338.
Kawano M, Reynolds AA, Miranda-Rios J, Storz G. 2005. Detection of 5'- and 3'-UTR-derived small RNAs and cis-encoded antisense RNAs in Escherichia coli. Nucleic Acids Res 33: 1040–1050.
Larson MH, Greenleaf WJ, Landick R, Block SM. 2008. Applied force reveals mechanistic and energetic details of transcription termination. Cell 132: 971–982.
Lynn SP, Kasper LM, Gardner JF. 1988. Contributions of RNA secondary structure and length of the thymidine tract to transcription termination at the thr operon attenuator. J Biol Chem 263: 472–479.
Massé E, Gottesman S. 2002. A small RNA regulates the expression of genes involved in iron metabolism in Escherichia coli. Genes Dev 16: 1305–1310.
Massé E, Escorcia FE, Gottesman S. 2003. Coupled degradation of a small regulatory RNA and its mRNA targets in Escherichia coli. Proc Natl Acad Sci USA 99: 4620–4625.
Morita T, Maki K, Aiba H. 2005. RNase E-based ribonucleoprotein complexes: mechanistic basis of mRNA destabilization mediated by bacterial noncoding RNAs. Genes Dev 19: 2176–2186.
Morita T, Mochizuki Y, Aiba H. 2006. Translational repression is sufficient for gene silencing by bacterial small non-coding RNAs in the absence of mRNA destruction. Proc Natl Acad Sci USA 103: 4858–4863.
Morita T, Ueda M, Kubo K, Aiba H. 2015. Insights into transcription termination of Hfq-binding sRNAs of Escherichia coli and characterization of readthrough products. RNA 21: 1490–1501.
Otaka H, Ishikawa H, Morita T, Aiba H. 2011. PolyU tail of rho-independent terminator of bacterial small RNAs is essential for Hfq action. Proc Natl Acad Sci 108: 13059–13064.
Panja S, Schu DJ, Woodson SA. 2013. Conserved arginines on the rim of Hfq catalyze base pair formation and exchange. Nucleic Acids Res 41: 7536–7546.
Papenfort K, Sun Y, Miyakoshi M, Vanderpool CK, Vogel J. 2013. Small RNA-mediated activation of sugar phosphatase mRNA regulates glucose homeostasis. Cell 153: 426–437.
Peters JM, Vangeloff AD, Landick R. 2011. Bacterial transcription terminators: the RNA 3'-end chronicles. J Mol Biol 412: 793–813.
Platt T. 1986. Transcription termination and the regulation of gene expression. Annu Rev Biochem 55: 339–372.
Ray-Soni A, Bellcourt MJ, Landick R. 2016. Mechanisms of bacterial transcription termination: all good things must end. Annu Rev Biophys 45: 319–347.
Reynolds R, Bermúdez-Cruz RM, Chamberlin MJ. 1992. Parameters affecting transcription termination by Escherichia coli RNA polymerase. I. Analysis of 13 rho-independent terminators. J Mol Biol 224: 31–51.
Rosenberg M, Court D. 1979. Regulatory sequences involved in the promotion and termination of RNA transcription. Annu Rev Genet 13: 319–333.
Santangelo TJ, Roberts JW. 2004. Forward translocation is the natural pathway of RNA release at an intrinsic terminator. Mol Cell 14: 117–126.
Sauer E, Weichenrieder O. 2011. Structural basis for RNA 3'-end recognition by Hfq. Proc Natl Acad Sci 108: 13065–13070.
Sauer E, Schmidt S, Weichenrieder O. 2012. Small RNA binding to the lateral surface of Hfq hexamers and structural rearrangements upon mRNA target recognition. Proc Natl Acad Sci U S A 109: 9396–9401.
Schu DJ, Zhang A, Gottesman S, Storz G. 2015. Alternative Hfq-sRNA interaction modes dictate alternative mRNA recognition. EMBO J 34: 2557–2573.
Uzzau S, Figueroa-Bossi N, Rubino S, Bossi L. 2001. Epitope tagging of chromosomal genes in Salmonella. Proc Natl Acad Sci 98: 15264–15269.
Vanderpool CK, Gottesman S. 2004. Involvement of a novel transcriptional activator and small RNA in post-transcriptional regulation of the glucose phosphoenolpyruvate phosphotransferase system. Mol Microbiol 54: 1076–1089.
Vanderpool CK, Gottesman S. 2007. The novel transcription factor SgrR coordinates the response to glucose-phosphate stress. J Bacteriol 189: 2238–2248.
Vogel J, Luisi BF. 2011. Hfq and its constellation of RNA. Nat Rev Microbiol 9: 578–589.
Wagner EG, Rompy B. 2015. Small RNAs in bacteria and archaea: who they are, what they do, and how they do it. Adv Genet 90: 133–208.
Waters LS, Storz G. 2009. Regulatory RNAs in bacteria. Cell 136: 615–628.
Yager TD, von Hippel PH. 1991. A thermodynamic analysis of RNA transcript elongation and termination in Escherichia coli. Biochemistry 30: 1097–1118.
Zuker M. 2003. Mfold web server for nucleic acid folding and hybridization prediction. Nucleic Acids Res 31: 3406–3415.