RESEARCH ARTICLE

Type IV pili promote early biofilm formation by *Clostridium difficile*

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One sentence summary: We demonstrate that type IV pili play an important role in the initial stages of *Clostridium difficile* biofilm formation, which has direct implications for future research into novel CDI therapies.

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ABSTRACT

Increasing morbidity and mortality from *Clostridium difficile* infection (CDI) present an enormous challenge to healthcare systems. *Clostridium difficile* express type IV pili (T4P), but their function remains unclear. Many chronic and recurrent bacterial infections result from biofilms, surface-associated bacterial communities embedded in an extracellular matrix. CDI may be biofilm mediated; T4P are important for biofilm formation in a number of organisms. We evaluate the role of T4P in *C. difficile* biofilm formation using RNA sequencing, mutagenesis and complementation of the gene encoding the major pilin *pilA1*, and microscopy. RNA sequencing demonstrates that, in comparison to other growth phenotypes, *C. difficile* growing in a biofilm has a distinct RNA expression profile, with significant differences in T4P gene expression. Microscopy of T4P-expressing and T4P-deficient strains suggests that T4P play an important role in early biofilm formation. A non-piliated *pilA1* mutant forms an initial biofilm of significantly reduced mass and thickness in comparison to the wild type. Complementation of the *pilA1* mutant strain leads to formation of a biofilm which resembles the wild-type biofilm. These findings suggest that T4P play an important role in early biofilm formation. Novel strategies for confronting biofilm infections are emerging; our data suggest that similar strategies should be investigated in CDI.

Keywords: *Clostridium difficile*; type IV pili; biofilm

INTRODUCTION

Clostridium difficile is a Gram-positive spore-forming obligate anaerobe, first isolated from the stool of newborn infants in 1935 (Hall and O'Toole 1935). *Clostridium difficile* infection (CDI) is the leading cause of infectious hospital-acquired gastrointestinal illness in the developed world (Musher *et al.* 2006; McFarland 2009). The disease is toxin mediated, ranging in severity from mild diarrhea to toxic megacolon, and can result in death (Bartlett and Gerding 2008). The frequency and severity of CDI are increasing rapidly in the United States and globally (Kelly and LaMont 2008); a recent report estimates that in 2011 CDI was responsible for almost half a million infections and approximately 29 000 deaths in the United States (Lessa *et al.* 2015). Improved methods for preventing and treating CDI will be crucial for reducing morbidity and mortality from this disease.

Bacteria commonly grow as biofilms, surface-associated communities embedded in a polysaccharide-rich substance; this mode of growth is critical to chronic and recurrent infections. Similarities between biofilm infections and CDI suggest that CDI is biofilm mediated. Both biofilm infections and CDIs are surface associated: while biofilms form on surfaces varied as prosthetic implants and the interiors of pipes, *C. difficile* infects the colon surface. Like many bacteria known to form biofilms, *C. difficile* employs quorum sensing as a means of cell–cell communication (Davies *et al.* 1998; Carter *et al.* 2005; Lee and Song 2005; Yang *et al.* 2014; Omer Bendori *et al.* 2015). Finally, biofilms are well known to allow their constituents to resist antibiotic treatment (Olsen 2015), and at times CDI does not respond to, or recurs after, appropriate antibiotic therapy. Although the precise mechanisms of CDI treatment resistance and infection relapse are currently unknown, they may be related to the formation of a *C. difficile* biofilm. Cells growing in a biofilm have a different phenotype compared to their planktonic counterparts. Some cells deep within a biofilm are relatively nutrient deprived and therefore slow or non-growing, which reduces susceptibility to antimicrobials (Becker *et al.* 2001). Determination of the phenotype of *C. difficile* in CDI may allow for better-targeted therapies. For example, compounds that disrupt the extracellular matrix of biofilms have demonstrated feasibility for reducing bacterial biofilms produced by clinically relevant organisms (Lu and Collins 2007). If *C. difficile* has growth characteristic of a biofilm in human infection, then study of these compounds could lead to enhancements in the efficacy of antimicrobials. Vaccines based on biofilm-specific antigens have been reported and used successfully in animal models of *Staphylococcus aureus* osteomyelitis and *Haemophilus influenzae* otitis media (Brady *et al.* 2011; Novotny *et al.* 2015).

Type IV pili (T4P) are a particular type of bacterial surface appendage that mediate adherence, colonization, biofilm formation, DNA transfer and twitching motility (Craig, Pique and Tainer 2004), among other functions; they have been well characterized in Gram-negative bacteria, and more recently described in Gram-positive bacteria. Gram-positive T4P were first recognized in *Ruminococcus albus* (Rakotoarivonina *et al.* 2002), and were subsequently observed in *C. perfringens* (Varga *et al.* 2006). Genes for T4P were identified in the *C. difficile* genome (Varga *et al.* 2006), although fimbrial structures on *C. difficile,* resembling T4P, were observed by electron microscopy nearly two decades previously (Borriello, Davies and Barclay 1988). The T4P structural subunits are called pilins; *C. difficile* T4P are composed primarily of PilA1 pilin (Piepenbrink *et al.* 2015). The *C. difficile* genome encodes genes for up to nine pilins, as well as the requisite assembly and scaffolding proteins, organized into three gene clusters as previously described by our lab (Maldarelli *et al.* 2014). A complex nanomachine assembles pilin monomers into filaments that can extend several micrometers from the bacterial cell (Craig, Pique and Tainer 2004). In many bacterial species, T4P can also retract; the energy for extension and retraction are provided by distinct ATPases (Whitchurch *et al.* 1991). T4P in Gram-negative organisms, such as enteropathogenic *Escherichia coli, Vibrio cholerae* and *Neisseria meningitidis,* are critical for colonization and pathogenesis (Nassif *et al.* 1994; Tacket *et al.* 1998; Humphries *et al.* 2009). In *N. meningitidis,* the binding of T4P to CD147 facilitates recruitment of beta-adrenergic receptors and leads to passage through the blood–brain barrier, a step absolutely critical to pathogenesis (Nassif *et al.* 1994; Bernard *et al.* 2014).

In addition to their role in colonization, T4P are also critical for development of biofilms in multiple organisms. In *Pseudomonas aeruginosa,* T4P-mediated motility is critical for early biofilm development (Klausen *et al.* 2003a), particularly in the attachment phase; T4P are also important in developing the architecture of the mature biofilm (Klausen *et al.* 2003b). In an *H. influenzae* strain, mutation of the major pilin results in thinner and less-stable biofilm as compared to the wild-type strain (Novotny *et al.* 2015). Recent studies also demonstrate the importance of T4P in Gram-positive biofilms. T4P genes are upregulated in *C. perfringens* biofilms grown at 37°C, and are required for biofilm formation (Obana, Nakamura and Nomura 2014). Recent work demonstrates cyclic-di-GMP regulation of *C. difficile* T4P gene expression and bacterial aggregation; Bordeleau *et al.* hypothesize that aggregation regulation is important in the development of *C. difficile* biofilms (Bordeleau *et al.* 2015; Purcell *et al.* 2016). Biofilm formation in *C. difficile* has been observed *in vitro* (Đapa *et al.* 2013), and putative bacterial aggregates implying biofilm formation have been observed *in vivo* (Buckley *et al.* 2011). Additionally, examination of the structure of PilA1, the major pilin subunit in *C. difficile,* shows a strong resemblance to the subunits of T4P known to mediate bacterial self-association, a prerequisite for formation of biofilms that depend on these structures (Piepenbrink *et al.* 2015).

Here, we investigate *C. difficile* biofilm gene expression and the impact of T4P deficiency, through knockout and complementation of the gene for the major *C. difficile* pilin, *pilA1,* on biofilm formation.

MATERIALS AND METHODS

Bacterial strains and growth conditions

The *Clostridium difficile* R20291 strain, isolated from a 2006 outbreak in the United Kingdom, was chosen for these experiments as its genetic sequence is known and mutants are available. Production of the R20291 *pilA1::ermB* and R20291 *pilA1::ermB* p*pilA1* *C. difficile* strains was previously described (Piepenbrink *et al.* 2015). *Clostridium difficile* was grown at 37°C within an anaerobic chamber (Coy Laboratory Products, Grass Lake, Michigan, US) with 10% H_2, 5% CO_2 and 85% N_2, unless otherwise stated.

RNA sample growth

For planktonic samples, overnight culture in brain heart infusion (BHI) media (Sigma-Aldrich, Saint Louis, Missouri, US) supplemented with 0.5% yeast extract and 0.1% L-cysteine brain heart infusion-supplemented (BHIS) was mixed 1:10 with fresh BHIS and grown with shaking for 6 h before mixing 1:1 with RNAProtect (Qiagen, Hilden, Germany). Biofilm

samples were grown for 1 week in glass jars containing 8 mm glass beads. The glass beads were added in order to increase surface area and maximize RNA yields. BHIS medium was changed daily and 6 h prior to cell harvest. To harvest adherent cells (biofilm), media was removed and the beads rinsed with fresh media before covering with RNAProtect and subjecting to vigorous shaking for 2 min. For plate growth samples, overnight culture was streaked onto Columbia blood agar plates (BD Biosciences, San Jose, California, US) and kept in the anaerobic chamber for 24 h prior to a 3-day room temperature growth in an anaerobic GasPak container (BD Biosciences). Plates were returned to the anaerobic chamber for cell harvesting and placement into RNAprotect. All samples were stored at $-80°$C until used for RNA sequencing.

Preparation of biofilm for imaging

Glass coverslips were placed in six-well (Costar, Corning, New York, US) tissue culture plates, and were inoculated with turbid overnight culture diluted to OD600 = 0.125 in pre-reduced BHI broth to a final volume of 4 mL, supplemented with thiamphenicol as needed for plasmid maintenance. Media was changed every other day. At day 1 or day 7, coverslips were washed with sterile PBS and stained with BacLight Live/Dead stain (Life Technologies, Carlsbad, California, US) according to manufacturer's instructions. Slides were fixed by incubation with 4% paraformaldehyde in PBS, mounted on glass slides, and stored in the dark at $-20°$C until imaged. Three slides per strain were grown for each strain per time point; two growths were conducted.

RNA sequencing

After thawing, cells were pelleted and RNAprotect removed. Cells were resuspended in 100 uL TES buffer with 35 000 units of Ready-Lyse lysozyme solution (Epicentre, Madison, Wisconsin, US) and incubated with shaking at room temperature for 90 min. The RNeasy RNA purification kit (Qiagen) was then used as per the manufacturer's instructions. Total RNA samples were treated with DNase I (Invitrogen, Carlsbad, California, US). The level of ribosomal RNA present in total RNA samples was reduced prior to library construction using the Ribo-Zero Gram-Positive Bacteria rRNA Removal Kit (Epicentre). Illumina RNAseq libraries were prepared with the TruSeq RNA Sample Prep kit (Illumina, San Diego, California, US), omitting the poly-A selection steps. Adapters containing six nucleotide indices were ligated to the double-stranded cDNA. The DNA was purified between enzymatic reactions, and the size selection of the library was performed with AMPure XT beads (Beckman Coulter Genomics, Danvers, Massachusetts, US). The libraries were sequenced on a 101 bp paired end run on the HiSeq 2000 (Illumina). Reads were processed for quality in Trimmomatic version 0.30 (Bolger, Lohse and Usadel 2014) using a leading and trailing minimum score of 10 and a four-base sliding window minimum score of 15; on average for all libraries 99.9% of reads survived. Bowtie2 version 2.1.0 (Langmead and Salzberg 2012) was used to map an average of 23.6 million paired reads to the *C. difficile* genome (NC_013316.1, NCBI) with an overall alignment rate of 96.2%. Samtools version 0.1.18 (Li *et al.* 2009) was used for file conversion to the binary BAM format. Transcript abundances were evaluated in Cufflinks version 2.1.1 (Li *et al.* 2009) using a ribosomal masking file for all 5S, 16S and 23S loci (NC_013316.gff annotation, NCBI), and fragments per kilobase per million were reported

(http://cufflinks.cbcb.umd.edu) (Trapnell *et al.* 2010). For comparative analysis of biofilm, plate and planktonic conditions, the Cuffdiff package in Cufflinks was used, and significant genes were reported. A cutoff false discovery rate-adjusted p-value (or q-value) of less than 0.05 was used to select significant differentially expressed genes. Heatmaps were generated using the gplots version 2.17.0 package (Warnes *et al.* 2015) in R (Team RC 2014). Mapped RNA-Seq reads were visualized using the Integrative Genomics Viewer (Robinson *et al.* 2011; Thorvaldsdóttir, Robinson and Mesirov *et al.* 2013).

RNA-sequencing data accession number

The data discussed in this publication have been deposited in NCBI's Gene Expression Omnibus (Edgar, Domrachev and Lash 2002) and are accessible through GEO Series accession number GSE69001 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE69001).

Confocal scanning laser microscopy (CLSM)

Biofilms were grown on 18×18 mm glass coverslips and prepared as above. Slides were read using a Zeiss 510 Meta confocal laser scanning microscope and accompanying software (Carl Zeiss AG, Oberkochen, Germany). Ten fields of view were imaged per slide. Live cells per field of view were enumerated with CellProfiler (Kamentsky *et al.* 2011). Z-stacks were acquired for a subset of these images; at least six z-stacks were analyzed per strain per time point. The structural organization of the biofilms was analyzed using the Comstat2 software package (http://www.comstat.dk) (Heydorn and Nielsen 2000; Vorregaard 2008). The 3D representations of the biofilms were generated using the 3D viewer plugin for ImageJ (http://3dviewer.neurofly.de) (Schmid *et al.* 2010).

Scanning electron microscopy (SEM)

Biofilms were grown on glass coverslips as described above and fixed in 100% ethanol. The samples were shipped to the Image and Chemical Analysis Laboratory, Department of Physics, Montana State University, where they were allowed to air dry and then taped to an SEM mount using double-sided conductive carbon tape. The samples were coated with iridium and then imaged with a Zeiss Supra 55VP Field Emission Scanning Electron Microscope using a 1 keV electron beam.

RESULTS

RNA sequencing reveals a unique *Clostridium difficile* transcriptome during biofilm growth

We performed RNA-Seq on *C. difficile* grown on plates under conditions favoring expression of PilA1 pili, on planktonic bacteria from BHIS broth culture, and on bacteria grown for 1 week in BHIS as a biofilm on glass beads. Of the 3652 *C. difficile* R20291 genes, 1604 were significantly differentially expressed in biofilm samples when compared to planktonic samples, and 2105 were significantly differentially expressed in biofilm samples when compared to plate-growth samples. We evaluated specific genes suspected to be growth phenotype related (Table 1). Not surprisingly, the flagellin gene *fliC* was significantly decreased in biofilm samples relative to planktonic samples, while there was no significant difference between biofilm and plate samples. Suspecting adhesins important in biofilm

Table 1. Differential expression of specific *C. difficile* genes during biofilm growth (pairwise comparison).

| Gene	Description	Biofilm/plate differential expression log$_2$(fold change)	Biofilm/planktonic differential expression log$_2$(fold change)
pilA1	Major pilin	− 0.90	0.87*
pilA2	Predicted pilin	− 3.12*	− 0.41
pilA3	Predicted pilin	− 2.17	0.37
pilJ	Minor pilin	0.43	0.07
pilK	Pilin-like protein	1.02*	1.18*
pilU	Pilin-like protein	1.04*	1.08*
pilV	Pilin-like protein	1.60*	0.96*
pilW	Pilin-like protein	− 0.27	− 0.60*
pilX	Pilin-like protein	− 2.21	− 0.26
fliC	Flagellin	− 0.25	− 1.35*
cwp66	Adhesin (cell-wall protein)	− 1.73*	− 0.18
cwp84	Cysteine protease important for surface layer maturation	0.37	− 0.31
CD0802	Adhesin (lipoprotein)	4.87*	− 0.87*
luxS	Quorum-sensing regulator	1.99	0.12
Spo0A	Transcription factor controlling spore formation	− 1.35	− 0.23
tcdA	Toxin A	− 1.70*	0.02
tcdB	Toxin B	− 1.53*	0.59*
sigB	Housekeeping gene	− 0.26	− 0.34

*Statistically significant (q-value < 0.05).

formation, we interrogated the adhesin genes *cwp66* (Waligora *et al.* 2001; Janoir *et al.* 2007) and CD0802, the equivalent of CD0873 in strain 630 (Kovacs-Simon *et al.* 2014). Unexpectedly, we found decreased expression of both in biofilm samples when compared to planktonic samples, although only CD0802 was significantly different. When comparing biofilm- and plate-growth samples, we found significantly decreased expression of *cwp66* and significantly increased expression of CD0802 in biofilm samples. Another adhesin, *cbpA*, encoding a collagen-binding adhesin, was recently identified in strain 630 (Tulli *et al.* 2013), but the gene is not present in strain R20291. *cwp84* encodes a cysteine protease that cleaves the highly expressed precursor surface layer protein SlpA into high-molecular-weight (HMW) and low-MW constituents; of the two, the HMW protein is more important for binding to human gastrointestinal tissues (Calabi *et al.* 2002). Interestingly, we find no differences in *cwp84* expression between all three growth conditions. This may reflect that our RNA-Seq biofilm samples were obtained after 1 week of growth, as a prior report documents that a *cwp84* mutant has a more dramatic decrease in biofilm formation on day 1 compared to days 3 and 5 (Đapa *et al.* 2013).

Cell-to-cell communication via a quorum-sensing system is vital for biofilm formation in multiple organisms (Davies *et al.* 1998; Yang *et al.* 2014; Omer Bendori *et al.* 2015). *Clostridium difficile* encodes the quorum-sensing regulator *luxS,* which is shown to regulate the expression of the genes encoding *C. difficile* toxin A and toxin B (Carter *et al.* 2005; Lee and Song, 2005). Mutation of *C. difficile luxS* results in a dramatic deficiency in biofilm formation, with inability to form even a bacterial monolayer on glass surfaces (Đapa *et al.* 2013). We observed no significant differences in expression of *luxS* between the three growth phenotypes examined. We did, however, find differential expression of toxin genes, suggesting that factors other than quorum sensing via the luxS regulator are important in regulation of toxin gene expression. The toxin B gene (*tcdB*), but not the toxin A gene (*tcdA),* was significantly increased in biofilm samples when compared to planktonic samples, while both *tcdA* and *tcdB* were significantly decreased in biofilm samples in comparison to plate-growth samples.

RNA sequencing shows differences in pilin gene expression

We previously demonstrated that there are far more sequence reads from *pilA1* than any other pilin gene when *C. difficile* is grown on blood agar under conditions that enhance the expression of pili (Fig. 1C and D), and that *pilA1* encodes the major pilin (Piepenbrink *et al.* 2015). Here, we also show that *pilA1* is the most highly expressed pilin gene during biofilm and planktonic growth (Fig. 1A, B and D), further supporting the designation of *pilA1*as the major pilin gene. We also find *pilJ* is the second most highly expressed pilin gene in all growth conditions. This is consistent with our prior work demonstrating PilJ incorporation into pili, indicating that PilJ is a minor pilin (Piepenbrink *et al.* 2014). When growth conditions are compared, significant expression differences are seen for multiple pilin genes (Fig. 1E). Expression of *pilA1* is significantly greater on plates than under planktonic conditions in broth (q < 0.05), as was expected from prior experience examining pili by electron microscopy (Piepenbrink *et al.* 2015). Relative expression of *pilA1* is also significantly greater when grown in BHIS as a biofilm on glass beads than when grown as planktonic cells. This finding is consistent with recent work using quantitative reverse transcriptase PCR that found a significant increase in *pilA1* expression in 48 h R20291 biofilms grown on plastic and compared to planktonic growth (Purcell *et al.* 2016). Although they reside in the same gene cluster as *pilA1*, our data indicate that expression of the pilin-like protein genes *pilK, pilU* and *pilV* may have different regulation. All three of these RNAs are expressed at significantly higher levels in biofilm conditions when compared to plate or planktonic growth, while *pilA1* is expressed at higher levels in biofilm relative to planktonic growth, but not relative to plate growth (Fig. 1E). Furthermore, while *pilA1* is expressed at a significantly higher level in plate growth in comparison to planktonic growth, this is not the case for *pilK, pilU* and *pilV* (Fig. 1E). Recent data demonstrate a predicted transcription terminator 36 bp downstream from the *pilA1* stop codon (Bordeleau *et al.* 2015), consistent with our finding of greater *pilA1* expression relative to *pil K, pil U* and *pil*V *expression*. Close inspection of the RNA-Seq data
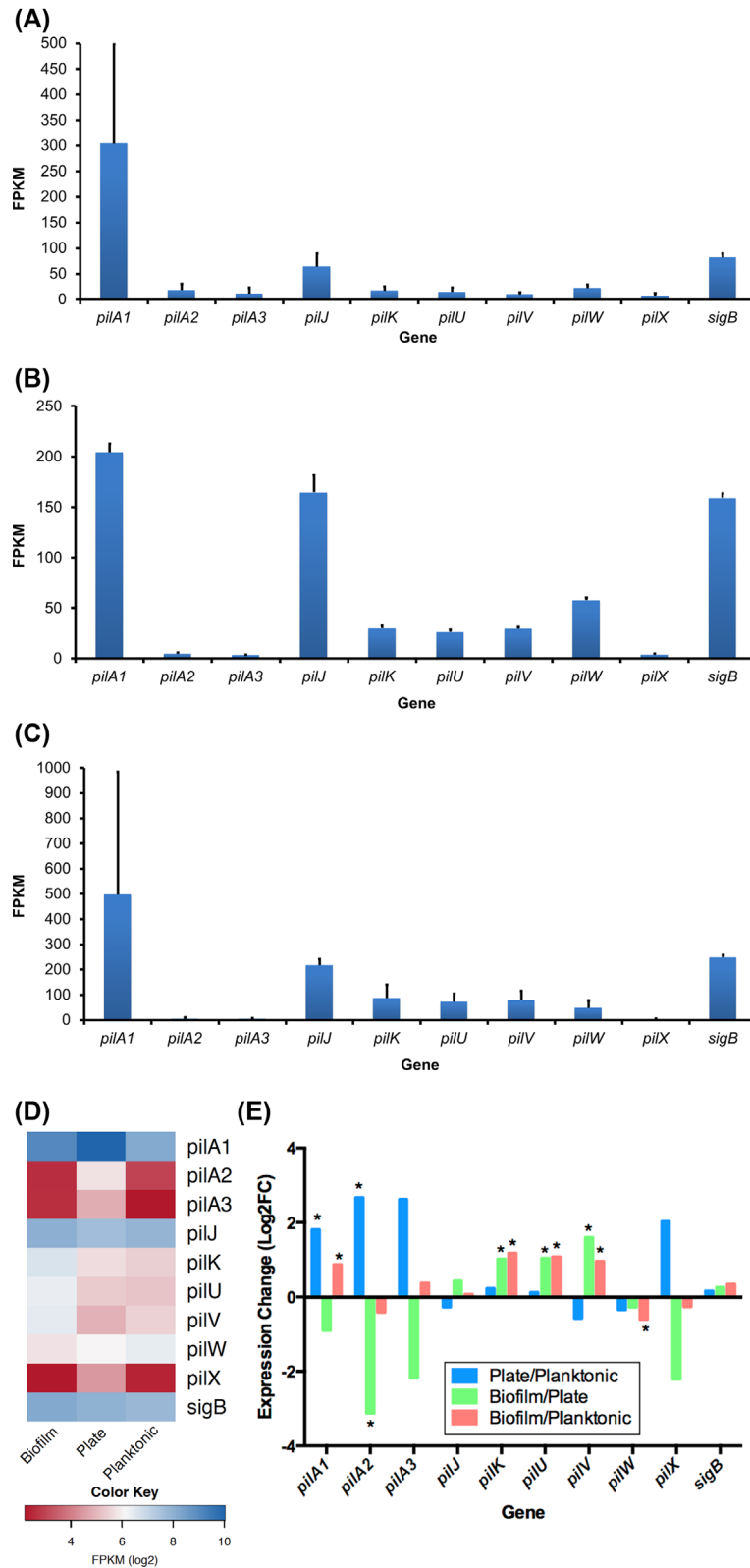
Figure 1. Pilin gene expression as determined by RNA sequencing of *C. difficile* samples grown on blood agar under conditions that express pili (A) as planktonic culture (B), and as a biofilm on glass beads (C). Using a three-way comparison, pilin gene expression is shown as a heatmap (D), and differential expression between the three growth conditions is shown using pairwise comparisons (E). *q value < 0.05. Please note that data included in panel A were previously published (Piepenbrink *et al.* 2015).

does not reveal evidence of a separate promoter or transcriptional terminator in this gene cluster. In contrast to the pilins from the largest gene cluster, *pilJ*, which is unlinked to the *pilA1* gene cluster, is expressed at equivalent levels under all three conditions.

A non-piliated mutant is deficient in initial biofilm formation

We previously demonstrated that PilA1 is the major *C. difficile* pilin, that the *pilA1* mutant produces no visible pili in conditions known to induce piliation in the wild-type strain and that piliation is restored with addition of a complementation plasmid (Piepenbrink *et al.* 2015). To investigate the role of T4P in biofilm formation, we grew wild-type *C. difficile*, the T4P-deficient mutant and the complemented mutant under biofilm-producing conditions.

If T4P are important in biofilm formation, we would expect a non-piliated *C. difficile* mutant to be delayed in biofilm formation as indicated by decreased biomass and decreased live cell count in early biofilms as compared to wild-type bacteria; indeed, we observe this to be the case. Analysis of the 3D biofilm structure demonstrates that biofilms formed by the *pilA1* mutant accumulate significantly less biomass at day 1 than the wild-type strain or the complemented mutant (Fig. 2A); correspondingly, the wild-type and the complemented *pilA1* mutant form significantly thicker biofilms than the *pilA1* mutant at day 1 (Fig. 2B). Quantification of CLSM images demonstrates that after 1 day of growth *pilA1* mutant biofilm appears to have fewer live

cells present per field of view than the parent wild-type strain (Figs 2C and 3A–C), although this finding is not statistically significant. The 3D reconstructions of biofilms illustrate the thicker biofilms formed by the wild-type and complemented mutant as compared to the *pilA1* mutant (Fig. 3G–J) Complementation of *pilA1* restores the count of live cells per field of view to levels slightly higher than seen in the wild-type strain.

The *pilA1* mutant strain's biofilm formation deficiency is not simply due to differences in growth rates between the wild-type and mutant strains. In broth culture, the *pilA1* mutant demonstrates similar growth as compared to the parent wild-type strain, indicating that the absence of T4P is the principal reason for the deficit in biofilm formation in the *pilA1* mutant strain. While the non-piliated *pilA1* mutant shows fewer live cells per field of view than the parent wild-type strain after 24 h of growth, this difference disappears by day 7 (Fig. 2C and H–J). As above, the 3D biofilm reconstructions demonstrate this phenotype (Fig. 3K–M). This phenotype, that of an early deficiency in biofilm formation, is consistent with previous observations of T4P-associated biofilms in *C. perfringens* and non-typeable *Haemophilus influenzae* (NTHI) (Carruthers *et al.* 2012; Novotny *et al.* 2015).

Along with total biomass and biofilm thickness, the measured surface-area-to-volume ratio provides a useful descriptor of a strain's tendency to form clumps or aggregates of bacteria; a lower surface-area-to-volume ratio indicates fewer isolated bacteria and more aggregates. Given that T4P are required for aggregation or microcolony formation by other bacteria, we would anticipate an increased surface-area-to-volume ratio in the *pilA1* mutant relative to the wild-type strain. Indeed, we find that the *pilA1* mutant has an increased ratio relative to the wild-type
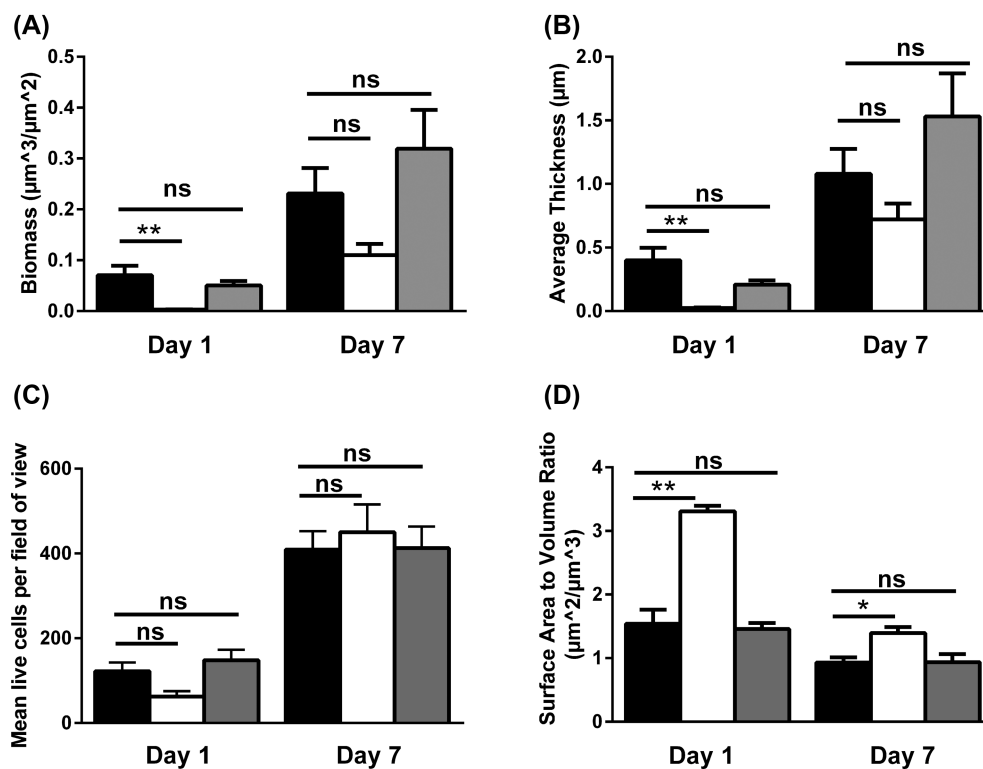

Figure 2. Assessment of lack of piliation on *C. difficile* biofilm formation. Biofilm biomass (**A**), thickness (**B**), live cell count (**C**) and surface-area-to-volume ratio (**D**) were calculated for wild-type *C. difficile* (black bars), the *pilA1* mutant strain (white bars) and the complemented mutant (gray bars) at 1 and 7 days of growth. After 1 day of growth, the *pilA1* mutant forms biofilms of significantly reduced biomass and thickness that contain fewer live cells as compared to the parent wild-type strain; the complemented mutant resembles the wild-type strain. These differences are still apparent after 7 days of growth, though these differences are no longer significant. The *pilA1* mutant strain also has a significantly elevated surface-to-volume ratio as compared to the wild type at 1 and 7 days of growth. * indicates $P < 0.01$, ** indicates $p < 0.001$. Groups were compared using a one-way ANOVA with Dunnett's post-test. Error bars indicate SEM.
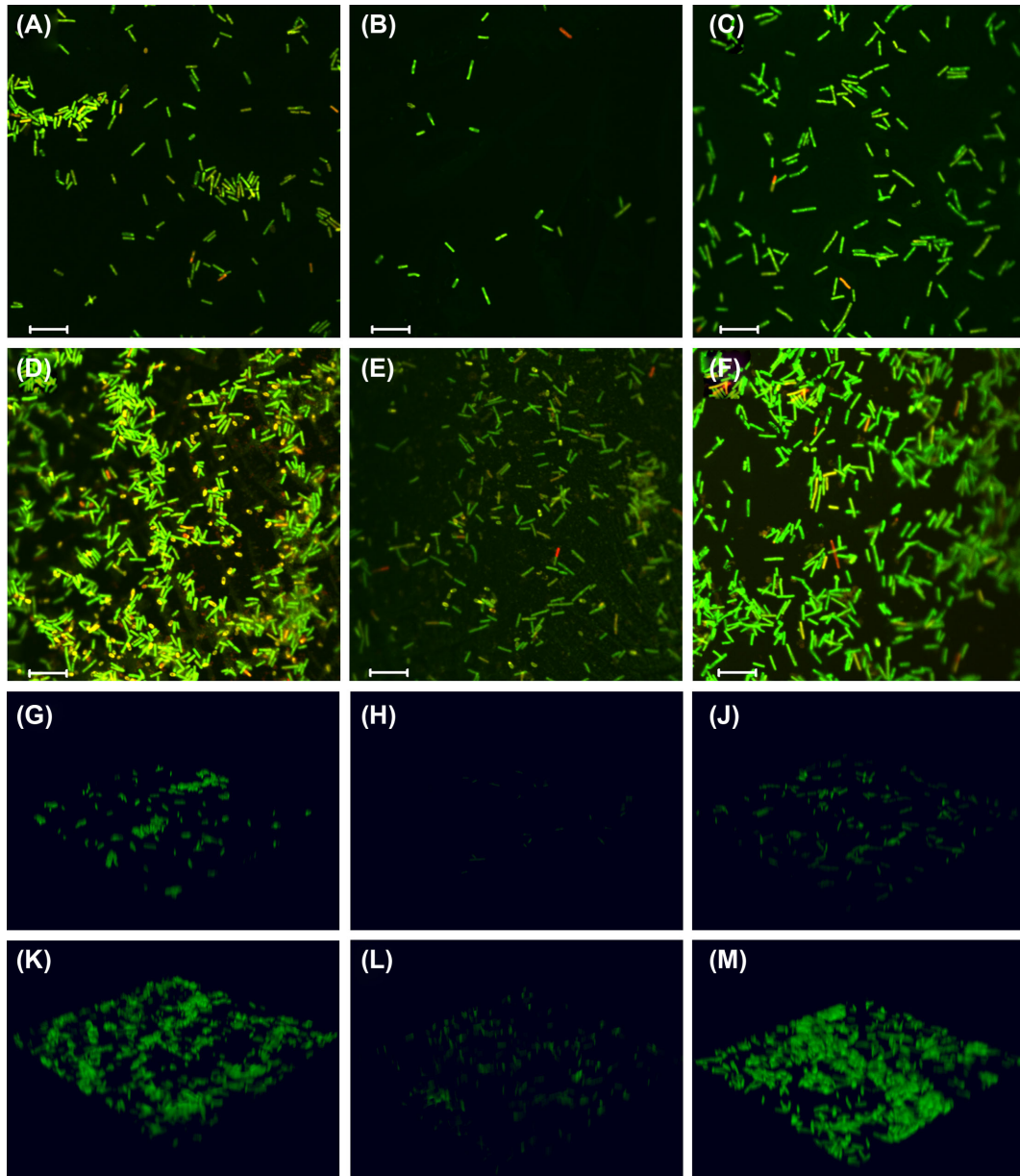
Figure 3. Visualization of *C. difficile* biofilms by CLSM. Representative images from each strain at both time points demonstrate the reduced biomass and thickness of the *pilA1* mutant biofilms as compared to the wild-type strain. Complementation of the mutation restores the wild-type phenotype. The 3D reconstructions of the imaged biofilms further illustrate these observations. Images demonstrate 1 day (**A-C, G-J**) and 7 days (**D–F, K-M**) of growth for wild type (**A, D, G and K**), the *pilA1* mutant (**B, E, H and L**) and the complemented mutant (**C, F, J and M**). Live cells are stained green, dead cells are stained red. The scale bar for images A–F indicates 10 μm.

strain, and complementation restores this ratio to wild-type levels (Fig. 2D). Importantly, these results clearly indicate a role for T4P in promoting bacterial aggregation rather than simply increasing adherence to an abiotic surface. However, CLSM images show aggregates of bacteria in all strains at both time points (Fig. 3A–G), although surface-area-to-volume ratios demonstrate that the wild-type strain has a greater propensity toward aggregate formation than the *pilA1* mutant (Fig. 3K–M). Similarly, SEM data show that both wild-type and *pilA1* mutants are capable of forming aggregates with an extracellular matrix (Fig. 4).

Given the presence of these aggregates in the *pilA1* mutant, these aggregates can form in the absence of T4P composed of PilA1. Thus, other adhesive proteins, for example CbpA or SlpA

(Merrigan *et al.* 2013; Tulli *et al.* 2013), may be involved in cell–cell adhesion alongside T4P; alternately, T4P composed of PilA2 or PilA3 may be involved in aggregate formation. As shown above, *pilA2* and *pilA3* are expressed at low levels in the *C. difficile* biofilm; however, no pili were observed on the *pilA1* mutant, suggesting their role, if any, is a small one.

In sum, T4P genes are differentially expressed during biofilm growth and critical for early biofilm formation in *C. difficile*: a non-piliated strain is deficient in initial biofilm formation, forms a biofilm of reduced thickness and biomass and has decreased tendency to aggregate in comparison to the parent wild-type strain. Complementation of the mutant restores the above measures to wild-type levels.
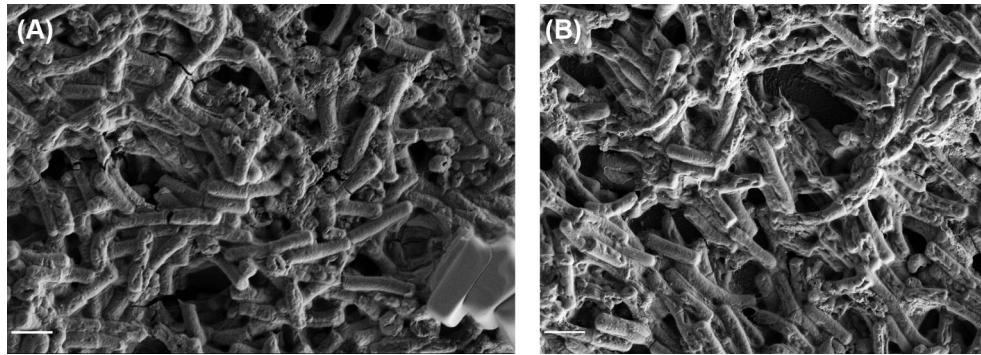
Figure 4. Formation of biofilm by *C. difficile*. SEM imaging demonstrates formation of cell aggregates linked by matrix by both wild-type (**A**) and *pilA1* mutant (**B**) strains of *C. difficile* after 7 days of growth. Scale bar indicates 2 μm.

DISCUSSION

Despite the human and medical costs of CDI, its mechanisms of colon colonization are poorly understood. Previous research demonstrates that *C. difficile* forms biofilms and expresses T4P (Đapa *et al.* 2013; Maldarelli *et al.* 2014; Bordeleau *et al.* 2015; Purcell *et al.* 2016). In this work, we sought to elucidate the role of T4P in *C. difficile* biofilm formation. We demonstrate major differences in gene expression when *C. difficile* is grown as a biofilm as compared to planktonic or plate growth, including differences in pilin gene expression. We also demonstrate the role of T4P in biofilm generation: a non-piliated mutant is slower to form biofilm and less likely to form aggregates than the parent wild-type strain. Together, our results demonstrate that biofilm is a distinct mode of *C. difficile* growth, one in which T4P play an important role. Furthermore, these data reinforce earlier work (Đapa *et al.* 2013) that emphasizes the importance of studying biofilms in this clinically relevant organism.

Our RNA-Seq studies provide valuable information regarding gene expression in *C. difficile* biofilms. We find that about half of all genes in our biofilm-growth samples are significantly differentially expressed when compared to either planktonic- or plate-growth samples. When comparing biofilm and planktonic gene expression, we expected to see increased expression of adhesin genes in the biofilm samples; however, we saw significantly lower expression of CD0802 and insignificantly lower expression of *cwp66*. These data lead us to hypothesize that these adhesins are not expressed at high levels once a biofilm is established, and this finding may reflect that samples were grown for 1 week before RNA extraction. Alternatively, as the ligands for these adhesins are unknown, and may be present on host tissue but not the surface of *C. difficile*, increased expression of these adhesins may rely on the presence of the host. Although others have shown a role for the quorum-sensing regulator *luxS*, the sporulation transcription factor *spo0A* and cysteine protease *cwp84* in biofilm formation (Đapa *et al.* 2013), we found no significant differences in expression of any of these genes in our samples. One limitation of our study is that we only have expression data for 1-week biofilm samples, and it is quite possible that differences occur at earlier time points.

Using quantitative biofilm formation assays, we show that a non-piliated mutant is deficient in early biofilm formation, which supports our hypothesis. The trend toward a decreased day 1 live-cell count in the non-piliated strain suggests that T4P are important in surface attachment, the first step of biofilm formation. While pili primarily composed of PilA2 and PilA3 may be expressed under other conditions, the genes encoding those

proteins may be remnants of other complete T4P gene clusters, and may no longer be expressed. Our RNA-Seq data suggest that *pilA2* and *pilA3* are poorly expressed under the conditions tested here, and no pili were observed in the *pilA1* mutant, indicating that, under the conditions used here, PilA2 and PilA3 pili are unlikely to contribute to biofilm formation. Deficiency in biofilm formation upon loss of piliation is common to other T4P-expressing, biofilm-forming organisms, including NTHI and *Pseudomonas aeruginosa*. NTHI strains lacking expression of the major pilin *pilA* form biofilms of reduced thickness and reduced biomass after 24 h of growth (Carruthers *et al.* 2012), results consistent with our findings in *C. difficile*. Similarly, experiments with cocultures of wild-type and non-piliated mutant *P. aeruginosa* strains demonstrate that non-piliated bacteria are less able to move between microcolonies, and consequently form less regular biofilms (Klausen *et al.* 2003a). The two T4P systems of the aquatic pathogen *Vibrio parahaemolyticus* have discrete roles in biofilm formation; the mannose-sensitive hemagglutinin pilus is involved in bacterial adhesion to surfaces, whereas the chitin-regulated pilus plays a role in cell–cell adhesion (Shime-Hattori *et al.* 2006). T4P in *C. difficile* may serve similar purposes, allowing bacteria in the nascent biofilm to adhere to each other, as well as to surfaces. *Clostridium difficile* encodes three putative major pilins, *pilA1, pilA2 and pilA3* (Merrigan *et al.* 2013). We demonstrate here that pili composed of PilA1 are involved in early biofilm formation, although this result does not exclude the possibility that pili composed primarily of PilA2 or PilA3 also contribute to biofilm formation and maintenance. As in *V. parahaemolyticus*, multiple types of pili may be involved in aggregation and biofilm formation; other mechanisms unrelated to pili are most likely involved as well.

Our observations regarding increased surface-area-to-volume ratio in the *pilA1* mutant relative to the wild-type strain support the hypothesis that T4P aid the bacteria in forming aggregates or microcolonies, as a lower relative surface-to-volume ratio is indicative of more cell aggregates. Microcolonies are a critical part of *P. aeruginosa* biofilm initiation, and in *Neisseria meningitidis,* T4P-reliant microcolony formation is critical for bacterial passage across the blood–brain barrier. Our results provide a potential explanation for *C. difficile* cell aggregates, that is linkage by hair-like structures, which have been seen by SEM in the ceca and colons of *C. difficile*-infected hamsters 24 and 36 h post-infection (Buckley *et al.* 2011). Taken together, our findings and the images acquired by Buckley and colleagues suggest a possible mechanism of colonic *C. difficile* colonization: T4P facilitate initial attachment to the colonic epithelium and

microcolony formation, which in turn leads to progression of disease. The involvement of T4P in initial *C. difficile* biofilm formation is dependent upon c-di-GMP levels, as was recently demonstrated by Purcell *et al.* (2016).

The ability to form biofilms may provide *C. difficile* a selective advantage. If T4P facilitate *C. difficile* adherence (both bacterium to bacterium and bacterium to epithelium), T4P-mediated adhesion could allow the bacteria to persist in the colon. Although spore formation may be the predominant mechanism underlying recurrent CDI, the presence of sessile communities in the colon may help to explain why symptoms of CDI can persist in some individuals despite recommended therapy and why infection recurs with high frequency. Bacteria in the biofilm mode of growth are more resistant to antimicrobials than bacteria grown on plates or in broth (Folsom *et al.* 2010). This phenomenon has been demonstrated *in vitro* in *C. difficile*; bacteria grown as a biofilm were demonstrably more resistant to killing by super-MIC concentrations of vancomycin when compared to equivalent planktonic cultures (Đapa *et al.* 2013). This advantage at super-MIC levels of vancomycin may indicate one mechanism by which *C. difficile* evades eradication during oral vancomycin treatment.

In conclusion, we demonstrate that the RNA expression profile of *C. difficile* in biofilm is distinct from that of *C. difficile* grown in broth or on agar plates, including significant differences in pilin, adhesin and toxin gene expression. We find that bacteria unable to produce T4P form thinner biofilms with decreased biomass and a trend toward fewer live cells than biofilms formed by T4P-producing, wild-type bacteria. This work leads us to a more general understanding of biofilm development and the role of T4P in that development.

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