IRON-DEPLETION PREVENTS BIOFILM FORMATION IN PSEUDOMONAS AERUGINOSA THROUGH TWITCHING MOTILITY AND QUORUM SENSING

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ABSTRACT

Influence of iron-depletion on twitching motility and quorum sensing (QS) system in P. aeruginosa was evaluated. The results demonstrated iron-depletion can retard biofilm formation and increase the twitching motility and expression of QS-related genes, suggesting a potential interaction between twitching motility and QS system in P. aeruginosa biofilm formation.

Key words: iron; biofilm; twitching motility; quorum sensing

Pseudomonas aeruginosa is one of the major causes of nosocomial infections (8). In addition, P. aeruginosa is a leading pathogen among patients with cystic fibrosis, diffuse panbronchiolitis, and chronic obstructive pulmonary disease (13, 22). In patients with these underlying diseases, it can cause chronic infections characterized by the formation of biofilms. Biofilms are composed of a group of bacteria attached to the surfaces and are encased in a hydrated polymeric matrix (4). Bacteria that form biofilms can withstand host immune responses and are much more resistant to antibiotics than their counterparts of nonattached, planktonic bacteria (20). Therefore, infections with biofilm-forming bacteria are persistent and difficult to treat with antibiotics.

P. aeruginosa possesses polar filaments called type IV pili that are involved in attachment and surface translocation by twitching motility (15). Twitching motility, mediated by pilus extension and contraction, is required for the formation of biofilms in P. aeruginosa (10, 21). Quorum-sensing (QS) system in bacteria detects cell density and regulates the expression of many genes at high cell densities. Many studies demonstrated that P. aeruginosa strains lacking functional QS system are less virulent than wild-type strains and form flat, undifferentiated biofilms that are less stable than the differentiated biofilms formed by wild-type P. aeruginosa. As a consequence, QS has been suggested as a potential target for new preventive and/or therapeutic strategies of P. aeruginosa infections (6).

Iron is essential for most pathogens because iron is an indispensable component of many proteins, especially some enzymes in bacteria. Therefore, iron acquisition from environment is important for the growth and metabolism of P. aeruginosa (5). Recently, many studies revealed that iron also play an important role in biofilm formation (1, 23). In vitro experiments showed both iron-depletion (<1µM) and iron-repletion (>100µM) retarded biofilm formation (14). Furthermore, some reports showed that the level of free iron is increased in airway secretions of cystic fibrosis patients, and this might be one of the possible reasons for the frequent identification of biofilms in the lungs of these patients (3).

To further determine the role of iron in P. aeruginosa
biofilm formation, we compared the ability of biofilm formation between type IV pilus mutant, QS mutant and wild-type strains under the iron-depletion condition. We select iron-depletion condition because the concentration of free iron in vivo would not exceed 100µM and thus the results are more relevant for control of biofilm infection.

*P. aeruginosa* wild-type strain PAO1, *P. aeruginosa* IV pilus mutant strain PAO-ΔpiliHIJK and *P. aeruginosa* lasR and rhlR mutant strain PAO-JP2 were used in this study. Iron-depletion condition in the medium was achieved by addition of iron-specific chelator 2, 2-dipyridyl (500µM) (DPD, Sigma, St. Louis, MO).

Static biofilm experiment was performed as described by Favre-Bonté *et al.* with some modifications (7). Briefly, PAO1, PAO-ΔpiliHIJK and PAO-JP2 were grown in M-H broth for 6 h with agitation at 37°C. Silica gel chips were immersed in the bacterial culture medium without agitation to allow bacteria to adhere and form biofilm. The size of all the chips were 1 cm² and culture medium were replaced every 48 h for 7 days of continuous cultivation. Silver staining method is that hydrated polymeric matrix of biofilm can be stained as black by AgNO₃ and the biofilm formation can be quantified by gray scale comparison (16). The gray scale of the chips was quantified by using Image pro plus (version 4.01, Media Cybernetics) system.

For twitching motility assay we used Petri dishes with LB agar (Sigma). DPD and FeCl₃ (Sigma) were added to the molten agar and the plates were dried overnight at room temperature. PAO1, PAO-ΔpiliHIJK and PAO-JP2 were point inoculated at the bottom of the agar plate. After 48 h, the twitching motility distance along the plastic–agar interface (at the bottom of the agar plate) was measured (2).

Total RNA from PAO1 and PAO-ΔpiliHIJK were isolated from bacterial pellet using a commercial kit (Trizol, Invitrogen, Carlsbad, CA) according to the manufacturer’s instructions. Reverse transcriptions were performed using 2µl of random hexamer primers, 2µl of 10mM dNTPmix 1µl of M-MLV (Promega, Madison, WI) and 1 µl of RNA in a total volume of 30 µl. Reverse transcription reaction was stopped after incubation at 37 °C for 60 min. Real-time PCR analyses were carried out by using the ABI Prism 7900HT SDS 2.2 software (ABI). The sequences of the genes were obtained from the gene bank of the National Library of Medicine and the primers were designed by using Primer Express software (version 2.0, ABI). The sequences of the primers: lasR (F: 5’-CTG CTC GAG CCG GCC TCG GCC TGT TCT-3’ R: 5’-CGG GAT CCG GAT GGC GCT CCA CTC CA-3’), rhlR (F: 5’-CAT GCG CGA GCA GGA GTT GCb-3’, R: 5’-TAG GGA TCC TAA TCG AAG CCC AGG CGC-3’), 16sRNA (F: 5’-GAT GAC GGT ACC GGA AGA ATA AGC-3’, R: 5’-CCA TGT CAA GGG TAG GTA AGG TTT-3’). Real-time PCR contained 2 µl of cDNA, 25µl of SYBR Green I (modified DNA polymerase, SYBR Green I, Optimized PCR buffer, 5mM MgCl₂, Dntp MIX including dUTP) and 1 µl of forward and reverse primers. Before amplification, PCR mixtures were heated at 50°C for 2 min to prevent carryover of PCR products and then to 95°C for 10 min to denature nucleic acids followed by 40 cycles of 95°C for 15 s and 60°C for 1 min. The mRNA level of lasR and rhlR genes in each strain was normalized on the basis of the 16sRNA levels.

Pictures of silica gel chips after silver staining were analyzed by the computer image analysis system (Figure 1). Under normal conditions, wild-type and PAO-ΔpiliHIJK formed similar level of biofilm. However, PAO-JP2 strain formed less biofilms. These results indicated that QS is required for the biofilm formation while pili are not required for biofilm formation under normal condition. Under the iron-depletion condition, biofilm formation of all the 3 strains decreased significantly and addition of iron increased biofilm formation although it is still lower than that under normal condition. These results demonstrated that iron is required for biofilm formation in *P. aeruginosa*.

Under normal condition, wild-type strain showed highest twitching motility compared with PAO-JP2 and PAO-ΔpiliHIJK strains suggesting that pili and QS are involved in twitching motility. Under iron-depletion condition, twitching motility of PAO1 and PAO-JP2 increased significantly and addition of iron decreased the ability of twitching motility. In
Biofilm formation by *P. aeruginosa*

contrast, twitching motility for PAO-ΔpilHIJK strain did not change significantly under iron-depletion or iron-repletion conditions (data not shown) suggesting that pili are required for the increased twitching motility under iron-depletion condition.

Previous studies suggested that twitching motility is necessary for microcolony formation and thus essential for normal biofilm development (10, 15, 21). However, our study showed that IV pili mutant strain PAO-ΔpilHIJK formed almost the same amount of biofilm compared to the wild-type strain, PAO1. The most distinct difference is that those studies focused on the initial stage of biofilm development (about 8–24 h), while we studied the mature biofilm formation (7 d). This may indicate that type IV pilus-mediated twitching motility mediates aggregation of cells in the initial stage of biofilm development, however given enough time (7 d in this study), twitching motility is not required for the formation of mature biofilm. Heydorn *et al.* (9) also reported that biofilm formed by the PAO-ΔpilHIJK strain contained a large number of distinct microcolonies that were regularly spaced and almost uniform in size. As the biofilms growing, the microcolonies of the PAO-ΔpilHIJK strain coalesced, leading to the formation of larger and larger microcolonies. They suggested that microcolonies formed by the PAO-ΔpilHIJK strain was due to the clonal growth of an individual bacterium in the biofilm because these bacteria were unable to move by twitching motility. Another study also reported that mutant strains defective in twitching motility were not impaired in the early stages of biofilm development and eventually formed mature biofilms, although the morphology were distinguishable from those of wild-type strains (12). Therefore, we hypothesize that another pathway for the formation of microcolonies might exist, and further studies on how PAO-ΔpilHIJK strain form a mature biofilm are necessary to test this hypothesis.

Biofilm formation by PAO1, PAO-ΔpilHIJK and PAO-JP2 strains were significantly decreased under iron-depletion condition compared with that under normal condition. We also showed that twitching motility of PAO1 and PAO-JP2 increased under iron-depletion condition. Singh *et al.* (19) showed that lactoferrin acted as a chelator to bind iron, leading to the twitching and wandering of bacteria across the surface instead of forming cell clusters and biofilms. These findings suggested that iron plays an important role at a critical juncture biofilm development in which bacteria stop roaming as individuals and aggregate into durable communities. Interestingly, as mentioned before, twitching motility is crucial in biofilm formation; however, too much twitching motility prevents biofilm development.

PAO-ΔpilHIJK strain lacks IV pili and the results showed that twitching motility did not change significantly under iron-depletion or iron-repletion conditions. Thus decrease of biofilm formation of PAO-ΔpilHIJK strain is not due to the decrease in twitching motility. We then determined whether iron-depletion has an effect on QS system, another important system involved in biofilm formation. We showed that upon iron depletion, expression of QS-related genes, *lasR* and *rhlR* were significantly increased consistent with the results obtained in other studies (5, 11, 23). One of the contradictory results is that iron-depletion decreased biofilm formation (Figure 1);

**Figure 1.** Comparison of biofilm formation under normal (without DPD), iron-depletion (with 500μM DPD) and partial iron-depletion (with 500μM DPD and 25μM FeCl₃) conditions. The bars represent the means for the 3 tests, and the error bars represent the standard errors of the means. One-factor analysis of variance was conducted. * means $P<0.05$, **means $P<0.01$, both as compared with the same strain in normal condition; # means $P<0.05$, as compared with PAO1 in normal condition.
however, it activated the expression of lasR and rhlR genes (Figure 2) suggesting an inhibitory role of QS on biofilm development. This could be explained by the results obtained by Patriquin et al. (17) showing that RhlIR/C4-HSL was required for low-Fe-stimulated twitching because rhlIR mutant strain was twitching defective on Fe-limited minimal medium and addition of C4-HSL restores twitching of the rhlI mutant strain. Therefore, under iron-depletion condition, increased rhlIR expression and concomitant C4-HSL production might upregulate a gene(s) leading to the upregulation of twitching and downregulation of biofilm formation.

Expression of QS-related genes, lasR and rhlR of PAO-DapiHIJK does not change significantly under the iron-depletion condition. However, in PAO1 strain, iron-depletion significantly enhanced the expression of these two genes (Figure 2). These results indicated that there is correlation between the expression of IV pili and QS-related genes. Pili are encoded by a gene cluster, pilGHIJK that may contain certain gene(s) responsible for the regulation of lasR or rhlR. Reid et al. (18) showed that response of clinical isolates to alterations of environmental iron is quite different to that of PAO1. Thus, we speculate that changes in genome might alter the response of QS to environmental factors.

Based on the results from this study, we concluded that twitching motility is not a prerequisite for the formation of mature biofilm and iron-depletion can decrease the formation of biofilm at least through both twitching motility and QS. The results also indicated there might be a correlation between QS and twitching motility. Further studies on the coordination and regulation of the two key pathways will enhance our understanding of P. aeruginosa biofilm formation and identify new targets to control biofilm formation.

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