Regulation of hierarchical carbon substrate utilization, nitrogen fixation, and root colonization by the Hfq/Crc/CrcZY genes in *Pseudomonas stutzeri*

Highlights

A1501 preferentially catabolizes succinate followed by citrate and ultimately glucose

The Hfq/Crc/CrcZY regulatory system orchestrates this preference

Hfq optimizes nitrogen fixation and root colonization

Glucose is shunted into an unidentified route to produce Psl exopolysaccharide
Regulation of hierarchical carbon substrate utilization, nitrogen fixation, and root colonization by the Hfq/Crc/CrcZY genes in Pseudomonas stutzeri

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SUMMARY
Bacteria of the genus Pseudomonas consume preferred carbon substrates in nearly reverse order to that of enterobacteria, and this process is controlled by RNA-binding translational repressors and regulatory ncRNA antagonists. However, their roles in microbe-plant interactions and the underlying mechanisms remain uncertain. Here we show that root-associated diazotrophic Pseudomonas stutzeri A1501 preferentially catabolizes succinate, followed by the less favorable substrate citrate, and ultimately glucose. Furthermore, the Hfq/Crc/CrcZY regulatory system orchestrates this preference and contributes to optimal nitrogenase activity and efficient root colonization. Hfq has a central role in this regulatory network through different mechanisms of action, including repressing the translation of substrate-specific catabolic genes, activating the nitrogenase gene nifH posttranscriptionally, and exerting a positive effect on the transcription of an exopolysaccharide gene cluster. Our results illustrate an Hfq-mediated mechanism linking carbon metabolism to nitrogen fixation and root colonization, which may confer rhizobacteria competitive advantages in rhizosphere environments.

INTRODUCTION
Bacteria can utilize numerous carbon substrates, and thus they can grow in a wide range of natural environments. Carbon substrates in the environment variable in type and abundance and can be utilized by bacteria either hierarchically or simultaneously.1 Hierarchical utilization refers to catabolite repression, in which preferred carbon sources prevent the utilization of other substrates, and a variety of mechanisms are used, depending on the bacterial genus2,3; the classic example of which is glucose-lactose diauxic growth of Escherichia coli.4 It is now well accepted that carbon catabolite repression controls the sequential catabolism of preferred carbon substrates, thereby enhancing the ability of bacteria to compete in their natural habitats.5 Bacteria of the genus Pseudomonas are considered generalists because they are isolated from a wide range of niches and have the remarkable property of metabolizing a wide range of compounds as a source of carbon and energy.6,7 Pseudomonas are metabolically versatile and behave as intelligent eaters that preferentially catabolize organic acids, especially tricarboxylic acid (TCA) cycle intermediates, rather than carbohydrates.5,6,7 Clearly, pseudomonads use a “reverse” carbon catabolite repression compared to enterobacteria, which may reflect better adaptation to specific environments due to the differential evolution that occurs in different niches.

Indeed, the molecular mechanisms responsible for carbon catabolite repression differ markedly among microorganisms. In particular, catabolite repression is mediated at the transcription level in E. coli (by the Crp transcriptional regulator) and Bacillus subtilis (by the CcpA transcriptional regulator) while in Pseudomonas, carbon catabolite repression operates at the translational level.3,5 The current model that was derived from studies of different Pseudomonas species showed that in this posttranscriptional repression system, the major players are the RNA chaperone Hfq, the catabolite repression control protein Crc and several regulatory ncRNAs.10–12 In the presence of preferred carbon sources, the translation of target genes involved in the utilization of non-preferred carbon sources is inhibited by Hfq by forming a repressive complex with Crc on specific target mRNAs.13,14 Emerging evidence further revealed that Hfq recognizes...
and binds to target mRNAs containing a common catabolite activity (CA) motif that is located close to the ribosome binding site,14 in contrast, Crc has no intrinsic RNA-binding activity15 but increases the stability of the Hfq/Crc/mRNA tripartite complexes.15 Sequences that resemble the CA motif are present in mRNAs of the oprB1, gtsABC, benR, benA, alkS, and alkB genes in Pseudomonas putida,16–18 the amiE, amIR, eisA, bkdR, and arfP2 genes in Pseudomonas aeruginosa,19–22 the xutR and xutA genes in Pseudomonas fluorescens,23,24 and oprB in Azotobacter vinelandii,25 which are involved in the utilization of non-preferred carbon sources. Several regulatory ncRNAs, such as CrcZ in P. aeruginosa,26 CrcZ/CrcY in P. putida or P. fluorescens,23,25 and CrcZ/CrcX in P. syringae,27 have been identified as antagonists of catabolite repression. Synthesis of these ncRNA is under the control of the two-component system CbrAB and the alternative sigma factor RpoN,20,22,27,28 with one exception for CrcY, in which the activation is thought to involve a different transcriptional regulator in P. putida.27

Although carbon catabolite repression in bacteria is one of the best-studied metabolic strategies, we are far from understanding the complexity of this regulatory system. Mechanisms that operate in carbon catabolite repression seem to control not only the expression of the genes involved in specific catabolic pathways but also the expression of the genes involved in microbe-plant interactions.5,29 For example, the control of catabolite repression was also found to be necessary for the growth and survival of P. syringae DC3000 during infection30 and the mineral phosphate-solubilizing/glycine betaine-producing ability of Acinetobacter sp. SK231 and P. aeruginosa.32,33 Similarly, the involvement of catabolite repression control in functions related to bacteria-plant interactions has been reported in Klebsiella pneumoniae34 and Rhizobium sp.,35 suggesting that this mechanism is widespread among rhizobacteria. In addition, the rhizosphere is a nitrogen-limiting environment that favors the proliferation of diazotrophs; the energy necessary to sustain root-associated nitrogen fixation is derived from the oxidation of carbon-rich exudates available in the rhizosphere.36,37 Therefore, another interesting emerging aspect is the connection between carbon catabolite repression and nitrogen fixation in root-associated diazotrophs.

Bacterial species belonging to the Pseudomonas genus are very efficient colonizers of the plant rhizosphere, resulting from the co-occurrence of multiple functions, such as chemotactic motility, surface attachment, biofilm formation, stress response, and nitrogen fixation.38–42 One of the most representative strains is Pseudomonas stutzeri A1501, a model diazotroph that was originally isolated from the rice rhizosphere.6 Due to its nitrogen-fixing ability, A1501 outcompetes other soil bacteria and proliferates in the carbon-rich but nitrogen-limiting environment of the rhizosphere to colonize root surface or inner cortex, and benefit host plant growth.43,44 Several genes (e.g., cbrAB, crc, and crcZ) involved in catabolite repression have been previously identified in P. stutzeri A1501.45 A subsequent transcriptional analysis of the P. stutzeri A1501 genome resulted in the identification of 53 ncRNAs, including crcZ and crcY.46 In addition, the chaperone gene Hfq was found to play a role in the control of nitrogen fixation, since an hfq mutant strain displayed reduced nitrogenase activity,46,47 although the actual mechanism of this regulation remains to be documented.

In this study, we focused on performing a detailed analysis of the genetic determinants involved in catabolite repression control, as well as some of the target genes involved in this mechanism, to better understand the regulatory mechanisms underlying hierarchical carbon substrate utilization in root-associated diazotrophic P. stutzeri A1501. We found that P. stutzeri A1501 preferentially catabolized succinate, followed by the relatively less favorable substrate citrate, and ultimately glucose. Genetic studies and microscale thermophoresis (MST) experiments showed that Hfq represses the translation of catabolic targets for non-preferred substrates by binding directly to mRNAs while Crc interacts with Hfq, enhancing the repression exerted by Hfq. Subsequent physiological and droplet digital PCR (ddPCR) analysis indicated that non-preferred carbon sources such as glucose and benzoate substantially increased the transcription of CrcZ, acting as a key antagonist of Hfq, by binding the protein and relieving the repression of the non-preferred catabolic pathways. Finally, our finding provides empirical evidence for the molecular mechanism by which Hfq contributes to optimal nitrogenase activity and efficient root colonization.

RESULTS
Carbon substrate utilization profile and diauxic growth
The hierarchy of carbon substrate utilization was determined by measuring the respiratory activity and growth ability of strain A1501 grown on single or double carbon sources. We first measured the respiratory activity on different carbon sources by using Biolog GENIII MicroPlates. Forty-three compounds that
supported metabolism were identified and grouped by their intensities of respiratory activity into two distinct groups (Table S1). Interestingly, many carbon sources from the MicroPlates are also present in plant root exudates, such as lactate, glucose, succinate, and citrate, with strong respiratory activities from 1694 to 1175 U (Table S1).

These data agree with the fact that organic acids are known to be preferentially used over sugars and aromatic compounds in Pseudomonas species. P. stutzeri A1501 displays better growth with lactate, succinate, or citrate than with glucose or benzoate, as shown in Figure 1A. The results from the growth assay were consistent with those of the Biolog assay (Table S1), except for glucose, supporting the high respiratory activity but poor growth of strain A1501. In addition, the carbon source preference is almost consistent with the capacity to support nitrogenase activity (Figure 1B), except for benzoate that does not support nitrogen fixation. In addition, strain A1501 preferentially utilized an organic acid until it was exhausted and then shifted to catabolism of glucose, exhibiting typical diauxic growth (Figures 1C–1E).

A1501 was previously shown to grow on 4 mM benzoate but rarely grew on 8 mM benzoate due to its toxic effect. Here we observed that in the presence of 4 mM benzoate plus succinate or lactate, A1501 was able to grow well but did not exhibit diauxic growth (Figures S1A and S1B). Surprisingly, A1501 grew very poorly on a citrate/benzoate mixture, and even more slowly on benzoate, implying that benzoate exerts a strong inhibitory effect on citrate utilization (Figure S1C). Conversely, A1501 co-utilized glucose and benzoate, exhibiting a much higher growth performance (the maximal OD600 of approximately 1.0) than on each individual source (the maximal OD600 of 0.30 and 0.45, respectively) (Figure 1F), as previously described for Pseudomonas sp. CSV86 and E. coli. Together, A1501 preferentially catabolized the top tier substrates succinate and lactate, followed by citrate at a relatively less favorable level, and ultimately the non-preferred substrates glucose and benzoate; however, the sophisticated mechanisms underlying this phenomenon remain to be investigated in depth.

Mutation of the hfq/crc/crcY genes and prediction of putative hfq-binding mRNAs
An analysis of the location of the putative regulatory genes controlling carbon catabolite repression in the P. stutzeri A1501 genome identified seven genes that are known to be involved in this process in other Pseudomonas species and revealed a cluster carrying the cbrAB genes of the two-component system linked to crcZ, while hfq, rpoN, crc, and crcY are not linked and belong to different clusters (Figure 2A and Table S2). Single and double mutants were constructed to obtain insights into their biological functions (Figure 2B). Growth with the preferred carbon source lactate or succinate was not impaired in any of the mutant strains, but a similar result was not observed for citrate. Although citrate is generally considered the preferred carbon source of Pseudomonas, the ΔcrcZΔcrcY double mutation strongly impaired growth on citrate as the sole carbon source (Figure 2B), suggesting that citrate utilization might be subject to carbon catabolite repression in strain A1501. Furthermore, we found that the ΔcbrA or ΔcbrB single mutants and the ΔcrcZΔcrcY double mutant did not grow on two non-preferred carbon sources, glucose or benzoate (Figure 2B), suggesting that these genes play key roles in carbon catabolite repression. The deletion of either crcZ or crcY had no effect on catabolite repression, suggesting that the two genes may be functionally complementary, as described in P. putida. The Δcrc mutant exhibited wild-type diauxic growth, whereas the Δhfq mutant exhibited the same phenotype defect in diauxie as the ΔhfqΔcrc double mutant, suggesting that Hfq has a stronger function than that of Crc in catabolite repression on glucose utilization.

Recent studies revealed that Hfq recognizes a common catabolite activity (CA) motif that contains an AANAANAA motif close to the AUG translation start codon of target mRNAs. An analysis of the phenotype of the hfq mutant helped identify the possible target genes that are involved in the utilization of non-preferred carbon sources by P. stutzeri A1501. Glucose is metabolized by the Entner-Doudoroff pathway in Pseudomonas species. The glnR gene encodes a transcriptional regulator that interacts with different promoters to regulate the expression of the oprB, gki, and edd genes located in different clusters in the A1501 genome (Figure 2A and Table S2), all of which are required for glucose transport and utilization. As shown in Figure 2C, a CA motif was identified in the mRNAs of these genes required for glucose metabolism. The characterization of the genes involved in benzoate utilization (Figure 2A) was previously reported in P. stutzeri A1501. An analysis of the mRNA sequences of these genes identified a CA motif in benR, benF, and benK (Figure 2C). Consistent with the hypothesis mentioned above that citrate utilization may also be controlled by catabolite repression, a CA motif was found in the mRNA of the citN gene.
encoding a citrate transporter (Figure 2C). These results suggest that Hfq has a major role in regulatory systems underlying hierarchical carbon substrate utilization, similar to those in other *Pseudomonas* species. It was previously reported that an *hfq* mutant strain displayed reduced nitrogenase activity.46 Most interestingly, a CA motif was also identified in the coding region of the *nifH* mRNA encoding a nitrogenase Fe
In P. aeruginosa, unlike Hfq, Crc was not found to bind CA motifs but to have an auxiliary function in Hfq-mediated repression. Thus, we used microscale thermophoresis (MST) to analyze the ability of both Hfq and Crc proteins to recognize predicted CA motifs in the 5’ untranslated region of the target genes, including the gltR, oprB, edd and zwf genes responsible for glucose utilization, the benR gene encoding a transcriptional activator for the benzoate degradation pathway, and the citN gene encoding a citrate transporter. As revealed in Figures 3E, 3G, 3I, and 3M (green curve), Hfq directly bound to each of the wild-type mRNAs. The same experiments were reproduced using mutated mRNAs devoid of the CA motif to support these data, and the ability to bind Hfq was lost (Figures 3E, 3G, 3I, and 3M, red curve). In the presence of Crc, the binding of Hfq to target mRNAs was significantly enhanced (Figures 3F, 3H, 3J, and 3M, green curve). These findings suggest that the presence of Hfq may be necessary and sufficient for carbon catabolite repression, as reported first in P. aeruginosa.20

CrcZ antagonizes hfq-mediated translational repression of non-preferred catabolic pathways

CrcZ and CrcY levels were reported to vary widely based on the carbon source used in P. aeruginosa and P. putida.53,54 In the present study, the level of crcZ is low under conditions causing strong catabolic
repression, but increases substantially in the presence of glucose or benzoate as the sole carbon source (Figure 4A). Notably, the level of \textit{crcZ} is also moderately increased during growth with citrate as a sole carbon source (Figure 4A), providing a possible explanation for the growth defect of the \textit{DcrcZDcrcY} double mutant on citrate (Figure 2B). During diauxic growth in the presence of succinate plus glucose, CrcZ levels increase sharply during the transition from the first exponential phase to the second (Figure 4B), suggesting that this ncRNA may be essential for abrogating Hfq-mediated translational repression. Unlike \textit{crcZ}, the level of \textit{crcY} is very low and remains virtually unchanged, regardless of the carbon source used under all growth conditions.

Figure 3. Hfq and Crc act as translational repressors of substrate-specific catabolic genes

(A) Absolute numbers of copies of the \textit{hfq} and \textit{crc} mRNAs during diauxic growth of WT A1501 on 4 mM succinate plus 20 mM glucose. The growth curve (circle) is shown together with the measured concentration of glucose (triangle) remaining in the medium. (B–D) Growth of the \textit{Dcrc} (B), \textit{Dhfq} (C), \textit{DhfqDcrc} (D) mutant strains (red) and their corresponding complemented strains (blue) that were grown on 4 mM succinate plus 20 mM glucose. (E–N) Determination of the binding affinity of labeled Hfq for the target mRNAs containing the WT or mutated CA motifs in the absence and presence of 400 nM Crc. Ligand-dependent changes in MST are plotted as normalized fluorescence (\(F_{\text{norm}}\)) values vs. ligand concentration in a dose-response curve. \(F_{\text{norm}}\) values are plotted as parts per thousand [%] for binding affinity analysis. N, ssRNA oligonucleotide; wt, wild type; mut, mutation. Data shown in (A–N) are representative of two independent experiments and are presented as means ± SEM (n = 6). See also Figures S2 and S3.

repression, but increases substantially in the presence of glucose or benzoate as the sole carbon source (Figure 4A). Notably, the level of \textit{crcZ} is also moderately increased during growth with citrate as a sole carbon source (Figure 4A), providing a possible explanation for the growth defect of the \textit{DcrcZDcrcY} double mutant on citrate (Figure 2B). During diauxic growth in the presence of succinate plus glucose, CrcZ levels increase sharply during the transition from the first exponential phase to the second (Figure 4B), suggesting that this ncRNA may be essential for abrogating Hfq-mediated translational repression. Unlike \textit{crcZ}, the level of \textit{crcY} is very low and remains virtually unchanged, regardless of the carbon source used under all growth conditions.
Figure 4. Functional analysis of the regulatory ncRNAs CrcZ and CrcY

(A) Absolute levels of the crcZ and crcY transcripts in strain A1501 grown on succinate and other carbon substrates. Bacteria were grown in minimal medium K containing 6.0 mM NH4\(^+\) and supplemented with 20 mM lactate (Lac), succinate (Suc), citrate (Cit), and glucose (Glu) and 4 mM benzoate (Ben). A 4 h incubation time was chosen for droplet digital PCR (ddPCR) assays. The absolute transcript level is reported as the number of copies per ng of total RNA. Means ± SEM (n = 6) are shown. Asterisks indicate statistical significance determined using two-way ANOVA with Tukey’s post hoc test: ****p < 0.0001, *p < 0.05, and ns: non-significant.

(B) Changes in absolute levels of the crcZ and crcY transcripts during diauxic growth. Bacterial cells were incubated in minimal medium K containing 6.0 mM NH4\(^+\) supplemented with a mixture of 4 mM succinate and 20 mM glucose, and samples were collected at regular intervals (0, 2, 4, 6, 8, 10, 12, and 14 h) for ddPCR assays. Data are representative of two independent experiments and are presented as the means ± SEM (n = 4).
conditions tested (Figures 4A and 4B), implying that CrcY does not appear to play a role in adjusting the strength of catabolite repression. However, the deletion of either crcZ or crcY led to a compensatory increase in the levels of the remaining ncRNA, as also observed in P. putida.57 For example, the levels of crcY in the strain lacking crcZ increased more than 3.0-fold compared to the wild-type strain (Figure 4C). This result may explain why the single deletion of crcZ or crcY had no effect on catabolite repression, but the double mutation resulted in constitutive catabolite repression that compromised the growth of citrate or glucose (Figure 2B). In particular, the level of CrcY was upregulated by about 13.4- and 4.5-fold at high temperature (45°C) or in the presence of saline stress, in contrast to CrcZ, whose expression was upregulated by approximately 2.2-fold under nitrogen fixation conditions (Figure S4), suggesting a CrcZY-mediated regulatory link between environmental stresses and carbon/nitrogen metabolism.

In addition, we found that the single deletion of the cbrA, cbrB, or rpoN genes caused a significant decrease in the expression levels of crcZ and crcY (Figure 4C), consistent with observations in other Pseudomonas strains.26,55 The position of the transcription start site for the promoters of both crcZ and crcY was determined by 5′RACE (Figures 4D and 4E). Further inspection of the DNA regions located upstream from the promoters of crcZ and crcY revealed the presence of conserved sequences (at position −24/−12) typical of a RpoN-dependent promoter and a putative CbrB binding site (Figures 4D and 4E), suggesting that both RpoN and CbrB controlled the expression of crcZ and crcY. This hypothesis was confirmed by DNase I footprinting assays (Figures 4F–4I). Both ncRNAs are predicted to contain five unpaired A-rich regions with putative CA motifs to which Hfq potentially binds (Figure 4D and 4E). A set of full-length CrcZ and CrcY ssRNA oligonucleotides containing the WT sequences (N-CrcZ-wt and N-CrcY-wt) and mutated sequences (the N-CrcZ-mut2 and N-CrcY-mut2 containing site-specific mutations in regions II and V, and the N-CrcZ-mut5 and N-CrcY-mut5 containing site-specific mutations in all five regions) were synthesized for the MST analysis to obtain experimental evidence for the predicted interactions between Hfq and CrcZ and CrcY individually (Figures 4D and 4E). As anticipated, both CrcZ and CrcY directly bind to Hfq, exhibiting Kd values of 23.7 ± 2.68 nM and 22.21 ± 4.68 nM, respectively (Figures S3A and S3D). The binding was significantly weakened or was even completely lost if the CA motif was mutated (Figures S3B, S3C, S3E, and S3F). Taken together, these experiments showed that the regulatory ncRNA CrcZ acts as a key decoy to antagonize Hfq-mediated translational repression of the catabolic genes necessary for the consumption of non-preferred carbon sources.

**Hfq optimizes root-associated nitrogenase activity via positive regulation of nifA and nifH expression at the transcriptional and translational levels**

Under laboratory conditions, nitrogenase activity associated with host seedlings or roots, which is usually low due to the limited carbon supply from root exudates, has been reported to increase after the addition of carbon sources.56,57 This result was also observed in rice roots inoculated with A1501 where no root-associated nitrogenase activity was detected in the absence of an additional carbon source; therefore, this condition was chosen as a negative control (Figure 5A). Interestingly, the root-associated nitrogenase activity of A1501 incubated with succinate was increased up to 12,000 U, which was approximately 3-fold higher than the nitrogenase activity of A1501 in the free-living state, and 6-fold higher than the root-associated nitrogenase activity of A1501 grown on glucose (Figure 5A), indicating that nitrogenase activity was significantly increased in the presence of roots, or more precisely certain root exudates. Next, acetylene reduction assays were performed with mutant strains using the same conditions as with the wild type. Inoculation with ΔcrcZ or ΔcrcY strains had little effect, while the inoculation of strains carrying Δcrc, Δhfq or...
DccZY mutations led to a significant decrease in root-associated nitrogenase activities, especially in the presence of mixed carbon sources (Figure 5A).

Knockout of hfq has also been reported to decrease the abundance of the nifH mRNA encoding a subunit of the nitrogenase, as well as the nifA mRNA, encoding a transcriptional activator of all nif operons in Azorhizobium caulinodans58 and Sinorhizobium meliloti.59 This is why we measured the nifA and nifH mRNA levels of wild-type and hfq mutant strains associated with roots using droplet digital PCR (ddPCR).

Figure 5. Hfq exerts a positive regulatory effect on root-associated nitrogenase activity at the transcriptional and translational levels

(A) Nitrogenase activity assays using wild-type and mutant strains associated with or without rice roots in the presence of 4 mM succinate (Suc) and 20 mM glucose (Glu) alone or in combination. Control 1: Rice roots without inoculation; Control 2: rice roots inoculated with A1501 but without an additional carbon source. (B and C) The absolute copy numbers of the nifA mRNA (B) and nifH mRNA (C) under the same conditions as in (A), which were determined using ddPCR.

(D and E) Determination of the binding affinity of labeled Hfq to nifH mRNA containing the WT (D) or mutated CA motif (E) using microscale thermophoresis. The CA motifs are shown in parentheses, and point mutations introduced into a synthesized ssRNA oligonucleotide are shown in red. N, ssRNA oligonucleotide; wt, wild type; mut, mismatch mutation.

(F) Analysis of the nifH mRNA half-life in the WT and hfq mutant strains. Rifampicin (400 μg/mL) was added at time 0. At the indicated times, an equal volume of frozen media was added to bring the temperature immediately to 4°C. RNA was extracted, followed by qRT-PCR.

Data shown in (A–C) are representative of two independent experiments and are presented as means ± SEM (n = 4). Asterisks indicate statistical significance determined using two-way ANOVA with Tukey’s post hoc test: ****p ≤ 0.0001, ***p ≤ 0.001, **p ≤ 0.01, p ≤ 0.05, and ns: non-significant. Data shown in (D–F) are presented as the means ± SEM (n = 6).
Consistent with the results shown in Figure 5A, an order of magnitude increases in the absolute copy number of both \(nifH\) and \(nifA\) mRNAs was observed when bacteria were grown on succinate or a mixture of glucose with rice roots compared with the level obtained without roots (Figures 5B and 5C, red asterisks indicate statistically significant differences). Mutation of \(hfq\) suppressed this stimulatory effect; for example, the absolute copy number of both \(nifH\) and \(nifA\) mRNAs in the \(hfq\) mutant decreased by more than 3.0- or 30.0-fold compared with the wild-type strain (Figures 5B and 5C), suggesting that Hfq exerts a positive regulatory effect on the expression of both genes in bacteria associated with roots. In addition, consistent with previous observations showing that the use of glucose as a sole carbon source poorly supports nitrogen fixation (Figure 1B), this non-preferred carbon source has no effect on the expression of both \(nifH\) and \(nifA\) mRNAs (Figures 5B and 5C). Moreover, \(nifH\) is the only gene among all the \(nif\) genes to contain the CA motif in its mRNA, suggesting that it is the target of Hfq (Figure 2C). As expected from MST experiments, Hfq directly binds to the CA motif located in the \(nifH\) coding region, while mutating this motif led to a complete lack of binding to Hfq (Figures 5D and 5E). We subsequently measured the half-life of the \(nifH\) transcript, which was 10.5 min for the WT and 7.0 min for the \(\Delta hfq\) mutant (Figure 5F), indicating that Hfq exerted a positive effect on the stability of the \(nifH\) mRNA.

**Involvement of \(hfq/crc/crcZY\) genes in root colonization via the transcriptional regulation of exopolysaccharide production**

A1501 exhibited poor growth on glucose but high respiratory activity (Figure 1F and Table S1); thus, we hypothesized that glucose is rapidly utilized not as a growth substrate but as a precursor to produce exopolysaccharides, which have been shown to be important for colony morphology and root colonization.\(^{60–62}\) Colony morphology was examined after seven days of growth with succinate and glucose alone or in combination. A1501 exhibited an opaque, wrinkled colony morphology after being grown on dual carbon sources, and this colony was much larger than the colony formed on each substrate alone (Figure 6A), which correlated with the amount of exopolysaccharides produced (Figure 6B). Wild-type phenotypes were totally abolished by the inactivation of the \(hfq/crc/crcZY\) genes (Figure 6B), suggesting their involvement in colony morphology and exopolysaccharide production. Consistent with the function of \(pslA\), the first gene in the \(psl\) operon that is involved in the synthesis of Psl exopolysaccharide in *P. aeruginosa*\(^{63}\) and *P. stutzeri*,\(^{64}\) the \(pslA\) mutant grown on mixed carbon sources not only produced fewer exopolysaccharides but formed colonies much smaller than those of the WT strain (Figures 6A and 6B). Unexpectedly, the \(\Delta glk\) mutant formed colonies very similar to the WT strain (Figure 6A) while producing more exopolysaccharides than the other mutants, suggesting that sugar nucleotide precursor synthesis in A1501 used a pathway independent of the glk gene. In addition, the production of exopolysaccharides is important for biofilm formation and root surface colonization.\(^{60,64}\) Here, \(pslA\) expression was significantly increased by approximately 33-fold after exposure to glucose in the presence of roots (Figure 6C). Sugar nucleotide precursors for exopolysaccharide production are generally synthesized from glucose-6-P dependent on glucokinase, which is encoded by the \(glk\) gene in *Pseudomonas*.\(^{65}\) When grown on a mixture of 3 mM glucose plus succinate, increasing the concentrations of glucose from 0.4 mM to 3.0 mM led to a continued increase in both \(pslA\) expression and exopolysaccharide production (Figures 6D and 6E). These results suggest that exopolysaccharide production depends on an adequate supply of glucose but is not inhibited by succinate.

Furthermore, we tested the effect of different concentrations of glucose on root surface colonization by A1501. As shown in Figure 6F, the level of colonization in the presence of 0.2 mM glucose was 2.7 times higher than that in the absence of glucose, indicating that glucose promotes root colonization. However, higher concentrations of glucose resulted in partial or total loss of the stimulatory effect (Figure 6F). We performed co-culture and co-inoculation assays using A1501 and its mutant derivatives to test the roles of the \(hfq/crc/crcZY\) genes. In both types of experiments, the individual mutants were recovered at significantly lower levels than A1501, suggesting a competitive disadvantage (Figures 6G and 6H). The \(\Delta hfq\) mutant was reduced to a minor fraction of the total bacteria (less than 0.1%) isolated from co-culture (Figure 6G) or from rice roots (Figure 6H), suggesting that Hfq is crucial for root surface colonization. Quantitative RT-PCR experiments were performed to compare the expression of the \(psl\) genes during root surface colonization. As shown in Figure 6I, the deletion of \(hfq\) resulted in a significant decrease in the expression of all \(psl\) genes, suggesting indirect Hfq-mediated regulation since the mRNAs of these genes do not contain the putative CA motif. These data indicate a crucial role for Hfq in root surface colonization through the positive regulation of \(psl\)-like genes that are required for Psl production.
Figure 6. The contribution of hfg/crc/crcZY genes to improving root colonization

(A) The colony morphology of WT and mutant strains was evaluated on minimal medium K that was supplemented with different carbon sources during the 7-day incubation period. Images were taken with the same magnification.

(B) Effect of gene mutations on exopolysaccharide production. Suc, succinate at 50 mM; Glu, glucose at 3 mM. Asterisks indicate statistical significance determined using two-way ANOVA with Tukey’s post hoc test: ****p ≤ 0.0001, ***p ≤ 0.001, **p ≤ 0.01, *p ≤ 0.05, and ns: non-significant.

(C) Relative expression levels of pslA in A1501 grown on various carbon substrates in the presence or absence of rice roots, as measured using qRT-PCR.

(D) Effect of different concentrations of glucose on the transcription of pslA in A1501 grown on minimal K medium supplemented with 6 mM NH4⁺ and 50 mM succinate. Asterisks indicate statistical significance determined using one-way ANOVA with Tukey’s post hoc test: ****p ≤ 0.0001.

(E) Effect of different concentrations of glucose on exopolysaccharide production by A1501 grown under the same conditions as in (D). Asterisks indicate statistical significance determined using one-way ANOVA with Tukey’s post hoc test: **p ≤ 0.01 and ns: non-significant.

(F) The effect of different glucose concentrations on root colonization by A1501 was determined by counting colony-forming units (CFUs) per g of root. Asterisks indicate statistical significance determined using one-way ANOVA with Tukey’s post hoc test: **p ≤ 0.01 and *p ≤ 0.05.

(G) Strain competition experiments. The fitness deficit of the test mutants was shown by co-culturing each mutant with wild-type A1501 mixed in a 1:1 ratio, inoculated at a starting OD₆₀₀ = 0.1 and grown for 24 h in LB. At 24 h, cultures were serially diluted and plated on LB plates containing the relevant antibiotics to enable counting of each strain. The wild-type strain was used as the competitor in all experiments. Asterisks indicate statistical significance determined using one-way ANOVA with Tukey’s post hoc test: ****p ≤ 0.0001, *p ≤ 0.05, and ns: non-significant.

(H) Competitive root colonization experiments. Rice seedlings were co-inoculated with the test mutant and wild-type A1501. After 24 h of incubation, root samples were collected, and the number of CFUs per g root was determined. The red bars represent the percentage of colonies recovered from the tested
Indeed, this posttranscriptional regulation has physiological advantages, which may allow of Hfq and CrcZ as a key translational repressor and antagonist pair in carbon catabolite repression. The rhizosphere is a hotspot for mutualistic interactions between microorganisms and host plants.67–69 In particular, plant roots secrete large amounts of carbon-rich compounds into the rhizosphere, including organic acids (succinic, citric, and lactic acids, among others), carbohydrates (glucose, fructose, and so forth), and phenolics (benzoate, benzoxazinoid, and so forth), which can be utilized as carbon and energy sources by root-colonizing pseudomonads.37,40 The metabolism of root exudates is controlled in a hierarchical manner by a complex regulatory system, including multiple RNA-binding proteins and ncRNAs at the posttranscriptional level.77 However, questions remain as why root-colonizing pseudomonads would adopt a complex strategy, as well as its physiological and adaptive advantages. In this study, ddPCR experiments quantitatively indicated that Hfq and CrcZ levels vary according to the nature of the carbon sources being used, a phenomenon that was also observed for A1501, which preferentially utilized citrate over glucose. The observation that citrate does not support the growth of the ΔcrcZΔcrcY double mutant remain puzzling; thus, citrate exhibits the characteristics of a typical non-preferred carbon source (Figure 2B). This result might be explained by the fact that Hfq binds to the A-rich motif of the citN mRNA encoding a citrate transporter (Figure 3K) while citrate moderately induces the expression of CrcZ (Figure 4A). The subject of most interest is benzoate, a derivative of lignin compounds and an intermediate in the catabolism of other aromatic compounds, which substantially inhibits growth and nitrogenase activity but increases the growth rate of glucose (Figure 1). The presumable explanation for this finding is that benzoate induced more CrcZ expression than glucose (Figure 4A), thus effectively relieving the inhibitory effect of Hfq on glucose utilization. Another explanation for our observations is that benzoate exerts a toxic effect on growth, and thus A1501 must evolve specifically adaptive regulatory mechanisms in a rhizosphere environment rich in aromatic carbon sources, as reported for the hegemonic preference of Cupriavidus pinatubonensis for benzoate.66 Therefore, this study provides a framework for understanding the molecular mechanism of hierarchical carbon substrate utilization in Pseudomonas. A complex model controlling this hierarchy is summarized in Figures 7 and S6.

The rhizosphere is a hotspot for mutualistic interactions between microorganisms and host plants.67–69 In particular, plant roots secrete large amounts of carbon-rich compounds into the rhizosphere, including organic acids (succinic, citric, and lactic acids, among others), carbohydrates (glucose, fructose, and so forth), and phenolics (benzoate, benzoazinoid, and so forth), which can be utilized as carbon and energy sources by root-colonizing pseudomonads.57,60 The metabolism of root exudates is controlled in a hierarchical manner by a complex regulatory system, including multiple RNA-binding proteins and ncRNAs at the posttranscriptional level.77 However, questions remain as why root-colonizing pseudomonads would adopt a complex strategy, as well as its physiological and adaptive advantages. In this study, ddPCR experiments quantitatively indicated that Hfq and CrcZ levels vary according to the nature of the carbon sources being used, a phenomenon that was also observed for P. aeruginosa,99 P. putida,77 and A. vinelandii.70 In particular, the abundance of Hfq increased rapidly under conditions that provoked strong repression by succinate (Figure 3A), whereas the synthesis of the ncRNA CrcZ was remarkably increased under non-repressing conditions (Figure 4A). These opposite expression patterns of Hfq and CrcZ, together with the fact that the Crc and CrcY levels are very low, regardless of the carbon source used, underscore the roles of Hfq and CrcZ as a key translational repressor and antagonist pair in carbon catabolite repression. Indeed, this posttranscriptional regulation has physiological advantages, which may allow Pseudomonas to respond to various signals in an extremely rapid and sensitive manner and ultimately improve their fitness and resource economy,71 as described in P. fluorescens23 and P. aeruginosa.26

In addition to specific genes subject to catabolite repression control, Hfq and CrcZ have multiple targets, which may confer regulatory plasticity and represent a new layer of hierarchical networks in response to nutritional and environmental cues. As shown in Figure 5, Hfq contributes to optimal root-associated nitrogenase activity via the indirect transcriptional regulation of nifA and direct posttranscriptional regulation of...
The expression of both crcZ and crcY is a RpoN-dependent process that is activated by the CbrA/CbrB two component sensor-regulator system (Figures 4F–4I), suggesting a carbon-nitrogen coupling regulation mode. Similar to other Pseudomonas strains, A1501 possesses two ncRNAs, CrcZ and CrcY, which can act additively or independently of each other. This functional redundancy may reduce the risk of loss of a critical component of a regulatory pathway. For example, the deletion of either crcZ or crcY led to a compensatory increase in the levels of the remaining sRNA (Figure 4C). Liu et al. developed a deterministic model and explored the evolutionary conditions for crcY and crcZ co-existence, proposing that both ncRNAs have different biological functions and confer distinct advantages or disadvantages under specific environmental conditions, e.g. on plant surfaces.72 Another earlier study showed that P. putida growing at low temperature shows increased levels of CrcZ and CrcY, implying their involvement in the cold stress response.72 We also found that in A1501, the expression of CrcZ and CrcY was associated with environmental stresses (Figure S4). Our data suggest that both ncRNAs might have different functions and consequently different contributions to different stress responses. A similar phenomenon has been described for the ncRNAs RsmY and RsmZ in P. stutzeri A1501, both of which have overlapping functions in the regulation of biofilm formation but distinctive roles in the regulation of nitrogen fixation.64

Another level of complexity might arise from the positive and negative regulations by Hfq through different modes of action.71 In one mode, Hfq acts as an RNA matchmaker, promoting base pairing between sRNAs...
and their mRNA targets. In a second mode, the chaperone Hfq directly binds the 5’ untranslated region or the coding region of mRNA targets, affecting their translation. Hfq is now recognized as a global regulator of cell physiology, which particularly affects the cell response to several stresses. For example, genetic, phenotypic, and proteomic data reveal wide-ranging effects of Hfq on the basic physiological traits of *P. putida* and *Pseudomonas protegens*. Moreover, another genome-wide analysis of Hfq-binding RNA targets in *S. meliloti* revealed that major stress response and symbiotic gene networks are extensively subject to Hfq-mediated posttranscriptional regulation. A previous study using strain A1501, the NifR and NifS, which are induced under nitrogen fixation conditions, optimize nitrogenase activity in a cooperative manner by increasing the stabilities of the nifD and nifK mRNAs, respectively. A reasonable speculation is that *P. stutzeri* A1501 may carry an unidentified ncRNA or employ an unidentified mechanism that specifically targets the nifH transcript, as previously described for sRNA154 in *Methanosarcina mazei*. The data presented here indicated that Hfq directly binds the nifH mRNA and ultimately enhances nitrogenase activity by increasing the half-life of the nifH mRNA, suggesting a novel Hfq-mediated regulatory mechanism underlying the efficient synthesis of the nitrogenase complex. Additionally, the Hfq mutation led to a significant decrease in the expression of nifA under nitrogen fixation conditions, as also previously reported. This effect is probably indirect, as no Hfq-binding site was identified in either the promoter or coding regions of nifA, thus markedly increasing the complexity of the regulatory circuitry.

Numerous investigations have demonstrated that when multiple carbon sources are available, bacteria display the following types of common growth behavior: hierarchical utilization or simultaneous utilization. Recent studies suggest that two utilization patterns can coexist during bacterial growth in nutritionally complex environments, allowing cells to allocate cellular resources to achieve optimal growth. Our data shown in Figures 7 and S6 provide empirical evidence of a resource allocation strategy that depends on Hfq-mediated opposing regulatory effects as follows: (i) succinate is catabolized as a good carbon and energy source for both growth and nitrogen fixation, while eliciting strong Hfq-mediated repression of glucose utilization that depends on glucokinase encoded by the glk gene; (ii) in addition, the intracellular glucose pool is shunted into an as yet unidentified route to produce the Psl exopolysaccharide, which is regulated positively by Hfq and independent of the glk gene. Obviously, Hfq-mediated opposing regulatory processes guarantee adequate supply of glucose as a raw material to produce exopolysaccharides, a key factor required for the early stage of both nitrogen-fixing biofilm formation and root colonization. We assumed that this strategy on Hfq-mediated opposing regulations may be the result of adaptive evolution within rhizosphere environments. As suggested by Liu et al., carbon catabolite repression confers a selective disadvantage at the plant surface, a nutritionally complex environment where various carbon sources are present simultaneously but fluctuate in their availability. Our findings further provide a fascinating example of sophisticated regulatory networks underlying carbon substrate utilization, nitrogenase activity, and exopolysaccharide production, which might be a common strategy employed by rhizobacteria to succeed in environments that are nutritionally complex and highly competitive.

**Limitations of the study**

Metabolically versatile pseudomonads are very competitive, as evidenced by their broad global distributions, for example as highly successful colonizers of the plant rhizosphere; however, the ecological rationale for the carbon catabolite repression strategy employed by rhizobacteria is still an open question. It is interesting to note that the carbon catabolite repression genes also participate in other important biological processes, such as biofilm formation, nutrient uptake, root colonization, and symbiotic nitrogen fixation, yet the underlying mechanisms are still far from being understood due to their complexities. A key challenge that remains is to develop various tools that will more rapidly and quantitatively characterize the genetic elements and regulatory pathways involved in diazotroph-plant interactions. Progress in this
area will drive the development of new strategies to ensure the compatibility of diazotrophs with host crops and rhizosphere environments, and exploit their potential to the maximum for improving soil fertility and crop growth. Therefore, future studies may provide opportunities to develop synthetic biology approaches that manipulate biofertilizer-producing diazotrophic strains widely used in sustainable agriculture.

STAR METHODS
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SUPPLEMENTAL INFORMATION
Supplemental information can be found online at https://doi.org/10.1016/j.isci.2022.105663.

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AUTHORS CONTRIBUTIONS
Y.Z., Y.Y., and M.L. established the research plan and interpreted the results. F.L. and M.L. designed the experiments. F.L. performed the majority of the experiments. Y.Z., Z.Y., and W.L. performed the MST experiments; X.K., S.J., L.S., and W.L. performed the RT-PCR and ddPCR experiments; Y.Z., Y-M.Y., and W.L. performed Biolog phenotyping and analyzed the growth characteristics. X.K., Y.S., Y.Y., S.J., L.S., Y.M., and J.Z. contributed to the construction of the mutants and vectors, protein purification, and the nitrogenase activity and root colonization assays. F.L., Y.Z., L.C., Y.Y., W.L., C.E., and M.L. analyzed data and prepared figures; C.E., Y.Y., and M.L. wrote the article with input from all co-authors.

DECLARATION OF INTERESTS
The authors have no competing interests to declare.
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## STAR METHODS

### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Bacterial strains** | | |
| Pseudomonas stutzeri A1501 | CGMCC 0351 | N/A |
| Pseudomonas stutzeri A1501 Δhfq | This study | N/A |
| Pseudomonas stutzeri A1501 Δhfq (pLhfq) | This study | N/A |
| Pseudomonas stutzeri A1501 Δcrc | This study | N/A |
| Pseudomonas stutzeri A1501 Δcrc (pLcrc) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔhfqΔcrc | This study | N/A |
| Pseudomonas stutzeri A1501 ΔhfqΔcrc (pLhfq) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔhfqΔcrc (pLcrc) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcbxA | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcbxB | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcbxA (pLcbxA) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcbxB (pLcbxB) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔrpoN | Shang et al. | N/A |
| Pseudomonas stutzeri A1501 ΔrpoN (pLrpoN) | Shang et al. | N/A |
| Pseudomonas stutzeri A1501 ΔcrcZ | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcrcZ (pLcrcZ) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcrcY | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcrcY (pLcrcY) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcrcYΔcrcZ | This study | N/A |
| Pseudomonas stutzeri A1501 ΔcrcYΔcrcZ (pLcrcZ) | This study | N/A |
| Pseudomonas stutzeri A1501 ΔjgK | This study | N/A |
| Pseudomonas stutzeri A1501 ΔpslA | Shang et al. | N/A |
| Pseudomonas stutzeri A1501 edd::lacZ | This study | N/A |
| Pseudomonas stutzeri A1501 gltR::lacZ | This study | N/A |
| **Escherichia coli DH5α** | NEB | Cat# C2987H |
| **Escherichia coli BL21** | NEB | Cat# C2527H |
| **Chemicals, peptides, and recombinant proteins** | | |
| D-Glucose | Sigma-Aldrich | Cat# G8270 |
| Succinic acid | Thermo Scientific | Cat# A12084.36 |
| Sodium lactate | Thermo Scientific | Cat# L14500.06 |
| Sodium benzoate | Thermo Scientific | Cat# 447802500 |
| Sodium citrate | Sigma-Aldrich | Cat# PHR1416 |
| Potassium phosphate monobasic | Aladdin | Cat# P104078 |
| Potassium hydrogen phosphate | Aladdin | Cat# P112219 |
| Sodium chloride | Aladdin | Cat# C111538 |
| Magnesium sulfate | Aladdin | Cat# M110771 |
| Manganese sulfate monohydrate | Aladdin | Cat# M111712 |
| Iron (III) sulfate hydrate | Aladdin | Cat# F112483 |

(Continued on next page)
| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Sodium molybdate dihydrate | Aladdin | Cat# S104871 |
| DMSO | Invitrogen | Cat# D12345 |
| EDTA | Aladdin | Cat# E116429 |
| EDTA-Ca | Sigma-Aldrich | Cat# 340073-100G |
| PBS | MP Biomedicals | Cat# PBS10X02 |
| Tris-HCl | Solarbio | Cat# TB230 |
| Yeast Extract Powder | Oxoid | Cat# LP0021B |
| Tryptone | Oxoid | Cat# LP0042B |
| Sulfuric acid | Sigma-Aldrich | Cat# 258105 |
| Ethanol | Thermo Scientific | Cat# 611050040 |
| Water-Saturated Phenol | Invitrogen | Cat# AM9712 |
| Gentamicin | Sigma-Aldrich | Cat# G1264-5G |
| Hygromycin B | Sigma-Aldrich | Cat# SBR00039 |
| Spectinomycin | Sigma-Aldrich | Cat# 59007-5G-9 |
| Kanamycin | Sigma-Aldrich | Cat# E004000-5G |
| Ampicillin | Sigma-Aldrich | Cat# A5354 |
| Tetracycline hydrochloride | Thermo Scientific | Cat# 233100025 |
| HEPES buffer | GIBCO | Cat# 15630080 |
| Trimethylamine | Sigma-Aldrich | Cat# 471283 |
| β-Mercaptoethanol | Sigma-Aldrich | Cat# 444203 |
| DTT | Thermo Scientific | Cat# R0861 |
| RNAlater solution | Invitrogen | Cat# AM7020 |
| Xba I | NEB | Cat# R0145S |
| Sph I-HF | NEB | Cat# R3182S |
| Nhe I-HF | NEB | Cat# R3131S |
| EcoR I-HF | NEB | Cat# R3101S |
| BamH I -HF | NEB | Cat# R3136S |
| Hind III-HF | NEB | Cat# R3104S |
| Nde I | NEB | Cat# R0111S |
| Spe I-HF | NEB | Cat# R3133S |
| Xho I | NEB | Cat# R0146S |
| Nco I-HF | NEB | Cat# R3193S |
| 4-Methylumbelliferyl-b-D-galactoside | Sigma-Aldrich | Cat# 4368814 |
| 7-Hydroxy-4-methylcoumarin | Sigma-Aldrich | Cat# 128724 |
| DNase I | Promega | Cat# M6101 |
| Biolog GENIII MicroPlate | BioLog | Cat# 1030 |
| Strep-Tactin column | IBA LifeSciences | Cat# 12-4013-001 |
| Chitin Beads | NEB | Cat# S6651V |
| Buffer W | IBA LifeSciences | Cat# 2-1003-100 |

Critical commercial assays

ClonExpress II One Step Cloning Kit | Vazyme | Cat# C112-01 |

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**Continued**

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| TIANamp Bacteria DNA Kit | TIANGEN | Cat# DP302 |
| M5 HiPer Multi-color Plasmid Miniprep Kit | Met5bio | Cat# MF031-plus-04 |
| CloneJET PCR Cloning Kit | Thermo Scientific | Cat# K1232 |
| High-Capacity cDNA Reverse Transcription Kit | ABI | Cat# 4368814 |
| T7 High Yield RNA Transcription Kit | Vazyme | Cat# TR101-01 |
| Quick Start Bradford Protein Assay Kit | Bio-Rad | Cat# 5000201 |
| InnuPREP RNA Mini Kit | Analytik Jena | Cat# 10755-390 |
| PowerUp SYBR Green PCR Master Mix | ABI | Cat# 01118369 |
| 5’ RACE System for Rapid Amplification of cDNA Ends | Invitrogen | Cat# 18374-058 |
| 2 × EvaGreen ddPCR SuperMix | Bio-Rad | Cat# 186-4034 |
| Ni-NTA Fast Start Kit | Qiagen | Cat# 30600 |
| Protein labeling Kit RED-NHS | NanoTemper | Cat# L001 |

**Experimental models: Organisms/strains**

| Organism | Source | Remarks |
|----------|--------|---------|
| Zhonghua 11 (Oryza sativa L. subsp. Japonica) | Chinese Academy of Agricultural Sciences | N/A |

**Oligonucleotides (Primers used in this study, see Table S4)**

**Recombinant DNA**

| Plasmid | Source | Remarks |
|---------|--------|---------|
| pK2013 | Lab collection Figurski and Helinski | N/A |
| pK18mob | Lab collection Schäfer et al. | N/A |
| pK18mobsacB | Lab collection Schäfer et al. | N/A |
| pUX-BF13 | Lab collection Choi et al. | N/A |
| pMD18-T | TaKaRa | Cat# 6011 |
| pLAFR3 | Lab collection Staskawicz et al. | N/A |
| pXY2 | Lab collection Liu et al. | N/A |
| pET30a | Novagen | Cat# 69909 |
| pET28a | Novagen | Cat# 69864 |
| pTWIN1 | NEB | Cat# N6951S |
| pK18mob-glk | This study | N/A |
| pKatCA75 | Lab collection | N/A |
| pKathPH4 | Lab collection | N/A |
| pKatAAD2 | Lab collection | N/A |
| pK18mobsacB-hfq | This study | N/A |
| pK18mobsacB-crc | This study | N/A |
| pK18mobsacB-crcZ | This study | N/A |
| pK18mobsacB-crcY | This study | N/A |
| pK18mobsacB-cbrA | This study | N/A |
| pK18mobsacB-cbrB | This study | N/A |
| pXY2-Pedd | This study | N/A |
| pXY2-PglrR | This study | N/A |
| pLhfxq | This study | N/A |
| pLcbnA | This study | N/A |
| pLcrCY | This study | N/A |

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RESOURCE AVAILABILITY

Lead contact
Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Min Lin (linmin@caas.cn).

Materials availability
This study did not generate new unique reagents.

Data and code availability
- All data reported in this paper will be shared by the lead contact upon request.
- This paper does not report original code.
- Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODELS AND SUBJECT DETAILS

Bacterial strains and growth conditions
*P. stutzeri* A1501 and its derivatives were grown on LB medium or minimal medium K (containing 0.4 g L\(^{-1}\) KH\(_2\)PO\(_4\), 0.1 g L\(^{-1}\) K\(_2\)HPO\(_4\), 0.1 g L\(^{-1}\) NaCl, 0.2 g L\(^{-1}\) MgSO\(_4\) \(\cdot\) 7H\(_2\)O, 0.01 g L\(^{-1}\) MnSO\(_4\) \(\cdot\) H\(_2\)O, 0.01 g L\(^{-1}\) Fe\(_2\)(SO\(_4\))\(_3\) \(\cdot\) H\(_2\)O, and 0.01 g L\(^{-1}\) Na\(_2\)MoO\(_4\) \(\cdot\) H\(_2\)O, pH 6.8) supplemented with the indicated carbon and nitrogen sources. Unless stated otherwise, growth experiments were conducted using minimal medium K containing NH\(_4\)\(^+\) (6.0 mM) and succinate (20 mM) as sole nitrogen and carbon sources with agitation at 200 rpm at 30°C. For diauxic measurements, minimal medium K was supplemented with an excess concentration of glucose (20 mM) plus a preferred carbon source at a concentration (4 mM) insufficient to support full growth, on which A1501 grow in a diauxic manner. If needed, *E. coli* and A1501 were grown in the presence of 100 μg mL\(^{-1}\) ampicillin (Amp), 10 μg mL\(^{-1}\) tetracycline (Tc), or chloramphenicol (Cm), 50 μg mL\(^{-1}\) spectinomycin (Spc), 50 μg mL\(^{-1}\) gentamicin sulfate (Gm), 50 μg mL\(^{-1}\) kanamycin (Km) and 50 μg mL\(^{-1}\) hygromycin (Hyg). The absorbance (OD\(_{600}\)) was measured with a U-3010 (Hitachi) spectrophotometer.

Plant materials and growth conditions
The rice seeds (zhonghua11 from a rice-breeding program at the Chinese Academy of Agricultural Sciences) were surface sterilized by a 2 min soak in 75% (v/v) ethanol, followed by 10 min in sodium hypochlorite solution (10% active chlorine). The seeds were rinsed five times (5 min each time) with sterile distilled water and germinated on humid sterile filter paper at 30°C. Three-day-old seedlings were transferred to the beaker containing sterilized MS medium (1.65 g L\(^{-1}\) NH\(_4\)NO\(_3\), 0.44 g L\(^{-1}\) CaCl\(_2\) \(\cdot\) 2H\(_2\)O, 0.37 g L\(^{-1}\) MgSO\(_4\) \(\cdot\) 7H\(_2\)O, 0.17 mg L\(^{-1}\) KH\(_2\)PO\(_4\), 1.9 g L\(^{-1}\) KNO\(_3\), 6.2 mg L\(^{-1}\) H\(_2\)BO\(_3\), 0.025 mg L\(^{-1}\) CoCl\(_2\) \(\cdot\) 6H\(_2\)O, 27.8 mg L\(^{-1}\) FeSO\(_4\) \(\cdot\) 7H\(_2\)O, 22.3 mg L\(^{-1}\) MnSO\(_4\) \(\cdot\) 4H\(_2\)O, 0.83 mg L\(^{-1}\) KI, 0.25 mg L\(^{-1}\) Na\(_2\)MoO\(_4\) \(\cdot\) 2H\(_2\)O, and 2.5 g L\(^{-1}\) sucrose).
The resulting plasmids were introduced into A1501 by triparental mating using pRK201383 as the helper plasmid of pXY2 was electroporated into chromosomally integrated upon first cross-over selection for Km and corresponding anti-biotic resistances. Excision of the vector in the second cross-over was obtained through enrichment of Km-sensitive cells on 10% (wt/vol) sucrose and corresponding antibiotic-supplemented plates. The Cm resistance gene was amplified by PCR from pKatCA75 for the construction of Δcrc, ΔcbrA, and ΔcbrB. The Hyg resistance gene was amplified from pKatHPH4 for the construction of Δhfq, ΔcrcZ, and ΔcrcY. The Spec resistance gene was amplified from pKatAAD2 for the construction of Δhfq, ΔcrcZ, and ΔcrcY. Correct recombination was confirmed by PCR followed by nucleotide sequencing of the amplicons obtained. After recombination in the host genome, genes located downstream of the insertion (when part of the same operon) were transcribed from the promoter of the resistance gene, and the expression levels of downstream genes in WT and mutant strains under normal growth conditions were compared using qRT-PCR.

The complemented strains were constructed using the broad host plasmid pLAFR3. A DNA fragment containing a WT gene with its promoter and terminator was amplified from genomic DNA of A1501 and cloned into pLAFR3. The resulting complementing plasmid was then introduced into the WT or mutant strain by triparental mating, generating complemented strains (Table S3). The gene expression levels in the overexpression strains were confirmed to be higher than those in the WT strain using qRT-PCR.

The ΔcrcZΔcrcY double mutant was constructed using the ΔcrcZ mutant as the parental strain. Briefly, a fragment that contained the Cm resistance cassette located between the upstream and downstream DNA fragments of crcY was generated by overlap extension PCR using the strategy of PCR-based fusions. The fusion PCR product was then cloned into the multiple cloning site of the pMD18-T vector (Takara, Japan). The resulting plasmid DNA was double digested with Xba I/Sph I and then cloned into the same site of pK18mobsacB. The resulting plasmid was introduced into the genome of ΔcrcZ by triparental mating and a double recombinant was selected based on sucrose resistance. Correct recombination in the resulting ΔcrcZΔcrcY mutant was checked by PCR using the primers M-crcY2(up)-F and M-crcY2(down)-R (Table S4), followed by nucleotide sequencing of the obtained PCR products. In the same way, we also constructed the ΔhfqΔcrc mutant using Δhfq as the starting strain. Similarly, a fragment containing the Spec resistance cassette located between the upstream and downstream DNA fragments of crc was cloned between the Nhe I and EcoRI sites of pK18mobsacB, generating the plasmid pK18crc, and then was introduced into the genome of Δhfq. Correct recombination in the resulting ΔcrcZΔcrcY double mutant was checked by PCR.

Construction of translational lacZ fusions and β-galactosidase assay
To construct translation gene fusions, the DNA fragments that carried the respective promoter of the target gene were amplified using the primers edd-lacZ F/edd-lacZ R2 and gtr-lacZ F/gtr-lacZ R2 (Table S4) and cloned into plasmid pXY2, a modified version of pUC18-mini-Tn7-Gm-lacZ. The recombinant plasmid of pXY2 was electroporated into P. stutzeri A1501 together with pUX-BF13, and the mini-Tn7 element carrying the lacZ reporter fusion was integrated into the unique Tn7 site located downstream of glmS. Specific β-galactosidase activity from bacterial suspensions growing in liquid cultures was measured by using 4-methylumbelliferyl-b-D-galactoside (4MUG) as the enzymatic substrate. The fluorescent product, 7-hydroxy-4-methylcoumarin (4MU), was detected at 460 nm after excision at 365 nm using a FlexStation 3 Plate Reader (Molecular Devices). Enzyme activity was reported as μmol 4 MU min⁻¹ mg⁻¹ protein.
Biolog phenotype profiling

The 96-well Biolog GENIII MicroPlate (Catalogue No.1030) was used to measure the respiratory activity of A1501 on different carbon sources. Briefly, the cultures were incubated at 30°C until mid-logarithmic phase, after which cells were harvested and suspended in a special “gelling” inoculating fluid (IF) at the recommended cell density. A total of 100 μL of the bacterial suspension was used to inoculate each well of the GENIII MicroPlate. The microplates were placed in an OmniLog instrument (Biolog Inc.) programmed to run at 30°C for 48 h. During the incubation, increased respiratory activity is observed in the wells, and cells utilize a carbon source and generate oxidants that then oxidize tetrazolium redox dye, forming a purple colour. The formation of formazan was recorded at 15 min intervals, and data were analysed using OmniLog Kinetic and Parametric software (Biolog Inc.).

Rhizosphere colonization and competition assays

The 12-day-old seedlings were transferred to 150 mL tubes (four seedlings per tube) that contained 30 mL of minimal K medium supplemented with the indicated carbon sources, and then inoculated with the indicated strains to obtain a final concentration of approximately 10⁵ total CFUs/mL. In competition experiments, strains were inoculated at a 1:1 ratio. To confirm starting ratios at a 1:1 ratio, colony counts were performed on the initial inoculum. After 24 h of incubation, seedlings were harvested and the leaves were cut off. Root tips were washed and resuspended in 10 mL phosphate-buffered saline (PBS: 137 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 2 mM KH₂PO₄). Root-attached bacteria were recovered by vortexing roots for 1.0 min and plating the appropriate dilutions on LB agar plates supplemented with the appropriate antibiotics. The plates were incubated overnight at 30°C. The amount of cells per g of root was then determined by colony forming unit (CFU) counting. The results were normalized to the fresh weight of the root.

Strain competition experiment

Freshly cultivated P. stutzeri wild-type strain A1501 and mutants were combined at a 1:1 ratio, inoculated at a starting OD₆₀₀ of 0.1 and grown for 24 h in LB. To confirm starting ratios at a 1:1 ratio, colony counts were performed on the initial inoculum. After 24 h of incubation, the cultures were serially diluted and plated on LB agar plates containing the relevant antibiotics to enable each strain to be counted. The wild-type strain A1501 was used as the competitor in all experiments. The mutant fitness score ratio was calculated as the ratio of CFUs of each mutant to that of A1501.

Nitrogenase activity assays

The nitrogenase activity of bacteria in pure cultures was evaluated using the acetylene reduction assay. Cells from an overnight culture in LB medium were centrifuged and resuspended in a 100 mL flask containing 10 mL of minimal medium K supplemented with the indicated carbon sources at an OD₆₀₀ of 0.1. The suspension was incubated for 24 h at 30°C with vigorous shaking under an argon atmosphere containing 0.5% oxygen, and then 10% acetylene was added. Gas samples (0.25 mL) were taken at 2-h intervals to determine the amount of ethylene produced on a poly divinylbenzene porous bead GDX-502 column using an SP-2100 gas chromatograph fitted with a flame ionization detector (Beijing Beifen-Ruili Analytical Instrument Co. Ltd.). The ethylene content in the gas samples was determined by reference to an ethylene standard. The nitrogenase activity was expressed as nmol ethylene min⁻¹ mg⁻¹ protein. Protein concentrations were determined using the Bio-Rad protein assay reagent kit (Bradford, Bio-Rad).

Root-associated nitrogenase activity was measured using the acetylene reduction assay. The 12-day-old seedlings were transplanted to 150 mL tubes (four seedlings per tube) that contained 30 mL of minimal K medium supplemented with the indicated carbon sources, and then inoculated with the indicated strains at a final OD₆₀₀ of 0.1. The tubes sealed with rubber septa were injected with 10% (vol/vol) pure acetylene, and ethylene concentrations were measured at 2-h intervals for 24 h at 30°C under static conditions. Ethylene production from uninoculated seedlings in the presence of 10% acetylene and inoculated seedlings in the presence of air was also assayed to determine the background ethylene emission of rice as a plant hormone.

Total RNA isolation and quantitative real time RT-PCR (qRT-PCR) analysis

To determine the expression of the indicated genes in pure bacterial cultures, total RNA was isolated with an innuPREP RNA Mini Kit (Analytik Jena) according to the manufacturer’s instructions. Total RNA was
reverse transcribed using random primers and the High Capacity cDNA Transcription Kit (Applied Biosystems) according to the manufacturer’s instructions. PCR was carried out with Power SYBR Green PCR Master Mix on an ABI Prism 7500 Sequence Detection System (Applied Biosystems) according to the manufacturer’s recommendations. The 16S rRNA gene was used as the endogenous reference control, and relative gene expression was determined using the comparative threshold cycle 2^ΔΔCT method. Data were analysed using ABI PRISM 7500 Sequence Detection System Software (Applied Biosystems). Primers were designed based on the full genome sequence of *P. stutzeri* A1501, and they are listed in Table S4.

Twelve-day-old seedlings were transferred to 150 mL tubes (four seedlings per tube) containing 30 mL of minimal K medium supplemented with the measured carbon sources, and then inoculated with the indicated strains at a final OD_{600} of 0.1 to determine the expression of genes in *P. stutzeri* A1501 associated with host rice roots. After 8 h of incubation, root tips were placed on 20 mL of 60% (v/v) RNAlater solution (Invitrogen) in PBS. Root-attached bacteria were removed from roots by vortexing for 1.0 min, harvested by quick centrifugation (18,900 × g, 1 min, 4°C), and re-suspended in cold lysis buffer (Analytik Jena). Total RNA was isolated with an innuPREP RNA Mini Kit (Analytik Jena), and qRT-PCR was performed using SYBR Premix Ex Taq (TaKaRa) and gene-specific primers (Table S4).

### 5' rapid amplification of cDNA ends (5' RACE) to determine transcriptional start sites

The transcriptional start of CrcZ and CrcY was determined using 5' RACE (Invitrogen) according to the manufacturer’s instructions. Briefly, first-strand cDNA was synthesized using the primer GSP1, which was specific for the target gene sequence. The purified cDNA was tailed with dCTP by terminal deoxynucleotidyl transferase. PCR amplification was performed using the sequence-specific primer GSP2 and the anchor primer AAP. The primers GSP1 and GSP2, which are specific for the target gene tested here, are listed in Table S3. The 5’ RACE products were cloned into the pGEM-T Easy vector (Promega) and sequenced to map the 5’ end of the transcript.

### Absolute quantification of the RNA copy number using droplet digital PCR (ddPCR)

Quantification by ddPCR was performed in 20 μL reaction mixtures containing 10 μL of 2 × EvaGreen ddPCR SuperMix (Bio-Rad) and primers at a final concentration of 0.2 μM were included. No template controls were used to monitor contamination and primer-dimer formation. Reactions were equilibrated for 3 min at room temperature and dispensed into each well of the droplet generator DG8 cartridge (Bio-Rad). Each oil compartment of the cartridge was filled with 70 μL of droplet generation oil for EvaGreen (Bio-Rad), and approximately 20,000 droplets were generated in each well using the droplet generator (Bio-Rad, QX200). The entire droplet volume (40 μL) was subsequently loaded onto a 96-well PCR plate and then heat sealed with a pierceable foil in the PX1 PCR Plate Sealer (Bio-Rad). The optimal thermal cycling conditions were used as follows: 95°C for 5 min; 40 cycles of 95°C for 30 s, 58°C for 1 min; and then 98°C for 10 min. The cycled droplets were read individually with the QX 200 droplet reader (Bio-Rad) and analysed with QuantaSoft droplet reader software.

### Microscale thermophoresis (MST) analysis

DNA templates that carried the wild or mutated crcZ (339 bp), crcY (315 bp), oprB (1254 bp), gltR (761 bp), zwf (1486 bp), citN (1344 bp), benR (984 bp), and nifH (882 bp) were synthesized by Sangon Biotech Co. All the synthesized DNA templates were sequenced to assure the absence of undesired mutations. For in vitro transcription from synthesized templates, the T7 High Yield RNA Transcription Kit (Vazyme) was used according to the manufacturer’s instructions. The resulting ssRNA oligonucleotides were purified by the RNAClean XP (Beckman Coulter Inc.) according to the manufacturer’s instructions. The purified Crc and Hfq proteins were labelled with a Monolith NT™ Protein labelling kit RED-NHS according to the manufacturer’s instructions (No. MO-L011, NanoTemper Technologies). For the measurements, the concentration of the labelled Hfq (20 nM) or Crc (200 nM) proteins was kept constant, while the concentrations of non-labelled ssRNA oligonucleotides varied from 0.3 μM to 10 μM. The dissociation constants (K_d) were calculated. Data analyses were performed with NanoTemper Analysis software (NanoTemper Technologies).

### Expression and purification of Hfq for MST measurements

The Hfq protein was expressed and purified using the Strept-Tactin system (IBA Life Sciences). To this end, a 255 bp DNA fragment containing the entire *hfq* ORF except the stop codon was amplified by PCR using the...
oligonucleotides str-hfq F/str-hfq R (Table S4). The resulting product was double-digested with Nde I/Xho I and then ligated into the corresponding sites of plasmid pET30a, generating the plasmid pET31a-hfq. The *E. coli Δhfq* strain (kindly provided by Yong Tao, the Institute of Microbiology of the Chinese Academy of Sciences, Beijing) containing the plasmid pET31a-hfq was incubated in LB medium containing the appropriate antibiotic at 37°C with shaking to reach an OD_{600} of 0.6. At this point, expression of the Hfq protein was induced by adding isopropyl β-D-1-thiogalactopyranoside (IPTG) to a final concentration of 0.5 mM. After 6 h of incubation at 30°C, cells were harvested by centrifugation and resuspended in buffer W (100 mM Tris-HCl, pH 8.0, 150 mM NaCl, 1 mM EDTA). The cells were disrupted by sonication after eliminating cell debris by centrifugation (10,000 rpm, 4°C) for 30 min, and the supernatant was loaded onto a Strep-Tactin column (IBA Life Sciences). The column was extensively washed with wash buffer (250 mM NaCl, 50 mM Tris-HCl, pH 7.5). The Hfq protein was eluted with wash buffer (100 mM Tris-HCl, 150 mM NaCl, pH 8.0, 1 mM EDTA, and 50 mM biotin) and analysed using SDS-PAGE. The protein concentration was determined using the Bio-Rad protein assay reagent kit (Quick Start Bradford, Bio-Rad). Final preparations were stored at −80°C in storage buffer (20 mM Tris-HCl, pH 8.0, 200 mM NaCl, and 30% glycerol) until use.

**Expression and purification of Crc and CbrB**

The Crc protein was expressed and purified using the IMPACT™ (Intein Mediated Purification with an Affinity Chitin-binding Tag) system according to the manufacturer’s instructions (New England Biolabs). A 777 bp fragment, which covers the entire coding region of the crc gene but excludes the termination codon, was amplified by PCR using pTW crc F/pTW crc R (Table S4). The resulting fragment was cloned between the Nde I and Xho I sites of the Pr7 expression vector pTWIN1, generating the plasmid pTWIN1-Crc, which carries an in frame fusion at the 3’-end of the crc gene with codons for the intein/chitin-binding domain to allow purification through a chitin column. The *E. coli Δhfq* strains containing the plasmid pTWIN1-Crc were incubated in LB medium containing the appropriate antibiotic at 37°C with shaking to reach an OD_{600} of 0.6. At this point, expression of the Crc protein was induced by the addition of IPTG to a final concentration of 0.5 mM. After 6 h of incubation at 30°C, the cells were harvested by centrifugation and resuspended in elution buffer (20 mM HEPES, 500 mM NaCl, 0.1 mM EDTA, pH 8.0). The cells were disrupted by sonication; cell debris was eliminated by centrifugation (10,000 rpm, 4°C) for 30 min, and the supernatant was loaded onto a chitin column. The column was extensively washed with wash buffer (20 mM HEPES, pH 8.0, 500 mM NaCl, and 0.1 mM EDTA). The Crc-intein fusion was cleaved by an overnight treatment at 4°C with elution buffer containing 30 mM dithiothreitol (DTT). The untagged Crc protein was eluted from the column, dialyzed against a buffer (500 mM NaCl and 20 mM HEPES, pH 8.0) and concentrated using 3K Amicon filtration units. The purity of the protein was as high as 90%, as judged using SDS-PAGE. The protein concentration was determined using the Bio-Rad protein assay reagent kit (Quick Start Bradford, Bio-Rad). Final preparations were stored at −80°C in storage buffer (20 mM Tris-HCl, pH 8.0, 200 mM NaCl, and 30% glycerol) until subsequent MST measurements.

The CbrB protein was expressed and purified using the IMPACT™ system for DNase I footprinting assays.

**Expression and purification of RpoN for DNase I footprinting assays**

The pET-28a expression system (Novagen) was used to produce C-terminally His-tagged RpoN within host *E. coli* BL21 (DE3) cells. A fragment of rpoN was amplified by PCR using the pET28a-rpoN-F/R primers (Table S4). The PCR product was digested with Nde I and Hind III and ligated into the protein expression vector pET28a, which had been digested with the same enzymes. The resulting plasmid (pET28a-RpoN) was introduced into the *E. coli* BL21 strain. An overnight culture of BL21 harbouring the expression plasmid was inoculated into LB medium containing the appropriate antibiotics. This cell culture was incubated with shaking at 37°C until the OD_{600} was 0.6 ~ 0.9, at which point production of His-tagged RpoN was induced by the addition of IPTG to a final concentration of 0.1 mM. His-tagged RpoN was purified using a Ni-NTA Fast Start Kit (Qiagen, Venlo, Netherlands) according to the manufacturer’s instructions. The purity of the protein was as high as 90%, as judged by SDS-Tris-glycine PAGE. His-tagged RpoN was quantitated using the Bio-Rad protein assay reagent kit (Quick Start Bradford, Bio-Rad) and stored at −80°C until further DNase I footprinting assays.

**DNase I footprinting assays**

DNase I footprinting assays were performed according to the manufacturer’s instructions (Tolo Biotech).92 The DNA probe was prepared by PCR amplification of the 264 bp CrcZ and 300 bp CrcY promoter regions...
using the primers FP crcZ F/FP crcZ R and FP crcY F/FP crcY R (Table S4), respectively. For each assay, 300 ng probes were incubated with 5 µg purified RpoN and 15 µg purified protein CbrB in a total volume of 40 µL. After an incubation for 30 min at 30°C, 10 µL of a solution containing ~0.015 units of DNase I (Promega) and 100 nmol of freshly prepared CaCl₂ were added and a subsequent incubation was performed at 37°C for 1 min. The reaction was stopped by adding 140 µL DNase I stop solution (200 mM unbuffered sodium acetate, 30 mM EDTA, and 0.15% SDS). Samples were first extracted with phenol/chloroform and then precipitated with ethanol. Pellets were dissolved in 30 µL of MilliQ water.

The preparation of the DNA ladder, electrophoresis, and data analysis were performed according to the manufacturer’s instructions,92 except that the GeneScan-LIZ600 size standard (Applied Biosystems) was used.

**Determination of nifH mRNA stability**

The stability of the nifH mRNA was determined in A1501 and the Δhfq or Δhfq (pLhfq) strains grown under nitrogen fixation conditions for 6 h followed by the addition of rifampicin (400 µg/mL, final concentration). Aliquots were added to two volumes of RNA protect (Sigma). After incubating at room temperature for 5 min, the samples were centrifuged for 5 min at 4°C and 12,000 rpm, and the pellets were quickly frozen in liquid nitrogen and stored at −80°C until they were ready for use. Total RNA was isolated with an innu-REP RNA Mini Kit (Analytik Jena) according to the manufacturer’s instructions. Full-length cDNAs were synthesized from total RNA using a First Strand cDNA synthesis kit (Takara Bio) and were used to estimate mRNA levels with qRT-PCR. The primers used to measure the mRNA half-life are listed in Table S4 (RT nifH F/RT nifH R). The relative mRNA concentration was calculated by the comparative threshold cycle (2⁻ΔΔCT) method with 16S rRNA as the endogenous reference. Data are presented as the percentage of nifH transcript levels relative to the amount of these transcripts at time point zero.

**Exopolysaccharide isolation and quantification**

The strains were grown in minimal medium K supplemented with 6 mM ammonium and different carbon sources at 30°C for 24 h. Then, the cultures were treated with 10% trichloroacetic acid (1:1), homogenized for 30 min, and centrifuged at 10,000 rpm for 30 min. Exopolysaccharides were isolated from culture supernatants by adding two volumes of chilled absolute ethanol. After further precipitation by cold absolute ethanol and lyophilization, purified exopolysaccharides were obtained.93 The exopolysaccharides were quantified using the phenol-sulfuric acid method94 and subsequently normalized to the total cellular protein level remaining after extraction, as determined using the Bio-Rad protein assay reagent kit (Quick Start™ Bradford, Bio-Rad). Final concentrations were expressed as mg exopolysaccharide per g bacterial protein.

**HPLC analysis of glucose consumption**

Culture samples (1.0 mL) were obtained at different time points on the growth curve and were centrifuged at 10,000 rpm for 30 min. The supernatant was collected and frozen at −80°C until HPLC analysis. The HPLC-RID system (Waters Corp.) equipped with a refractive index (RI) detector was used for each sample.95 The injection volume was 10 µL and the mobile phase was a 50-mg/L EDTA-Ca water solution delivered at a flow rate of 0.5 mL/min. The column temperature was set at 90°C. The chromatography running time was 15 min. The HPLC peaks were identified to be glucose by comparing the retention time between them and the standard compound.

**QUANTIFICATION AND STATISTICAL ANALYSIS**

Unless stated otherwise, experiments were performed three times with similar results. Each bar on graphs represents a mean of biological replicates and error bars indicate the SEM (standard error of the mean), as mentioned in figure legends for each experiment. For all assays, n represents the number of biological replicates. Statistical analysis was performed using GraphPad Prism 9.0. A two-tailed unpaired Student’s t-test with a 95% confidence interval was used to evaluate the difference between two groups. For more than two groups, one-way ANOVA was used. A probability value of p ≤ 0.05 was considered significant. Data are presented as averages ± SEM. ****: p ≤ 0.0001; ***: p ≤ 0.001; **: p ≤ 0.01; *: p ≤ 0.05; and ns: non-significant.