



Quorum sensing signal autoinducer-2 promotes root colonization of *Bacillus velezensis* SQR9 by affecting biofilm formation and motility

Qin Xiong¹ · Di Liu¹ · Huihui Zhang² · Xiaoyan Dong¹ · Guishan Zhang¹ · Yunpeng Liu¹ · Ruifu Zhang^{1,2}

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Abstract

Root colonization of beneficial rhizobacteria is critical for their beneficial effects. Quorum sensing (QS) has been reported to affect the colonization of many plant pathogens. However, how QS signals regulate root colonization of beneficial rhizobacteria is unclear. In this study, the QS signal AI-2 synthetase-encoding gene *luxS* was completely deleted from the genome of the plant beneficial rhizobacterium *Bacillus velezensis* SQR9, and bioluminescence experiments showed that AI-2 production was blocked. Deletion of *luxS* reduced biofilm formation, motility, and root colonization of *B. velezensis* SQR9, while addition of exogenous AI-2 to the mutant restored this phenomenon. These results indicated that AI-2 positively affects the root colonization of *B. velezensis* SQR9. This study provided new insights for enhancing the colonization of beneficial rhizobacteria.

Key points

- *LuxS* participated in the synthesis of the quorum sensing signal AI-2 in *B. velezensis*.
- AI-2 enhanced motility, biofilm formation, and root colonization of *B. velezensis*.
- AI-2 stimulated the production of γ -polyglutamic acid by *B. velezensis*.

Keywords AI-2 · *Bacillus velezensis* · Biofilm formation · Motility · Quorum sensing · Root colonization

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✉ Guishan Zhang
zhangguishan@caas.cn

✉ Yunpeng Liu
liuyunpeng@caas.cn

¹ Key Laboratory of Microbial Resources Collection and Preservation, Ministry of Agriculture, Institute of Agricultural Resources and Regional Planning, Chinese Academy of Agricultural Sciences, Beijing 100081, People's Republic of China

² Jiangsu Key Lab and Engineering Center for Solid Organic Waste Utilization, National Engineering Research Center for Organic-based Fertilizers, Nanjing Agricultural University, Nanjing 210095, People's Republic of China

Introduction

Plant growth-promoting rhizobacteria (PGPRs) have been widely used in agriculture for their environmentally safe and plant growth-promoting properties (Lugtenberg and Kamilova 2009). Efficient root colonization of PGPRs is necessary for the bacteria to exert their beneficial effects on plants. Chemotactic motility towards roots (chemotaxis) and the formation of biofilms on the rhizoplane are the major processes of root colonization by PGPRs (Cao et al. 2011; Li et al. 2013). In the biofilm formation process, extracellular matrix components, such as extracellular polysaccharides (EPSs) and γ -polyglutamic acid (γ -PGA), play an important role in clustering the cells together. Motility and biofilm formation are regulated by a variety of factors, including both exogenous and endogenous signals. Quorum sensing (QS), which is associated with bacterial density and generally determined by endogenous signals, has been widely studied for its effects on the motility and biofilm formation of bacteria (Huber et al. 2001), especially pathogenic bacteria.

Therefore, we speculated that QS would be an important factor that influences the root colonization of PGPRs.

QS is a process for regulating specific behaviors in a cell density-dependent manner (Bassler 1999). QS is accomplished by the release, detection, and transport of small signaling molecules called autoinducers that accumulate in proportion to cell density (Wai-Leung and Bassler 2015). With some exceptions, QS signals mainly include *N*-acyl-homoserine lactones (AHLs), cholera autoinducer 1 (CAI-1), autoinducer-2 (AI-2), and oligopeptides (Vendeville et al. 2005). The QS systems in gram-negative and gram-positive bacteria are different. While AHLs are the most common and well-studied class of autoinducers in gram-negative bacteria, these molecules have not been detected or are not functional in gram-positive bacteria. In gram-positive bacteria, QS is usually mediated by AI-2, and this molecule could be shared by both gram-negative and gram-positive bacteria (Xavier and Bassler 2003).

AI-2 is a 2-methyl-2,3,3,4-tetrahydroxytetrahydrofuran (THMF) formed through spontaneous rearrangements of the precursor 4,5-dihydroxy 2,3-pentanedione (DPD), which is synthesized by S-ribosylhomocysteinase LuxS (Schauder et al. 2001). AI-2 was first identified in the marine bacterium *Vibrio harveyi* and is part of a complex multilayered QS system responsible for regulating bioluminescence and other virulence-associated traits (Bassler et al. 1994). In addition, AI-2 has been proposed as a chemokine in *Escherichia coli* K12 (Hegde et al. 2011). The effect of AI-2 on biofilm formation varies in different bacterial species, and the *Staphylococcus aureus luxS* mutant strain formed stronger biofilms than the wild-type strain under static, flowing, and anaerobic conditions (Yu et al. 2012). *Bacillus* species are the most important group of PGPRs; however, how AI-2 and the QS system affect the root colonization of *Bacillus* species is not clear.

Bacillus velezensis SQR9 is a beneficial plant bacterium isolated from the rhizosphere of cucumber plants (Cao et al. 2011). This PGPR strain has shown several beneficial effects on plants, including promoting growth (Liu et al. 2016), inducing plant resistance (Wu et al. 2018), and enhancing plant tolerance to abiotic stress (Chen et al. 2017). In this study, the potential AI-2 synthetase-encoding gene *luxS* was deleted, and the mutant was assessed for AI-2 production, biofilm formation, motility, and root colonization. We demonstrated that AI-2 plays positive roles in biofilm formation, γ -PGA production, motility, and root colonization of *B. velezensis* SQR9.

Materials and methods

Strains and culture conditions

The strains and plasmids used in this study are shown in Table 1. *B. velezensis* strain SQR9 (CGMCC accession no. 5808, China General Microbiology Culture Collection Center,

NCBI accession NO. CP006890) was cultured at 37 °C in Luria-Bertani medium (1% (w/v) tryptone, 0.5% (w/v) yeast extract, and 0.5% (w/v) NaCl). *V. harveyi* (from the Han lab of Shanghai Veterinary Research Institute, CAAS) was cultured at 30 °C in AB medium (0.05 M MgSO₄, 0.3 M NaCl, 1% (w/v) glycerol, 10 mM K₂HPO₄, 10 mM L-arginine, and 0.02% (w/v) casamino acids). To measure biofilm formation and extracellular polymeric substances, *B. velezensis* SQR9 and its derived mutants were cultured in MSgg medium (100 mM 3-(*N*-morpholino) propane sulfonic acid (MOPS), 5 mM potassium phosphate, 2 mM MgCl₂, 700 μ M CaCl₂, 50 μ M MnCl₂, 50 μ M FeCl₃, 1 μ M ZnCl₂, 2 μ M thiamine, 0.5% (w/v) glycerol, 0.5% (w/v) glutamate, 50 μ g/mL tryptophan, 50 μ g/mL phenylalanine, and 50 μ g/mL threonine, pH 7.0) (Branda et al. 2001). When necessary, antibiotics were added to the growth media at the following concentrations: erythromycin, 1 μ g/mL; spectinomycin, 100 μ g/mL; and zeocin, 20 μ g/mL for *B. velezensis* strains.

luxS deletion and complementation

To disrupt the target gene in the SQR9 genome, an erythromycin resistance gene was used to replace the *luxS* gene to obtain the *luxS* mutant. The upstream and downstream regions that flanked the *luxS* gene were amplified from the SQR9 genome. The primers used to knock out the *luxS* gene in SQR9 are shown in Table S1. The upstream homologous fragment, the erythromycin gene fragment with the promoter, and the downstream homologous fragment were fused by using the two-step overlapping PCR method described by Shao et al. (2015), and the resulting product was transformed into *B. velezensis* SQR9 to generate the *luxS* mutant strain of *Bacillus velezensis* SQR9, $\Delta luxS$. Complementation of the *luxS* gene was performed by inserting the *luxS* operon (including its promoter) into the $\Delta luxS$ mutant at the amylase gene locus (*amyE*) and inserting spectinomycin (*Spc*, amplified from plasmid p7S6) as the screening marker. All these mutants were verified by PCR amplification and sequencing.

Detection of AI-2

Detection of AI-2 was performed following the method reported by Shao et al. (2007). AI-2 detection was carried out using *Vibrio harveyi* BB 170 (sensor 1⁻, sensor 2⁺) as a reporter strain. *V. harveyi* BB170 was cultured in AB medium overnight at 30 °C and 170 rpm. The cell suspension with an OD₆₀₀ of 0.2–0.3 was diluted with AB medium at 1:5000 as the reporter solution. At the same time, supernatants of wild-type SQR9, $\Delta luxS$, and $\Delta luxS/luxS$ cultures were collected at exponential phase, filtered through a 0.22- μ m membrane and filtered through a 3-kDa exclusion filter (Millipore). Subsequently, the filtrate was lyophilized, and the powder was dissolved in PBS solution (pH 7.0). In addition, 4 μ M

Table 1 Strains and plasmids used in this study

Strains or plasmids	Genotype	Source/reference
Plasmids		
pAX01	Em ^R	Li et al. (2018)
p7S6	Spc ^R , pMD18-T ligated with <i>spc</i> gene	Yan et al. (2008)
Strains		
<i>B. velezensis</i>		
SQR9	Wild-type isolate	Cao et al. (2011)
$\Delta luxS$	Complete deletion of the <i>luxS</i> gene, Em ^R	This work
SQR9 $\Delta luxS$ /- <i>uxS</i>	Genome complementation of <i>luxS</i> gene with original promoter at the locus of <i>amyE</i> , Spc ^R	This work
<i>V. harveyi</i>		
BB170	Kan ^R , sensor1 ⁻ , sensor2 ⁺ , LuxPQ (AI-2 receptor), <i>lux</i> system of luminescence	Han lab (Shanghai Veterinary Research Institute, CAAS)

pure AI-2 (4,5-dihydroxy 2,3-pentanedione (DPD)) (Omm Scientific Inc, TX, USA) was included as a positive control. Both the culture medium (AB medium) for the reporter strain and PBS solution were included as negative controls. Then, 20 μ L of each sample was added to 180 μ L of reporter solution and incubated at 30 °C and 200 rpm. Bioluminescence was measured and recorded every hour.

Plants and growth conditions

The seeds of the “jingtian” maize cultivar were soaked in 75% ethanol for 40 s, washed with sterile water, and soaked in 0.2% (w/v) NaClO solution for 5 min. The seeds were placed on sterile wet filter paper for germination at 25 °C. The germinated seeds were transferred to sterile vermiculite and cultured at a suitable temperature for 3 days. Then, maize seedlings were transplanted to 100 mL flasks containing 75 mL of sterile liquid 1/4 Hoagland medium. The plants were grown with a 16-h light/8-h dark photoperiod. The seedlings were ready for the colonization experiment after culturing for an additional 15 days.

Biofilm formation assay

The biofilm assay was carried out in MSgg medium, as described by Hsueh et al. (2006). *B. velezensis* SQR9 and its derived strains were cultured in LB medium to an OD₆₀₀ of 1.0. The cells were collected by centrifugation at 8000 rpm. The pellets were washed with sterile water three times and suspended in an equal volume of MSgg medium. The concentrations were adjusted to an OD₆₀₀ of 1.0. Then, for qualitative analysis, 10 μ L of each cell suspension was inoculated into a 48-well plate containing 1 mL of MSgg medium in each well. Exogenous AI-2 was added to the corresponding wells to a

final concentration of 4 μ M. The 48-well plate was cultured at 37 °C without shaking. Biofilm formation was evaluated at 16 h postinoculation. For quantitative analysis, 100 μ L of each cell suspension was inoculated into 6-well plates containing 10 mL of MSgg medium in each well. Sterile biofilm filters 15-1040 (Biologixb, Changzhou, China) were placed in each well to collect the biofilm for weighing. After 16 h, the biofilm was weighed. For each treatment, four replicates were included.

Motility assay

The swimming and swarming assay was performed following the method of Sperandio and Inoue (Sperandio et al. 2002; Inoue et al. 2007). *B. velezensis* SQR9 and its derived strains were grown in LB medium until the OD₆₀₀ reached 0.8. The cultures were inoculated on petri dishes containing semisolid LB medium (0.5% (w/v) glucose and 0.6% (w/v) Eiken agar (Eiken, Nogi-machi, Japan) for the swarming assay, and 0.3% (w/v) Eiken agar for the swimming assay) by sterilized toothpicks. The petri dishes were incubated at 37 °C for 4–6 h to allow the bacteria to swim. Then, the water content of the medium in the petri dishes was reduced with dry air to terminate the swimming process. Subsequently, the petri dish was incubated overnight at room temperature.

Detection of γ -PGA

The viscosity assay was performed as described by Feng et al. (2014) to determine the production of γ -PGA. For each strain, 1 mL of cell suspension with an OD₆₀₀ of 1.0 was inoculated into a 250-mL flask containing 100 mL of MSgg medium. Exogenous AI-2 was added to the corresponding wells to a final concentration of 4 μ M. The bacteria were cultured with

shaking at 170 rpm at 37 °C. At 24 h postinoculation, the viscosity of 20 mL of fermentation broth was measured by an NDJ-8S rotatory viscometer (Ping Xuan, Shanghai, China) with an adapted No. 0 rotator. The detection speed was set at 60 rpm for samples with a viscosity under 10 mPa s and 30 rpm for samples with a viscosity over 10 mPa s. For each treatment, three replicates were included.

The cetyltrimethylammonium bromide (CTAB) detection assay was performed as described by Ashiuchi (2011). In solutions with high ionic strength, CTAB forms complexes with proteins and non-acid polymers, so the concentration of γ -PGA can be calculated by the turbidity of the complexes. For each strain, 1 mL of cell suspension with an OD₆₀₀ of 1.0 was inoculated into a 250-mL flask containing 100 mL of MSgg medium. Exogenous AI-2 was added to the corresponding wells to a final concentration of 4 μ M. The bacteria were cultured with shaking at 170 rpm at 37 °C. At 24 h postinoculation, γ -PGA was purified using a previously described method (Goto and Kunioka 1992), and pure γ -PGA was dissolved in an equal volume of ddH₂O for further detection. At the same time, standard γ -PGA (Yuan Ye, Shanghai, China) solutions were prepared with concentrations of 0.0391, 0.0781, 0.1563, 0.3125, 0.625, 1.25, 2.5, and 5 g/L. Then, 2 mL of each sample solution or standard γ -PGA solution was mixed with 2 mL of CTAB solution (2% NaOH, 5 g/L CTAB); subsequently, the OD₂₅₀ was measured by Infinitr M200PRO (TECAN, Männedorf, Switzerland). The production of γ -PGA was calculated based on the standard curve. For each treatment, three replicates were included.

Colonization assay

The maize seedlings were grown in a hydroponic system as described above and divided into three groups for treatment. *B. velezensis* SQR9 and its derivative strains were cultured in LB medium to an OD₆₀₀ of 1.0. The cells were collected by centrifugation at 8000 rpm. The pellets were washed with sterile water three times and resuspended in the same volume. The bacterial cells were inoculated into the maize seedling rhizosphere to a final concentration with an OD₆₀₀ of 0.1. Exogenous AI-2 was added to the corresponding treatment at a final concentration of 4 μ M. After incubation for 3 days, the population of the cells colonizing the rhizoplane was determined by plate counting as described previously (Liu et al. 2017). For each treatment, six replicates were included.

Result

LuxS is necessary for AI-2 production in *B. velezensis* SQR9

The *luxS* gene encoding *S*-ribosylhomocysteinase, which catalyzes the production of AI-2 from *S*-ribosylhomocysteine

(SRH), was identified in the *B. velezensis* SQR9 genome. Deletion of *luxS* did not affect the growth of *B. velezensis* SQR9 (Figure S1). To verify its function in AI-2 production, the SQR9 wild-type strain and the *luxS* mutant were tested for AI-2 production by bioluminescence assay. For this experiment, both the culture medium (AB medium) for the reporter strain and the PBS solution were included as negative controls. Pure AI-2 was applied as a positive control. The results showed that deletion of *luxS* significantly decreased AI-2 production while reintroducing *luxS* into the mutant restored AI-2 production (Fig. 1). These results indicated that *luxS* is responsible for AI-2 signal production in SQR9.

AI-2 signal promotes biofilm formation of *B. velezensis* SQR9

To test the effect of AI-2 on biofilm formation of *B. velezensis* SQR9, both qualitative and quantitative analysis of biofilm formation by the wild-type strain and the *luxS* deletion mutant in MSgg medium were performed. At 16 h postinoculation, wild-type SQR9 formed more wrinkles than $\Delta luxS$, and the wet weight of the pellicle biofilm of the wild-type strain was 1.6-fold that of the $\Delta luxS$ strain. Reintroducing *luxS* into the mutant with its original promoter or adding 4 μ M AI-2 restored biofilm formation. Moreover, the addition of exogenously AI-2 further increased the biofilm of the wild-type strain (Fig. 2). These results showed that AI-2 could enhance biofilm formation of *B. velezensis* SQR9.

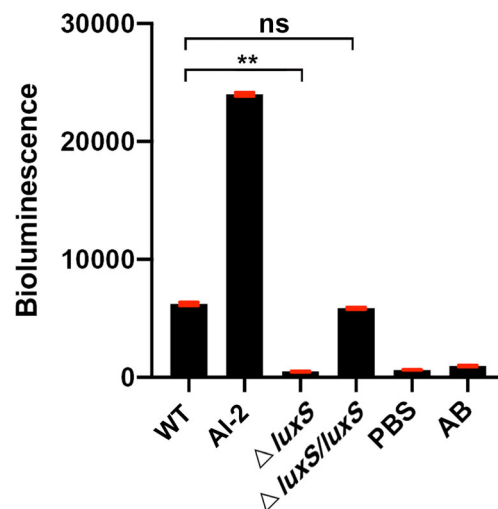


Fig. 1 Measurement of AI-2 production by bioluminescence assay. AI-2 in the culture medium of the SQR9 wild-type, $\Delta luxS$ and $\Delta luxS/luxS$ strains was tested using a reporter strain *V. harveyi* BB170, whose luminescence was enhanced with increasing concentrations of AI-2 in the environment. Pure AI-2 was applied with the final concentration of 4 μ M as a positive control, and AB medium was included as a negative control. Error bars indicate the standard deviations based on six independent replicates

AI-2 signal increases the synthesis of γ -PGA by *B. velezensis* SQR9

The extracellular matrix is important for biofilm formation. To investigate how AI-2 stimulates the biofilm formation of strain SQR9, the effect of AI-2 on the *Bacillus* biofilm matrix, including exopolysaccharides (EPSs), the amyloid-like protein TasA, and γ -polyglutamic acid (γ -PGA), were tested. As the extracellular matrix can be analyzed by the viscosity of the supernatant, we first measured the viscosity of the wild-type strain and $\Delta luxS$ mutant. The results showed that the SQR9 wild-type strain had a higher culture supernatant viscosity than the $\Delta luxS$ mutant. The viscosity of the culture supernatant of the wild-type strain was 7.48 mPa s, while that of $\Delta luxS$ was decreased to 2.32 mPa s. Complementation with *luxS* and adding 4 μ M AI-2 restored the viscosity of the cultures to 5.32 mPa s and 6.33 mPa s, respectively. In addition, exogenously adding AI-2 to wild-type SQR9 cultures resulted in a viscosity of 11.64 mPa s (Fig. 3a).

Fig. 2 Biofilm formation of *B. velezensis* SQR9. WT, wild-type; WT+AI-2, wild-type strain supplied with exogenous AI-2; $\Delta luxS$, *luxS* mutant; $\Delta luxS$ /AI-2, $\Delta luxS$ supplied with exogenous AI-2; $\Delta luxS$ /*luxS*, $\Delta luxS$ complemented with *luxS* gene at the amylase gene locus (*amyE*). The final concentration of exogenous AI-2 was 4 μ M. **a** Biofilm formation in 48-well plates. **b** The weight of biofilms at 16 h. Error bars indicate the standard deviations based on four independently replicated experimental values

The synthesis of EPS and TasA by the SQR9 wild-type and $\Delta luxS$ mutant strains was similar (Figure S2 and Figure S3), and we speculate that biofilm matrix γ -PGA synthesis may be different between the SQR9 wild-type and $\Delta luxS$ mutant strains. CTAB can bind γ -PGA specifically to form a water-insoluble, highly dispersed micelle-like complex, which can be measured on the basis of the turbidity to determine the concentration of γ -PGA. Therefore, we used CTAB solution to react with the fermentation supernatant of each treatment. The results showed that the concentration of γ -PGA in the supernatant of the SQR9 wild-type strain was 0.92 g/L, and adding exogenous AI-2 to wild-type SQR9 increased the concentration of γ -PGA in the supernatant to 2.50 g/L. When *luxS* was deleted from the wild-type strain, the concentration of γ -PGA in the supernatant was decreased to 0.08 g/L, and complementation of the mutant with *luxS* and adding 4 μ M AI-2 restored the concentration of γ -PGA to 0.40 g/L and 0.62 g/L, respectively (Fig. 3b). SDS-PAGE of the γ -PGA samples from each treatment had the same tendency

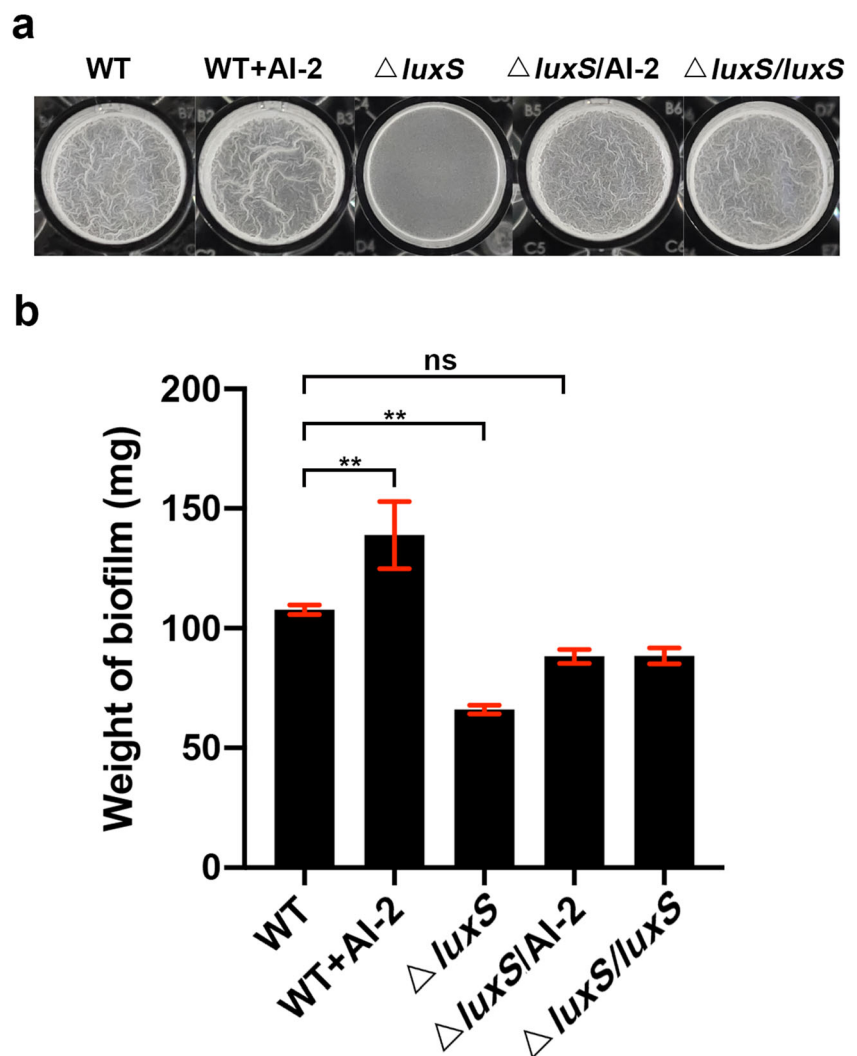
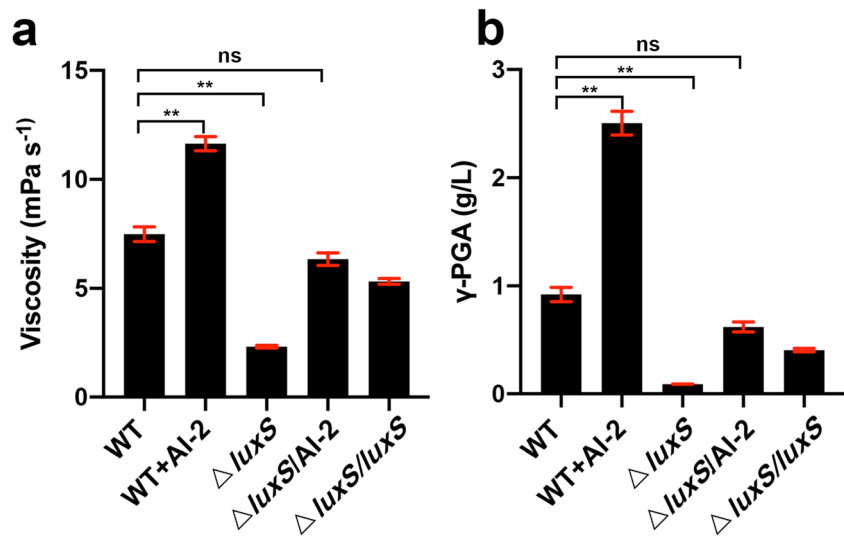


Fig. 3 The γ -PGA yield of *B. velezensis* SQR9. WT, wild-type; WT+AI-2, wild-type strain supplied with exogenous AI-2; $\Delta luxS$, *luxS* mutant; $\Delta luxS$ /AI-2, $\Delta luxS$ supplied with exogenous AI-2; $\Delta luxS/luxS$, $\Delta luxS$ complemented with *luxS* gene at the amylase gene locus (*amyE*). The final concentration of exogenous AI-2 was 4 μ M. **a** The viscosities of fermentations. **b** The yield of γ -PGA. Error bars indicate the standard deviations based on three independently replicated experimental values



(Figure S4). The results showed that AI-2 affected the synthesis of γ -PGA in *B. velezensis* SQR9.

AI-2 signal promotes motility of *B. velezensis* SQR9

Bacterial motility is critical for PGPR to colonize plant roots. To investigate the effect of AI-2 signaling on the motility of SQR9, swimming analysis of the SQR9 wild-type strain, $\Delta luxS$ mutant strain, and *luxS* gene complementation strain was performed; the results were compared and showed that the diameter of the motility circle of $\Delta luxS$ was significantly decreased compared with that of SQR9. Complementation with *luxS* or adding 4 μ M AI-2 to the medium restored the motility circle of $\Delta luxS$ mutant strain to the wild-type level (Fig. 4). This result indicates that AI-2 has a positive effect on the motility ability of SQR9.

AI-2 signal enhances the root colonization of *B. velezensis* SQR9

Biofilm formation and motility are important factors in determining the root colonization of PGPRs. As shown in this study, the effects of LuxS and AI-2 on biofilm formation and motility are both positive. We proposed that LuxS/AI-2 should have a positive effect on root colonization. Quantitative analysis of colonization of maize roots was performed at 3 days postinoculation using the plate counting method. The results showed that the root colonization of $\Delta luxS$ decreased significantly by 35%. When *luxS* was introduced into the mutant, the colonization of the resulting strain on maize roots was restored to the same level as that of the wild-type strain. When exogenous AI-2 was added to the rhizosphere of maize, colonization of SQR9 on roots significantly increased by 16% (Fig. 5). Exogenous AI-2 also rescued the colonization decreasing caused by *luxS* mutation. These

results indicated that AI-2 could promote root colonization of SQR9.

Discussion

Quorum sensing is an important system that regulates specific functions in bacteria. *Bacillus* is the most widely used plant beneficial bacteria in agriculture, but knowledge on how QS signals regulate the rhizospheric behavior of *Bacillus* is lacking. In this study, the QS signal AI-2 synthetase-encoding gene *luxS* was deleted from the *B. velezensis* SQR9 genome. AI-2 production was completely blocked in the *luxS* mutant, and biofilm formation, motility, and root colonization of the *luxS* mutant were significantly reduced. These results indicated that AI-2 could act as a quorum sensing signal and promote root colonization of *B. velezensis*. This finding would provide new insights into the colonization of PGPR.

Quorum sensing of gram-positive bacteria is regulated by few kinds of oligopeptides and AI-2. The signaling oligopeptides are generally produced by specific cleavage of proteins. For example, bacterial pheromones, which play roles in sporulation and the production of certain secondary metabolites (Rosenberg et al. 2012), are cleaved from the ComX protein. Nevertheless, knowledge of the QS signal in gram-positive bacteria is still very poor. In comparison with other QS signals, AI-2 more widely exists in both gram-negative and gram-positive bacteria, and the structure of AI-2 is highly conserved among species, giving it the capability to act as an interspecies QS signal. However, pathways regulated by AI-2 among different bacteria are varied. Merritt et al. found that deletion of the AI-2 synthetase LuxS could affect the biofilm morphology of *Streptococcus mutans* (Merritt et al. 2003). Similarly, the addition of AI-2 promoted the biofilm formation of *Staphylococcus epidermidis* (Xue et al. 2015) and *Bacillus cereus* (Auger et al. 2006). This study also showed similar results

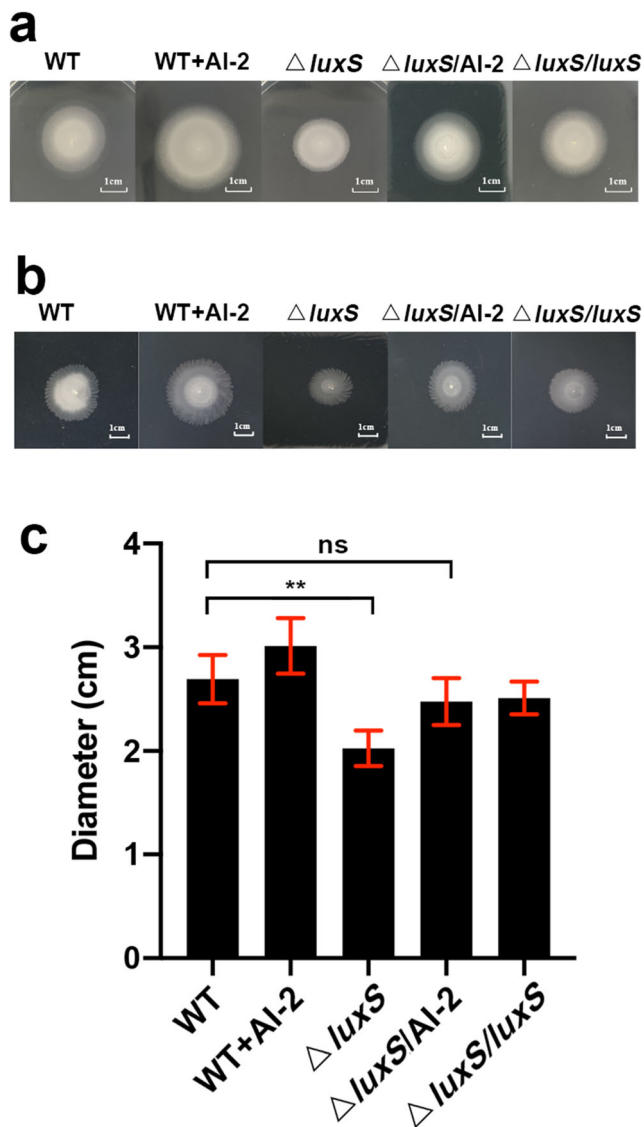


Fig. 4 Motility analysis of *B. velezensis* SQR9. WT, wild-type; WT+AI-2, wild-type strain supplied with exogenous AI-2; $\Delta luxS$, *luxS* mutant; $\Delta luxS/AI-2$, $\Delta luxS$ supplied with exogenous AI-2; $\Delta luxS/luxS$, $\Delta luxS$ complemented with *luxS* gene at the amylase gene locus (*amyE*). The final concentration of exogenous AI-2 was 4 μ M. **a** Swimming of cells on LB swim medium solidified with 0.3% agar. **b** Swarming of cells on LB swarming medium solidified with 0.6% agar and 0.5% glucose. **c** The diameters of cell spots inoculated on swarming plates that incubated at 37 °C for 4 h. Error bars indicate the standard deviations based on six independently replicated experimental values

that AI-2 could positively affect biofilm formation of *B. velezensis* SQR9. It has been reported that deletion of *luxS* from *E. coli* K-12 led to enhanced motility (Ling et al. 2010). However, in *Bacillus subtilis*, deletion of *luxS* led to reduced motility (Lombardía et al. 2006). Similarly, in this study, AI-2 also showed a positive effect on the motility of *B. velezensis* SQR9.

Bacillus biofilm formation is a complex process regulated by a range of regulation systems, such as Spo0A, SinI-SinR (Xu et al. 2017), and the DegSU system (Verhamme et al. 2009). In this study, AI-2 increased the production of γ -

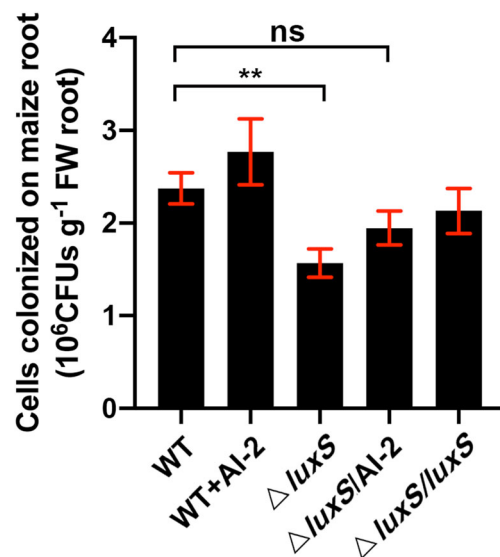


Fig. 5 Root colonization of *B. velezensis* SQR9. WT, wild-type; WT+AI-2, wild-type strain supplied with exogenous AI-2; $\Delta luxS$, *luxS* mutant; $\Delta luxS/AI-2$, $\Delta luxS$ supplied with exogenous AI-2; $\Delta luxS/luxS$, $\Delta luxS$ complemented with *luxS* gene at the amylase gene locus (*amyE*). The final concentration of exogenous AI-2 was 4 μ M. Error bars indicate the standard deviations based on six independently replicated experimental values

PGA. The synthesis of γ -PGA is regulated by phosphorylated DegU (Stanley and Lazazzera 2005). Interestingly, the pheromone synthesized from ComX is also involved in the regulation of the DegSU system. The pheromone could promote the transcription of *degQ*, which facilitates the transfer of an activating phosphate from DegS to DegU. It is proposed that AI-2 might have a similar pathway as the ComX pheromone and regulate biofilm formation of *B. velezensis*. Further study on how AI-2 participates in biofilm formation regulation would provide insight into the AI-2 and ComX pheromone signaling pathways.

This study revealed that AI-2 had the same effect on the synthesis of γ -PGA and biofilm formation. Therefore, it is presumed that AI-2 could affect biofilm formation in *Bacillus* spp. by influencing the synthesis of γ -PGA. γ -PGA is a natural polymer with many good properties, such as being nontoxic, edible, degradable, water soluble, and modifiable. γ -PGA can be used in fertilizers to enhance their effectiveness (Hoppensack et al. 2003), as well as biological flocculants for sewage treatment (Ritchie et al. 1999). γ -PGA is also used in food and cosmetics (Shyu et al. 2008), and it is widely used in medicine; it can be used as a drug carrier to improve drug solubility (Liang et al. 2006). γ -PGA is mainly produced by microbial fermentation; therefore, it is very important to improve the efficiency of microbial synthesis of γ -PGA. Our study found that the addition of AI-2 can significantly increase γ -PGA production in SQR9, which is important for the production of γ -PGA in industry.

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Authors' contributions RZ and GZ conceived and designed research. QX, DL, and HZ conducted experiments. XD and YL analyzed data. QX and YL wrote the manuscript. All authors read and approved the manuscript.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval This article does not contain any studies with human participants or animals performed by any of the authors.

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