The *Xanthomonas axonopodis* pv. *citri* flagellum is required for mature biofilm and canker development

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*Xanthomonas axonopodis* pv. *citri* (*Xac*) is the causative agent of citrus canker. This bacterium develops a characteristic biofilm on both biotic and abiotic surfaces. To evaluate the participation of the single flagellum of *Xac* in biofilm formation, mutants in the *fliC* (flagellin) and the *flgE* (hook) genes were generated. Swimming motility, assessed on 0.25 % agar plates, was markedly reduced in *fliC* and *flgE* mutants. However, the *fliC* and *flgE* mutants exhibited a flagellar-independent surface translocation on 0.5 % agar plates. Mutation of either the *rpfF* or the *rpfC* gene, which both encode proteins involved in cell–cell signalling mediated by diffusible signal factor (DSF), led to a reduction in both flagellar-dependent and flagellar-independent surface translocation, indicating a regulatory role for DSF in both types of motility. Confocal laser scanning microscopy of biofilms produced in static culture demonstrated that the flagellum is also involved in the formation of mushroom-shaped structures and water channels, and in the dispersion of biofilms. The presence of the flagellum was required for mature biofilm development on lemon leaf surfaces. The absence of flagellin produced a slight reduction in *Xac* pathogenicity and this reduction was more severe when the complete flagellum structure was absent.

INTRODUCTION

Biofilms are bacterial communities in which cells are embedded in an extracellular polysaccharide (EPS) matrix that can be attached to a surface (Branda et al., 2005; Southey-Pillig et al., 2005). Life in these communities provides protection to the organisms from deleterious conditions (Davey & O'Toole, 2000), and biofilm formation is considered to be important for the disease cycle of bacterial pathogens of both animals and plants. We are interested in understanding the role of biofilm formation in the development of canker disease by *Xanthomonas axonopodis* pv. *citri* (*Xac*), one of the most devastating diseases of citrus species. *Xac* is a foliar pathogen that enters the plant leaves through stomata and can infect both fruits and leaves (Brunings & Gabriel, 2003). Confocal laser scanning microscopy (CLSM) of citrus cankers, using *Xac* bacteria expressing green fluorescent protein (GFP), showed the occurrence of structured arrangements of cells (Rigano et al., 2007b). Our aim here was to identify bacterial factors involved in the development of such three-dimensional structures.

The development of bacterial biofilms is generally a multistep process, which is initiated when the bacteria reach a surface. First, the bacteria attach to a surface reversibly, where they can move freely across it until they become immobilized (Stoodley et al., 2002). To achieve a mature biofilm, new individuals appear in the community either by recruitment from planktonic bacteria or through replication of cells already present in the biofilm. Finally, some grouped cells disperse to develop new structures elsewhere (Heydorn et al., 2000; Tolker-Nielsen et al.,

**Abbreviations:** CLSM, confocal laser scanning microscopy; CV, crystal violet; DSF, diffusible signal factor; EPS, extracellular polysaccharide; *Xac*, *Xanthomonas axonopodis* pv. *citri*; *Xcc*, *Xanthomonas campestris* pv. *campestris*.

Two supplementary figures are available with the online version of this paper.
Flagella have been implicated in surface attachment in several bacteria (Lemon et al., 2007; O’Toole & Kolter, 1998a, b). For instance, flagella are necessary for swarming motility in Pseudomonas aeruginosa, which in turn is important in determining the final structure of the biofilm (Merritt et al., 2007). Similarly, Escherichia coli is not able to form an organized structure when flagella are lost. It has been proposed that E. coli flagella serve to overcome surface repulsion (Van Houdt & Michiels, 2005) and also allow attached cells to migrate along the abiotic surface (Pratt & Kolter, 1998).

However, flagella are important but not essential for surface attachment in Vibrio cholerae El Tor (Watnick & Kolter, 1999). Previous work from our laboratory has established that synthesis of the EPS xanthan contributes to the formation of structured biofilms in Xac and that both synthesis of xanthan and cell–cell signalling involving diffusible signal factor (DSF) contribute to structured biofilm formation in the related Xanthomonas campestris pv. campestris (Xcc) (Rigano et al., 2007a; Torres et al., 2007). Synthesis and perception of DSF require proteins coded in the rpf cluster (for regulation of pathogenicity factors). RpfF directs DSF synthesis whereas RpfC is involved in DSF perception (Barber et al., 1997; Slater et al., 2000). Mutants in the rpfF (DSF-minus) and rpfC (DSF overproducer) genes in Xcc can only form unstructured arrangements of bacteria (Torres et al., 2007). Synthesis of xanthan is directed by genes from the gum operon (da Silva et al., 2002), which is highly conserved in Xanthomonas spp. Disruption of the first gene, gumB, leads to complete loss of xanthan production (Vojnov et al., 1998); strains of Xac or Xcc with mutations in gumB cannot form structured biofilms (Rigano et al., 2007b; Torres et al., 2007). Beyond DSF signalling and xanthan production, relatively little is known about factors influencing biofilm formation in Xanthomonas spp. and the role that the flagellum plays in this process.

Both Xcc and Xac bear a single polar flagellum. Flagellar gene clusters encode all the structural proteins of the flagellum in Xac (da Silva et al., 2002). In addition, it has been shown that several genes encoding auxiliary protein subunits are involved in regulation of the flagellar assembly (Khater et al., 2007). As in many other bacterial species, FliC and FlgE proteins are the flagellin (Vonderviszt et al., 1998) and the hook components (Aizawa, 1996), respectively, of the flagellar structure. This structure is formed by a thin filament that protrudes from the cell body bound to a basal body that includes a rotor and a stator (Chevance & Hughes, 2008).

Here we address the role of the single flagellum in biofilm formation in Xac. Our initial approach was to test the effects of mutations in fliC and flgE on motility and biofilm structure. Our results indicate that the flagellum and flagellum-dependent motility are important for initial adherence to surfaces, for the development of a mature structured biofilm and for biofilm dispersion. Unexpectedly, our observations reveal a second type of motility that is flagellum-independent and EPS-dependent. Both types of motility are regulated by the DSF signalling system. Thus, these findings indicate possible additional roles for cell–cell signalling and EPS in the biofilm formation process.

METHODS

Bacterial strains. Xanthomonas strains were cultured at 28 °C with shaking in PYM (Cadmus et al., 1976) or in Y minimal medium (YMM) (Sherwood, 1970). To examine biofilm development, bacteria were grown in YMM containing 1 % (w/v) glucose as the carbon source (Rigano et al., 2007b). E. coli was grown at 37 °C in Luria–Bertani medium (Sambrook et al., 1989). Bacterial growth was measured in a Spectronic 20 Genesys spectrophotometer (Thermo Electron) at 600 nm. When required, the antibiotics ampicillin (100 μg ml⁻¹), kanamycin (50 μg ml⁻¹), spectinomycin (100 μg ml⁻¹), tetracycline (10 μg ml⁻¹) or gentamicin (20 μg ml⁻¹) were added to the growth media.

Construction of fliC and flgE mutants. Molecular techniques and extraction of genomic DNA from Xac were performed according to Rigano et al. (2007b). To obtain the XacfliC mutation strain (flic), the flic gene was partially deleted. First, two fragments of 408 and 484 bp were amplified from the region encoding the wild-type Xac fliD and flic genes by using primers flic1/flic2 and flic3/flic4, respectively (Table 1). The fragments were digested with restriction enzyme Xbal, purified and ligated to one another. A PCR was then performed using this ligation as template and primers flic1/flic4. The PCR product was resolved by 0.8% agarose gel electrophoresis and a fragment of 892 bp was excised and purified from the gel using the Wizard SV Gel and PCR Clean-Up System (Promega). The product was cloned into pGEM-T easy (Promega) and subcloned into the suicide vector pK18mobacB (Katen et al., 1999), producing the pKΔfliC construct.

The Xac flegE::Ω strain (flegE) was constructed by allelic exchange. Two regions, upstream and downstream from flegE, were amplified using fleg1/fleg2 (546 bp) and fleg3/fleg4 (547 bp) primers, respectively (Table 1). The two fragments were digested with HindIII, purified and ligated to one another, and a PCR was then performed using this ligation as template and primers fleg1/fleg4. The PCR product of 1093 bp was resolved and purified as described above. The product was cloned into pGEM-T easy (Promega) and subcloned into the suicide vector pl8mobacB (Katen et al., 1999), producing the pKΔflgE construct.

The two constructs, pKΔfliC and pKΔflgE, were used to transform the wild-type strain by electroporation (do Amaral et al., 2005). Transformed bacteria were selected on PYM agar medium supplemented with kanamycin (Müller et al., 1993). Sucrose-sensitive clones were grown in the absence of antibiotics and double recombination events were selected on PYM agar plates supplemented with 5 % sucrose and spectinomycin in the case of the flegE mutant. Km²/sucrose² candidates were screened by PCR followed by sequencing of the amplified fragments. Isolated mutants were confirmed by flagellum staining as described by Kearns & Losick (2003) and motility was further analysed.

For genetic complementation of the flic mutants, the entire gene and 245 bp from the upstream region were amplified by PCR using...
primers c-Flic-sense and Flic4 (Table 1). The amplified fragment was cloned into pLAFR3 (Staskawicz et al., 1987) to obtain pLAFR-fliC. Similarly, the flgE gene was amplified using primers c-Flge-sense and c-Flge-antisense (Table 1); the amplicon was cloned into pLAFR3 to obtain pLAFR-flgE. Both constructs were confirmed by DNA sequencing and electroporated into fliC and flgE mutant strains.

### Bacterial adhesion and biofilm quantification

The crystal violet (CV) technique was used to quantify biofilm development of the different strains on an abiotic surface (O'Toole & Kolter, 1998b). Bacterial strains were grown overnight in PYM nutrient medium and then inoculated into YMM to a final OD 600 of 0.1. Aliquots of 150 μl were used to fill the wells of a 96-well polystyrene plate and incubated at 28 °C for 24, 48 or 72 h. To confirm similar bacterial growth, the OD600 was measured before the adhesion assay was performed.

To analyse the capacity of the strains to adhere to the abiotic surface, bacteria from an overnight culture growth in YMM at 28 °C and 200 r.p.m. were used. Aliquots of 150 μl were dispensed into the wells of a 96-well plate and bacteria were incubated as previously described for 1, 3, 6 and 24 h. The OD 600 and cell attachment were measured as described above.

For quantification of biofilm development and adhesion of cells to an abiotic surface, the medium was gently removed using a pipette, and the 96-well plate was washed using 0.9 % NaCl and stained with 0.1 % CV solution. After 30 min incubation the unbound CV stain was

### Table 1. Strains, plasmids and primers used in this study

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*Km’, Ap’, Gm’, Sp’ and Tc’ indicate kanamycin, ampicillin, gentamicin, spectinomycin and tetracycline resistance, respectively. Sites for restriction enzymes are underlined: *EcoRI, §XbaI, §BamHI, ||HindIII.*
removed and the wells were washed with distilled water. The CV in each well was solubilized by adding 150 µl 70% ethanol and was quantified by absorbance at 570 nm.

The adhesion value was normalized according to the number of cells. This value was termed relative adhesion (A570/OD600).

**In vitro and in vivo analysis of biofilm by CLSM.** For in vitro experiments each strain expressing the autofluorescent protein GFP was grown at 28 °C on PYM supplemented with kanamycin. The OD600 was adjusted to 0.004 in YMM. Aliquots of 500 µl were transferred to chambered coverglass slides containing a 1 µm thick borosilicate glass (no. 155411; Lab-Tek, Nunc), as described by Russo et al. (2006).

Biofilm formation was monitored with a Zeiss LSM 510/Axiovert 100M confocal laser scanning microscope at 1, 2, 3, 4 and 5 days post-inoculation (days p.i.) of the bacteria on the chambered coverglass slides. The assay was performed in triplicate and three-dimensional images were generated by the program Zeiss LSM Image Browser, version 3.2.0.

For in vivo studies, leaves of lemon (*Citrus limon*) were infected by spraying with a 1 x 10^6 c.f.u. ml⁻¹ suspension of GFP-labelled wild-type *Xac* or the *fliC* or *flgE* mutants. After 6 days of incubation, areas of approximately 1 cm² were cut from the leaves and mounted on the adaxial leaf surface under glass coverslips. The samples were observed with a Nikon C1 confocal laser scanning microscope. The simulated images obtained were analysed with the program Nikon EZ-C1. Assays were performed in triplicate.

**Quantification of biofilm structures.** Quantifications of biofilm volume and thickness of three-dimensional CLSM biofilm image stacks were performed by using the COMSTAT program (Heydorn et al., 2000). For comparative analysis of the wild-type, *fliC* and *flgE* mutant strains, three independent biofilm experiments were performed. In each round, seven image stacks were acquired at the following time points: 1, 2, 3, 4 and 5 days p.i.

**RNA purification and RT-PCR.** Total RNA of *Xac* cells was isolated by using an RBC kit, according to the manufacturer’s instructions (RBCbioscience). The RNA was treated with RNase-free DNase (Promega) and the integrity of the nucleic acid was checked by agarose gel electrophoresis. Total RNA (1 µg) was used to synthesize the cDNA using MMLV-RT (Promega) and the oligonucleotide dN6. Expression of the *gumB*, *fliC* and 16S genes was determined by RT-PCR using the primers listed in Table 1.

The density corresponding to the RT-PCR bands was quantified by using ImageJ 1.41 software from the National Institutes of Health (http://rsbweb.nih.gov/ij/download.html). Each gene was normalized using 16S rRNA as the housekeeping gene. The ratio between normalized *fliC* or *gumB* in *rpfC* and *rpfF* mutants and in wild-type *Xac* was determined. Data are expressed as the mean ± SEM (n=3).

**Motility assays.** Motility assays were performed as described by Rashid & Kornberg (2000). Briefly, bacteria were grown overnight in PYM medium, then 3 µl of bacterial cultures with normalized OD600 were used to inoculate plates of 0.25% agar NYGB medium (Barber et al., 1997) (swimming) and 0.5% agar PYM. After 72 h, motility was assessed qualitatively by examining the circular halo formed by the growing bacterial cells. The assays were performed in triplicate.

**Flagellar staining with CV.** Flagellar staining was done based on a modification of the method described by Mayfield & Inniss (1977). The staining solution was prepared by mixing 10 vols solution A (2 g tannic acid, 10 ml 5% phenol, 10 ml AlKO₂S₂.12H₂O) with 1 vol. 12% (w/v) CV in ethanol. Then 3 µl of bacterial culture grown overnight in PYM (OD₆₀₀ 2) was placed over a coverglass with a coverslip and 10 µl of the solution was used. Observations were made with an optical microscope (Carl Zeiss).

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**Fig. 1.** Effects of mutation of the *fliC*, *flgE* and *gumB* genes on motility in *Xac*. (a) Flagellum-dependent swimming motility on 0.25% NYGB agar plates. (b) Motility on 0.5% PYM agar plates with 1% glucose.
**Plant material and inoculations.** *Citrus limon* ‘Eureka’ was used as the host plant for *Xac*. Plants were kept in a controlled temperature room at 20–25 °C and with a 16 h photoperiod. Bacteria were grown in PYM with the appropriate antibiotics. Bacterial suspensions were diluted in 10 mM MgCl₂ to a final concentration of $1 \times 10^6$, $1 \times 10^5$ or $1 \times 10^4$ c.f.u. ml⁻¹ and were inoculated by spraying as previously described (Rigano et al., 2007b). Bacterial growth in the host plant was quantified as previously described (Rigano et al., 2007b), using $1 \times 10^6$ c.f.u. ml⁻¹ to spray the leaves. Statistical significance of differences between bacterial populations was determined by Student’s unpaired two-tailed t-test. P-values <0.05 were considered statistically relevant. Cankers from 20 inoculated leaves were quantified and their areas were calculated using the program ImageJ version 1.41.

**RESULTS**

**FliC and FlgE are required for swimming motility**

The genomic arrangement near the *fliC* and *flgE* genes in *Xac* is shown in Supplementary Fig. S1(a), available with the online version of this paper. *Xac* strains with either a *fliC* deletion or a disruption in the *flgE* gene with a spectinomycin resistance cassette were created as described in Methods (Supplementary Fig. S1b). The mutations were confirmed by PCR (Supplementary Fig. S1c) and DNA sequence analysis (data not shown). The absence of the flagellar structure in these mutants was confirmed by CV staining (Kearns & Losick, 2003). A typical single polar flagellum was observed in wild-type *Xac*, but was absent in the two mutants (Supplementary Fig. S1d). This observation confirms the expected and fundamental role for FliC and FlgE in the morphogenesis of the flagellum in *Xac*.

The ability of the different *Xac* strains to swim in low-percentage agar was assayed to evaluate the role of the flagellum in motility. NYGB 0.25 % agar plates were inoculated with the wild-type strain and the *fliC* and *flgE* mutants. After incubation at 28 °C for 72 h, both mutants showed a considerable reduction in their motility compared with the wild-type (Fig. 1a). This was expected because the same phenotype has been reported for these kinds of mutations in other bacteria (Murray & Kazmierczak, 2006; Young et al., 1999). The complemented mutant strains showed a partial restoration of their motility, confirming that no other genes had been affected in the mutants (Supplementary Fig. S2).
In Xcc, DSF signalling elevates the expression of a wide range of genes, including those involved in flagellum synthesis (He et al., 2006). This prompted us to evaluate whether DSF signalling regulates swimming motility in Xac. The reduced motility of rpfE and rpfC mutants (Fig. 2a) and the reduced expression levels of fliC in both mutants (Fig. 2b) were indicative that flagellar-mediated swimming in Xac depends on cell–cell signalling. The gumB mutant showed a wild-type motility, indicating that xanthan has no role in Xac swimming motility (Fig. 1a). The gumB gene, as reported previously (Vojnov et al., 2001), is also regulated by the DSF/Rpf system (Fig. 2d).

**A second type of Xac motility is independent of the flagellum**

The different Xac strains were also tested for motility on 0.5% agar PYM swarm plates. Both fliC and flgE mutants showed a comparable surface translocation to the wild-type (Fig. 1b) although the gumB (Fig. 1b) and rpf (Fig. 2c) mutants showed a reduction in motility. Given these findings, under the conditions here, motility was not dependent on the presence of a flagellum, but was dependent on xanthan production and DSF signalling. Such motility has been termed ‘sliding’ (Horng et al., 2002), and it has been proposed that EPS acts as a surfactant or surface-wetting agent to facilitate this type of movement (Murray & Kazmierczak, 2008).

**Flagellar mutants show reduced attachment to abiotic surfaces and biofilm development**

Adhesion to an abiotic surface and biofilm formation was assayed using 96-well polystyrene plates and CV staining. The fliC and flgE mutants showed reduced adherence throughout a 24 h experiment, suggesting that the intact flagellum is required for this stage (Fig. 3a). To follow biofilm development, a lower bacterial concentration (OD$_{600}$ 0.1) was used to inoculate the 96-well plates. Staining of the biofilm was performed at 24, 48 and 72 h. The wild-type Xac showed an increased biofilm biomass as revealed by CV staining while the biofilm biomass of the fliC and flgE mutants was reduced. However, these strains were still able to attach and develop a biofilm at 72 h (Fig. 3b).

**Role of the flagellum in the structural development of Xac biofilms**

We previously observed by CLSM that GFP-tagged Xanthomonas strains develop three-dimensional biofilm structures in static cultures in minimal medium, by using chambered cover slides (Rigano et al., 2007b; Torres et al., 2007). During the initial attachment Xac first contacted the glass surface in a polar fashion before becoming attached laterally. The colony of attached bacteria then spread over the entire well (Fig. 4a, day 3 and data not shown). After this first step, Xac developed microcolonies that increased in number (Fig. 4a). A more structured biofilm was formed from these microcolonies after 4 days (Fig. 4a). At higher concentrations of Xac, biofilms formed by the fliC and flgE mutants were reduced. However, these strains were still able to attach and develop a biofilm at 72 h (Fig. 3b).
magnifications, it was possible to observe that the bacteria were linked predominantly through lateral interactions. After 5 days, wild-type Xac developed mushroom-type structures separated by extensive water channels (Fig. 4e). At this time, some groups of cells (pioneer cells) spread from the biofilm to other locations (Fig. 4e).
Although the fliC and flgE mutants showed no apparent alteration in the first steps of cell attachment, they were not able to occupy the whole surface area of the chamber, in contrast to the wild-type strain (Fig. 4a, d). After 3 days, both mutants were able to develop aggregates (Fig. 4a, day 4) that later formed structures similar to those of the wild-type but thinner (Fig. 4d). Furthermore, these mutants failed to form a mature biofilm (Fig. 4e). These visual observations were confirmed by quantification of two variables, mean thickness and biofilm volume, by using the COMSTAT program (Fig. 4b, c).

The flagellum is involved in mature biofilm formation on leaf surfaces and is required for bacterial growth and canker development

To investigate if the flagellar mutants were affected in virulence, canker development, bacterial growth and biofilm formation, we evaluated the mutant strains in the lemon leaves infection model. The plants were spray-inoculated with Xac wild-type, fliC and flgE mutant strains. Fig. 5(a) shows canker symptoms on lemon leaves 22 days p.i. at a concentration of $1 \times 10^5$ and $1 \times 10^4$ c.f.u ml$^{-1}$. Quantification of canker lesions in lemon leaves was done at 22 days p.i. The flgE mutant was severely compromised in its ability to cause canker disease in citrus leaves (Fig. 5b). Comparison of bacterial numbers in planta revealed no significant statistical difference between Xac wild-type and the fliC mutant. However, the flgE mutant showed a statistically significant reduction in growth compared with the Xac wild-type (Fig. 5c).

When biofilm was analysed at 6 days p.i. on lemon leaves, the wild-type strain showed aggregation within a typical biofilm, as observed previously (Rigano et al., 2007b) (Fig. 5d). However, biofilms from the fliC and flgE mutants developed in a more dispersed fashion with either few or no aggregated structures. The flgE mutant was more affected than the fliC mutant (Fig. 5d).
DISCUSSION

In the present study we have addressed the importance of motility and the role of the flagellum in the development of mature biofilm structures in Xac. Previous work from our group showed that a wild-type Xac strain was able to attach and develop a biofilm with an organized structure on both abiotic and biotic surfaces. Similar structures were also seen within the cankers of infected plants (Rigano et al., 2007b). In addition, it has been shown that the xanthan polysaccharide in Xac is essential in the formation of microcolonies and the subsequent development of more complex structures (Rigano et al., 2007b). In the present work, we show evidence that the flagellum is essential for the establishment of a mature biofilm in Xac, that EPS is required for a flagellum-independent type of motility (‘sliding’) and that both sliding motility and swimming motility (flagellum-dependent) are regulated by the rpf/DSF system.

In addition to swimming motility in low-percentage agar media, many bacteria exhibit swarming motility on plates with higher (0.5–0.6% w/v) agar concentrations. Swarming motility is associated with changes in flagella number and/or placement, quorum sensing, and also with the production of wetting agents (Henrichsen, 1972; Kaiser, 2007; Overhage et al., 2007). Both flagella and type IV pili contribute to swarming in Pseudomonas aeruginosa (Harshey, 1994; Rashid & Kornberg, 2000) and probably in other bacteria (Köhler et al., 2000; Overhage et al., 2007). In contrast, Xac is capable of flagellum-independent surface translocation on 0.5% agar PYM. We consider that this type of motility is more consistent with sliding rather than swarming. Sliding motility has been defined as ‘a kind of translocation produced by expansive forces in a growing culture in combination with special surface properties of the cells resulting in reduced friction between cell and substrate’ (Murray & Kazmierczak, 2008). Unlike swarming, sliding motility does not require flagella (Henrichsen, 1972). It is not known how DSF signalling and EPS impinge on the proposed sliding motility, although it is possible that EPS plays a role in modifying the properties of the agar surface to promote motility.

Xac does not exhibit twitching motility (unpublished observations), and consequently it is tempting to speculate that the type of motility described here as sliding has a role in the formation of initial microcolonies which occurs in a flagellum-independent manner. This remains an open question however, since the factors that we demonstrate to influence sliding are DSF signalling, which has a very pleiotropic effect in Xanthomonas spp., and the EPS xanthan, which probably also has a number of different roles in biofilm formation (Dow et al., 2003).

Bacterial attachment, the formation of aggregates or microcolonies, and finally biofilm maturation have been recognized as the three stages of biofilm development that are responsive to diverse environmental signals (Stanley & Lazazzera, 2004). Mutations of the fliC or flgE gene significantly reduced, but did not eliminate, initial attachment to both abiotic and biotic surfaces, suggesting that other genes may be involved in this process. CLSM revealed that in chambered slides, fliC and flgE mutants do not develop the structured organization seen in wild-type Xac. A typical biofilm contains bacteria tightly packed in hexagonal arrays separated by water-filled channels that allow nutrient flux between the towers of cells, and develops multicellular mushroom-shaped structures. The biofilms of the flagellar mutants lack water channels, which may explain why these mutants produce thinner structures that are reflected in the low CV staining. Overall, the data suggest that the flagellum plays a major role in the formation of the mature Xac biofilm. However, it remains unclear whether this deficiency is due to the absence of the flagellum structure or the absence of motility.

Similarly to the findings described here, a study in Xcc reported that the absence of flagellin affected the swimming motility of the bacteria as expected but did not alter virulence (Lee et al., 2003). However, results obtained with other vascular pathogens such asRalstonia solanacearum, Pantoea stewartii and Dickeya dadantii showed that the flagellum-dependent motility was required for biofilm formation and systemic colonization of the host plant (Herrera et al., 2008; Jahn et al., 2008; Trans-Kersten et al., 2001). Although the mutation in the fliC gene in Xac affects the formation of biofilm structures, these strains are not significantly less virulent. The mutation in the flgE gene, in contrast, affects both the ability to form biofilms and virulence. As far as we are aware, this is the first study to establish the participation of FlgE in biofilm formation and virulence in a Xanthomonas species.

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