The Transcriptional Activator *rfiA* Is Quorum-Sensing Regulated by Cotranscription with the *luxI* Homolog *pcoI* and Is Essential for Plant Virulence in *Pseudomonas corrugata*

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The gram-negative phytopathogen *Pseudomonas corrugata* has an acyl-homoserine lactone (AHL) quorum-sensing (QS) system called PcoI/PcoR that is involved in virulence on tomato. This work identifies, downstream of *pcoI*, a gene designated *rfiA*, which we demonstrate is directly linked to QS by cotranscription with *pcoI*. The deduced RfiA protein contains a DNA-binding domain characteristic of the LuxR family but lacks the autoinducer-binding terminus characteristic of the QS LuxR-family proteins. We also identified, downstream of *rfiA*, an operon designated *pcoABC*, encoding for the three components of a tripartite resistance nodulation-cell-division (RND) transporter system. The expression of *pcoABC* is regulated by RfiA. We found that lipodepsipeptide (LDP) production is cell density dependent and mutants of *pcoI*, *pcoR*, and *rfiA* are unable to inhibit the growth of the LDP-sensitive microorganisms *Rhodotorula pilimanae* and *Bacillus megaterium*. *P. corrugata* *rfiA* mutants were significantly reduced in their ability to cause necrosis development in tomato pith. In addition, it was established that PcoR in the absence of AHL also played a role in virulence on tomato. A model for the role of PcoI, PcoR, and RfiA in tomato pith necrosis is presented.

Pseudomonas corrugata (Roberts and Scarlett 1981 emend. Sutra et al. 1997) is a ubiquitous bacterium isolated from a wide variety of sources (Catara 2007). It was first described by Scarlett and associates (1978) as the causal agent of tomato pith necrosis (TPN) and was later isolated in association with TPN in all the tomato-growing areas of the world. *P. corrugata* has also been identified occasionally as causal agent of pepper pith necrosis (Lopez et al. 1988) and has been reported once in chrysanthemum (Fiori 1992) and geranium (Magyarosy and Buchanan 1995). The characteristic symptom of the disease is stem pith necrosis; the pith may appear as necrotic and frequently disaggregated in the core. The disease is commonly observed in the field and greenhouse where conditions seem

more favorable: free water on leaves and stem surfaces and high soil nitrogen levels, which lead to more tender vegetation, promote *P. corrugata* infection (Carroll et al. 1992; Naumann et al. 1989; Scarlett et al. 1978). *P. corrugata* has been isolated from nondiseased plants, mainly as an endophyte from the rhizosphere, and has also been recovered from bulk soil (Catara 2007). Interestingly, *P. corrugata* has an antimicrobial activity in vitro against a large number of microorganisms (including gram-negative and gram-positive bacteria, Chromista, yeast, and fungi) and has been successfully tested as a biological control agent against plant pathogens in different pathosystems (Catara 2007; Choi et al. 2003; Schmidt et al. 2004; Zhou and Paulitz 1993).

P. corrugata produces the phytotoxic and antimicrobial cationic lipodepsipeptides (LDP), corpetin A and corpeptin B, and the lipodepsinona peptide cormycin A, which are thought to act as major virulence factors (Emanuele et al. 1998; Scaloni et al. 2004). *P. corrugata* has also been reported as producing alginate (Fett et al. 1996) and a siderophore, corrugatin, which could be involved in pathogenicity and colonization (Risse et al. 1998). Antimicrobial activity is probably a result of the production of LDP (Emanuele et al. 1998; Scaloni et al. 2004) and other substances such as inhibitory volatiles (Fernando et al. 2005), hydrogen cyanide (Ramette et al. 2003), and pyrrolnitrin (Garbeva et al. 2004).

Recently, we demonstrated that quorum sensing (QS) is intimately linked to *P. corrugata* virulence (Licciardello et al. 2007). QS allows bacterial populations to coordinate the expression of some traits in a cell-density-dependent manner and relies on the production and response to signal molecules (Fuqua et al. 1994). *N*-acyl homoserine lactones (AHL) are most commonly used as QS signal molecules in gram-negative bacteria (Fuqua and Greenberg 2002). The paradigm of the production and response to AHL is the *Vibrio fischeri* LuxI/R system, consisting of an AHL synthase member of the LuxI family and a transcriptional regulator belonging to the LuxR family. As the density of the population increases, so does the concentration of AHL; and, once the latter reaches a critical threshold concentration, the AHL signal molecule binds the cognate LuxR-family sensor/regulator, which activates or represses target gene expression (Fuqua and Greenberg 2002).

In plant-pathogenic bacteria, expression of virulence factors is often dependent on AHL QS; as, for example, conjugation in *Agrobacterium tumefaciens*, cell-wall-degrading enzyme pro-

The nucleotide sequence data is available in the GenBank/EMBL/DBJ database under accession number EF189721.

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*The e-Xtra logo stands for “electronic extra” and indicates that Figure 4 appears in color online.

duction in *Erwinia carotovora*, extracellular polysaccharide production in *Pantoea stewartii* subsp. *stewartii*, and traits involved in survival, host invasion, and virulence in *Pseudomonas syringae* pv. *syringae* (Barnard and Salmond 2007; Dulla and Lindow 2008; Quinones et al. 2004, 2005; von Bodman and Farrand 1995; von Bodman et al. 2003; Whitehead et al. 2001; Zhang et al. 1993). Generally, phytotoxin production is not regulated by QS (Bender et al. 1999). Nevertheless, recently, it has been demonstrated that the biosynthesis and export of toxoflavin by the rice pathogen *Burkholderia glumae* requires the regulator ToxJ, the expression of which is regulated by the TofI/R QS system (Kim et al. 2004).

Our previous studies demonstrated that *P. corrugata* CFBP 5454 produces *N*-hexanoyl-L-homoserine lactone (C6-HSL), 3-oxo-C6-HSL, and C8-HSL and this ability is well conserved within the species because strains with different origins and phenotypic and genetic characters produced the same AHL at comparable levels (Licciardello et al. 2007). The AHL QS system of *P. corrugata* was designated PcoI/PcoR, consisting of an AHL synthase, PcoI, and a transcriptional sensor/regulator belonging to the LuxR family protein, PcoR (Licciardello et al. 2007). The contribution of QS to TPN and antimicrobial activity was previously demonstrated (Licciardello et al. 2007). Interestingly, only the *P. corrugata* *pcoR* mutant and not the *pcoI* mutant induced significantly attenuated disease symptoms in tomato, suggesting an intricate network of regulation. To our knowledge, the identification and analysis of *P. corrugata* QS is the only available molecular study on this bacterial pathogen. The importance and conservation of QS in *P. corrugata* as well as the absence of further genetic and molecular information prompted us to investigate the DNA sequence adjacent to the QS locus because, often, QS-regulated genes are genetically linked. Downstream of *pcoI*, we identified a gene that encodes a transcriptional regulator of the LuxR family, which we designated *rfiA*. Importantly, *rfiA* is cotranscribed with *pcoI*. Further sequencing of the downstream region revealed a 5,611-bp operon encoding for a tripartite resistance nodulation-cell-division (RND) transporter system. The role of RfiA, QS, and the transporter system in LDP regulation and tomato virulence was investigated.

RESULTS

Sequencing and analysis of genes adjacent to the QS *pcoI/pcoR* locus.

Our previous studies determined that *P. corrugata* possesses an AHL QS system designated PcoI/R which plays a role in *P. corrugata* virulence (Licciardello et al. 2007). To further characterize the DNA region linked to the QS *pcoI/R* locus within a previously sequenced 2.7-kb region, we sequenced an additional 6.7-kb DNA region adjacent to and downstream of the *pcoI* gene (Fig. 1).

DNA sequence analysis of the newly sequenced region revealed four potential open reading frames (ORF) downstream of the *pcoI/R* locus. The first ORF was closely associated with the *pcoI* locus, separated by only 32 bp of DNA. This ORF shared 91% identity with the *rfiA* gene sequence of *P. corrugata* 2140R, which is predicted to encode a transcriptional activator involved in the biosynthesis of antimicrobial compounds (this gene has only been annotated as a possible regulator of fungal inhibition without experimental evidence; GenBank accession number AF199370).

In addition, three ORF, designated *pcoA*, *pcoB*, and *pcoC*, were identified downstream of *rfiA*, transcribed in the opposite direction. The stop codon (TGA) of *pcoA* overlapped the start coding region of *pcoB* by 58 bp and, similarly, the stop codon (TGA) of *pcoB* overlapped the start coding region of *pcoC* by

4 bp, which indicates that *pcoA*, *pcoB*, and *pcoC* are most probably organized as an operon. The *pcoA*, *-B*, and *-C* genes are 1,428, 1,161, and 3,084 bp in size, respectively, and composition analysis revealed that the G+C contents range from 62 to 66%. Nucleotide sequence analysis revealed that these three genes encode proteins, which constitute a multidrug efflux system (discussed below).

The *rfiA* gene encodes a regulator belonging to the LuxR family.

Nucleotide sequence analysis revealed that *rfiA* encoded a predicted 288-amino-acid protein. RfiA displayed 94% sequence identity (96% homology) with a putative transcriptional activator of *P. corrugata* 2140R (AAG28559) and 51% identity (65% homology) with the protein SalA of *P. syringae* MIS38 (BAF76155). RfiA also showed 45% identity (62% homology) with the well-characterized SalA protein of *P. syringae* B301D (Lu et al. 2002; Wang et al. 2006). RfiA contains a DNA-binding domain with a helix-turn-helix motif characteristic of the LuxR family near its carboxy-terminus (amino acids 215 to 265). It consists of four helix bundles in which the central helices form the HTH motif (Fuqua and Greenberg 2002).

Unlike PcoR, RfiA does not contain the five highly conserved amino acids (Trp59, Tyr69, Asp79, Pro80, and Gly121) in the *N*-terminus characteristic of the autoinducer-binding subfamily of the QS LuxR-family proteins (Fuqua et al. 1996). Moreover, RfiA lacks the characteristic "acid pocket" composed of four highly invariant residues (Asp11, Asp12, Asp55, and Lys105) of members of the response regulator subfamily domains (Parkinson and Kofoed 1992).

The *rfiA* gene is part of the same transcriptional unit as *pcoI*.

The first putative triplet codon for the *rfiA* ORF was located only 32 bp downstream of the stop codon of the *pcoI* AHL synthase gene; therefore, it was of interest to determine whether the *rfiA* was part of the same transcriptional unit of *pcoI*. We used reverse transcription with three sets of oligonucleotides (Table 1) to determine whether the *pcoI* and *rfiA* genes are cotranscribed. Specifically, primer set PCR1 was used to amplify a segment of the *pcoI* gene, PCR2 to capture an intervening region between *pcoI* and *rfiA*, and PCR3 to amplify a region of the *rfiA* gene. As expected, the PCR1 primer set yielded the expected 101-bp amplification product in the wild-type (WT) strain but not in the *pcoI* transposon insertion mutant GL1 (data not shown). Primer set PCR2 yielded a 614-bp amplicon in the WT strain, indicating that *pcoI* and *rfiA* are cotranscribed. The *rfiA* PCR3-specific amplicon was absent in the GL1 *pcoI* mutant strain, indicating that the transposon inserted in the *pcoI* gene had a polar effect on the cotranscribed *rfiA* gene. Therefore, it was concluded that GL1 is a double *pcoI/rfiA* mutant.

PcoR and AHL activate *pcoI* and *rfiA* expression.

In order to determine whether PcoR regulates *pcoI* expression, the promoter region of *pcoI* was cloned in the broad-host-range low-copy-number β -galactosidase promoter probe vector pMP220, yielding pMPPcoI. Promoter activity was determined in response to the growth phase of the WT strain CFBP 5454 and QS mutant derivatives. *PcoI* promoter activity was undetectable in the *pcoR* or *pcoI* *P. corrugata* mutants, indicating that it was under positive autoregulation, as is often the case with the *luxI/R* systems (Fig. 2A). The presence of a *lux*-box consensus sequence in the upstream region of *pcoI* further supported the idea that PcoR regulated *pcoI* expression. Exogenous addition of C6-HSL, C8-HSL, and 3-oxo-C6-HSL (1 μ M), the three signal molecules produced by *P. corrugata*

CFBP 5454 (Licciardello et al. 2007), to the *pcol* GL1 mutant harboring pMPPcoI restored promoter activity (Fig. 2B). Therefore, it seems that all three AHL are able to interact and activate PcoR; however, C8-HSL had the highest inducing activity ($P < 0.05$). Because *pcol* and *rflA* are part of the same transcriptional unit, it was concluded that *rflA* was also under the control of *pcol* promoter region.

Fig. 1. Gene map of a 9.8-kb DNA region of *Pseudomonas corrugata* CFBP 5454. The *rflA* gene and *pcoABC* operon are localized downstream of the *pcol*/R quorum-sensing system. *rflA*, oriented in the same direction as *pcol*, encodes for an LuxR transcriptional activator protein. Genes *pcoA*, *pcoB*, and *pcoC* form the *pcoABC* operon, which is divergently oriented, and encode for a resistance nodulation-cell-division transporter system.

The *pcoABC* operon encodes an RND transporter system and is regulated by RflA.

The *pcoABC* operon encodes three components of a tripartite RND transporter system. This is composed of an outer membrane protein (PcoA), a periplasmic membrane fusion protein (PcoB), and a cytoplasmic RND transporter (PcoC) (Saier and Paulsen 2001). The PcoABC efflux system is highly homologous to the PseABC RND efflux system described in *P. syringae* pv. *syringae* B301D. In *P. syringae*, this system is localized in the *syr-syp* genomic island, which is involved in the secretion of lipopeptide toxins such as syringomicin and syringopeptins (Kang and Gross 2005). The PcoA-predicted protein (475 amino acids long) showed 67% identity and 82% similarity with PseA (ABN45752) and contains the TolC conserved domain. The putative PcoB protein, composed of 386 amino acids, showed 66% identity (80% similarity) with PseB (ABN45753) and contains the AcrA conserved domain, anchored to the inner membrane. The 1,027-amino-acid PcoC protein showed the highest homology (77% identity, 87% similarity) with PseC (ABN45754) and contains the domain AcrB, composed of 12 transmembrane α -helices and two large hydrophilic loops. The activity of a *pcoA::lacZ* promoter probe reporter construct expressed in GL1 (*pcol/rflA* mutant), GL2

Table 1. Bacterial strains, plasmids, and oligonucleotides used in this study

Strains, plasmids, or oligonucleotides	Characteristics or sequences ^a	References or sources ^b
<i>Pseudomonas corrugata</i>		
CFBP 5454	Wild type, source of <i>pcol</i> and <i>pcoR</i>	CFBP
GL1	<i>pcol</i> 362::Tn5, Km ^r	Licciardello et al. 2007
GL2	<i>pcoR</i> 76::Tn5, Km ^r	Licciardello et al. 2007
GLRflA	<i>rflA</i> :: pKnock, Km ^r	This study
GLPCOA	<i>pcoA</i> :: pKnock, Km ^r	This study
<i>Escherichia coli</i>		
pLc3.34	pLAFR3 containing <i>P. corrugata</i> CFBP 5454 DNA, Tc ^r	DISTEF
DH5 α	F2 f80dlacZDM15 D(lacZYA-argF)U169 endA1 recA1 hsdR17 deoR gyrA96 thi-1 relA1 supE44	Sambrook et al. 1989
CC118 λ pir	Δ (ara, leu)7697 araD139 Δ lacX74 galE galK phoA20 thi-1 rpsE rpoB (Rf)	Herrero et al. 1990
Plasmids		
pCR2.1	Cloning vector TA, Amp ^r	Invitrogen
pRK2013	Km ^r Tra ⁺ Mob ⁺ ColE1 replicon	Figurski and Helinski 1979
pMOSBlue	Cloning vector, Amp ^r	Amersham-Pharmacia
pMP220	Promoter probe vector, IncP Tc ^r	Spaink et al. 1987
pBBR1MCS-5	Broad-host-range vector, Gm ^r	Kovach et al. 1975
pKNOCK-Km ^r	Mobilizable suicide vector, Km ^r	Alexeyev, 1999
pGEM-T	Cloning vector; Amp ^r	Promega
pMOPcoI	<i>pcol</i> promoter cloned in pMOSblue; Amp ^r	This study
pGEPcoA	<i>pcoA</i> promoter cloned in pGem-T; Amp ^r	This study
pMPPcoI	<i>pcol</i> promoter cloned in pMP220	This study
pMPPcoA	<i>pcoA</i> promoter cloned in pMP220	This study
pKMRflA	pKNOCK containing an internal fragment of <i>P. corrugata</i> CFBP 5454 <i>rflA</i> gene	This study
pKMPcoA	pKNOCK containing an internal fragment of <i>P. corrugata</i> CFBP 5454 <i>pcoA</i> gene	This study
pBBR-RflA	pBBR1MCS-5 containing the full length <i>P. corrugata</i> CFBP 5454 <i>rflA</i> gene	This study
Oligonucleotides		
PrpcoI-fw	5'-CGGTACCCACTGTGCATCAGTGAAAGC-3'	This study
PrpcoI-rew	5'-GTCTAGAGGGATTTTCGAGTAAGATG-3'	This study
PrpcoA-fw	5'-GCTCTAGATTGAAGACATGCATAGG-3'	This study
PrpcoA-rew	5'-GGGTACCTATCGCGAGCAGGCTCG-3'	This study
RflAKn-fw	5'-CTCTAGAACGACCTGATACTTACCG-3'	This study
RflAKn-rew	5'-GGGTACCTTATGGCAGCAGCTTCAG-3'	This study
PcoAKn-fw	5'-CAGGATCCTGGGAGATCGACCTGTTCG-3'	This study
PcoAKn-rew	5'-GAAAGCTTTTGAACGACTCACGAAAGCC-3'	This study
RflA-fw	5'-AGAAGCTTCTATAACACCAAGACTCTG-3'	This study
RflA-rew	5'-TTGGATCCAACGTTCTATACGGCTTGG-3'	This study
PCR1-Fw	5'-CATCAGGCCTCACTCACACACT-3'	This study
PCR1-Rew	5'-GTTGGCCTGGGTGACTTCTC-3'	This study
PCR2-Fw	5'-CATCAGGCCTCACTCACACACT-3'	This study
PCR2-Rew	5'-ATGCGTTGCATCGACAGGTAT-3'	This study
PCR3-Fw	5'-CTCTAGAACGACCTCATACTTACCG-3'	This study
PCR3-Rew	5'-GGGTACCTTATGGCAGCAGCTTCAG-3'	This study

^a Km^r, Rf^r, Tc^r, Amp^r, and Gm^r indicate resistance to kanamycin, rifampicin, tetracycline, ampicillin, and gentamicin, respectively.

^b CFBP = Collection Francaise de Bacteries Phytopathogenes, Angers, France; DISTEF = Dipartimento di Scienze e Tecnologie Fitosanitarie, Catania, Italy.

(*pcoR* mutant), and GLRFIA (*rflA* mutant) was 10-fold lower compared with the activity measured in the parent strain, indicating that *pcoA* is positively regulated by QS or RfIA. We then introduced a plasmid constitutively expressing *rflA* (i.e., pBBRRfIA, where *rflA* is expressed from the *lac* promoter) in both GL1 and GL2 *P. corrugata* mutants harboring the *pcoA::lacZ* transcriptional fusion. The expression of *rflA* in

trans restored *pcoA* promoter activity in both mutants to the parent strain levels, whereas exogenous addition of the three AHL produced by *P. corrugata* had no effect on the promoter activity (Fig. 2C). This data indicates a hierarchical regulatory scenario in which the PcoR-AHL complex regulates the *pcoI/rflA* operon and, in turn, RfIA activates *pcoABC*. Therefore, it was concluded that the expression of the *pcoABC* transporter system is indirectly controlled by QS via RfIA.

RfIA regulates LDP secretion.

Our previous studies demonstrated that the antimicrobial activity of *P. corrugata* was altered in QS mutants (Licciardello et al. 2007). Because *P. corrugata* is known to produce the antimicrobial and phytotoxic LDP cormycin A and corpeptines A and B (Emanuele et al. 1998; Scaloni et al. 2004), we investigated whether LDP production is population-density dependent and whether it is regulated by the PcoI/R QS system or by RfIA.

We performed the production profile during bacterial growth of AHL and LDP in *P. corrugata* CFBP 5454 as described in the Materials and Methods section. AHL abundance was indirectly quantified by measuring the amount of violacein produced by the biosensor *Chromobacterium violaceum* CV026 (Fig. 3). LDP production, on the other hand, was assayed by inhibition of in vitro growth of *Rhodotorula pilimanae* (cormycin) and *Bacillus megaterium* (cormycin and corpeptines). AHL and LDP production were first detected in culture after 16 h of growth with cell concentration as low as 1.6×10^8 CFU ml⁻¹. Culture filtrates displayed inhibition zones of 4.5 and 2.5 mm against *R. pilimanae* and *B. megaterium*, respectively (Fig. 3). However, LDP and AHL concentration increased rapidly at higher bacterial densities; maximum LDP concentration was observed at cell densities over 6×10^9 CFU ml⁻¹ (late exponential growth phase) when inhibition zones against *R. pilimanae* and *B. megaterium* were 7 and 5 mm, respectively (Fig. 3). Thus, it was concluded that LDP synthesis in *P. corrugata* CFBP 5454 occurs in a cell-density-dependent manner, with a trend similar to AHL production.

GL1 (*pcoI/rflA* double mutant), GL2 (*pcoR* mutant), and GLRFIA (*rflA* mutant) did not inhibit the growth of LDP-sensitive indicator microbes, suggesting they were unable to produce and/or secrete LDP (Fig. 4A). Expression in trans of the

Fig. 2. *pcoI* and *pcoA* promoter activities in the parent strain *Pseudomonas corrugata* CFBP 5454 and mutant derivatives. Values are averages of at least three independent experiments; the standard deviation of the mean are indicated by the bars. **A**, *pcoI* promoter activities in the parent strain and quorum-sensing (QS) mutant derivatives obtained by using the *pcoI-lacZ* reporter construct. *pcoI* gene expression was under positive feedback QS regulation. **B**, Effect of exogenous *N*-hexanoyl-L-homoserine lactone (C6-HSL), C8-HSL, and C6-3-oxo-HSL on *pcoI* transcription as measured using GL1 (pMPPcoI). The three lactones, and particularly C8-HSL, are effective in restoring *pcoI* promoter activity in the GL1 mutant (*pcoI*⁻) in presence of the *pcoI-lacZ* reporter construct. **C**, *pcoABC* operon promoter activities in the parent strain *P. corrugata* 5454 and QS mutant derivatives by using the *pcoA-lacZ* reporter construct. Expression of the *pcoABC* operon requires both the acyl-homoserine lactone (AHL)-QS system and the RfIA activator but promoter activities were fully restored in the QS mutant derivatives only by the presence of pBBRRfIA, which carried the intact *rflA* gene, and not by exogenous AHL.

Fig. 3. Lipopeptide and acyl-homoserine lactone (AHL) production in *Pseudomonas corrugata* CFBP 5454 during growth phase. Lipopeptide production and secretion was evaluated by antimicrobial activity of the culture filtrates (10×) recovered at 8, 16, 24, 48, and 72 h after inoculation on inducing conditions against *Rhodotorula pilimanae* and *Bacillus megaterium* indicator microorganisms on potato dextrose agar medium. The values of the inhibition zones (mm) are indicated by the bars. Values are the means of three replications. Results presented are representative of three different experiments. AHL biosynthesis was indirectly measured as violacein produced by a bioassay using the biosensor *Chromobacterium violaceum* CV026. Values are the means of 10 replications.

Fig. 4. Biological activity of the parent strain *Pseudomonas corrugata* CFBP 5454 and mutant derivatives. **A**, Lipopeptide (LDP) production. Culture filtrates (10 $\times$) were tested on potato dextrose agar medium overlaid with *Rhodotorula pilimanae* and *Bacillus megaterium* indicator microorganisms: 1 = wild type (WT), 2 = GL1 (*pcoI/rfiA* double mutant), 3 = GL1 (*pcoI/rfiA* double mutant) + pBBRRfiA, 4 = GLPCOA (*pcoA* mutant), 5 = GLRFIA (*rfiA* mutant), 6 = GL2 (*pcoR* mutant). No LDP were detected in quorum-sensing and RfiA mutants. The complementation of the GL1 (*pcoI/rfiA* double mutant) with the intact *rfiA* gene through the pBBRRfiA plasmid could restore LDP production. The inhibition zone produced by the GLPCOA (*pcoA* mutant) culture filtrate was slightly reduced compared with the WT. Results presented are representative of two different experiments with three replicates for each. **B**, Mean disease index (DI) of tomato plants inoculated with the parent strain *P. corrugata* CFBP 5454 (WT) and GL1 (*pcoI/rfiA* double mutant), GL2 (*pcoR* mutant), GLRFIA (*rfiA* mutant), and GLPCOA (*pcoA* mutant) mutant derivatives, GL1 with the addition of exogenous acyl-homoserine lactones (GL1+AHL), and GL1 and GL2 complemented in trans with intact *rfiA* (GL1+RfiA and GL2+RfiA, respectively). Error bars represent the standard deviation of the mean DI obtained from 20 individual disease rating scores. Below the graphic, the presence of AHL (either produced by the bacterial strain or provided exogenously) and the regulators PcoR and RfiA in the interaction with tomato plant are reported. **C**, Stem pith symptoms in tomato plants prick inoculated with bacterial cells of the parent strain and GLRFIA (*rfiA* mutant) and GLPCOA (*pcoA* mutant). Plants inoculated with the GLRFIA mutants did not show necrosis or hollowing of the pith, whereas the trace of the wound caused by inoculation is still visible.

rfiA gene in the GL1 mutant by introducing pBBRRfiA restored the antimicrobial activity to the WT strain level, which suggested that RfiA was regulating LDP production (Fig. 4A). A mutation in the *pcoA* transporter gene led only to the partial loss of antimicrobial activity because GLPCOA (*pcoA* mutant) culture filtrates ($P < 0.05$) (Fig. 4A) displayed inhibition zones against *R. pilimanae* and *B. megaterium* of approximately 48 and 46% respectively, compared with those of the parental strain CFBP 5454. This data indicated that the PcoABC efflux system had a role in LDP secretion but it was not essential because, most probably, *P. corrugata* possesses other efflux pumps able to transport LDP.

The RfiA and PcoR regulators are involved in *P. corrugata* tomato virulence.

The contributions of RfiA, PcoABC, and the PcoI/R QS system were examined in the formation of stem pith necrosis. The parent strain was compared with the mutants on the basis of its ability to induce symptoms in artificially inoculated tomato stem pith. Disease rating scores were considered and used to calculate a mean disease index (DI). The summary of these in planta infection studies is presented in Figure 4B and C. Plants inoculated with GL1 (*pcoI/rfiA* double mutant) and GLPCOA (*pcoA* mutant) displayed symptoms and DI similar to that of the parent strain (Fig. 4B). Stem-pith-infected tissues appeared as necrotic, partially dehydrated, and hollow (Fig. 4C). Most of the plants inoculated with GL2 (*pcoR* mutant) and GLRFIA (*rfiA* mutant) showed only a dark discoloration along the pin puncture and were scored 1 on the susceptibility scale (Fig. 4B). On the remaining plants, lesions were significantly reduced in length compared with those caused by the parent strain and were only discolored (Fig. 4C). The typical necrosis of the TPN syndrome was not observed (score 2). Expression of *rfiA* in trans in the GL2 mutant restored the virulence nearly to the same levels of that of the parent strain whereas, in GL1, virulence was maintained. Therefore, it was concluded that RfiA is of pivotal importance in virulence. In order to further assess the role of AHL, we inoculated GL1 with 5 μ l of C6-HSL and C8-HSL (5 μ g ml⁻¹) by pipetting into the inoculation wounds. GL1 exposed to exogenously added AHL showed symptoms similar to those induced by the GL2 and GLRFIA mutants (i.e., reduction of lesion extension and necrosis) (Fig. 4B). These results indicated that there were two conditions in which *P. corrugata* was virulent toward tomato: presence of PcoR in the absence of AHL or with RfiA. Therefore, it was concluded that, in the presence of AHL, PcoR activates gene expression of virulence factors via RfiA. In the absence of AHL signals, PcoR, however, can induce virulence gene expression via another yet unknown mechanism (Fig. 5). Finally, it was observed that the *pcoABC* transport system was not pivotal for virulence because a *pcoA* mutant retained its ability to be pathogenic (Fig. 4B and C). Most probably, therefore, other LDP transport efflux systems are operational in *P. corrugata*.

DISCUSSION

In this study, we have further demonstrated that, in *P. corrugata*, virulence is regulated by QS. Injection of high-density inoculum within the stem of a wide range of plant species, including members of *Solanaceae* (egg plant and bell pepper) and species in other families (cucumber, zucchini, bean, and cauliflower) led to pith necrosis (Siverio et al. 1993; Sutra et al. 1997; Catara et al. 1997, 2002). In nature, however, the disease is only widespread in tomato, with just a few cases of infections being reported on pepper and one report on chrysanthemum and geranium (Catara 2007). Therefore, it is tempting to speculate that, in some plants, *P. corrugata* is not able to reach the bacte-

rial density necessary for a synchronous concerted attack through activation of effectors or virulence factors.

P. corrugata has a conserved AHL QS system designated PcoI/R which plays a significant role in tomato virulence (Licciardello et al. 2007). In this study, we sequenced the DNA region downstream of the AHL QS locus, and this led to the identification of genes of considerable phytopathological interest. A regulatory gene, *rfiA*, located at the right border of the *pcoI* gene, was identified as a member of the LuxR family of regulatory proteins. RfiA contains an HTH DNA motif in the C-terminus characteristic of the LuxR family of bacterial regulatory proteins but lacks the conserved residues at the N-terminus characteristic of the two major LuxR subfamilies (Finney et al. 2002). Due to high sequence similarity and domain homologies, RfiA could belong to the novel LuxR subfamily described for the SalA and SyrF regulators in *P. syringae* pv. *syringae* B301D (Lu et al. 2002). SalA and SyrF are localized in the *syr-syp* genomic island and regulate syringomycin and syringopeptins production (Wang et al. 2006). However, in *P. syringae* pv. *syringae*, the production of LDP is not regulated by QS but is directly controlled by a complex regulatory cascade which involves the two-component sensor/regulator GacS/GacA and the transcriptional activators SalA and SyrF (Wang et al. 2006). Interestingly, we found that, in *P. corrugata*, the newly characterized *rfiA* forms an operon with the *pcoI* AHL-synthase gene. To our knowledge, this is the first report of a *luxI* homolog co-transcribed with a regulatory gene. We demonstrated that PcoR activates *pcoI* expression in the presence of exogenous AHL via a typical positive-feedback regulatory loop. Because *pcoI* and *rfiA* constitute an operon, the expression of *rfiA* is directly regulated by the PcoR-AHL complex.

Further sequencing revealed the presence of an RND-type efflux system next to *rfiA*, designated *pcoABC*. The RND superfamily of multidrug transporters is composed of eight phylogenetic families involved in the export of several compounds, including heavy metals, multiple drugs, and lipo-oligosaccharides (Murakami and Yamaguchi 2003). The predicted PcoABC system was highly homologous to the PseABC efflux system of *P. syringae* pv. *syringae* B301D, where it has a role in the secretion of syringomycin and syringopeptins (Kang and Gross 2005). We have shown that the *pcoABC* operon is under positive regulation by RfiA and, indirectly, by the PcoI/R system. Therefore, the regulation of the *P. corrugata* *pcoABC* operon occurs according to a hierarchical model. In late exponential growth, the bacterium activates AHL production, which results in the formation of the AHL-PcoR complex that most probably binds the *lux*-box element in the *pcoI* promoter and activates AHL-synthase expression as well as *rfiA* transcription. RfiA, in turn, activates the transcription of the *pcoABC* operon either directly or indirectly. Further experiments are needed to molecularly confirm this working model.

P. corrugata produces the LDP corpeptin A and corpeptin B, two isoforms consisting of 22-amino-acid residues, which are toxic when inoculated into tobacco leaves and are also antimicrobial against *B. megaterium* (Emanuele et al. 1998). Moreover, some of the strains also produce cormycin A, a lipodepsinonapeptide which shows in vitro inhibition of not only *B. megaterium* but also *R. pilimanae*, and exhibited phytotoxic activity (Scaloni et al. 2004). Mutants unable to produce toxins did not inhibit growth of the two target microorganisms and did not induce necrotic lesions in tomato leaves or stems (Chun and Leary 1989). Phenotype analysis of the *pcoI/rfiA*, *pcoR*, and *rfiA* mutants revealed that they were impaired in LDP secretion because their culture filtrates were unable to inhibit the growth of the *R. pilimanae* and *B. megaterium* target organisms. Moreover, time-course monitoring of antimicrobial activity of *P. corrugata* CFBP 5454 showed that LDP are not produced until high popu-

lation densities are reached. The importance of RfiA in the development of the disease symptoms in tomato plants was demonstrated by the reduction of lesions and the absence of necrosis within the stem pith tissues inoculated with the *rfiA* mutant compared with those inoculated with the WT strain CFBP 5454. Similarly, a *P. syringae* pv. *syringae* *salA* mutant failed to produce syringomycin and was significantly less virulent than the WT strain B301D (Lu et al. 2002). SalA is essential for the expression of *syrB1*, a biosynthesis gene for syringomycin (Lu et al. 2002). The PcoABC and PseABC RND transport systems of *P. corrugata* and *P. syringae* pv. *syringae*, respectively, share considerable homology. Both systems were shown to be involved in LDP secretion; however, both *pcoABC* and *pseABC* mutants were as virulent as their WT strains (Kang and Gross 2005; this study). In *P. syringae* pv. *syringae*, two additional ABC transporters, encoded by *syrD* and *sypD*, are also involved in the secretion of syringomycin and syringopeptins (Quigley et al. 1993). When the *pcoA* mutant was grown in laboratory culture media, it produced and secreted LDP, indicating that other LDP secretion mechanisms also exist in *P. corrugata*.

Even in *Burkholderia glumae*, phytotoxin production and transport is regulated indirectly by AHL QS via the ToxJ regulator. ToxJ is a transcriptional regulator similar to RfiA, which is regulated by the ToxI/R AHL QS system and, in turn, regulates toxoflavin production in a cell-density-dependent manner (Kim et al. 2004). ToxJ simultaneously activates the expression of the *toxABCDE* operon responsible for toxoflavin biosynthesis and the *toxFGHI* operon responsible for its transport (Kim et al. 2004). RfiA, SalA, SyrF, and ToxJ belong to a new emerging subfamily of LuxR-family regulators which, thus far, have been shown to be involved in virulence and the regulation of the RND-type transporters in toxin secretion in phytopathogenic bacteria. It is likely that these regulators are also involved in regulating other virulence loci, including LDP structural genes as well as other LDP secretion systems.

Plant pathogenicity studies revealed that PcoR and RfiA play a role in bacterial virulence in tomato (Fig. 4B). As shown by Licciardello and associates (2007), the GL1 mutant (which is a double *pcoI/rfiA* mutant) was as virulent in planta as the WT, suggesting that the sole presence of PcoR in the cell is sufficient for *P. corrugata* to be pathogenic. However, when PcoR is in the cell with AHL and no RfiA, *P. corrugata* is less virulent, as in the case of the addition of exogenous AHL to the *pcoI/rfiA* mutant. The presence of RfiA is crucial for bacterial virulence. Like other LuxR-type regulators, RfiA does not require AHL to

Fig. 5. Working model for acyl-homoserine lactone (AHL) quorum-sensing (QS) and RfiA involvement in *Pseudomonas corrugata* virulence. In the presence of AHL QS, virulence is regulated via RfiA. In absence of AHL, PcoR does not activate transcription of the *pcoI/rfiA* locus and virulence is regulated by PcoR.

be active; therefore, its complementation in trans is sufficient to restore pathogenicity in the *pcrR* mutant in absence of AHL. A working model of how the PcoI/PcoR/RfIA system could be involved in virulence, in which either QS regulates the production of RfIA or PcoR regulates virulence independently of AHL, is shown in Figure 5. Therefore, it is hypothesized that PcoR acts directly or indirectly in the absence of AHL as an activator or repressor of different virulence-associated genes. There are several reports of QS-LuxR family proteins which are active independently of AHL, as is the case for CarR of *Serratia* spp. (Cox et al. 1998), SmaR of *Erwinia* spp. (Slater et al. 2003), and EsaR of *Pantoea stewartii* (von Bodman et al. 1998). Further molecular studies of this working model may shed light on the mechanisms by which PcoR and RfIA contribute to virulence gene expression.

MATERIALS AND METHODS

Bacterial strains, plasmids, and media.

Pseudomonas corrugata strains, plasmids, and primers used in this study are listed in Table 1. *P. corrugata* strains were routinely grown at 28°C in either nutrient agar (Oxoid, Milan, Italy) plus 1% dextrose (NDA) or in Luria-Bertani (LB) agar. Transcriptional fusion plasmids for the various gene promoters based on the pMP220 promoter probe vector were constructed as follows. The 695-bp fragment containing the *pcrI* promoter region was amplified by polymerase chain reaction (PCR) by using Vent DNA polymerase (New England Biolabs, Frankfurt, Germany) according to the instructions of the supplier, genomic DNA of *P. corrugata* CFBP 5454 as the template, and oligonucleotides PrpcrI-fw and PrpcrI-rew (Table 1). The fragment was then cloned in pMOSBlue (Amersham Biosciences, Amersham, U.K.), yielding pMOPcrI, and verified by DNA sequencing. The *pcrI* promoter was then removed as a *KpnI-XbaI* fragment and cloned in the corresponding sites in pMP220, yielding pMPpcrI. Similarly, the *pcoA* promoter was amplified as a 220-bp fragment by using oligonucleotides PrpcoA-fw and PrpcoA-rew (Table 1) and cloned in pGEM-T, yielding pGEMpcoA. The *pcoA* promoter was then removed as a *KpnI-XbaI* fragment and cloned in pMP220, yielding pMPpcoA. The full-length *rflA* gene (912 bp) was amplified by PCR by using *Taq* DNA Polymerase Recombinant (Invitrogen, Milan, Italy) and primers RFIA-fw and RFIA-rew and cloned in pCR2.1 vector (Invitrogen). The fragment was then removed as a *BamHI-HindIII* and cloned in the corresponding sites in pBBR1MCS-5 (Kovach et al. 1995) according to the instructions of the supplier, yielding pBBR-RfIA. Antibiotics were added as required at the following final concentrations: ampicillin, 100 µg ml⁻¹; tetracycline, 15 µg ml⁻¹ (*Escherichia coli*) or 40 µg ml⁻¹ (*Pseudomonas* spp.); gentamicin, 10 µg ml⁻¹ (*E. coli*) or 30 µg ml⁻¹ (*Pseudomonas* spp.); and kanamycin, 50 µg ml⁻¹ (*E. coli* and *C. violaceum*) or 100 µg ml⁻¹ (*Pseudomonas* spp.).

Recombinant DNA techniques.

DNA manipulations, including digestion with restriction enzymes, agarose gel electrophoresis, purification of DNA fragments, ligation with T4 ligase, DNA hybridization, radioactive labeling by random priming, and transformation of *E. coli*, were performed as described by Sambrook and associates (1989). Southern hybridizations were performed by using N+Hybond membranes (Amersham Biosciences); plasmids were purified using Jet star columns (Genomed GmbH, Löhne, Germany) or by the alkaline lysis method (Birnboim 1983); total DNA from *Pseudomonas* spp. was isolated by Sarkosyl-pronase lysis as described previously (Better et al. 1983) or by a Puregene DNA isolation kit (Gentra Systems, Minneapolis, MN, U.S.A.), according to the manufacturer's instructions. Triparental matings

from *E. coli* to *P. corrugata* were carried out with the helper strain *E. coli* (pRK2013) (Figurski and Helinski 1979).

DNA sequencing and sequence analysis.

The cosmid pLC3.34 DNA insert was partially sequenced on both strands (Macrogen, Inc, Seoul, Korea). Sequencing for verification of cloned DNA inserts was determined by BMR-CRIBI (University of Padua, Italy). Homology searches of nucleotide and protein sequences were performed using the BLAST searching program (Altschul et al. 1990).

Cloning and construction of the genomic *rflA* and *pcoA* null mutants of *P. corrugata* CFBP 5454.

The *P. corrugata* CFBP 5454 *rflA* and *pcoA* genes were amplified, in part, by PCR in order to generate independent genomic mutations in *P. corrugata* CFBP 5454. The central part of the *rflA* gene was amplified by PCR as a 678-bp fragment using primers RfIAKn-fw and RfIAKn-rew (Table 1) and cloned as a *KpnI-XbaI* fragment in the corresponding sites in pKNOCK-Km, generating pKMRfIA. The central part of the *pcoA* gene was amplified by PCR as an 818-bp fragment using primers PcoAKn-fw and PcoAKn-rew (Table 1) cloned as a *BamHI-HindIII* fragment in the corresponding sites in pKNOCK-Km (Alexeyev 1999), generating pKMPcoA. The pKMRfIA and pKMPcoA plasmids were then used as a suicide delivery system in order to create *rflA* and *pcoA* knockout mutants through homologous recombination in strain CFBP 5454, as described previously (Alexeyev 1999), generating GLRFIA and GLPCOA, respectively. The fidelity of the marker exchange events was confirmed by Southern analysis (data not shown).

Reporter gene fusion assay.

β-Galactosidase activities were determined during growth in LB medium essentially as described by Miller (1972), with the modifications of Stachel and associates (1985). All experiments were performed in triplicate. β-Galactosidase activities were determined after a 20-ml LB medium culture started with an initial inoculum of 1.6 × 10⁸ CFU.

Reverse-transcriptase PCR analysis.

P. corrugata CFBP 5454 and GL1 and GLRFIA mutant derivatives were grown in LB medium to the exponential growth phase (14 h after inoculation); total RNA was isolated using a commercial RNA extraction kit (Purescript, Gentra) as recommended by manufacturer. For this purpose, triplicate samples (1 ml) were removed from *P. corrugata* cultures (optical density at 600 nm of 0.2 to 0.4) grown overnight. RNA samples were quantitatively analyzed by agarose gel electrophoresis. Following a DNase purification step by DNase I (Invitrogen), 200 ng of total RNA was used in each 20-µl reaction containing one unit of Superscript III reverse transcriptase (Invitrogen), 10 pmol random examer (Invitrogen), and 10 pmol dNTPs. Samples in which reverse transcriptase was not added were used as negative controls. The reverse transcription reaction was performed at 50°C for 1 h followed by 15 min at 65°C, according to the manufacturer's instructions. PCR reactions were performed using a Gene Amp PCR system 9700 (PE Applied Biosystem, Milan, Italy) under the following conditions: an initial 94°C for 3 min; followed by 35 cycles of 94°C for 30 s, 58°C for 30 s, and 72°C for 1 min; and a final extension of 72°C for 10 min. Three sets of primers were used to amplify a partial *pcrI* gene region (PCR1) (101 bp), a *pcrI-rflA* overlapping region (PCR2) (614 bp), and a partial *rflA* gene region (PCR3) (678 bp) (Table 1). *P. corrugata* CFBP 5454 DNA was used as a positive control. As a negative control, PCR reactions with the same primer sets were performed using RNA samples that had not been reverse transcribed.

Culture filtrate preparation.

P. corrugata culture filtrates were used to assess AHL and LDP production. Culture filtrates were prepared according to Cirvilleri and associates (2005), with minor modifications. Bacterial strains were grown in IMM (Surico et al. 1988) at 28°C for 4 days. The time course production was determined by sampling aliquots of 20 ml of culture incubated in a orbital shaker from triplicate flasks immediately after inoculation and after 4, 8, 16, 24, 32, 48, 72, and 96 h. The mutant and WT strain activities were also assessed in 4-day-old still cultures. After centrifugation (9000 × g, 20 min), the supernatant was passed through a 0.22-µm Millipore filter (Millipore, Billerica, MA, U.S.A.) to obtain cell-free culture filtrates. Aliquots of samples at time (T) 8, 16, 24, 48, 72, and 96 were 10× concentrated by using the Vacufuge concentrator 5301(Eppendorf, Milan, Italy) for LDP bioassay.

Bioassay for AHL production.

AHL were quantified by adding the culture filtrates of *P. corrugata* CFBP 5454 or its mutant derivatives to CV026 liquid cultures and measuring violacein formation, which is dependent upon the external addition of medium-chain-length lactones (Martinelli et al. 2004). Experiments were conducted in micro-titer-plate-based assay and absorbance values read using the Bioscreen C instrument (LabSystems, Espoo, Finland).

P. corrugata culture filtrates were added to freshly grown CV026 (1×10^9 CFU ml⁻¹) at a ratio of 1:10 to a final volume of 300 µl (10 replicates for each cultural filtrate) and incubated in the Bioscreen C system for 16 h at 27°C with continuous shaking to allow induction of violacein formation. A 10-replicate lane loaded only with CV026 was also included. Turbidity at 600 nm was used for growth control. After incubation, the plates were dried at 60°C until all medium had evaporated (approximately 6 h or overnight). The violacein was resolubilized by adding 100 µl of dimethyl sulfoxide to each well and incubating the plates in the Bioscreen C system for 2 h with continuous shaking. The amounts of violacein measured at 590 nm by adding the culture filtrates were used as an indirect measure of AHL production. These values were subtracted by those obtained in the wells inoculated only with CV026.

In vitro bioassay for LDP production.

Cultural filtrates of *P. corrugata* and mutant derivatives were evaluated for LDP production by a bioassay based on antimicrobial activity against *R. pilimanae* and *Bacillus megaterium* performed essentially as previously described (Cirvilleri et al. 2005). Antimicrobial activity was assessed by well-diffusion assay in plates containing solidified potato dextrose agar (PDA) (Oxoid) overlaid with 6 ml of PDA added with 1 ml of the test microorganism (approximately 10^6 CFU ml⁻¹). Wells, each 7 mm in diameter, were made in the agar using a cork borer, and 20 µl of culture filtrate were transferred into each well. The plates were incubated up to 4 days at 27°C, after which they were examined for clear inhibition zones around the well. Tests were carried out twice with triplicate wells each time.

Plant inoculations.

P. corrugata CFBP 5454 and the derivative mutants were tested for pathogenicity on tomato cv. Marmande plants grown in nursery flats, 1 month after germination. During the trials, plants were maintained in a growth chamber with a photoperiod of 16 and 8 h and 26°C temperature. Tomato plants were pin-prick inoculated on the stem at the axil of the first true leaf (20 plants per strain) with bacterial cells from a 48-h culture on NDA. After inoculation, plants were enclosed in polyethylene bags to maintain 100% relative humidity for 3 days; the bags were then removed until the end of the experiment. Ex-

periments were performed at least twice. Host–pathogen interaction phenotype was evaluated approximately 10 days after inoculation using an empirical 0-to-4 increasing susceptibility scale based on the range of symptoms. Asymptomatic plants were scored as 0; score 1 was assigned to those plants showing only a dark discoloration along the pin puncture and was considered to be negative. Scores from 2 to 4 focused on symptom progressions and the presence of typical dark necrotic lesion within the stem (TPN): 2, light discoloration of the stem >5 mm; 3, typical TPN between 0.5 and 1 cm; 4, TPN > 1. A point each was added if hollowing of the pith or presence of brown discoloration of the xylem beyond the TPN-affected areas were observed. Results were expressed as mean DI.

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