Two (p)ppGpp Synthetase Genes, \textit{relA} and \textit{spoT}, Are Involved in Regulating Cell Motility, Exopolysaccharides Production, and Biofilm Formation of \textit{Vibrio alginolyticus}

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The stringent response mediated by the signal molecule (p)ppGpp is involved in response to multiple environmental stresses and control of various physiological processes. Studies have revealed that (p)ppGpp strongly affects the formation and maintenance of several bacterial biofilms. However, the specific regulatory roles of (p)ppGpp in biofilms, especially in the expression of genes related to cell motility and exopolysaccharides (EPSs) production, remain poorly understood.

We recently reported two (p)ppGpp synthetase genes \textit{relA} and \textit{spoT} from the epizootic pathogen \textit{Vibrio alginolyticus}. Herein, we found that the (p)ppGpp synthetase genes of \textit{V. alginolyticus} contributed to biofilm formation at low cell density and biofilm detachment at high cell density, respectively, in polystyrene microtiter plates. Quantitative reverse transcription PCR (qRT-PCR) analysis revealed that the expression levels of both EPSs and motility associated genes were consistent with the development of biofilms. Besides, the (p)ppGpp synthetase gene \textit{spoT} was found to be closely involved in the regulation of flagellum, smooth/translucent colony morphology and spotty pellicle at the air-liquid interface. Interestingly, pleiotropic phenotypes of $\Delta$\textit{relA}$\Delta$\textit{spoT} were similar to that of the \textit{rpoN} (\textit{\(\sigma^{54}\)}) deletion mutant. Meanwhile, the absence of (p)ppGpp synthetase genes significantly reduced the expression levels of \textit{rpoN} at low cell density, suggesting that (p)ppGpp may mediate the formation via positively affecting the alternative sigma factor RpoN. These findings allow us to propose (p)ppGpp as a crucial regulator for biofilm development in \textit{V. alginolyticus}, in view of the regulatory roles of \textit{relA} and \textit{spoT} in cell motility and EPSs production.

\textbf{Keywords: Vibrio alginolyticus, stringent response, (p)ppGpp, biofilm formation, motility, EPSs, rpoN}
INTRODUCTION

Vibrio alginolyticus is a halophilic, facultative anaerobic, and Gram-negative opportunistic pathogen that inhabits coastal waters and estuaries. V. alginolyticus has been reported as a notorious causative agent of vibriosis in many marine animals, including fish, shellfish, and shrimp (Lee et al., 1996; Gömez-León et al., 2005; Kahla-Nakbi et al., 2006; Austin, 2010). Besides, it can cause ear infections, gastroenteritis and septicemia in humans through wounds or ingestion of contaminated food (Mustapha et al., 2013; Jacobs Slifka et al., 2017). The robust response abilities to environmental stresses and stringent control of virulence genes are thought to be essential for the pathogenicity of this bacterium (Wang et al., 2007; Rui et al., 2008).

Biofilm formation is one of the most effective strategies for pathogenic Vibrio spp. to confront harsh environmental conditions (Faruque et al., 2006; Crozatto et al., 2007; Hall and Mah, 2017). Initially, flagella and other motility factors accelerate bacterial attachment to the surface and initiate microcolony formation (O’Toole and Kolter, 1998; Utada et al., 2014). Exopolysaccharides (EPSs), the most prevalent component of extracellular matrix, are produced to promote the development of mature biofilms (Yildiz and Visick, 2009; Teschler et al., 2015) and responsible for the transition of colony morphology between opaque/rugose (Op) and translucent/smooth (Tr) (Chen et al., 2009). EPSs production depends on transcription of the Vibrio polysaccharide synthesis (vps) genes (Yildiz and Schoolnik, 1999). Deletion of vpsM or other vps genes lead to a reduction in biofilm formation and EPSs production in V. chOLERa, indicating vps genes have important roles in biofilm formation (Fong et al., 2010). A complex intracellular regulatory network, including transcriptional activators, alternative sigma factors and other factors, could directly regulate the vps genes (Zhu et al., 2002; Teschler et al., 2015). VpsT, a member of the UhpA (FixJ) family of transcriptional regulators, is found to be required for vps genes expression and development of rugose colonial morphology in V. chOLERa O1 El Tor (Casper-Lindley and Yildiz, 2004; Beyhan et al., 2007). The alternative sigma factor-54 (RpoN) has been reported to be strongly involved in affecting vps genes expression of Vibrio spp. (Yildiz and Visick, 2009; Cheng et al., 2018). Interestingly, RpoN could indirectly promote the vps genes expression at low cell density in V. chOLERa (Herzog et al., 2019). Finally, bacteria stimulated by nutrient stress or extracellular signals would downregulate vps genes, degrade polysaccharides, and reorientate in direction to escape from the biofilm matrix mesh (Fong and Yildiz, 2007; Pratt et al., 2009; Hay and Zhu, 2015; Bridges et al., 2020).

The stringent response mediated by the bacterial alarmones pppGpp and ppGpp [collectively termed (p)ppGpp] is considered as an important adaptive response to stressful conditions, whether living in the environment or in the host (Srivatsan and Wang, 2008). In many Gram-negative bacteria, such as V. alginolyticus, the metabolism of (p)ppGpp is controlled by two conserved enzymes: monofunctional synthetase RelA specialized for the synthesis of (p)ppGpp and bifunctional synthetase/hydrolase SpoT (Magnusson et al., 2005; Gaca et al., 2015). As a global regulator, (p)ppGpp is able to activate or repress the transcription of many genes by directly regulating RNA polymerase in cooperation with the RNA polymerase-binding transcription factor DksA or indirectly interacting with σ-factors (Dalebroux and Swanson, 2012). It has been reported that (p)ppGpp synthetases are involved in the regulation of biofilm formation in several bacterial species (Sugisaki et al., 2013; Azriel et al., 2016; Díaz-Salazar et al., 2017; Colomer-Winter et al., 2019). The lack of (p)ppGpp synthetase genes in most pathogens showed reduced biofilm formation (Renier et al., 2011; He et al., 2012; Sugisaki et al., 2013), while some (p)ppGpp-deletion mutants could form significantly enhanced biofilms compared with the wild type, such as Actinobacillus pleuropneumoniae (Li et al., 2015) and Pseudomonas putida KT2440 (Liu et al., 2017). This means that (p)ppGpp synthetase genes play significant regulatory roles in biofilm formation among bacterial species.

Although a close link between (p)ppGpp synthetase genes and biofilm formation has been found, the regulation mechanism of (p)ppGpp synthetase genes on biofilm formation remains unclear. In this manuscript, we constructed the (p)ppGpp-synthetase deletion mutants (ΔrelA, ΔrelAΔspoT) and observed the alteration of biofilm formation, EPSs production, colonial morphology, and flagellum by comparison with the wild type (WT) and complemented strains (ΔrelA-pRelA, ΔrelAΔspoT-pSpoT). Finally, we elaborated the potential regulatory role of (p)ppGpp synthetase genes in biofilm formation by analyzing these phenotypes and the expression levels of several key genes related to biofilm formation and motility. These findings help us better understand the complex regulatory network of biofilm formation in V. alginolyticus.

MATERIALS AND METHODS

Bacterial Strains and Growth Conditions

Vibrio alginolyticus HN08155 and its derivative strains were cultured at 30°C and 180 rpm in 2216E medium consisting of 5 g/L tryptone, 1 g/L yeast extract, and 0.01 g/L FePO4. Escherichia coli β2163 was grown at 37°C and 180 rpm in standard Luria-Bertani (LB) medium containing 0.3 mM diaminopimelic acid (DAP) (Luo et al., 2015). Antibiotics were used at the following concentrations: 100 µg/mL ampicillin (Amp) was used for the growth of Vibrio spp., 10 µg/mL chloramphenicol (Cm) was additionally added to complemented strains for integrated plasmids. All reagents were purchased from Solarbio (Beijing, China).

Construction of Deletion Mutants and Complemented Strains

All strains and plasmids used are described in Supplementary Table 1. The (p)ppGpp synthetase gene mutants of V. alginolyticus HN08155 were generated by allelic exchange (Milton et al., 1996). This method has been described in detail in our previous work (Yin et al., 2021), and the oligonucleotides used in these procedures are listed in Supplementary Table 2. In brief, the sucrose-sensitive suicide plasmid pDM4-relA containing the fusion product of a 542 bp upstream fragment
and a 542 bp downstream fragment of relA coding sequence was introduced into E. coli β2163 cells by heat shock. Subsequently, the recombinant plasmid pDM4-relA was integrated into wild type through conjugation, and the relA internal coding sequence was deleted after homologous recombination occurred on both homologous arms. Single crossover mutants with plasmids integrated into specific chromosomal loci were obtained on 2216E agar plates supplemented with Amp and Cm. The second crossover mutants were screened on 2216E agar plates supplemented with Amp and 10% sucrose. The in-frame deletion mutant ΔrelA was confirmed by PCR and DNA sequencing. For generation of the double deletion strain ΔrelA ΔspoT, the suicide plasmid pDM4-spoT containing the fusion product of a 563 bp upstream fragment and a 500 bp downstream fragment of spoT coding sequence was transferred into the single deletion mutant ΔrelA, and the spoT internal coding sequence was deleted after double cross-over recombination. High cellular (p)ppGpp level in strains would inhibit cell growth, due to the lack of hydrolysis of (p)ppGpp. The opening reading frame of tetracycline resistance gene were amplified, respectively. The open reading frame of tetracycline resistance gene) were amplified, respectively. The open reading frame of tetracycline resistance gene) were amplified, respectively. The open reading frame of tetracycline resistance gene) were amplified, respectively. The open reading frame of tetracycline resistance gene), designated ΔrelA-ΔspoT. For colonial morphology assay, the cultures were adjusted to an OD_{600} of 0.6, diluted 1:100 into 50 mL fresh LB medium without antibiotics. Each 200 µL sample was taken from the diluted medium into the 96-well polystyrene microplate and cultivated statically at 30°C. After culturing for 2, 4, 6, 8, 10, 12, 14, 18, 24, and 36 h respectively, bacterial cells were removed, rinsed once with phosphate buffer saline (PBS) and 220 µL of 0.1% crystal violet was added to stain biofilm for 30 min. The residual stains were then removed, and the stained biofilms were washed twice with PBS. Finally, the amount of biofilm was determined at 570 nm using a microplate reader (Epoch2, BioTek) after dissolving crystal violet with 200 µL of 95% ethanol for 1 h. Three separate experiments were performed with biological triplicates each. To exclude growth effects on biofilm formation, the cell density was measured at 600 nm, and biofilm formation was normalized by dividing total biofilm by the cell density.

### Table 1 | Primers used for qRT-PCR.

| Gene name | Gene function | Primer sequence (5′ to 3′) |
|-----------|---------------|--------------------------|
| vpsH      | Polysaccharides biosynthesis protein | F: 5′-CGACTACGGATGACACCACTCG-3′<br>R: 5′-AACCCTGATGTTGAACAACCTG-3′ |
| vpsT      | Helix-turn-helix transcriptional regulator | F: 5′-GCAGACAGAAGAGGCAATCGT-3′<br>R: 5′-GCCGAGAACCACTCATCCAC-3′ |
| vpsM      | Polysaccharides biosynthesis protein | F: 5′-GCCGAGAACCACTCATCCAC-3′<br>R: 5′-GCCGAGAACCACTCATCCAC-3′ |
| cheR      | Chemotaxis protein | F: 5′-ATACCTTGCTCAGTACGCCA-3′<br>R: 5′-TTGGTTAGTCGATGC-3′ |
| flgE      | Flagellar hook protein | F: 5′-ACCGAAGGCTAGATGTTTCTC-3′<br>R: 5′-ACCGAAGGCTAGATGTTTCTC-3′ |
| flgH      | Flagellar basal body L-ring protein | F: 5′-GCCGAGAACCACTCATCCAC-3′<br>R: 5′-GCCGAGAACCACTCATCCAC-3′ |
| rpoN      | Regulator | F: 5′-CGCAACAGAAAGATACGCTCG-3′<br>R: 5′-CGCAACAGAAAGATACGCTCG-3′ |
| 16S       | 16S ribosomal RNA | F: 5′-GCAGTCACGTATCAGCACCT-3′<br>R: 5′-GCAGTCACGTATCAGCACCT-3′ |

### Assays of Biofilm, Spotty Pellicle, and Colonial Morphology

The biofilm assay was performed following a previously described protocol with modification (O’Toole and Kolter, 1998). After overnight incubation, the cultures were adjusted to an OD_{600} of 0.6, diluted 1:100 into 50 mL fresh LB medium without antibiotics. Each 200 µL sample was taken from the diluted medium into the 96-well polystyrene microplate and cultivated statically at 30°C. After culturing for 2, 4, 6, 8, 10, 12, 14, 18, 24, and 36 h respectively, bacterial cells were removed, rinsed once with phosphate buffer saline (PBS) and 220 µL of 0.1% crystal violet was added to stain biofilm for 30 min. The residual stains were then removed, and the stained biofilms were washed twice with PBS. Finally, the amount of biofilm was determined at 570 nm using a microplate reader (Epoch2, BioTek) after dissolving crystal violet with 200 µL of 95% ethanol for 1 h. Three separate experiments were performed with biological triplicates each. To exclude growth effects on biofilm formation, the cell density was measured at 600 nm, and biofilm formation was normalized by dividing total biofilm by the cell density.

In spotty pellicle assay, 5 mL above diluted medium was added in each glass tube and incubated without shaking for 16 h at 30°C. For colonial morphology assay, the cultures were adjusted to an OD_{600} of 0.6, and collected by centrifuging at 12,000 rpm for 2 min. After removing the supernatant, bacteria were resuspended in 10 µL LB medium. Subsequently, 10 µL mixed medium is spotted on a trypticase soy sheep blood agar plate (Huankai Microbial, Guangdong, China) and cultivated statically at 30°C for 24 h, after which the colonial morphology was observed and photographed.

### Extracellular Polysaccharides Quantitative Assays

The modified phenol-sulfuric acid method was used to determine the polysaccharides production (Liu et al., 2021). In the beginning, 8 mg glucose was dissolved into the 100 mL volumetric flask, and then 0, 0.2, 0.4, 0.6, 0.8, and 1 mL glucose solution were added to each centrifuge tube, respectively. Double-distilled water was added to the centrifuge tube containing less than 1 mL glucose solution. Subsequently, 1 mL phenol solution and 5 mL concentrated sulfuric acid were added into each tube, after 30 min of reaction, the absorbance at 490 nm was respectively measured to draw the standard glucose curve. Subsequently, 1 mL phenol solution and 5 mL concentrated sulfuric acid were added to the tube, after 30 min of reaction, the absorbance at 490 nm was respectively measured to draw the standard glucose curve. Each overnight culture was adjusted to an OD_{600} of 0.6, diluted 1:100 into 50 mL LB medium and 6 mL broth was incubated to the 6-well polystyrene microplate statically at room temperature for 6 and 24 h, respectively. 5 mL bacterial culture was collected and centrifuged at 5,000 rpm and 4°C for 5 min, and 1 mL supernatant was collected and transferred into a new tube. Afterward, the supernatant, phenol solution (9%, v/v) and concentrated sulfuric acid were added to the tube sequentially in the ratio of 1:1.5 (v:v:v). After 30 min of reaction, the absorbance at 490 nm was measured, and the
FIGURE 1 | Normalized biofilm formation (total amount of biofilm/growth) of WT, ΔrelA, ΔrelAΔspoT, ΔrelA-pRelA, and ΔrelAΔspoT-pSpoT in 96-well polystyrene microtiter plates at different cell densities (A). The normalized biofilm differences between WT and ΔrelA, ΔrelAΔspoT, ΔrelA-pRelA, and ΔrelAΔspoT-pSpoT were shown at 6 (B) and 24 h (C), respectively. Three separate experiments were performed with biological triplicates each, and results were presented as mean ± SD (n = 3). * indicates p ≤ 0.05, and ** indicates p ≤ 0.01.

amount of polysaccharides was further calculated according to the above standard glucose curve.

Flagella Observation and Swarming Ability Assays
Vibrio containing Amp antibiotic (OD_{600} = 0.6) were fixed in 2.5% glutaric dialdehyde, and then bacterial flagella were observed by using the transmission electron microscope (TEM, JEM-2100, Japan). To evaluate swarming ability, fresh cultures of V. alginolyticus strains containing antibiotics (OD_{600} = 0.6) were centrifuged, precipitated, and resuspended in 10 µL 2216E medium, and then 1 µL culture was spotted on 0.9% (w/v) 2216E agar plates at room temperature.

RNA Extraction and Quantitative Reverse Transcription PCR Analysis
The WT and mutants were cultured in LB medium containing Amp overnight, and then diluted 1:100 into fresh medium and

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grown to the exponential phase (OD_{600} = 0.6) and stationary phase (OD_{600} = 1.2). RNA was extracted with a Bacteria Total RNA Extraction Kit (Promega, Madison, WI, United States), and then HiScript II Q RT SuperMix (+gDNA wiper) (Vazyme, Nanjing, China) was used for reverse transcription after determining the RNA quality by Nanophotometer NP80 (IMPLEN, München, Germany). The final cDNA samples were analyzed by Quantitative Reverse Transcription PCR (qRT-PCR) using ChamQ Universal SYBR qPCR Master Mix (Vazyme, Nanjing, China). Primers used in the qRT-PCR analysis are listed in Table 1. Expression of the genes encoding helix-turn-helix transcriptional regulator VpsT, polysaccharides biosynthesis protein VpsM and VpsH, chemotaxis protein CheR, flagellar hook protein FlgE, flagellar basal body L-ring protein FliD, and alternative sigma transcription factor RpoN was determined in triplicate. The 16S rRNA gene was used as an endogenous control for normalization of expression values. The 2^{ΔΔCt} method was used to quantify and compare each gene expression (Livak and Schmittgen, 2001).

Statistical Analysis

Three separate experiments were performed with biological triplicates each, and results were presented as mean ± standard deviation (SD; n = 3). Analysis of statistical differences was conducted on GraphPad Prism version 7.01 for Windows (GraphPad Software, SanDiego, CA, United States) through one-way analysis of variance (ANOVA) with Dunnett’s post-test. Differences were considered significant by calculated p value (*p ≤ 0.05, **p ≤ 0.01), and p value ≥ 0.05 was regarded as no significant.

RESULTS

The Lack of relA and spoT Affects Biofilm Formation at Different Cell Densities

The amount of biofilm formation of WT, ΔrelA, ΔrelAΔspoT, ΔrelAΔrelA, and ΔrelAΔspoTΔrelA was evaluated by staining with crystal violet. Afterward, biofilm formation was normalized by dividing total biofilm by the cell density (Supplementary Figure 1) to exclude growth effects (Figure 1A). Overall, the trend of biofilm formation in WT was to continuously increase at low cell density and then reduce at high cell density. The increasing amount of biofilm formation reached a peak at 6 h (OD_{570}/OD_{600} = 3.61), and kept a high level for a period of time until 10 h. Interestingly, the second small peak of biofilm formation appeared at 12 h (OD_{570}/OD_{600} = 1.04) and decreased again to a low level. The amount of biofilm formation in ΔrelA was significantly lower than that of the WT at low cell density (Figure 1B, p ≤ 0.01). However, it maintained a high level (OD_{570}/OD_{600} = 2.61–3.36) during 8–14 h and then began to decompose slowly until close to the level of WT (Figure 1C). In terms of the biofilm formation in ΔrelAΔspoT, its absorbance value was significantly lower than that of WT at low cell density (Figure 1B, p ≤ 0.05). Surprisingly, the amount of biofilm formation reached its peak at 8 h (OD_{570}/OD_{600} = 4.66), but the biofilm did not decompose within 36 h, even maintained at a high level (Figure 1C). The biofilm formation of ΔrelAΔrelA and ΔrelAΔspoTΔrelA were similar to WT and ΔrelA, respectively. An obvious crystal violet-stained circular ring appeared at the air-liquid interface (data not shown), suggesting that bacteria might tend to accumulate in oxygen-rich areas. However, there was additional biofilm staining at the bottom of the microtiter plates in ΔrelAΔspoT at high cell density (8–36 h), exhibiting robust bacterial self-aggregation ability. These results revealed that the deletion of two (p)ppGpp synthetase genes led to postponed biofilm formation at low cell density, and the absence of relA and spoT resulted in delayed and failed biofilm disassembly, respectively.

The Effects of relA and spoT on Spotty Pellicle and Colony Morphology

It has been reported that the formation of spotty pellicle strongly depends on oxygen level (Hare et al., 1981). After static incubation at 30°C for 16 h in the glass tube, the liquid solutions of WT, ΔrelA, and ΔrelAΔspoTΔrelA (OD_{600} = 0.8–1.0) were more turbid than that of ΔrelAΔspoT (OD_{600} = 0.6–0.8), without apparent bacterial flocculation and transparent air-liquid surface layer (Figure 2A). However, a spotty, unattached floating and thicker pellicle of ΔrelAΔspoT was observed at the air-liquid surface layer (Figure 2A). This result implied that spoT might participate in sensing oxygen and regulation of spotty pellicle.

A surprising but expected phenomenon was discovered in the hemolysis test when we determined whether the hemolytic ability of V. alginolyticus HN08155 could be affected by the absence of (p)ppGpp synthetase genes. In the beginning, there was no obvious difference in colony morphology among all strains (data not shown), whereas the colony morphology of ΔrelAΔspoT began to wrinkle gradually during cultivation. As
FIGURE 3 | The effects of relA and spoT on EPSs production. The content of EPSs was measured by phenol-sulfuric acid method in WT, ΔrelA, ΔrelAΔspoT, ΔrelA-pRelA, and ΔrelAΔspoT-pSpoT at 6 (A) and 24 h (B), respectively. Three separate experiments were performed with biological triplicates each, and results were presented as mean ± SD (n = 3). ** indicates p ≤ 0.01.

The Effects of relA and spoT on Exopolysaccharides Production at Low and High Cell Densities

Exopolysaccharides are an indispensable component of biofilm formation in V. alginolyticus (Chen et al., 2009). Based on our previous experiments, 6 and 24 h were chosen as the representatives of the low cell density and high cell density, respectively (Zhang et al., 2021). Our results indicated that the EPSs content of WT was the highest at low cell density (Figure 3A), but it gradually decreased with the disruption of (p)ppGpp synthetase genes, which was almost 3- and 5-fold of that in ΔrelA and ΔrelAΔspoT, respectively (Figure 3B). Complementation of the relA or spoT gene into mutant strains ΔrelA-pRelA and ΔrelAΔspoT-pSpoT respectively increased the synthesis of EPSs, although these contents were relatively lower in comparison with that in WT (p ≤ 0.01). At high cell density, WT had the lowest EPSs content, followed by ΔrelA, while ΔrelAΔspoT had the highest EPSs content. Notably, the content of EPSs in ΔrelAΔspoT was almost 1.5 times that of WT and ΔrelA. The EPSs content in ΔrelA was almost the same as that in WT (p ≥ 0.05), and supplementation of the spoT gene in ΔrelAΔspoT could reduce the secretion of EPSs to some extent. Overall, both relA and spoT favored the synthesis of EPSs at low cell density, and only spoT helped the degradation of it at high cell density.

The Effects of relA and spoT on Flagella Synthesis and Swarming Ability

Transmission electron microscope observation showed that V. alginolyticus HN08155 was short and rod-shaped, with a
single curved polar flagellum that was 2–3 times longer than that of the bacterium (Figure 4A). The flagellum of ΔrelA and ΔrelAΔspoT-pSpoT was identical to WT (Figures 4B,D), but the lack of spoT resulted in the loss of flagellum (Figure 4C). There was no doubt that spoT played a critical role in synthesizing flagellum. Previous study revealed that the swimming ability of ΔrelAΔspoT decreased sharply (Yin et al., 2021). Here we further investigated whether the absence of spoT would affect its swarming ability. After 12 h of continuous culture on 0.9% 2216E agar plates at room temperature, the swarming halo diameter of the WT, ΔrelA, and ΔrelAΔspoT-pSpoT averaged 6.80 ± 0.3, 7.65 ± 0.5, and 5.75 ± 0.4 mm, respectively, while those of ΔrelAΔspoT averaged 2.10 ± 0.8 mm only (Figures 5A,B). The diameters of the swarming rings of ΔrelAΔspoT were significantly smaller than that of the WT, ΔrelA and complemented strains (Figure 5B, p ≤ 0.01), demonstrating that the swimming ability of ΔrelAΔspoT was sharply diminished. Taken together, the disruption of spoT in V. alginolyticus not only negatively affected the production of flagellum, but also significantly impaired its swarming ability.

Deletion of relA and spoT Affects Expression Levels of Exopolysaccharides- and Motility-Related Genes

Although the content of extracellular polysaccharides can be determined by the phenol-sulfuric acid method by which the product concentration is qualified by observing the absorbance value (Liu et al., 2021). However, we could not rule out the interference of the residual nucleic acid and other sugars in the samples. Considering that vps-related genes affect the synthesis and transport of bacterial polysaccharides (Casper-Lindley and Yildiz, 2004; Chang et al., 2010), and cheR, fliG, and fliD are closely related to bacterial flagella synthesis and motility (Echazarreta and Klose, 2019; Liu et al., 2020), the expression levels of some functional genes were further
determined by qRT-PCR. As shown in Figures 6A–C, at low cell density, the expression level of vpsH in WT was 1.92- and 6.14-fold higher than that of ΔrelA and ΔrelAΔspoT, respectively. Conversely, at high cell density, the transcription of vpsH in ΔrelAΔspoT was 5- and 3.5-fold higher than that of WT and ΔrelA, respectively. Similarly, the expression levels of the other EPSs-related genes, vpsT and vpsM, in WT and ΔrelA showed a trend of first increasing and then decreasing, which was similar to the trend of biofilm formation and EPSs production in the culture medium under the same conditions. As for motility-related genes, the expression level of cheR and flgE in ΔrelA was significantly lower than that of WT, whereas the expression levels of flgH in ΔrelA were not significantly different from that of WT (Figure 6D). The transcriptional profiles of these motility-related genes in ΔrelAΔspoT exhibited lower levels compared with WT, which was consistent with the results of the swarming ability test. These results will further help us understand the effects of (p)ppGpp synthetase gene-mediated alteration in EPSs and motility on biofilms.

**Deficiency of relA and spoT Reduces the Expression of rpoN**

Previous study has shown that the deletion of rpoN leads to failure of biofilm detachment, loss of flagellum, reduction of spoty pellicle, transparent surface rugose colonies, and decreased swimming motility (Zhang et al., 2021). To explore the relationship between (p)ppGpp synthetase genes and rpoN, we measured the expression level of rpoN at low and high cell densities. At low cell density, the expression levels of rpoN in ΔrelA was significantly higher than that in WT, but that in ΔrelAΔspoT was significantly reduced by almost 3-fold compared with WT (Figure 7). At high cell density, the transcription levels of rpoN in ΔrelA and ΔrelAΔspoT was lower than that of WT even if the difference was
not significant. It was likely that (p)ppGpp synthetase genes had no obvious effect on rpoN at high cell density. These results implied that the deletion of (p)ppGpp synthetase genes had negative effects on the expression of rpoN at low cell density.

DISCUSSION

To date, many internal factors and external stimuli are known to regulate biofilm formation (Martin-Rodriguez and Romling, 2017; Ranieri et al., 2018; Bridges et al., 2020). Recently, the roles of nucleotide-based second messenger (p)ppGpp in biofilm regulation have attracted more and more attention (Díaz-Salazar et al., 2017; Colomer-Winter et al., 2019; Salzer et al., 2020). However, little information is available on the (p)ppGpp-mediated regulation of biofilm formation at different cell densities. In this study, we provide evidence by polystyrene plate assays that the (p)ppGpp synthetase genes, relA and spoT, favor biofilm formation at low cell density and biofilm detachment at high cell density, respectively, in V. alginolyticus.

The initial adhesion mediated by flagella is essential for biofilm formation (Utada et al., 2014). Our study indicated that the lack of (p)ppGpp synthetase gene spoT rather than relA led to loss of flagella in V. alginolyticus. Meanwhile, we found that the biofilm of WT and other strains formed more rapidly than that of ΔrelAΔspoT within 6 h, which may depend on the flagella-mediated aggregation. This result was similar to that of Vibrio cholerae O139 mutant strain, which was deficient in flagella synthesis and failed to form an obvious biofilm stained by crystal violet (Watnick et al., 2001).

Nevertheless, ΔrelAΔspoT could form biofilm, which might be due to other motility-related mechanisms such as type IV pili (Klausen et al., 2003). Flagella or other motility-related organelles also play an indispensable role in the biofilm detachment (Bridges et al., 2020). Interestingly, both ΔrelA and ΔrelAΔspoT showed a higher level of biofilm formation in comparison with WT between 8 and 18 h, then the biofilm in ΔrelA instead of in ΔrelAΔspoT began to decompose, implying that flagella in ΔrelA could play an important role in promoting biofilm detachment. Considering that bacterial motility, including swimming and swarming, is beneficial to the biofilm development (Bridges et al., 2020). We thus speculated the motility ability might be negatively affected in ΔrelAΔspoT. As expected, the swarming motility of WT and ΔrelA was not affected (p ≥ 0.05), while that of ΔrelAΔspoT significantly decreased relative to WT (p ≤ 0.01). qRT-PCR analysis further demonstrated that only the lack of spoT in V. alginolyticus significantly decreased the expression level of cheR, flgH, and flgE. In summary, the flagella synthesis and motility ability mediated by spoT would contribute to biofilm formation at low cell density and is necessary for the decomposition of biofilms at high cell density.

The most prevalent extracellular matrix component in biofilms is EPSs (Teschler et al., 2015). To better understand the mechanism of biofilm formation, we first measured the content of EPSs at low and high cell densities. Our results showed that the EPSs content of WT was higher than those of other strains at low cell density, whereas lower than those of ΔrelAΔspoT at high cell density. Both relA and spoT promoted EPSs production at low cell density and only spoT inhibited EPSs at high cell density. Similarly, the biofilm in WT was the highest at low cell density and the lowest at high cell density compared with that of other strains. Therefore, we propose that the content of EPSs is positively correlated with the amount of biofilm formation at low and high cell densities in V. alginolyticus. qRT-PCR analysis further showed that two (p)ppGpp synthetase genes relA and spoT increased the expression level of vpsT and vpsH at low cell density, whereas only spoT negatively regulated them at high cell density. Unlike vpsH and vpsT, only spoT contributed to the expression level of vpsM at low cell density, and two (p)ppGpp synthetase genes had negative effects on the expression of vpsM at high cell density. Our results indicated that (p)ppGpp synthetase genes could positively regulate transcription of some EPSs-related genes at low cell density, but negatively regulate expression level of them at high cell density. Besides, ΔrelAΔspoT was discovered to change the colony morphology from smoothness to rugosity, and form a spotty pellicle at the air-liquid interface. Yip et al. (2006) showed that induction of polysaccharide biosynthetic genes resulted in wrinkled colonies, pellicle formation and matrix production in Vibrio fischeri. We speculated that the higher expression levels of vpsT, vpsM, and vpsH in ΔrelAΔspoT than that in WT and ΔrelA were key elements to produce these biofilm phenotypes. In addition, the degradation of the biofilm in ΔrelA was delayed for a long time (8–18 h) might be explained, at least in part, by the higher expression of vpsM in ΔrelA than that in WT at high cell density.
Many studies have found that (p)ppGpp is closely related to the expression of rpoS and biofilm formation including in Pseudomonas aeruginosa and Pseudomonas putida KT2440 (van Delden et al., 2001; Liu et al., 2017). Previous study revealed that rpoN of V. alginolyticus played essential roles in controlling biofilm mediated by flagellum and EPSs (Zhang et al., 2021). In this study, we focused on the alteration of rpoN and rpoS expression in (p)ppGpp-deficient mutants. Although no obvious regular changes mediated by rpoS was observed (data not shown), the expression level of rpoN in ΔrelAΔspoT was lower than that in WT at low cell density, suggesting that (p)ppGpp synthetase genes could promote the biofilm formation by increasing the transcription level of rpoN. But unexpectedly, (p)ppGpp likely does not decompose the biofilm via rpoN-related pathways at high cell density (Figure 8). Lange et al. (1995) found that ppGpp played an important role in transcriptional elongation of rpoS. We speculated (p)ppGpp might have a similar effect on rpoN. In addition, the reason why ΔrelA had a higher gene expression level of rpoN than that of WT, but kept a notable decreased biofilm was that low concentration of (p)ppGpp might not be beneficial to the accumulation and stability of the alternative sigma factors in cell (Bougdour and Gottesman, 2007).

![Figure 8](image-url)
CONCLUSION

In summary, our results revealed the impact of (p)ppGpp-mediated regulation on biofilm, including biofilm formation in multi-well plates, the spotty pellicle at air-liquid interface and colony morphology. Combined with the results of cell motility, EPS production, and the expression of related functional genes, we found that (p)ppGpp synthetase genes may facilitate biofilm formation at low cell density and biofilm detachment at high cell density through flagella-mediated motility, EPSs production and rpoN-regulated pathways.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

WY and ZX designed the study and analyzed the data. WY, ZX, WR, HL, XZ, XC, and AH performed the experiments. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2022.858559/full#supplementary-material
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