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Aminoglycoside resistance of *Pseudomonas aeruginosa* biofilms modulated by extracellular polysaccharide

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Summary

Pseudomonas aeruginosa is an opportunistic pathogen that produces sessile communities known as biofilms that are highly resistant to antibiotic treatment. Limited information is available on the exact role of various components of the matrix in biofilm-associated antibiotic resistance. Here we show that the presence of extracellular polysaccharide reduced the extent of biofilm-associated antibiotic resistance for one class of antibiotics. Minimal bactericidal concentration (MBC) for planktonic and biofilm cells of *P. aeruginosa* PA14 was measured using a 96 well microtiter plate assay. The MBC of biofilm-grown $\Delta pelA$ mutant, which does not produce the Pel polysaccharide, was 4-fold higher for tobramycin and gentamicin, and unchanged for $\Delta bifA$ mutant, which overproduces Pel, when compared to the wild type. Biofilms of *pelA* mutants in two clinical isolates of *P. aeruginosa* showed 4- and 8-fold higher MBC for tobramycin as compared to wild type. There was no difference in the biofilm resistance of any of these strains when tested with fluoroquinolones. This work forms a basis for future studies revealing the mechanisms of biofilm-associated antibiotic resistance to aminoglycoside antibiotics by *P. aeruginosa*.

Keywords

Pseudomonas aeruginosa; exopolysaccharides; biofilms; aminoglycosides; minimal bactericidal concentration

Introduction

Pseudomonas aeruginosa is a ubiquitous gram-negative bacterium readily isolated from water and soil environments. It is an opportunistic human pathogen frequently causing septicemia in burn patients, chronic infections in individuals with cystic fibrosis, and urinary tract infections in catheterized patients [7,9]. This organism can adopt a planktonic (free swimming) or a biofilm mode of life. In response to environmental cues, planktonic cells of *P. aeruginosa* adhere to surfaces and adopt a sessile existence. Sessile cells proliferate on surfaces, eventually differentiating to form structured communities called biofilms that are encased in a self-produced matrix comprised of a variety of macromolecules, including exopolysaccharides (EPS) [16].

Two different loci that contribute to the EPS component of the matrix in non-mucoid *P. aeruginosa* strains have been identified. The *pel* locus (referring to *pel*licle, a biofilm formed

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at the air-medium interface), containing the genes *pelA-G*, is responsible for synthesis of a glucose-rich component of the matrix, whereas *psl* (polysaccharide synthesis locus), containing the *pslA-O* genes, is responsible for a mannose and galactose rich EPS [4–6,10,12]. It has been suggested that exopolysaccharide matrix that surrounds the cells in the biofilm prevents diffusion of antimicrobial agents to the microbes. While this reduced diffusion may be the case for some antimicrobial agents, for others it has been shown that antimicrobial agents can penetrate the matrix but still cannot kill the cells in the biofilm [1,19]. In this report we address the role of the Pel EPS in biofilm-associated antibiotic resistance of *P. aeruginosa* PA14 grown in a microtiter well system.

Materials and methods

Bacterial strains, plasmids and media

Strains, plasmids and primers used in the study are listed in Table 1. *P. aeruginosa* PA14 and *Escherichia coli* were cultured on lysogeny broth (LB) medium [2] or minimal M63 salts supplemented with arginine (0.4%) and MgSO₄ (1 mM) [11].

Molecular techniques

PCR amplification for 16SrRNA gene was carried out for the two clinical isolates, isolated from an infected wound from a tertiary care hospital of Pakistan, using primers listed in Table 1. The resulting sequences were analyzed using the BLAST software and percentage similarity to other sequences was determined. Presence of the *pel* genes was assayed in the clinical isolates by PCR using primers listed in Table 1. The *pelA* mutation in two clinical isolates was generated by a single-crossover (SCO) insertion using the pMQ89 suicide vector as described by Kuchma et al. [8].

Antibiotic resistance assay

Ninety-six-well microtiter plate-based assay was used to determine biofilm-specific antibiotic resistance as described by Mah et al. [11], with slight modification. The minimal bactericidal concentrations for biofilm grown cells (MBC-B) was determined by exposing 24 h-old biofilms to various concentrations of antibiotic for 24 h, followed by a single wash of biofilms with sterile medium to remove the antibiotics. Bacteria that survived antibiotic treatment were allowed to grow for 24 h in fresh medium lacking antibiotics, and these bacteria subsequently were transferred to LB agar plate using a multipronged device to assess growth. The MBC of planktonic bacteria (MBC-P) was determined by adding antibiotic to bacteria at the time of inoculation, incubating for 24 h followed by plating for viability. Both planktonic and biofilm populations were comprised of ca. 1×10^7 colony forming units (CFU)/well.

Biofilm formation assay and assessment of EPS production

Biofilm formation was measured using a microtiter dish system described previously [3,15]. Congo red assays were performed as reported [3,5,6]. Aliquotes of 2.5 µl of LB grown cultures were spotted onto the plates for 24 h at 37°C followed by 3–4 days of incubation at room temperature.

Results and Discussion

Loss of Pel EPS in lab strains does not lower biofilm antibiotic resistance

For *P. aeruginosa* PA14 (referred to as PA14), the primary polysaccharide component of the biofilm matrix *in vitro* is synthesized by the enzymes encoded by the *pel* locus. The Pel EPS is the cellular component bound by the dye Congo Red (CR) [4,5]. Previous studies have

shown that strains of PA14 with mutations in *sadCh*, *pelA* and *roeA* genes, as well as a double *bifAroeA* mutant (Table 1) show a decrease in Pel-dependent polysaccharide production compared to the wild type [13,14], while the Δ *bifA* mutant shows increased *pel*-dependent polysaccharide synthesis [8]. We tested biofilms formed by these mutants and the wild type for their resistance to two different classes of antibiotics using a 96 well dish biofilm assay.

The MBC of biofilm (MBC-B)- and planktonically (MBC-P)-grown wild type and mutants was determined for gentamicin (an aminoglycoside antibiotic) and ciprofloxacin (a fluoroquinolone antibiotic). No difference in the planktonic MBC was observed between wild type and the mutant strains for either of these antibiotics (Tables 2 and 3).

Wild type biofilms showed a 4-fold higher MBC for gentamicin when compared to planktonic cells. The biofilm formed by the Δ *bifA* mutant had the same MBC for gentamicin as that of wild type, indicating that increased *pel*-polysaccharide production in Δ *bifA* does not result in an increase in resistance (Table 2, left). Although Pel polysaccharide produced by the Δ *pelA*, Δ *sadC*, Δ *bifA* Δ *roeA* and Δ *roeA* mutants was reduced as compared to wild type, the MBC of these strains increased by 4-fold, 2-fold, 8-fold and 2-fold, respectively, for gentamicin.

For ciprofloxacin, the MBC-B was 8-fold higher for wild type as compared to planktonic cells. In contrast to the studies using gentamicin, all the mutant strains tested for ciprofloxacin showed the same MBC-B as that of wild type (Table 3, left), indicating that the presence or absence of Pel polysaccharide does not affect the resistance to ciprofloxacin.

Loss of Pel EPS results in increased MBC for aminoglycoside antibiotics in biofilms

To test whether loss of Pel EPS has the same role in biofilm antibiotic resistance for other aminoglycoside and fluoroquinolone antibiotics, as described above, we tested tobramycin (an aminoglycoside) and ofloxacin (a fluoroquinolone). There was no difference in MBC-P of wild type and the mutant strains when tested against tobramycin and ofloxacin (Table 2 and 3, right).

A 4-fold higher MBC was observed for wild type biofilm as compared to planktonic cells when tested for tobramycin. As observed for the other aminoglycoside (gentamicin), the MBC of biofilm-grown Δ *bifA* was the same as that of wild type when tested for tobramycin. This confirmed that increased production of Pel polysaccharide does not result in an increase in antibiotic resistance to aminoglycosides. The biofilm formed by the Δ *pelA*, Δ *sadC*, Δ *bifA* Δ *roeA* and Δ *roeA* mutants, all of which have lower Pel polysaccharide production than the wild type, showed an increase in MBC for tobramycin as compared to wild type (Table 2). Like ciprofloxacin, the MBC-B for ofloxacin was the same for the wild type and all mutants tested (Table 3). These data suggest that at least for aminoglycoside antibiotics, loss of Pel polysaccharide enhanced biofilm resistance of *P. aeruginosa*.

Clinical isolates defective for Pel polysaccharide production display the same biofilm antibiotic resistance patterns as the PA14 lab strain

In order to assess whether biofilm antibiotic resistance patterns observed in lab strains was also observed in clinical isolates, we repeated the assays described above with two clinical, non-mucoid infected wound isolates of *P. aeruginosa*, designated SMC4685 and SMC4686. 16SrRNA sequence analysis was used to confirm that these isolates were *P. aeruginosa* using primers listed in Table 1. C>99% similarity to other well characterized strains of *P. aeruginosa* in Genbank. The *pelA* mutation was generated by a single-crossover (SCO) insertion in the two clinical isolates. The SCO insertion was confirmed by PCR, and these mutants showed a decrease in CR binding and biofilm formation (Fig. 1).

The MBC for planktonic and biofilm cells were measured for the clinical isolates and for their *pelA* mutants for ciprofloxacin, ofloxacin and tobramycin. Since the single crossover insertion was generated using the suicide vector pMQ89, which carries a gentamicin resistance marker, we could not determine the MBC for gentamicin in the mutants of the clinical isolates. For all the antibiotics tested, there was no difference in the planktonic MBC of the *pelA* mutant and their respective wild type for any of the antibiotics (Tables 2 and 3). Biofilms of the two clinical isolates SMC4685 and SMC4686, respectively, showed a 4-fold and 8-fold higher MBC for tobramycin as compared to planktonically grown cells. The MBC of biofilm grown derivatives of the clinical isolates, SMC4685 *pelA*::SCO and SMC4686 *pelA*::SCO, was 8-fold and 4-fold higher for tobramycin when compared to their respective wild type parent.

For these two clinical isolates, loss of Pel polysaccharide production resulted in only a modest decrease in biofilm formation in the microtiter dish assay, which was not statistically significant (Fig. 1B). Thus, these strains showed increased biofilm resistance despite a small, non-significant decrease in total biofilm formation. Like the PA14 lab strain, both the clinical isolates showed no difference in the MBC-B of *pelA* mutants and their respective wild type when tested for fluoroquinolone resistance.

In conclusion, here we present evidence that presence of Pel polysaccharide is not important for biofilm antibiotic resistance of *P. aeruginosa* under the conditions tested in this report, that is, in biofilms grown in microtiter dishes. Our data show that biofilm antibiotic resistance increases for aminoglycosides when the Pel polysaccharide is reduced in both lab and clinical isolates of *P. aeruginosa*. Furthermore, the degree of increased biofilm resistance measured in these Pel-deficient mutants is likely somewhat of an under-estimation given that these mutants typically show some decrease in biofilm formation [13,14]. It is possible that the *pelA* mutants produce a second EPS molecule, which might help explain our findings. Alternatively, it is possible that decreasing *pel*-dependent polysaccharide might result in a compensatory up-regulation of other factors that either directly or indirectly cause the observed resistance. A more detailed study of the role of Pel polysaccharide in biofilm antibiotic resistance will be necessary to address questions arising from this work.

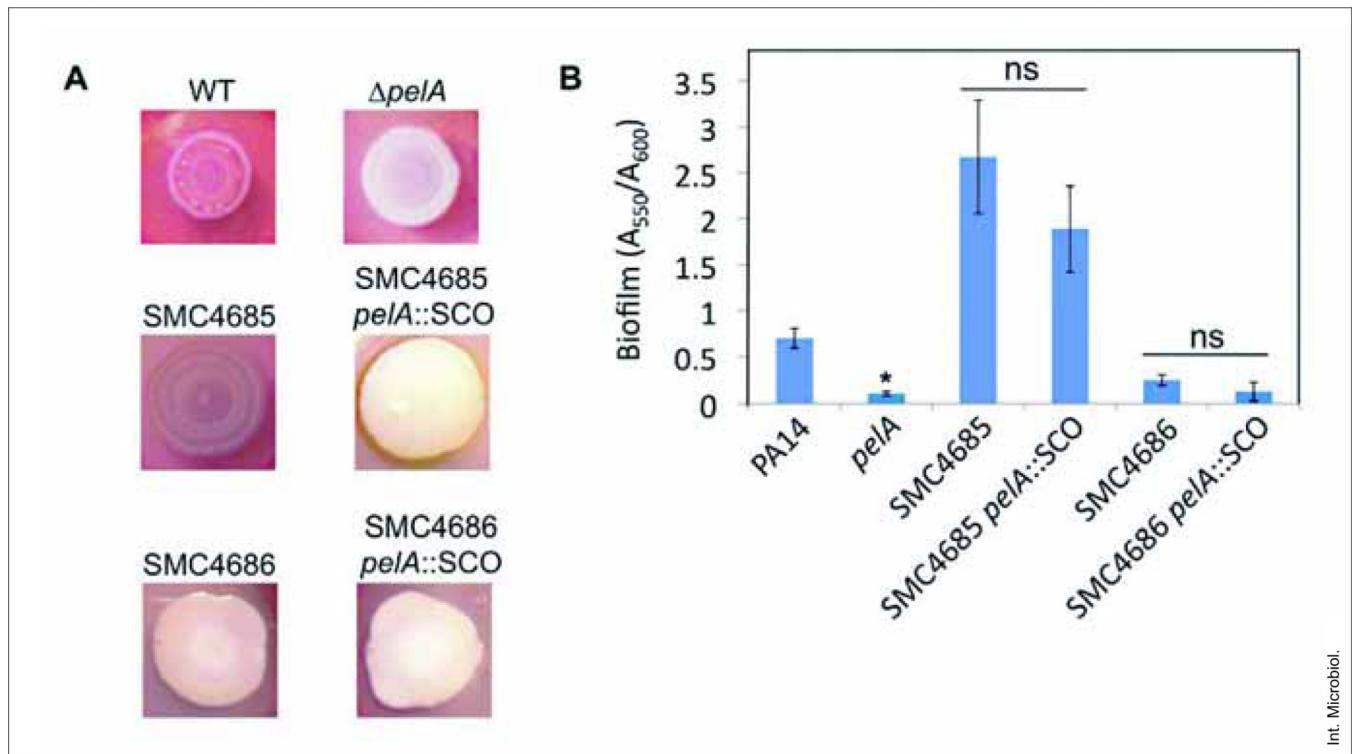
Acknowledgments

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**Fig. 1.**

Requirement of *pelA* for CR binding and biofilm phenotype of wild type. (A)

Representative images of CR binding by the wild type, $\Delta pelA$, clinical wound isolates SMC4685 and SMC4686 and their *pelA::SCO* derivatives. Plates were incubated for 24 h at 37°C, followed by 4 days at room temperature. (B) Quantification of CV-stained biofilms (A_{550}) normalized to their OD (A_{600}). Strains were grown as described in Methods for 24 h prior to CV staining. Asterisks (*): significantly different from the wild type PA14 strain (Student's t test, $P < 0.05$); ns, difference not significant from corresponding parental strain ($P > 0.05$). The average A_{600} value (\pm SD) for each of the strains from three independent experiments, which serves as a measure of planktonic growth, is as follows: PA14, 0.223 (\pm 0.024); *pelA*, 0.281 (\pm 0.045); SMC4685, 0.214 (\pm 0.020); SMC4685 *pelA::SCO*, 0.208 (\pm 0.044); SMC4686, 0.204 (\pm 0.037); SMC4686 *pelA::SCO*, 0.198 (\pm 0.077). There is no significant difference in planktonic growth between the wild type and their respective mutants for any of these strains ($P > 0.05$).

Table 1

Strains, plasmids and primers used in this study

Strain, plasmid, or primer	Relevant genotype or sequence (5'-3')	Source or reference
<i>Pseudomonas aeruginosa</i> strains		
PA14	WT	[17]
SMC3335	PA14Δ <i>sadC</i>	[13]
SMC2893	PA14Δ <i>pelA</i>	[4,5]
SMC3351	PA14Δ <i>bifA</i>	[8]
SMC3812	PA14Δ <i>roeA</i>	[14]
SMC3814	PA14Δ <i>bifA</i> Δ <i>roeA</i>	[14]
SMC4685	Clinical isolate #99 from infected wound	This study
SMC4687	SMC4685 <i>pelA</i> ::pMQ89	This study
SMC4686	Clinical isolate #160 from infected wound	This study
SMC4688	SMC4686 <i>pelA</i> ::pMQ89	This study
Plasmids		
pMQ89	Suicide cloning vector	[18]
Primers		
<i>pelA</i> -F	GCTACGTGCCGTTTCGTCAGCA	
<i>pelA</i> -R	CAGGCCCGCAGGTAGACGTG	
<i>pelG</i> -F	TATTGCTGGCGACCCTGTTCGATG	
<i>pelG</i> -R	ATGAAACGCAGCAGGTAGGCACAG	
16SrRNA-F	AGAGTTTGATCCTGGCTCAG	
16SrRNA-R	ACGGGCGGTGTGTRC	

Table 2
Resistance of planktonically and biofilm-grown strains to aminoglycoside antibiotics

Strains	Aminoglycosides							
	Gentamicin			Tobramycin				
	MBC-P ^a	Fold Change	MBC-B ^a	Fold Change	MBC-P ^a	Fold Change	MBC-B ^a	Fold Change
PA14	39 ^b	-	156 ^b	-	19.5 ^b	-	78 ^b	-
$\Delta pelA$	39 ^b	1X ^c	625 ^b	4X ^c	19.5 ^b	1X ^c	312.5 ^b	4X ^c
$\Delta bifA$	39 ^b	1X ^c	156 ^b	1X ^c	19.5 ^b	1X ^c	78 ^b	1X ^c
$\Delta bifA \Delta roeA$	39 ^b	1X ^c	1250 ^b	8X ^c	19.5 ^b	1X ^c	625 ^b	8X ^c
$\Delta sadC$	39 ^b	1X ^c	312.5 ^b	2X ^c	19.5 ^b	1X ^c	625 ^b	8X ^c
$\Delta roeA$	39 ^b	1X ^c	312.5 ^b	2X ^c	19.5 ^b	1X ^c	625 ^b	8X ^c
SMC4685 ^d	39 ^b	-	312.5 ^b	-	39 ^b	-	156 ^b	-
SMC4685 <i>pelA</i> ::SCO	ND ^e	ND	ND	ND	39 ^b	1X ^c	1250 ^b	8X ^c
SMC4686 ^d	39 ^b	-	312.5 ^b	-	39 ^b	-	312.5 ^b	-
SMC4686 <i>pelA</i> ::SCO	ND	ND	ND	ND	39 ^b	1X ^c	1250 ^b	4X ^c

^aMBC-P ($\mu\text{g/ml}$), MBC of planktonically grown cells; MBC-B ($\mu\text{g/ml}$), MBC of biofilm-grown cells.

^bValues are from at least three separate experiments.

^cFold-change compared to WT grown under the same conditions.

^dClinical, non-mucoid *P. aeruginosa* isolate.

^eND, not determined. As outlined in the text, the construct used to make this mutation confers gentamicin resistance and thus we could not determine biofilm resistance to this antibiotic for these mutants.

Table 3

Resistance of planktonically and biofilm-grown strains to fluoroquinolones antibiotics

Strains	Ciprofloxacin				Ofloxacin			
	MBC-P ^a	Fold Change	MBC-B ^a	Fold Change	MBC-P ^a	Fold Change	MBC-B ^a	Fold Change
PA14	3.9 ^b	-	31.25 ^b	-	62.5 ^b	-	250 ^b	-
$\Delta pelA$	3.9 ^b	1X ^c	31.25 ^b	1X ^c	62.5 ^b	1X ^c	250 ^b	1X ^c
$\Delta bifA$	3.9 ^b	1X ^c	31.25 ^b	1X ^c	62.5 ^b	1X ^c	250 ^b	1X ^c
$\Delta bifA \Delta roeA$	3.9 ^b	1X ^c	31.25 ^b	1X ^c	62.5 ^b	1X ^c	250 ^b	1X ^c
$\Delta sadC$	3.9 ^b	1X ^c	31.25 ^b	1X ^c	62.5 ^b	1X ^c	250 ^b	1X ^c
$\Delta roeA$	3.9 ^b	1X ^c	31.25 ^b	1X ^c	62.5 ^b	1X ^c	250 ^b	1X ^c
SMC4685 ^d	0.9 ^b	-	31.25 ^b	-	7.8 ^b	-	62 ^b	-
SMC4685 <i>pelA</i> ::SCO	0.9 ^b	1X ^c	31.25 ^b	1X ^c	7.8 ^b	1X ^c	62 ^b	1X ^c
SMC4686 ^d	1.8 ^b	-	31.25 ^b	-	7.8 ^b	-	250 ^b	-
SMC4686 <i>pelA</i> ::SCO	1.8 ^b	1X ^c	31.25 ^b	1X ^c	7.8 ^b	1X ^c	250 ^b	1X ^c

^aMBC-P (μ g/ml), MBC of planktonically grown cells; MBC-B (μ g/ml), MBC of biofilm-grown cells.^bValues are based from results from at least three separate experiments.^cFold-change compared to WT grown under the same conditions.^dClinical, non-mucoid *P. aeruginosa* isolate.