The *HopPtoF* Locus of *Pseudomonas syringae* pv. *tomato* DC3000 Encodes a Type III Chaperone and a Cognate Effector

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Type III secretion systems are highly conserved among gram-negative plant and animal pathogenic bacteria. Through the type III secretion system, bacteria inject a number of virulence proteins into the host cells. Analysis of the whole genome sequence of *Pseudomonas syringae* pv. *tomato* DC3000 strain identified a locus, named *HopPtoF*, that is homologous to the avirulence gene locus *avrPphF* in *P. syringae* pv. *phaseolicola*. The *HopPtoF* locus harbors two genes, *ShcF_{Pto}* and *HopF_{Pto}*, that are preceded by a single *hrp* box promoter. We present evidence here to show that *ShcF_{Pto}* and *HopF_{Pto}* encode a type III chaperone and a cognate effector, respectively. *ShcF_{Pto}* interacts with and stabilizes the *HopF_{Pto}* protein in the bacterial cell. Translation of *HopF_{Pto}* starts at a rare initiation codon ATA that limits the synthesis of the *HopF_{Pto}* protein to a low level in bacterial cells.

Many gram-negative bacterial pathogens of plants and animals harbor a specialized secretion system, termed type III secretion system (Galan and Collmer 1999). In plant bacterial pathogens, genes encoding the type III secretion system are called the hypersensitive response and pathogenicity (*hrp*) genes, because mutations in these genes abolish the induction of hypersensitive response (HR) on nonhost plants or resistant host plants and pathogenicity on host plants (He 1998; Lindgren 1997). Type III secretion system can secrete a group of proteins, termed Hrp-dependent outer (Hop) proteins, to the outside of the bacterial cell. Hop proteins bear the secretion signals at the N-terminus. The secretion signal sequences of different Hop proteins do not share significant homology but have unique biophysical properties that may lead the protein through the type III secretion pathway (Guttman et al. 2002; Petnicki-Ocwieja et al. 2002). Some Hop proteins are secreted into intercellular spaces that may act as the type III accessory proteins, whereas others are delivered inside the host cells (Galan and Collmer 1999). Hop proteins that are delivered into the host cells are designated type III effectors (Hueck 1998). Many type III effectors of plant bacterial pathogens were identified as avirulence (Avr) proteins because they induce HR and disease resistance on plants carrying the corresponding disease resistance (*R*) genes (Collmer 1998). The Avr proteins interact directly or indirectly with the corresponding *R* proteins

in the host cells, leading to the activation of HR and disease resistance (Kim et al. 2002; Mackey et al. 2002; Scofield et al. 1996; Shao et al. 2003; Tang et al. 1996). A number of Avr proteins exhibit virulence functions on susceptible plants (White et al. 2000).

The function of certain type III effectors requires a partner protein termed type III chaperone (Bennett and Hughes 2000). Type III chaperones do not share sequence similarities, but they show common structural characteristics such as a small molecular mass, an acidic isoelectric point, and a predominantly helical secondary structure (Aldridge and Hughes 2001; Bennett and Hughes 2000). Most of the type III chaperones characterized to date are specific for a single effector protein, but some are associated with two or more effectors (Page and Parsot 2002). Chaperones often bind directly to the N-terminal portion of the secreted proteins. The loss of the chaperone often results in reduced stability of the partner protein in the bacterial cytoplasm (Fu and Galan 1998; Niebuhr et al. 2000) and sometimes aberrant secretion and translocation of the Hop proteins (Cambronne et al. 2000; van Dijk et al. 2002).

The complete genome sequence of *Pseudomonas syringae* pv. *tomato* DC3000 strain, a bacterial pathogen of tomato and *Arabidopsis thaliana*, enabled the identification of more than 30 putative type III effector genes (Collmer et al. 2002). Among them, the *HopPtoF* locus shows homology to the *avrPphF* locus of *P. syringae* pv. *phaseolicola*, a bean pathogen (Fouts et al. 2002; Zwiesler-Vollick et al. 2002). *avrPphF* is a plasmid-borne *avr* gene conferring specific resistance on bean cultivars carrying the *R1* disease resistance gene (Jackson et al. 1999; Tsiamis et al. 2000). *avrPphF* also confers cultivar-specific virulence on bean and soybean plants. In certain bean cultivars, *avrPphF* suppresses the HR caused by other *avr* gene. The *avrPphF* locus comprises two open reading frames (ORFs) that both are required for avirulence and virulence functions. The two ORFs are led by a single promoter containing the *hrp* box motif (Tsiamis et al. 2000). The proteins encoded by the *avrPphF* locus have not been characterized and, therefore, it is unclear how the two ORFs coordinate to condition the avirulence and virulence functions.

We report here that the DC3000 *HopPtoF* locus harbors two ORFs that are controlled by a single *hrp* box promoter. The first ORF encodes a molecular chaperone that interacts with the effector protein encoded by the second ORF and stabilizes the effector protein. We name the effector protein *HopF_{Pto}* to reflect its nature as a Hop protein, homology to *AvrPphF*, and

origin from *P. syringae* pv. *tomato*. The chaperone protein is designated ShcF_{Pto} to indicate it as a specific Hop chaperone for HopF_{Pto}. The *HopF_{Pto}* gene carries a rare initiation codon ATA that was reported to serve as a translation initiation site in mitochondria but not in bacterial cells (NCBI website). The use of ATA as the initiation codon limits the protein synthesis to a low level, and mutation of ATA to ATG of *HopF_{Pto}* in the *HopPtoF* locus drastically increases the HopF_{Pto} protein synthesis in bacterial cells.

RESULTS

Sequence characterization of the *HopPtoF* locus.

The *HopPtoF* locus (Fig. 1) of *P. syringae* pv. *tomato* DC3000 consists of two ORFs, *ShcF_{Pto}* and *HopF_{Pto}*, that encode proteins homologous to ORF1 and ORF2 encoded by the *P. syringae* pv. *phaseolicola* *avrPphF* locus (Tsiamis et al. 2000). *ShcF_{Pto}* is preceded by a Shine-Dalgarno (SD) sequence (–13GAGGAG–7) and a promoter carrying the *hrp* box motif. The sequence of *ShcF_{Pto}* predicts a protein of 15 kDa with an

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GGGAACCTGATGCTGCTCAGTGACCCTCATAACCTTAAATTATGAGGATA
T GAGGAGATGCATTATGAAAAACGCATTTGACCTGCTTGTGGAAGGGCTGG
ShcF M K N A F D L L V E G L A
CTAAGGACTACAACATGCGCCCTTGCTGACAAAGAAACATATCGATGAAG
K D Y N M P P L P D K K H I D E V
TCTATTGCTTTGAGTTTCAAAGTGGTATGAACGTAAAAGTATACCAAGACG
Y C F E F Q S G M N V K V Y Q D E
AATTTCGCTGGGTATATTTACCGCTGACGTTGGGACATTTCAAGATAGCA
F R W V Y F T A D V G T F Q D S S
GTATTGACACATTAACTACGCGCTCCAGCTGAACAACCTTAGCCTTAGAA
I D T L N Y A L Q L N N F S L R K
AACCTTTCTGACCTTCGGAATGACGAGGAGAGAAAATGGTGTATTGCATA
P F L T F G M T K E K N G V L H T
CACGCAACCCCTTGATTGAGGTAGACAACGTGCAATGCGCAGGATATTTG
R T P L I E V D N V Q M R R I F E
AGGAGCTTATAGGCGTGGCAGGTGAATCAGAAAACTAAACTCAAAT
E L I G V A G T A E I R K T L K L K *
AGTAGCGAGGAAGTTTTTAGCCTTGTGAACCTCTGGTAATGTACATAAAT
GCATCAATATTACATGGGCACTGTACATACACCAGAGCGTCGGGCATAAAA
GAAGCAATCTGCAGTTTTTAAAAAGAGGAACCATATATAGGTAATATTTCG
HopF M G N I C
GGCACCTCGGGCTCACGTCATGTGTATAGCCCATCCCATACACAAACGATA
G T S G S R H V Y S P S H T Q R I
ACTTCAGCTCCCTCTACATCCACTCATGTTGGTGAGATACACTGACATCC
T S A P S T S T H V G G D T L T S
ATTCACTAGCTTTTCGATAGTCAGAGAGAGCATTTCTGAACATGCATGAT
I H Q L S H S Q R E Q F L N M H D
CCAATGAGAGTAATGGGACTTGACCATGATACCGAGCTTTTCAGAACGACG
P M R V M G L D H D T E L F R T T
GATAGTCGCTATATAAAAAACGATAAACTCGCGGCAATCCACAATCCATG
D S R Y I K N D K L A G N P Q S M
GCGAGTATCCTTATGATGAAGAACTGCGCCCAATCGTTTTGCCAGCCAT
A S I L M H E E L R P N R F A S H
ACAGGTGCCCAACACAGCAAGCAAGGCGTACGTTCCGAAAAGAATAAAA
T G A Q P H E A R A Y V P K R I K
GCCACCGATCTAGAGTTCCATCACTGAACGTAATGACTGGCTCGCTAGCG
A T D L G V P S L N V M T G S L A
CGAGACGGAATTAGAGCTTATGATCAGATGATGATAATCAGGTCTCTGTG
R D G I R A Y D H M S D N Q V S V
AAAATGCGAGTCGGGAGATTTCTCGAAAGGGGTGGCAAGGTCTATGCCGAC
K M R L G D F L E R G G K V Y A D
GCTTCGTCTGTAGCTGACGATGGGGAACATCACAAGCTCTGATTGTCACA
A S S V A D D G E T S Q A L I V T
TTGCCAAAGGACAGAAAGTGCCCGTGAAGAGGTCATGA
L P K G Q K V P V E R V *
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Fig. 1. Nucleotide sequence of the *HopPtoF* locus (AY321312) and deduced amino acid sequences of ShcF_{Pto} and HopF_{Pto}. The amino acid sequence is presented in single-letter code below the nucleotide sequence. The *hrp* box sequence is in boldface, Shine-Dalgarno sequences in bold italics, termination codons in asterisks, and ATA₁ and GTG₁₃ of *HopF_{Pto}* are underlined.

isoelectric point 6.5. Secondary structure analysis using the PSIPRED program predicts a predominantly α -helical structure throughout the ShcF_{Pto} protein and a C-terminal α -helical structure in particular (McGuffin et al. 2000). These characteristics often are observed in type III chaperones (Bennett and Hughes 2000). Compared with the *avrPphF* ORF2 gene, a standard initiation codon ATG was not found in the *HopF_{Pto}* gene. Instead, it has ATA at the predicted initiation site (Fig. 1). Gene annotation using the GLIMMER program (Salzberg et al. 1998) suggests that HopF_{Pto} starts at GTG₁₃, 13 codons downstream of ATA₁. But GTG₁₃ is not preceded by a SD sequence. In addition, homology between HopF_{Pto} and the *avrPphF* ORF2 extends to the region between GTG₁₃ and ATA₁. Seven bases upstream of ATA₁ is a typical SD sequence (–13AGAGGA–7). ATA can be used as the initiation codon in mitochondria but was not reported to act as a start codon in any bacteria. Downstream of GTG₁₃, there are eight in-frame ATGs.

ShcF_{Pto} is not secreted in culture.

Many bacterial Avr proteins characterized to date are secreted through the type III secretion system (Galan and Collmer 1999). The similarity between the *HopPtoF* and *avrPphF* loci led us to test the secretion of the *HopPtoF*-encoded proteins in culture. The ShcF_{Pto} protein was tagged at the C-terminus with hemagglutinin (HA) epitope. The ShcF_{Pto} protein was highly expressed in the *hrp*-inducing minimum medium (MM) (Huynh et al. 1989) but not in King's B (KB) broth (King et al. 1954) (Fig. 2). In MM, ShcF_{Pto} was detected only in the cell-bound fraction. In contrast, AvrPto, a type III effector, was detected in both bacterial cells and supernatant, indicating that ShcF_{Pto} is not secreted or the secretion of ShcF_{Pto} is undetectable under the experimental conditions. We also tagged HopF_{Pto} at its C-terminus with HA in the native context (i.e., *hrp* box promoter, *ShcF_{Pto}*, the spacer DNA between *ShcF_{Pto}* and *HopF_{Pto}*, and *HopF_{Pto}*). However, we were unable to detect the HopF_{Pto} protein in either bacterial cells or supernatant. We then replaced the *hrp* box promoter with the strong constitutive *nptII* promoter for the expression of the *ShcF_{Pto}*-*HopF_{Pto}*-HA genes. The *nptII* promoter enabled the detection of a small amount of HopF_{Pto}-HA protein in *P. syringae* pv. *tomato* DC3000 cells but not in the supernatant (data not shown).

HopF_{Pto} but not ShcF_{Pto} is translocated into plant cells.

The failure to detect the secretion of ShcF_{Pto} in culture might be caused by experimental conditions that did not favor the secretion of this protein. Similar result was reported for the *P. syringae* AvrB protein that showed poor secretion in culture (van Dijk et al. 1999). Therefore, we used an alternative approach to test the secretion of ShcF_{Pto} by fusing a partial AvrRpt2 protein (AvrRpt2⁴¹⁻²⁵⁵) to the C-terminus of ShcF_{Pto} and testing the HR-inducing activity of the fusion protein on

Fig. 2. ShcF_{Pto} is not secreted in liquid culture. DC3000 (*pLK-hrp-ShcF-hemagglutinin* [HA]) was cultured in King's B broth (KB) and the hyper-sensitive response and pathogenicity (*hrp*)-inducing minimum medium (MM). The total protein (T) of the DC3000 (*pLK-hrp-ShcF-HA*) culture in MM was fractionated into pellet (P) and supernatant (S), separated by sodium dodecyl sulfate-gel electrophoresis, and probed with the anti-HA antibody. The pellet (P) and supernatant (S) fractions of DC3000 (*pLK-hrp-ShcF-HA*) in MM also were probed with anti-AvrPto antibodies as positive control of the type III secretion. Untransformed DC3000 was cultured in MM as negative control for ShcF_{Pto}-HA.

Arabidopsis plants carrying the *RPS2* gene. AvrRpt2 is an Avr protein that induces the HR on *Arabidopsis* plants carrying the *RPS2* gene (Bent et al. 1994). Upon secretion into the plant cells, AvrRpt2 is cleaved by a plant protease between the amino acid residues Gly71 and Gly72, exposing a potential myristoylation site for the C-terminal peptide that has the avirulence activity in *RPS2* plants (Mudgett and Staskawicz 1999). AvrRpt2⁴¹⁻²⁵⁵ lacks the N-terminal type III secretion signal and, therefore, is unable to travel through the type III secretion system into the host cells. However, if the test protein carries the type III secretion signal, the fusion protein can be secreted into the host cells to induce the HR on plants carrying the *RPS2* gene. This strategy has been used successfully to demonstrate the secretion of type III effectors (Guttman and Greenberg 2001; Guttman et al. 2002; Mudgett et al. 2000; Petnicki-Ocwieja et al. 2002). In *P. syringae* pv. *phaseolicola*, *pLK-hrp-ShcF_{Pto}-avrRpt2⁴¹⁻²⁵⁵*, which gives rise to a fusion protein of ShcF_{Pto}-AvrRpt2⁴¹⁻²⁵⁵, did not cause HR on *RPS2* plants (Fig. 3). This result further indicated that ShcF_{Pto} is not secreted. In contrast, *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵*, which produces the ShcF_{Pto} protein and the HopF_{Pto}-AvrRpt2⁴¹⁻²⁵⁵ fusion protein, elicited an *RPS2*-dependent HR (Fig. 3A), indicating that the *HopF_{Pto}* gene encodes a type III effector. Because ShcF_{Pto} and HopF_{Pto} are encoded by a single operon, ShcF_{Pto} has the structural characteristics of a molecular chaperone, and the secretion of ShcF_{Pto} was not detectable, we speculated that ShcF_{Pto} serves as the type III chaperone for HopF_{Pto}.

We further tested whether the translocation of HopF_{Pto} is dependent on type III secretion system. The HR-inducing activity of pLK-hrp-ShcF_{Pto}-HopF_{Pto}-AvrRpt241-255 was examined in a DC3000 type III secretion deficient mutant, the *hrcC* deletion strain (Wei et al. 2000). No HR was induced by the *hrcC* deletion strain carrying *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-AvrRpt2⁴¹⁻²⁵⁵* (Fig. 3B), indicating that the translocation of HopF_{Pto} relies on the type III secretion system.

Translation of HopF_{Pto} starts at ATA.

Type III secretion signals are confined to the very N-terminal portion of the effector proteins. Thus, determination of the translation initiation site is necessary to understand the HopF_{Pto} secretion signal. Compared with the *avrPphF* *ORF2* sequence that has the predicted start codon ATG, *HopF_{Pto}* has ATA at position 1, a codon unknown for translation initiation in bacteria. Other potential start codons include GTG at position 13 and eight in-frame ATGs downstream of GTG₁₃ (Fig. 1). To unequivocally determine the translation initiation site of *HopF_{Pto}*, we first tested whether one of the in-frame ATGs serves as the initiation codon. Serial deletions from the 3' end of *HopF_{Pto}* were created that sequentially removed the ATGs, the remaining DNA fragments (including the native promoter, complete *ShcF_{Pto}*, the spacer DNA, and partial *HopF_{Pto}*) were fused to *avrRpt2⁴¹⁻²⁵⁵*, and the resulting plasmids were introduced into *P. syringae* pv. *phaseolicola* for the translocation assay. All the deletion constructs, including the one (*pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{Δ151}-avrRpt2⁴¹⁻²⁵⁵*) that has only the N-terminal

Fig. 3. HopF_{Pto} but not ShcF_{Pto} shows type III-dependent secretion in planta. **A**, Translocation assay of ShcF_{Pto} and HopF_{Pto}. *ShcF_{Pto}* and *HopF_{Pto}* were fused in frame with the partial *avrRpt2⁴¹⁻²⁵⁵* gene in the plasmids *pLK-hrp-ShcF-avrRpt2⁴¹⁻²⁵⁵* and *pLK-hrp-ShcF-HopF^{ATA}-avrRpt2⁴¹⁻²⁵⁵*, respectively. *Pseudomonas syringae* pv. *phaseolicola* NPS 3121 (*PspH*) was used as the carrier strain for the plasmids. Test strains were infiltrated into *Arabidopsis* Col-0 (*RPS2*) and Col-0 *rps2-201* (*rps2*) plants for the hypersensitive response (HR) assay. The HR was induced by *PspH* strains expressing the wild-type AvrRpt2 protein (*pavrRpt2*) and HopF_{Pto}-AvrRpt2⁴¹⁻²⁵⁵ fusion protein (*hrp-ShcF-HopF^{ATA}-avrRpt2⁴¹⁻²⁵⁵*). **B**, Type III-dependent secretion of HopF_{Pto} in planta. *pLK-hrp-ShcF-HopF-avrRpt2⁴¹⁻²⁵⁵* was introduced into *P. syringae* pv. *tomato* DC3000 and DC3000 *hrcC*⁻ mutant strains. Bacteria were infiltrated into *Arabidopsis* Col-0 plants (*RPS2*). The HR was induced by DC3000 carrying *pLK-hrp-ShcF-HopF-avrRpt2⁴¹⁻²⁵⁵*. N = no visible response.

portion of the *HopF_{P10}* gene without any in frame ATGs, still retained the ability to cause HR on *RPS2* plants (Fig. 4A), suggesting that the translation initiation site of *HopF_{P10}* is further upstream of ATG₅₄, the first ATG from the 5' end.

GTG₁₃ was predicted by the GLIMMER program to be the translation start codon for *HopF_{P10}*. To test this prediction, we constructed a plasmid, *pML-nptII-HopF_{P10}^{GTG13}-HA*, to express

the *HopF_{P10}-HA* protein starting from GTG₁₃. This plasmid carries the *HopF_{P10}* gene starting at GTG₁₃ and a synthetic SD sequence placed upstream of GTG₁₃. As control, we also constructed a plasmid, *pML-nptII-ShcF_{P10}-HopF_{P10}-HA*, to express the *HopF_{P10}-HA* protein encoded by the native *HopPtoF* locus. The gene transcription in both plasmids was controlled by the strong *nptII* promoter. We examined the *HopF_{P10}-HA* protein

Fig. 4. ATA is the translation initiation codon of *HopF_{P10}*. **A**, *pLK-hrp-ShcF-HopF^{ATA}Δ35-avrRpt2⁴¹⁻²⁵⁵*, *pLK-hrp-ShcF-HopF^{ATA}Δ107-avrRpt2⁴¹⁻²⁵⁵*, and *pLK-hrp-ShcF-HopF^{ATA}Δ151-avrRpt2⁴¹⁻²⁵⁵* carry deletions of 35, 107, and 151 amino acid residues, respectively, at the C-terminus of *HopF_{P10}*. The plasmids were introduced into *Pseudomonas syringae* pv. *phaseolicola* NPS 3121 (*Psph*) and the resulting bacterial strains were infiltrated into *Arabidopsis* Col-0 plants (*RPS2*) and Col-0 *rps2-201* (*rps2*) plants. **B**, DC3000 strains carrying *pML-nptII-ShcF-HopF^{ATA}-hemagglutinin (HA)* and *pML-nptII-HopF^{GTG13}-HA* plasmids were cultured in minimal medium. Protein from the bacterial pellet was assayed with Western blot using the HA antibody. The *HopF_{P10}* protein encoded by the native *HopF_{P10}* gene in *pML-nptII-ShcF-HopF^{ATA}-HA* (lane 1) is ≈23 kDa and slightly larger than the protein translated from GTG₁₃ by the *pML-nptII-HopF^{GTG13}-HA* plasmid (lane 2). **C**, *pLK-hrp-ShcF-HopF^{ATA}-avrRpt2⁴¹⁻²⁵⁵*, *pLK-hrp-ShcF-HopF^{CTA}-avrRpt2⁴¹⁻²⁵⁵* (mutation of ATA₁ to CTA), and *pLK-hrp-ShcF-HopF^{GCG13}-avrRpt2⁴¹⁻²⁵⁵* (mutation of GTG₁₃ to GCG) were introduced into *Psph*, and the resulting bacterial strains were infiltrated into *Arabidopsis* Col-0 plants (*RPS2*) and Col-0 *rps2-201* (*rps2*) plants for the hypersensitive response (HR) assay. The HR was disrupted by mutation of ATA₁ to CTA but not affected by mutation of GTG₁₃ to GCG. **D**, *RPS2* *Arabidopsis* leaves were injected with *Psph* strains carrying *avrRpt2⁴¹⁻²⁵⁵*, *pavrRpt2*, *pLK-hrp-ShcF-HopF^{ATA}-avrRpt2⁴¹⁻²⁵⁵*, and *pLK-hrp-ShcF-HopF^{ATG}-avrRpt2⁴¹⁻²⁵⁵* (mutation of ATA₁ to ATG). The HR caused by *Psph*(*pLK-hrp-ShcF-HopF^{ATG}-avrRpt2⁴¹⁻²⁵⁵*) was visible ≈16 h after inoculation, significantly earlier than the HR caused by *Psph*(*pLK-hrp-ShcF-HopF^{ATA}-avrRpt2⁴¹⁻²⁵⁵*), which was visible ≈22 h after inoculation. **E**, *Psph* strains carrying *pML-nptII-ShcF-HopF^{ATG}-HA* (ATG), *pML-nptII-ShcF-HopF^{CTA}-HA* (CTA), *pML-nptII-ShcF-HopF^{ATA}-HA* (ATA), and *pML* empty vector (EV) were cultured in minimal medium. Protein from equal amount of bacterial cells was assayed with Western blot using the HA antibody. The inoculation experiments for **A**, **C**, and **D** were repeated numerous times with the same results.

produced by the two plasmids in DC3000 using Western blot analysis with the anti-HA antibodies. If GTG₁₃ is the start codon, the two plasmids were expected to yield the HopF_{Pto}-HA protein of the same size. However, the protein produced by *pML-nptII-HopF_{Pto}^{GTG13}-HA* (Fig. 4B, lane 2) was slightly smaller than the protein produced by *pML-nptII-ShcF_{Pto}-HopF_{Pto}-HA* (Fig. 4B, lane 1), indicating that GTG₁₃ is not the start codon. To further test this possibility, we created point mutation in the construct *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵* that converted GTG₁₃ to GCG. GCG was not reported to act as a translational start codon in any organisms. Mutation of GTG₁₃ to GCG did not affect the HR-inducing ability of *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵* (Fig. 4C), further supporting the theory that GTG₁₃ is not the initiation codon for HopF_{Pto}. The authentic translation initiation site of HopF_{Pto} must be further upstream of GTG₁₃ because the HopF_{Pto} protein encoded by the native *HopPtoF* locus is larger than the protein starting from GTG₁₃.

We speculated that ATA₁ functions as the start codon of HopF_{Pto}, because the homology between HopF_{Pto} and AvrPphF ORF2 extends to the ATA₁ codon, and the protein translated from ATA₁ exhibits the features of type III secretion signal sequence at its N-terminus (Guttman et al. 2002; Petnicki-Ocwieja et al. 2002). We tested this possibility by converting ATA₁ to CTA and ATG in the plasmid *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵*. CTA never was reported to act as an initiation codon in any organisms. Interestingly, mutation of ATA₁ to CTA in HopF_{Pto} completely disrupted the HR-inducing activity (Fig. 4C), whereas mutation of ATA₁ to ATG accelerated the HR (Fig. 4D). In addition, mutation of ATA₁ to ATG in HopF_{Pto} in the *pML-ShcF_{Pto}-HopF_{Pto}-HA* plasmid resulted in the detection of a large amount of the HopF_{Pto}-HA protein produced by *P. syringae* carrying the plasmid, while no protein was detected when ATA₁ was converted to CTA (Fig. 4E). The HopF_{Pto}-HA proteins produced by *pML-ShcF_{Pto}-HopF_{Pto}^{ATG}-HA* (indicated by ATG in Fig. 4E) is the same size as that encoded by the native *HopPtoF* locus (indicated by ATA in Fig. 4E). Because the *HopF_{Pto}^{ATA}*

gene in the native *HopPtoF* locus showed the HR-inducing activity in the *avrRpt2⁴¹⁻²⁵⁵*-mediated translocation assay and also could direct the synthesis of a small amount of protein in bacteria, we believed that ATA₁ serves as the translation initiation codon of HopF_{Pto}.

ShcF_{Pto} interacts with HopF_{Pto} in yeast and in protein pull-down assay.

Type III chaperones often interact physically with the cognate Hop proteins (Aldridge and Hughes 2001; Bennett and Hughes 2000). To determine that ShcF_{Pto} indeed functions as the type III chaperone for HopF_{Pto}, we tested the interaction of ShcF_{Pto} and

Fig. 6. HopF_{Pto} is unstable but can be secreted in the absence of ShcF_{Pto}. 1, Untransformed *Pseudomonas syringae* pv. *phaseolicola* NPS 3121 (*Psph*); 2, *Psph* carrying both *pML-nptII-HopF^{ATG}-FLAG* and *pHM2-hrp-ShcF-hemagglutinin (HA)* plasmids; and 3, *Psph* carrying *pML-nptII-HopF^{ATG}-FLAG* alone were cultured in minimum medium and diluted to the same concentration with the same medium before protein analysis. Equal volumes of bacterial cultures were fractionated into pellet and supernatant and examined with antibodies against FLAG, HA, and NPTII for HopF_{Pto}-FLAG, ShcF_{Pto}-HA, and the control NPTII protein (for bacterial lysis). T, total bacterial protein in culture; P, pellet; S, supernatant. Arrows indicate the corresponding proteins. Equal loading of protein samples is shown by the nonspecific bands above the HopF_{Pto}-FLAG and ShcF_{Pto}-HA proteins. The weak bands of the same size as ShcF_{Pto}-HA in lanes 1/T and 3/P are from nonspecific hybridizations.

Fig. 5. ShcF_{Pto} interacts with HopF_{Pto}. **A**, Yeast two-hybrid analysis. ShcF_{Pto} was cloned into the bait vector pEG202; the full-length HopF_{Pto} gene and HopF_{Pto} lacking the first 13 codons were cloned into the prey vector pJG4-5. The LexA yeast two-hybrid system was used for protein interaction assay. **B**, Protein pull-down assay. The GST-AvrPto and GST-HopF_{Pto} fusion proteins were purified with glutathione-conjugated cellulose beads. Washed beads carrying 50 µg of GST-AvrPto (lane 1) and GST-HopF_{Pto} (the fusion proteins) (lane 2) were incubated with the bacterial lysate of *Pseudomonas syringae* pv. *phaseolicola* NPS 3121 strain containing ShcF_{Pto}-hemagglutinin (HA). The beads were washed with phosphate-buffered saline plus 0.1% Tween-20 and assayed with Western blot and the anti-HA antibody.

HopF_{Pto} using the yeast two-hybrid assay. Expression of neither ShcF_{Pto} nor HopF_{Pto} alone in yeast activated the *lacZ* reporter gene, as indicated by the white yeast colonies (Fig. 5A). However, expression of ShcF_{Pto} and HopF_{Pto} together in the same yeast cell activated the *lacZ* reporter gene, indicating the interaction of the two proteins. HopF_{Pto} carrying deletion of the N-terminal 13 amino acid residues still showed interaction with ShcF_{Pto} (Fig. 5A). The interaction of HopF_{Pto} with ShcF_{Pto} was further confirmed by the protein pull-down assay. GST-HopF_{Pto} fusion protein, but not GST-AvrPto, specifically pulled down the ShcF_{Pto}-HA protein from the protein extract of *P. syringae* pv. *phaseolicola* expressing the *ShcF_{Pto}-HA* gene (Fig. 5B).

ShcF_{Pto} stabilizes HopF_{Pto} in bacterial cytoplasm.

The interaction of a type III chaperone with the corresponding Hop protein often stabilizes the effector protein in bacterial cytoplasm or facilitates the secretion of the effector through the type III pathway. To determine how ShcF_{Pto} affects HopF_{Pto}, we tagged the C-terminus of HopF_{Pto}^{ATG} with FLAG epitope and expressed the protein using the *nptII* promoter. The HopF_{Pto} gene with ATG as start codon was used in this experiment to facilitate the protein detection. We investigated the stability and secretion of HopF_{Pto}-FLAG in the presence or absence of ShcF_{Pto} in *P. syringae* pv. *phaseolicola* strains. Upon induction in MM, over 50-fold more HopF_{Pto}-FLAG protein was detected in *P. syringae* pv. *phaseolicola* carrying both *pHM-hrp-ShcF_{Pto}-HA* and *pML-HopF_{Pto}^{ATG}-FLAG* than in the strain carrying *pML-HopF_{Pto}^{ATG}-FLAG* alone (Fig. 6). In both strains, the HopF_{Pto} protein was detected in the supernatant as well as in the bacterial pellet. The HopF_{Pto} protein in the supernatant was not caused by cell lysis because the ShcF_{Pto} protein and the control NPTII protein were not detected in the supernatant (Fig. 6). These results indicated that ShcF_{Pto} is required for the stability but dispensable for the secretion of the HopF_{Pto} protein.

DISCUSSION

In this study, we showed that the *HopPtoF* locus of DC3000 encodes two proteins, ShcF_{Pto} and HopF_{Pto}. The HopF_{Pto} gene carries a rare initiation codon ATA that limits the HopF_{Pto} protein synthesis in *P. syringae* to a low level. ShcF_{Pto} serves as the type III chaperone that interacts with and stabilizes the type III effector protein HopF_{Pto}.

The ShcF_{Pto} protein exhibits several structural and functional characteristics supporting its nature as the chaperone of HopF_{Pto}. Structurally, ShcF_{Pto} is predicted to be a small cytoplasmic protein (molecular mass = 15 kDa) with a relatively low isoelectric point (pI = 6.5) and a secondary structure predominantly helical. These characteristics are shared by other type III chaperones (Aldridge and Hughes 2001; Bennett and Hughes 2000; Page and Parsot et al. 2002). Functionally, ShcF_{Pto} is required for the stability of HopF_{Pto} in bacterial cytoplasm. The interaction of ShcF_{Pto} with HopF_{Pto} in the yeast two-hybrid system and in a protein pull-down assay provided further evidence that ShcF_{Pto} is the chaperone of HopF_{Pto}. Unlike ShcA and ShcM, two chaperones of *P. syringae* pv. *syringae* that facilitate secretion but are dispensable for the stability of their partner effectors HopPsyA and HopPtoM, respectively (Badel et al. 2003; van Dijk et al. 2002), ShcF_{Pto} is dispensable for the secretion but required for the stability of HopF_{Pto}. The requirement of type III chaperone for effector protein stability also was reported for the *Erwinia amylovora* type III chaperone DspB/F which is required for the stability of its cognate effector DspA/E (Gaudriault et al. 2002). In addition, several animal bacterial type III chaperones also protect their corresponding effectors from proteolysis but are not required for the effector secretion (Parsot et al. 2003).

Unlike the *avrPphF* ORF2 that has an in-frame ATG downstream of the SD sequence, the HopF_{Pto} gene does not have a standard translation initiation site. However, this locus is able to direct the synthesis of the HopF_{Pto} protein in both HA tagging and AvrRpt2⁴¹⁻²⁵⁵ tagging experiments. Gene annotation using the GLIMMER program (Salzberg et al. 1998) suggested that the translation of HopF_{Pto} starts at GTG₁₃, a codon that often is used as an alternative initiation codon in bacteria (Osawa et al. 1992). However, this codon is not preceded by a typical SD sequence for ribosomal binding. In addition, mutagenesis of GTG₁₃ to GCG (GCG does not serve as the start codon in any organisms) in *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵* did not affect the HopF_{Pto}-AvrRpt2⁴¹⁻²⁵⁵ protein synthesis, as indicated by the HR-inducing activity of this plasmid in *RPS2* plants. Furthermore, the protein synthesized from GTG₁₃ was smaller than the protein encoded by the native gene. These results strongly argue against GTG₁₃ as the initiation codon of HopF_{Pto}.

In contrast, several pieces of evidence support the theory that ATA₁ is the translation initiation site for HopF_{Pto}, although ATA was not reported to act as the start codon in bacteria. First, we could detect a small amount of the HopF_{Pto} protein encoded by the native *HopPtoF* locus in the HA tagging experiment when the strong constitutive *nptII* promoter was used to drive the expression of the genes. Second, in the AvrRpt2-mediated translocation assay, the fusion of the HopF_{Pto}^{ATA} gene in the native *HopPtoF* locus with *avrRpt2⁴¹⁻²⁵⁵* (in the *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}-avrRpt2⁴¹⁻²⁵⁵* plasmid) resulted in the induction of HR on *RPS2* plants, suggesting that HopF_{Pto}^{ATA} could direct the synthesis of the fusion protein. Additional evidence came from the mutagenesis studies of ATA₁ in the plasmids *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2⁴¹⁻²⁵⁵* and the *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-HA*. We found that mutation of ATA₁ to CTA in the HopF_{Pto}^{ATA} gene (in *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}-avrRpt2⁴¹⁻²⁵⁵* plasmid) abolished the HR-inducing activity, whereas mutation of ATA₁ to ATG enhanced the HR-inducing activity of this plasmid. Consistently, we could not detect the HopF_{Pto} protein when ATA₁ was changed to CTA, but mutation of ATA₁ to ATG in *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-HA* dramatically increased the HopF_{Pto} protein synthesis in *P. syringae* bacteria. Furthermore, mutation of the ATA₁ to ATG resulted in a protein the same size as that encoded by the native gene. Taken together, these results strongly supported the theory that ATA₁ is the start codon of HopF_{Pto}^{ATA}. The direct evidence supporting ATA₁ as the start codon of HopF_{Pto}^{ATA} can be provided by the protein sequence analysis. However, the low abundance of the HopF_{Pto} protein encoded by the native *HopPtoF* locus deterred our effort to obtain enough protein for sequence analysis.

The use of ATA as the start codon does not appear to be general for the *HopF* loci in *Pseudomonas* bacteria. Among the three reported homologous genes, only the HopF_{Pto} gene from DC3000 starts with ATA. The other two genes, one from *P. syringae* pv. *phaseolicola* and the other from *P. syringae* pv. *delphinii*, both started with ATG (Deng et al. 2003; Tsiamis et al. 2000). The *HopPtoO* gene of DC3000, which encodes a protein with the N-terminal 10 amino acid residues identical to those of HopF_{Pto}, also starts with ATG (Buell et al. 2003). This suggests that the ATA₁ codon of the DC3000 HopF_{Pto} gene might be derived from mutation of ATG.

The use of a nonstandard codon for translation initiation is not uncommon in bacteria and other organisms. When a non-ATG codon is used, it often results in low translation efficiency (Osawa et al. 1992). Consistently, we were able to detect only a small amount of the HopF_{Pto}-HA protein with the HopF_{Pto}-HA gene carrying ATA₁ as the start codon. The finding that ATA can act as a translation initiation codon is significant to bacterial genomics in prediction of putative ORFs.

MATERIALS AND METHODS

Bacterial strains and growth conditions.

Escherichia coli strain XL1-Blue was used for gene cloning. *P. syringae* pv. *tomato* DC3000 and *P. syringae* pv. *phaseolicola* NPS 3121 were used for secretion and translocation assays. Antibiotics were used at the following concentrations: rifampicin, 100 µg/ml; ampicillin, 100 µg/ml; gentamicin, 10 µg/ml; kanamycin, 50 µg/ml for *E. coli* and 20 µg/ml for *P. syringae*; spectinomycin, 50 µg/ml.

Construction of plasmids.

Three broad host range plasmids, pLK, pML123 (Petnicki-Ocwieja et al. 2002), and pHM2 vector were used to construct plasmids listed in Table 1. pLK plasmid was derived from pPTE6 (Ronald et al. 1992) by removing the *avrPto* gene with a *HindIII* digestion. pHM2 plasmid was modified from pHM1 (Zhu et al. 1998) by replacing the *Bam*HI fragment with DNA containing the cloning sites of *Bam*HI, *Eco*RI, *Xho*I, *Kpn*I, *Hind*III, and *Xba*I (X. Tang, unpublished data). Construction of *pavrRpt2*⁴¹⁻²⁵⁵ was previously described (Petnicki-Ocwieja et al. 2002).

To construct *pLK-hrp-ShcF_{Pto}-HA*, a DNA fragment spanning 200 bp upstream of the *hrp* box promoter and the complete *ShcF_{Pto}* (excluding the stop codon) was polymerase chain reaction (PCR)-amplified from DC3000 using an upstream primer 5'-TTGAATTCGCCCCCTTCGTTACCTTCCAGC-3', (*Eco*RI site in boldface) and a downstream primer 5'-TTGTCGACTCAGACGTAGTCTGGGACGTCGTATGGGTAGGATCCTTTGAGTTTTAGTGTCTTCTG-3' (preceding the *ShcF_{Pto}* 3' end sequence are a *Sal*I site [bold], the HA sequence plus a stop codon [underlined], and a *Bam*HI site [italics]). The PCR products were cloned into the *Eco*RI and *Sal*I sites of pMODTM-2<MCS> plasmid (Epicentre, Madison, WI, U.S.A.), confirmed by sequence analysis, and further cloned into the *Eco*RI and *Sal*I sites of the pLK plasmid, resulting in *pLK-hrp-ShcF_{Pto}-HA*. The DNA fragment also was cloned into the *Eco*RI and *Sal*I sites of the pBluescript SK(-) plasmid (Stratagene, La Jolla, CA, U.S.A.). The resulting pBS-*hrp-ShcF_{Pto}-HA* plasmid was digested further by *Eco*RI and *Kpn*I and cloned into the same sites in pHM2 vector, resulting *pHM2-hrp-ShcF_{Pto}-HA*. The DNA fragment spanning the *hrp* box promoter and *ShcF_{Pto}* in *pLK-hrp-ShcF_{Pto}-HA* was released by *Eco*RI and *Bam*HI digestion and cloned into the *Eco*RI and *Bam*HI sites of *pavrRpt2*⁴¹⁻²⁵⁵, resulting in *pLK-hrp-ShcF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵.

To generate *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-HA* and *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵ plasmids, the same upstream

primer for *pLK-hrp-ShcF_{Pto}-HA* and the downstream primer 5'-AAGGATCCGACCCCTTTCGACCGGCA-3', corresponding to the 3' end of *HopF_{Pto}* plus the *Bam*HI site (bold), were used to PCR amplify the DNA fragment encompassing the *hrp* box promoter, *ShcF_{Pto}*, spacer between *ShcF_{Pto}* and *HopF_{Pto}*, and the complete *HopF_{Pto}* from DC3000. The PCR product was cloned into the *Eco*RI and *Bam*HI sites of pMODTM-2<MCS> plasmid for sequence analysis. The confirmed sequence was further cloned into the *Eco*RI and *Bam*HI sites of *pLK-hrp-ShcF_{Pto}-HA* and *pLK-hrp-ShcF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵, replacing *ShcF_{Pto}* and resulting in *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-HA* and *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵, respectively.

The serial deletions of the *HopF_{Pto}* gene in plasmids *pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵ were generated by PCR using the same upstream primer for *pLK-hrp-ShcF_{Pto}-HA* and one of the following downstream primers containing the *Bam*HI site (bold): 5'-CAGGATCCACCCCTTTCGAGAAAATC-3' for deletion of 35 amino acid (aa) residues, 5'-CAGGATCCTTCATGCATAAGGATACT-3' for deletion of 107 aa residues, and 5'-CAGGATCCGTTTCAGAAACTGCTCTCT-3' for deletion of 151 aa residues from the *HopF_{Pto}* C-terminus. The PCR products were digested with *Eco*RI and *Bam*HI and cloned into pMODTM-2<MCS> for sequence analysis. The confirmed sequences were cloned into the *Eco*RI and *Bam*HI sites of *pLRP-ShcF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵, replacing *ShcF_{Pto}*.

A QuickChange site-directed mutagenesis kit (Stratagene) was used to generate point mutations in the *HopF_{Pto}* gene in pMOD::*hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵ and pMOD::*hrp-ShcF_{Pto}-HopF_{Pto}-HA* with the following three pairs of complementary primers: 5'-GAGGAAACCATTCTAGGTAATATTTGCGGC-3' and 5'-GCCGCAAATATTACCTAGAAATGGTTTCCTC-3' (ATA₁ to CTA); 5'-CGGGCTCACGTCATGCGTATAGCCCATCCC-3' and 5'-GGGATGGGCTATACGCATGACGTGAGCCCG-3' (GTG₁₃ to GCG); and 5'-GAGGAAACCA TTATGGGTAATATTTGCGGC-3' and 5'-GCCGCAAATATT ACCCTAAATGGTTTCCTC-3' (ATA₁ to ATG). The mutants were confirmed by sequence analysis and subsequently cloned into the *Eco*RI and *Bam*HI sites of *pLK-hrp-ShcF_{Pto}-avrRpt2*⁴¹⁻²⁵⁵ and *pLK-hrp-ShcF_{Pto}-HA* plasmids, resulting in *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{CTA}-avrRpt2*⁴¹⁻²⁵⁵, *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{GCG}-avrRpt2*⁴¹⁻²⁵⁵, and *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATG}-avrRpt2*⁴¹⁻²⁵⁵, *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{CTA}-HA*, and *pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATG}-HA*.

To generate *pML-nptII-ShcF_{Pto}-HopF_{Pto}-HA* and *pML-nptII-HopF_{Pto}^{GTG13}-HA* plasmids, we used the forward primers 5'-AAAAAGCAGGCTTCGAAGGAGATAGAACCATGA

Table 1. Plasmids

Designation	Characteristics
<i>pavrRpt2</i> ⁴¹⁻²⁵⁵	Broad host plasmid pLK carrying partial <i>avrRpt2</i> gene with the N-terminal 40 codons deleted
<i>pavrRpt2</i>	Full length <i>avrRpt2</i> gene in broad host plasmid
<i>pLK-hrp-ShcF_{Pto}-avrRpt2</i> ⁴¹⁻²⁵⁵	<i>ShcF_{Pto}</i> under native promoter and fused in frame with partial <i>avrRpt2</i> , in <i>pLK</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}-avrRpt2</i> ⁴¹⁻²⁵⁵	<i>HopPtoF</i> operon under the native promoter fused with partial <i>avrRpt2</i> , in <i>pLK</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}Δ35-avrRpt2</i> ⁴¹⁻²⁵⁵	Deletion of 35 amino acids (aa) from the C-terminus of <i>HopF_{Pto}</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}Δ107-avrRpt2</i> ⁴¹⁻²⁵⁵	Deletion of 107 aa from the C-terminus of <i>HopF_{Pto}</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}Δ151-avrRpt2</i> ⁴¹⁻²⁵⁵	Deletion of 151 aa from the C-terminus of <i>HopF_{Pto}</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{CTA}-avrRpt2</i> ⁴¹⁻²⁵⁵	Mutation of ATA ₁ of <i>HopF_{Pto}</i> to CTA in <i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2</i> ⁴¹⁻²⁵⁵
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{GCG13}-avrRpt2</i> ⁴¹⁻²⁵⁵	Mutation of GTG ₁₃ of <i>HopF_{Pto}</i> to GCG in <i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2</i> ⁴¹⁻²⁵⁵
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATG}-avrRpt2</i> ⁴¹⁻²⁵⁵	Mutation of ATA ₁ of <i>HopF_{Pto}</i> to ATG in <i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}-avrRpt2</i> ⁴¹⁻²⁵⁵
<i>pLK-hrp-ShcF_{Pto}-HA</i>	Hemagglutinin (HA)-tagged <i>ShcF_{Pto}</i> under native promoter, in <i>pLK</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}-HA</i>	<i>HopF_{Pto}</i> in the <i>HopPtoF</i> operon tagged by HA, in <i>pLK</i>
<i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}^{ATA}-HA</i>	ATA ₁ of <i>HopF_{Pto}</i> replaced by ATG in <i>pLK-hrp-ShcF_{Pto}-HopF_{Pto}-HA</i>
<i>pML-nptII-HopF_{Pto}^{GTG}-HA</i>	HA-tagged <i>HopF_{Pto}</i> starting at GTG ₁₃ in <i>pML123</i> plasmid
<i>pML-nptII-ShcF_{Pto}-HopF_{Pto}^{ATA}-HA</i>	<i>HopPtoF</i> locus driven by <i>nptII</i> promoter, <i>HopF_{Pto}</i> is HA-tagged, in <i>pML123</i>
<i>pML-nptII-ShcF_{Pto}-HopF_{Pto}^{ATA}-HA</i>	ATA ₁ of <i>HopF_{Pto}</i> replaced by ATG in <i>pML-nptII-ShcF_{Pto}-HopF_{Pto}^{ATA}-HA</i>
<i>pML-nptII-ShcF_{Pto}-HopF_{Pto}^{CTA}-HA</i>	ATA ₁ of <i>HopF_{Pto}</i> replaced by CTA in <i>pML-nptII-ShcF_{Pto}-HopF_{Pto}^{CTA}-HA</i>
<i>pHM2-hrp-ShcF_{Pto}-HA</i>	HA-tagged <i>ShcF_{Pto}</i> under native promoter, in broad host plasmid pHM2
<i>pML-nptII-HopF_{Pto}^{ATG}-Flag</i>	Flag-tagged <i>HopF_{Pto}^{ATG}</i> driven by <i>nptII</i> promoter, in <i>pML123</i>

AAAACGCATTT-3' and 5'-AAAAAGCAGGCTTCGAAGG AGATAGAACCTGTGTATAGCCCATCC-3' containing the attB2 sequence (underlined) and SD sequence (italic) and the same reverse primer 5'-AGAAAGCTGGGTAACAGACCCTTTCGAC-3' containing the attB1 sequence (underlined) in PCR with the DC3000 genomic DNA as template. The PCR products were cloned into pDONR201 (Invitrogen, Carlsbad, CA, U.S.A.) using a BP reaction of Gateway system (Invitrogen). The genes then were cloned into pCPP5040, the pML123-based destination vector containing the *nptII* promoter, a gateway site, and the HA tag (Petnicki-Ocwieja et al. 2002).

To generate *pML-nptII-HopF_{Pto}^{ATG}-Flag*, the *HopF_{Pto}^{ATG}* gene was amplified by PCR using the forward primer 5'-**CGCTCGAGAAGCTT**AAAAAGAGGAAACCATT**ATGG**-3' (*Xho*I, bold; *Hind*III, italic; ATG, underlined) and the reverse primer 5'-**GCTTCGAAGACCCTTTCGACCGGCAC**-3' (*Csp*45I, bold) and pMOD::hrp-ShcF_{Pto}-*HopF_{Pto}^{ATG}-avrRpt2⁴¹⁻²⁵⁵* as template. The PCR products were digested by *Xho*I and *Csp*45I and cloned into *pBS-Flag* (J. Chen and X. Tang, unpublished), resulting in *pBS-HopF_{Pto}^{ATG}-Flag*. This plasmid then was digested with *Hind*III and *Xba*I, and the insert was cloned into the same sites in pML123, resulting in *pML-nptII-HopF_{Pto}^{ATG}-Flag*.

Secretion assays.

P. syringae bacteria were grown in KB medium (King et al. 1954) containing appropriate antibiotics. The bacterial cells were spun down at 4,000 rpm, washed twice with the hrp-inducing minimal medium (pH 5.8) (Huynh et al. 1989), diluted in MM to an optical density at 600 nm = 0.2, and grown at room temperature for another 16 h. The bacterial culture was centrifuged at 4,000 rpm for 5 min, and the supernatant was transferred into a clean microcentrifuge tube and spun at 14,000 rpm for 5 min to remove all residual bacteria. The protein in bacterial pellet and supernatant was separated by sodium dodecyl sulfate (SDS)-polyacrylamide gel electrophoresis and examined by immunoblotting using anti-HA, anti-FLAG, and anti-AvrPto antibodies.

Translocation assays.

The *avrRpt2⁴¹⁻²⁵⁵* fusion plasmids were introduced into *P. syringae* pv. *phaseolicola* NPS 3121 or *P. syringae* pv. *tomato* DC3000 and the DC3000 *hrcC* mutant strains by electroporation. Bacteria were grown overnight in liquid KB medium with appropriate antibiotics, washed twice with 10 mM MgCl₂, and resuspended in the same solution. Bacteria at the concentration of 2 × 10⁸ cells/ml were infiltrated into *Arabidopsis* Col-0 and Col-0 *rps2-201* plants with a needle-less syringe.

Interaction assay in yeast two-hybrid system.

ShcF_{Pto}, *HopF_{Pto}^{ATG}*, and *HopF_{Pto}^{GTG13}* (*HopF_{Pto}* with the N-terminal 13 codons deleted) were PCR amplified using the following three pairs of primers: 5'-**GCGAATTCATGAAAA**CGCATTTGAC-3' and 5'-**GCCTCGAGCTATTTGAGTTTT**AGTGT-3' for *ShcF_{Pto}*; 5'-**GCGAATTCATGGGTAATATT**TGCGGCACCTCG-3' and 5'-**TACTCGAGTCAGACCCTTT**CGACCGG-3' for *HopF_{Pto}*; and 5'-**GCGAATTCATGTATAG**CCCATCCCAT-3' and 5'-**TACTCGAGTCAGACCCTTTTCG**ACCGG-3' for *HopF_{Pto}^{GTG13}* (*Eco*RI and *Xho*I sites are in boldface). The PCR products were digested with *Eco*RI and *Xho*I and cloned into *pEG202* and *pJG4-5* plasmids (Golemis et al. 1997), resulting in *pEG202::ShcF_{Pto}*, *pJG4-5::HopF_{Pto}*, and *pJG4-5::HopF_{Pto}^{GTG13}*. Procedures described by Golemis and associates (1997) were followed to test the interaction of *ShcF_{Pto}/HopF_{Pto}* and *ShcF_{Pto}/HopF_{Pto}^{GTG13}* in yeast.

In vitro protein pull-down assay.

HopF_{Pto} was released from *pJG4-5::HopF_{Pto}* by *Eco*RI and *Xho*I digestion and inserted into *pGST-AvrPto* plasmid (Shan et al. 2000) for production of the glutathione-S-transferase (GST)-*HopF_{Pto}* recombinant protein. Procedures described by Guan and Dixon (1991) were followed for purification of the GST-*HopF_{Pto}* protein with the glutathione-conjugated cellulose beads (Sigma, St. Louis). The GST-AvrPto fusion protein was produced for control using the same procedure. The GST fusion proteins coupled with the cellulose beads were used in protein pull-down assay with *ShcF_{Pto}* tagged with HA.

P. syringae pv. *phaseolicola* carrying *pLK-hrp-ShcF_{Pto}-HA* was grown in 2 ml of KB medium containing kanamycin and rifampicin for overnight and then transferred into 10 ml of hrp-inducing MM. Bacteria cells were harvested 16 h after culture in MM, resuspended in 5 ml of TBS (50 mM Tris-HCl, 150 mM NaCl, pH 7.4), and sonicated on ice. The cell lysate was centrifuged at 4°C, and the supernatant was incubated with beads coupled with 50 µg of GST-*HopF_{Pto}* and GST-AvrPto, respectively, overnight at 4°C with constant shaking. The beads were washed five times with phosphate-buffered saline plus 0.1% Tween-20, boiled in 1× SDS sample buffer, and analyzed with Western blot and the anti-HA antibody.

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AUTHOR-RECOMMENDED INTERNET RESOURCE

NCBI Taxonomy Browser Genetic Codes webpage: www3.ncbi.nlm.nih.gov/Taxonomy/Utils/wprintgc.cgi?mode=c