Two Host Cytoplasmic Effectors Are Required for Pathogenesis of Phytophthora sojae by Suppression of Host Defenses

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Phytophthora sojae encodes hundreds of putative host cytoplasmic effectors with conserved FLAK motifs following signal peptides, termed crinkling- and necrosis-inducing proteins (CRN) or Crinkler. Their functions and mechanisms in pathogenesis are mostly unknown. Here, we identify a group of five P. sojae-specific CRN-like genes with high levels of sequence similarity, of which three are putative pseudogenes. Functional analysis shows that the two functional genes encode proteins with predicted nuclear localization signals that induce contrasting responses when expressed in Nicotiana benthamiana and soybean (Glycine max). PsCRN63 induces cell death, while PsCRN115 suppresses cell death elicited by the P. sojae necrosis-inducing protein (PsojNIP) or PsCRN63. Expression of CRN fragments with deleted signal peptides and FLAK motifs demonstrates that the carboxyl-terminal portions of PsCRN63 or PsCRN115 are sufficient for their activities. However, the predicted nuclear localization signal is required for PsCRN63 to induce cell death but not for PsCRN115 to suppress cell death. Furthermore, silencing of the PsCRN63 and PsCRN115 genes in P. sojae stable transformants leads to a reduction of virulence on soybean. Intriguingly, the silenced transformants lose the ability to suppress host cell death and callose deposition on inoculated plants. These results suggest a role for CRN effectors in the suppression of host defense responses.

Many plant pathogens, including bacteria, fungi, oomycetes, and nematodes, secrete distinct proteins into different cellular compartments of their hosts to modulate host defense circuits and benefit parasite colonization (Bhavsar et al., 2007; Hogenhout et al., 2009; Tyler, 2009). These pathogen-secreted proteins are named effectors (Hogenhout et al., 2009). Through coevolution, plants have developed effective surveillance systems to recognize particular effectors and induce defense pathways. This host response is called effector-triggered immunity (ETI; Chisholm et al., 2006; Jones and Dangl, 2006). A major function of pathogen effectors is believed to be the suppression of host defense responses through their interaction with critical host targets, including signal transduction pathways involved in pathogen-associated molecular pattern (PAMP)-triggered immunity (PTI) and ETI (Chisholm et al., 2006; Jones and Dangl, 2006). For example, many characterized effectors of bacterial or oomycete plant pathogens, such as Pseudomonas syringae AvrPtoB (Abramovitch et al., 2003), Phytophthora sojae Avr1b (Dou et al., 2008a), and Phytophthora infestans Avr3a (Bos et al., 2006), can suppress host cell death and/or the hypersensitive response (HR), which are involved in PTI and/or ETI.

P. sojae, causing soybean (Glycine max) root and stem rot, is a major threat to soybean cultivation and leads to annual losses of $1 to $2 billion worldwide (Tyler, 2007). Besides P. sojae, the genus Phytophthora contains over 90 species, almost all of which are destructive pathogens of a huge range of agriculturally and ornamentaly important plants (Erwin and Ribiero, 1996). Phytophthora belongs to the fungus-like oomycetes, which are evolutionarily related to algae in the kingdom Stramenopila (Kamoun, 2003; Tyler et al., 2006). Oomycetes are distinct from fungi in many aspects, including biochemical pathways and infection strategies, management approaches used for fungal pathogens are often not effective for controlling oomycete diseases (Erwin and Ribiero, 1996; Kamoun, 2003). The study of effectors and relevant functional analyses is important for understanding oomycete pathogenesis and for developing novel and effective strategies to manage disease (Hogenhout et al., 2009; Tyler, 2009). For instance, several potato (Solanum
Furthermore, strong host cell death and callose deposition were only observed in plants inoculated with the CRN-silenced transformants. Therefore, we propose a role for CRN effectors in the suppression of host defense responses.

RESULTS
Identification of a Group of P. sojae-Specific Effectors Containing the FLAK Motif and NLS

The P. sojae CRN family, containing 202 genes, has been extensively analyzed (Tyler et al., 2006; Haas et al., 2009). In this study, we selected one group of genes for functional analysis for three reasons: first, the predicted gene models encode proteins with a predicted NLS (Fig. 1; Cokol et al., 2000); second, the group shows high sequence similarity with P. infestans CRN2 (PiCRN2), which has the ability to elicit cell death in N. benthamiana and induce the expression of defense responses in tomato (Solanum lycopersicum; Torto et al., 2003); and third, selected genes within the group are the most highly expressed among all P. sojae RXLR and CRN effectors based upon transcriptional profiling of various life cycle and infection stages (W. Ye, X. Wang, D. Dou, and Y. Wang, unpublished data). Five genes in this group are PsCRN63, PsCRN77, PsCRN84, PsCRN86, and PsCRN115 (Haas et al., 2009). Among these, PsCRN77, PsCRN84, and PsCRN86 are apparently pseudogenes, because they possess one or more stop codons in their predicted open reading frames that would result in truncated proteins (Supplemental Fig. S1).

We determined the actual transcription units for PsCRN63 and PsCRN115 by reverse transcription (RT)-PCR from total RNA of the cultured mycelium of P. sojae isolate P6497 (Förster et al., 1994). Primers were designed to amplify the predicted open reading frame shifts by deleting one or two nucleotides, the theoretical protein sequences encoded by PsCRN63 and 499 amino acids (PsCRN63) and 499 amino acids (PsCRN115; Fig. 1; Supplemental Fig. S1).

After removing the stop codons or revising the frame shifts by deleting one or two nucleotides, the theoretical protein sequences encoded by PsCRN77, PsCRN84, and PsCRN86 were obtained and analyzed together with PsCRN63 and PsCRN115. They share a high level of sequence similarity and contain identical sequence segments corresponding to a NLS motif (Fig. 1; Supplemental Fig. S1). In total, the sequences of PsCRN63 and PsCRN115 are 95.7% identical at the amino acid level. The C-terminal portion of PsCRN63, which is sufficient to trigger cell death in N. benthamiana (described below; Fig. 1; Supplemental Fig. S1),
differs by only four amino acids from that of PsCRN115. To identify potentially orthologous genes in other Phytophthora species, we searched the P. infestans and P. ramorum genomes using BLASTN, with a cutoff of E \( \leq 10^{-2} \). As a result, five genes were obtained from P. infestans and one gene was obtained from P. ramorum. All of the predicted protein sequences contain the conserved signal peptides and FLAK motif but lack the NLS (Fig. 1; Supplemental Fig. S1).

Phylogenetic relationships among the 11 CRN proteins from three species of Phytophthora were modeled by using MEGA 4.1 with the neighbor-joining method (Tamura et al., 2007). On the phylogenetic tree, the 11 CRNs were sorted into three distinct groups according to species of origin (Fig. 1). The recent expansion of this class of CRN genes in the genomes of P. sojae and P. infestans, and the presence of pseudogenes in P. sojae, suggest that the genes are subject to high selective pressure and are consistent with a “rapid birth-and-death” mechanism (Tyler, 2009). Among this group of 11 closely related proteins, the P. sojae CRNs are exceptional because they are the only ones with clearly predicted NLS motifs (Fig. 1).

**Expressional Analysis of Genes in the P. sojae Group**

To investigate the relative expression level of each gene in the P. sojae group, we designed specific primers and performed quantitative RT-PCR using RNA from growing hyphae. Figure 2A shows that PsCRN63 expression is the highest among the five genes tested. Expression levels of PsCRN115 and PsCRN86 were 38.1% and 10.4% of that of PsCRN63, respectively. Transcripts of pseudogenes PsCRN77 and PsCRN84 were nearly undetectable in this assay. We also determined the expression patterns of PsCRN63 and PsCRN115 in different stages of development, including mycelium, germinated cysts, and infection stages at 1.5, 3.0, 6.0, 12.0, and 24.0 h post infection (hpi). PsCRN63 transcripts are slightly (approximately 1.5-fold) induced during the late infection stages (12.0 and 24.0 hpi), whereas the highest levels of PsCRN115 RNA are found at the mycelium stage. Overall, these results show that transcripts of PsCRN63 and PsCRN115 are detectable in all the tested stages (Fig. 2B). In addition, relative transcript levels of PsCRN63 and PsCRN115 were 12.4% to 21.8% and 3.4% to 7.1% of that of the P. sojae actin gene, respectively.

Contrasting Effects of PsCRN63 and PsCRN115 on Cell Death

We tested whether PsCRN63 or PsCRN115 proteins could induce necrosis in N. benthamiana using infiltration of Agrobacterium tumefaciens cells with a potato virus X (PVX) vector carrying each of the mature genes (the predicted signal peptides were removed). These methods have been widely used for transient expression of genes in N. benthamiana and for determining whether particular genes can induce or suppress cell death (Torto et al., 2003; Bos et al., 2006; Dou et al., 2008a; Haas et al., 2009). A known elicitor of cell death, the P. sojae necrosis-inducing protein (PsojNIP; Qutob et al., 2002), was used as positive control. Figure 3A shows that PsCRN63 triggered cell death in N. benthamiana. In contrast, only mosaic symptoms due
to PVX were observed following infiltration with PsCRN115 or gfp under the same conditions. The necrotic symptoms triggered by PsCRN63 developed about 5 d after infiltration and showed light-colored lesions, whereas PsojNIP triggered dark necrotic lesions around the site of infiltration and the symptoms developed about 4 d after infiltration (Fig. 3A). Considering that the cell death-inducing activity of PsCRN63 might possibly be caused by gene overexpression in the PVX system, we performed additional plant transformation experiments by transient expression of PsCRN63 in N. benthamiana leaves by agroinfiltration. The leaves infiltrated with PsCRN63 exhibited cell death at 5 d after infiltration. In contrast, no obvious cell death was found when a control gene, gfp, was expressed under the same conditions (Supplemental Fig. S2).

To further explore the functions of PsCRN115, we tested whether it could suppress cell death like other bacterial and oomycete effectors (Abramovitch et al., 2003; Bos et al., 2006; Dou et al., 2008a). Several cell death inducers were used in this assay: BAX (triggering HR-mimicking cell death in plants; Lacomme and Santa Cruz, 1999), PsojNIP (Qutob et al., 2002), a combination of P. infestans Avr3a and potato R3a as an HR inducer (Armstrong et al., 2005), a P. sojae RXLR effector identified as an elicitor of cell death (Avh241; Q. Wang, C. Han, and Y. Wang, unpublished data), and PsCRN63. Cell death symptoms were observed when A. tumefaciens cells carrying the above genes were infiltrated into N. benthamiana leaves (Fig. 3, B and C). However, when PsCRN115 was infiltrated in N. benthamiana leaves 12 h prior to infiltration of each of the inducers, this pretreatment blocked cell death normally triggered by PsojNIP and PsCRN63. In contrast, PsCRN115 could not protect N. benthamiana tissue from cell death triggered by BAX, Avh241, or Avr3a/R3a (Fig. 3, B and C). Furthermore, prior infiltration with A. tumefaciens cells containing the gfp gene, a negative control, did not protect against cell death. A previously identified cell death suppressor, Avh331 (Dou et al., 2008a), was used as a positive control and could block cell death triggered by all the tested effectors (Fig. 3, B and C).

To independently confirm the activities of PsCRN63 and PsCRN115 in the host soybean, we performed cobombardment assays using a double-barreled attachment for the Bio-Rad Gene Gun, which enables us to shoot two different DNA samples side by side into a leaf simultaneously. This method improves the reproducibility of the results (Dou et al., 2008a, 2008b; Kale et al., 2010). The GUS gene was used as a reporter with the indicated combination of tested genes (Table I). Cell death induced or suppressed by the test gene results in elimination or restoration of GUS reporter gene expression, respectively. Table I shows that expression of the PsCRN63 gene reduced the number of GUS-positive blue patches by 52% (compared with a gfp control gene), while expression of the PsCRN115 gene did not significantly alter reporter gene expression. These results indicate that PsCRN63 is a cell death inducer in soybean cells and that PsCRN115 is not. When PsCRN115 was coexpressed with PsCRN63, the number of GUS-positive blue patches was doubled, indicating that PsCRN115 could partially suppress PsCRN63-induced cell death. This result was obtained when the cobombardment of PsCRN63 and PsCRN115 was compared directly with cobombardment of PsCRN63 and GFP using the double-barreled bombardment (direct assay) and when each mixture was separately compared with a reference consisting of GUS plus empty vector (indirect assay; Table I).
Furthermore, the PsojNIP-induced cell death inhibition activity of PsCRN115 was also tested by the direct and indirect assays described above. All of the results indicate that PsCRN115 can inhibit PsojNIP-induced cell death in soybean and thus confirm the observations from the N. benthamiana assays (Table I).

Deletion Analysis of PsCRN63 and PsCRN115

To determine the functional regions in PsCRN63 and PsCRN115, we created seven deletion mutants for PsCRN63 and tested their ability to induce cell death using the above-introduced methods. Figure 4 demonstrates that positions from 133 to 411 of PsCRN63 can trigger a weak cell death, while positions from 133 to 425 are required for full cell death activity. We also tested the corresponding positions of PsCRN115 for cell death suppression activity. The region from 132 to 424 was sufficient to inhibit cell death induced by PsCRN63 or PsojNIP, but the region from 132 to 410 was insufficient for the suppressive activity of PsCRN115. The characterized regions do not contain the conserved FLAK motif, indicating that this motif is not required for the activity when the protein is expressed inside plant cells. Thus, the FLAK motif, as well as RXLR-dEER motifs in oomycete RXLR effectors (Whisson et al., 2007; Dou et al., 2008b) and RXLXE(Q) motifs in Plasmodium (Hiller et al., 2004; Marti et al., 2004), might serve as host translocation signals and deliver proteins into host cells. The full-length PsCRN63 protein including the putative signal peptide also triggered cell death (Fig. 4). This result supports the hypothesis that the FLAK motif performs a host-targeting function similar to the RXLR-dEER motif. However, further experiments are required to validate this hypothesis.

The Contradictory Roles of the Predicted NLS Motif in PsCRN63 and PsCRN115

Although orthologous genes to this group of P. sojae CRN effectors are present in other Phytophthora species, the NLS was only found in the P. sojae orthologs (Fig. 1). To test whether this signal is required to induce cell death, we first made mutations of this NLS [PsCRN63 (pNLSAAAA)] and then expressed it in N. benthamiana leaves by infiltration methods. Figure 5 shows that mutations of the NLS abolished the ability to trigger cell death. Furthermore, when we added a synthetic NLS to the C terminus of the mutant PsCRN63 gene [PsCRN63 (pNLSAAAA)], we found that this complemented the mutation and led to restoration of cell death-triggering activity. These results indicate that a functional NLS is required for the activity. To further confirm this, we sequestered PsCRN63 either in the nucleus or the cytoplasm by attachment of a NLS or a nuclear exclu-
sion signal (NES; Shen et al., 2007) to the C terminus, respectively. Cell death symptoms were not observed when the NES was attached (Fig. 5). To exclude the possibility that the NES could interfere with activity, a nonfunctional NES (nes) was fused to the C terminus of PsCRN63. In the nonfunctional nes, the second and third Leu residues and the first Ile residue were all replaced with Ala residues (Shen et al., 2007). This construct triggered the same symptoms as wild-type PsCRN63. Considering these findings together, we infer that PsCRN63-mediated cell death is triggered in the plant nucleus.

To test whether the predicted NLS of PsCRN115 is required to suppress cell death, we made mutations of

| Experiment | Barrel 1a | Barrel 2a | Direct Ratiob | Indirect Ratio c | ρd |
|------------|-----------|-----------|---------------|-----------------|----|
| 1          | GFP       | GFP       | 0.99 ± 0.01   |                 |    |
| 2          | GFP       | PsCRN63   | 0.48 ± 0.02   | b/a = 0.48 <0.001 |    |
| 3          | GFP       | PsCRN115  | 1.10 ± 0.04   | c/a = 1.11 >0.1  |    |
| 4          | GFP       | PsCRN63 + GFP | 0.54 ± 0.03 |                 |    |
| 5          | PsCRN63 + GFP | PsCRN63 + PsCRN115 | 1.01 ± 0.02 | e/d = 1.87 <0.001 |    |
| 6          | GFP       | PsojNIP   | 0.04 ± 0.00   |                 |    |
| 7          | GFP       | PsojNIP + PsCRN115 | 0.38 ± 0.03 | h/g = 9.50 <0.001 |    |
| 8          | PsojNIP + GFP | PsojNIP + PsCRN115 | 1.84 ± 0.40 |                 |    |

aBarrels 1 and 2 are physically identical. Half of all the replicates were conducted using the configuration of DNA samples indicated plus GUS, and half were conducted with the samples reversed between barrels 1 and 2. In all cases, the mass of DNA for each barrel was identical. bRatios between the numbers of spots produced by each barrel. Geometric averages and st were calculated from log ratios obtained from 10 to 16 pairs of shots. cComparison of the two averaged ratios from the experiments indicated by the lowercase letters. dP values for the indirect comparisons were calculated from the log ratios using the Wilcoxon rank sum test.

Figure 4. Deletion analysis of PsCRN63 and PsCRN115 defines their functional fragments. Deletion mutants of PsCRN63 (A; induction of cell death) and PsCRN115 (B; suppression of cell death) were expressed by agroinfiltration in N. benthamiana. A schematic view of the different deletion mutant constructs is shown on the left. The typical symptoms for each infiltration site are shown on the right. Photographs of symptoms were taken 5 d after infiltration. The numbers show the ratio of necrotic responses and the total number of infiltrated sites. In B, cell death inducers (PsCRN63 and PsojNIP) are indicated.
this NLS [PsCRN115(pNLSAAAA)] and then expressed it in *N. benthamiana* to test its activity. Figure 5 shows that the mutations did not abolish the ability of PsCRN115 to suppress PsCRN63- or PsojNIP-derived cell death, indicating that the predicted NLS is not required for its function. Significantly, in the region of PsCRN63 (positions 133–411) that is required for triggering cell death, the PsCRN115 sequence differs by only four residues from PsCRN63 (Supplemental Fig. S1). Despite the fact that PsCRN63 and PsCRN115 are nearly identical in sequence, the two proteins have contrasting activities and differential requirements for the NLS. We propose that PsCRN115 and PsCRN63 may share the same molecular host targets that are involved in the cell death signal transduction pathway and that their differential activities are dependent on plant nuclear localization or not.

### Generation of Transformants with Silenced PsCRN63 and PsCRN115

To obtain transformants that do not express PsCRN63 and PsCRN115, we used a gene-silencing strategy based on polyethylene glycol-mediated protoplast stable transformation of *P. sojae* (Judelson et al., 1991; Dou et al., 2008a; McLeod et al., 2008). The construct pH709 that carries the selectable marker gene for genetin resistance was used for cotransformation with the construct containing the open reading frame of PsCRN63 driven by the constitutive Ham34 promoter (Judelson et al., 1991). Using established procedures (Dou et al., 2008a, 2008b), three independent silenced transformants were identified from 25 putative transformants that could grow on selection medium containing 50 μg mL⁻¹ genetin (Shanghai Sangon BS723). In the three transformants, T3, T20, and T21, expression levels of PsCRN63 were 26.49%, 29.35%, and 16.32% relative to the wild type, respectively (Fig. 6). Meanwhile, we assessed the expression levels of PsCRN115 because it shares high sequence similarity with PsCRN63. Figure 6 shows that the expression level of PsCRN115 was also impaired, and the silencing of this gene was strongly correlated to that of PsCRN63. These results indicate that the transformants T3, T20, and T21 are deficient in mRNA accumulation for both PsCRN63 and PsCRN115.

### Silenced Lines Exhibit Reduced Virulence on Soybean

To examine the pathogenicity of the silenced transformants, we inoculated etiolated soybean seedlings (Dong et al., 2009) of cv Williams, which is susceptible to most strains of *P. sojae*, including P6497. The expression deficiency of these two CRN effectors resulted in a significant reduction of virulence (Fig. 7). Slow-spread necrotic lesions (0.5 ± 0.08 to 0.9 ± 0.29 cm) developed on plant hypocotyls inoculated with the CRN-gene silenced transformants, whereas rapidly spreading water-soaked lesions (3.45 ± 0.42 cm) were observed on plants inoculated with the wild type (Fig. 7).

### Reduction of the Ability in Silenced Lines to Suppress Host Cell Death and Callose Deposition

To analyze which stage of pathogen development is affected, we used an inverted microscope to visualize hyphae in infected tissue. At 6 hpi, hyphae of silenced line T21 and of the wild type could be equally well detected in epidermal cells (Fig. 8A). Infectious hyphae of silenced line T21 and of the wild type could be equally well detected in epidermal cells (Fig. 8A). Infectious hyphae of silenced line T21 and of the wild type could be equally well detected in epidermal cells (Fig. 8A).
phae of the wild type grew actively and occupied neighboring primary infected cells by 24 hpi. However, infectious hyphae of the silenced lines were mostly restricted to primary infected cells, and only a few infectious hyphae extended into neighboring cells, where cell death was observed (Fig. 8A). Callose deposition was also observed under the same conditions. At 6 hpi, soybean epidermal cells infected by T21 and the wild type showed similar low or undetectable levels of callose deposition. However, a strong signal of callose deposition was only found at 24 hpi in soybean epidermal cells inoculated with T21 (Fig. 8B).

Since the silenced lines apparently lost the ability to suppress host cell death and defense responses, we further determined whether the silencing of the two CRN effectors interfered with ETI. The wild-type P. sojae strain (P6497) is virulent on soybean cultivars without any known resistance genes (rps) or those carrying Rps1b, but it is avirulent on soybean cultivars carrying Rps1a, Rps1c, Rps1d, Rps1k, Rps2, Rps3a, Rps3b, Rps3c, Rps4, Rps5, or Rps6 (Table II). Two silenced lines (T3 and T21) were determined to be virulent on Rps1b- or rps-containing soybean seedlings, although development of the susceptible phenotype was delayed compared with plants inoculated with the wild type. In other incompatible reactions, the silenced transformants and wild-type strains remained avirulent (Table II). These results indicate that a reduction in expression of the genes of PsCRN63 and PsCRN115 does not interfere with ETI, at least for the interactions of the tested Avr-Rps genes.

DISCUSSION

We have examined two CRN effectors of P. sojae with high sequence similarity and high expression levels. We demonstrate that these two proteins cause opposite effects when expressed in plant cells: PsCRN63 induces cell death, while PsCRN115 suppresses cell death induced by PsojNIP or PsCRN63. Next, we showed that the C-terminal fragment of each protein was necessary and sufficient for their activities when the proteins were transiently expressed in N. benthamiana, supporting the hypothesis that oomycete CRN effectors are translocated into host cells. We further showed that the predicted NLS was required for PsCRN63 to trigger cell death but not for PsCRN115 to suppress cell death. We obtained P. sojae transformants with expression deficiencies in both genes and showed that the silenced lines had significantly reduced infection ability. The silenced lines could penetrate host cells but had lost the abilities to suppress cell death and callose deposition. From these results,

Figure 7. Reduced virulence in silenced lines on soybean. A, Etiolated seedlings of the susceptible soybean cv Williams were inoculated with the wild type (WT) and the silenced line (SL) T21. The inoculated seedlings were photographed at 48 hpi. The experiments were repeated four times in all mutants with similar results, and only one silenced line (T21) is shown as an example. B, Lesion lengths of inoculated sites measured 48 h after inoculation.

Figure 8. Silenced lines could not suppress host cell death and callose deposition. A, Microscopic observations of invasive hyphae (top panels) and cell death (bottom panels) in soybean root epidermal cells. Bars = 20 μm. B, Callose deposition in soybean root epidermal cells detected with aniline blue staining. Bars = 100 μm. In A and B, the experiments were repeated four times in all mutants with similar results; only one silenced line (T21) is shown as an example, and the average number of callose deposits per microscopic field of 1 mm² was calculated using the ImageJ software. SL, Silenced line; WT, wild type.
we inferred that the two effectors were critical to pathogenesis by modulating host defenses. 

*P. sojae*, as well as other oomycete pathogens, contains a vast repertoire of effectors predicted to act in the host cytoplasm (Tyler et al., 2006; Haas et al., 2009). RXLR and CRN effectors are two major classes of such effectors and are both identified by conserved motifs following the signal peptide (Kamoun, 2007). So far, all effectors and are both identified by conserved motifs (Jiang et al., 2008; Haas et al., 2009). Each encodes over 350 RXLR factor for *P. sojae* physiological races; in parentheses are the genes carrying resistance. Values shown refer to numbers of plants killed/numbers of all inoculated plants. 

### Table II. The phenotypes of different soybean cultivars inoculated with the silenced lines (T3 and T21) and the wild type

| Soybean Cultivar<sup>a</sup> | T3<sup>b</sup> | T21<sup>c</sup> | Wild Type |
|-----------------------------|-------------|--------------|-----------|
| Harlon (Rps1a)              | 1/8<sup>b</sup> | R<sup>c</sup> | 0/10      | R         | 2/10 | R |
| L77-1863 (Rps1b)            | 15/20       | S<sup>d</sup> | 13/20     | S         | 19/20 | S |
| Williams79 (Rps1c)          | 0/10        | R            | 0/9       | R         | 0/11 | R |
| PI103091 (Rps1d)            | 1/10        | R            | 0/10      | R         | 1/9  | R |
| Williams82 (Rps1k)          | 0/8         | R            | 0/10      | R         | 0/10 | R |
| L76-1988 (Rps2)             | 1/8         | R            | 1/9       | R         | 1/10 | R |
| Chapman (Rps3a)             | 1/10        | R            | 0/9       | R         | 1/9  | R |
| PRX146-36 (Rps3b)           | 0/9         | R            | 0/10      | R         | 0/9  | R |
| PRX145-48 (Rps3c)           | 1/8         | R            | 0/9       | R         | 1/8  | R |
| L85-2352 (Rps4)             | 3/18        | R            | 1/16      | R         | 4/17 | R |
| L85-3059 (Rps5)             | 0/10        | R            | 0/10      | R         | 0/9  | R |
| Harosoy62xx (Rps6)          | 1/10        | R            | 0/10      | R         | 1/10 | R |
| Williams (ips)              | 16/20       | S            | 11/17     | S         | 21/21 | S |

<sup>a</sup>Soybean cultivars contain 14 soybean differentials of *P. sojae* physiological races; in parentheses are the genes carrying resistance. <sup>b</sup>Values shown refer to numbers of plants killed/numbers of all inoculated plants. <sup>c</sup>R (resistance) ratio below 40%. <sup>d</sup>S (susceptible) ratio more than 60%.

CRN family effectors are named such because they can induce necrosis and leaf crinkling in plants (Torto et al., 2003). Surprisingly, we demonstrated that PsCRN115 could block cell death induced by PsojNIP and PsCRN63, indicating that this family of effectors also has similar abilities to RXLR effectors in suppressing plant defense (Bos et al., 2006; Dou et al., 2008a). Wild-type *P. sojae* can successfully invade compatible soybean cultivars, and only weak or no cell death symptoms are observed in the early infection stage (Chen et al., 2008). Since infection of PsCRN63/115-deficient transformants was hindered by premature host cell death under the same conditions, it appears that cell death suppression was severely weakened in the silenced lines. Considering that PsCRN63 is a cell death inducer and that PsCRN115 is a cell death suppressor, and that both genes are constitutively expressed, we hypothesize that these two effectors serve as essential suppressors of cell death mediated by PAMPs such as PsojNIP (Qutob et al., 2002). However, we cannot exclude other possibilities that may cause the altered host responses we observed in plants infected with the CRN-silenced transformants.

We show that PsCRN63 is a cell death inducer. Our results, together with deletion analysis of PsCRN2
and the RXLR effectors (Bos et al., 2006; Dou et al., 2008a) and PsojNIP (Qutob et al., 2002), proteins secreted into the host apoplast, such as elicinins, have been identified as modulators of host cell death. Pro-Phytophthora
tiptions and discoveries contribute to the understanding of how Phytophthora colonies fit the concept of accessibility. 

Many secreted proteins from Phytophthora have been identified as modulators of host cell death. Proteins secreted into the host apoplast, such as elicinins (Huitema et al., 2005) and PsojNIP (Qutob et al., 2002), can trigger cell death and defense, while selected RXLR effectors (Bos et al., 2006; Dou et al., 2008a) and the P. infestans SNE1 protein (Kelley et al., 2010) have the ability to suppress cell death and defense. The P. infestans RXLR effector appears to play both roles (Bos et al., 2010), as is required for full virulence. In analogous fashion, PsCRN63 and PsCRN115 have contrasting roles in manipulating plant cell death. Expression of PsCRN63 and PsCRN115 is jointly required for full pathogenesis. Our functional investigations and discoveries contribute to the understanding of how Phytophthora and other oomycete pathogens invade plants and defeat their immune systems.

In addition, the concept of accessibility was proposed to describe host cellular conditioning toward a symbiotic relationship induced by compatible pathogens (Ouchi, 1983, 2006), based on the observations that the susceptibility of a plant to an incompatible pathogen and nonpathogen could be induced by prior inoculation with a compatible pathogen (Ouchi et al., 1991). Thus, it has been assumed that compatible pathogens have evolved a variety of strategies to establish a symbiotic relationship with the host, such as suppressing defense responses in the host cell (Ouchi, 1983, 2006). Our findings that CRN effectors, as well as RXLR effectors (Bos et al., 2006; Dou et al., 2008a; Sohn et al., 2007; Kelley et al., 2010), can modulate host defense circuitry and benefit parasite colonization fit the concept of accessibility.

Materials and Methods

Sequence Search, Structure Prediction, and Phylogenetic and Molecular Evolution Analyses

The P. sojae CRN effectors (Tyler et al., 2006; Haas et al., 2009) were renamed by the order of localization within the different scaffolds (Tyler et al., 2006). The PsCRN63, PsCRN77, PsCRN84, PsCRN86, and PsCRN115 genes were obtained by a TBLASTN search (Altschul et al., 1997) using the query sequence PICRN2 (Torto et al., 2003) against the gene models of P. sojae. Then, PsCRN63 was used in TBLASTN searches against the gene models of Phytophthora ramorum and Phytophthora infestans (Tyler et al., 2006; Haas et al., 2009) with an E value cutoff of E < 10^{-4}. To obtain the theoretical protein sequences encoded by their nonseed conjugate progenitors, the sequences of three pseudogenes, PsCRN77, PsCRN84, and PsCRN86, were adjusted by removal of one or more stop codons or by correcting frame shifts by deleting one or two nucleotides.

Signal peptide scores for each gene were predicted by the SignalP3.0 server ( Bendtsen et al., 2004), and NLS was analyzed by the Predict NLS server ( Ouchi, 1983, 2006). Our findings that CRN effectors, as suppressing defense responses in the host cell (Ouchi et al., 1991), have been traced to the nucleus, we hypothesized that CRNs could modulate host defense circuitry and benefit parasite development. This hypothesis was supported by the work of others who have demonstrated that an intact NLS is required for cell death-inducing processes of PsCRN63, suggesting that the signal involved in this process and the role of PsCRN115 are currently under investigation.

Plasmids and Strain Construction

The oligonucleotides used for the following plasmid constructions are documented in Supplemental Table S1. For the PVX assay, fPsCRN63 (full length of PsCRN63), PsCRN63, PsNIP, and PsCRN115 were amplified using combinations of oligonucleotide primers PsCRN63-F1 and PsCRN63-R, PsCRN63-F2 and PsCRN63-R, PsNIP-F and PsNIP-R, and PsCRN115-F2 and PsCRN115-R, respectively. The amplicons were cloned using appropriate restriction enzymes (Supplemental Table S2) into the PVX vector pGRI07 ( Lu et al., 2003). For the PsCRN63 deletions, the mutants were amplified using combinations of oligonucleotide primers (Supplemental Tables S1 and S2).

To make the sense construct of pHamPsCRN63 (driven by the Ham34 promoter; Judelson et al., 1991), we used fPsCRN63 in the vector of pGRI07 as template to amplify PsCRN63 using PrimeSTAR HS DNA Polymerase (Takara code DR010A) with the primers PsCRN63-F1 and PsCRN63-R. The PCR product was inserted into Smal-digested pTH210 (Wang et al., 2009; Judelson et al., 1991). The constructs were screened and confirmed by sequencing. The other constructs are shown in Supplemental Table S2. All the above plasmids were validated by sequencing by GenScript.

SYBR Green Real-Time RT-PCR Assay

Total RNA was isolated from the hyphae and the germinating cysts using NucleoSpin RNA II (Macherey-Nagel) following the manufacturer’s protocol. The integrity of total RNA was confirmed by agarose gel electrophoresis. The RNA was quantified using a spectrophotometer (Nanodrop ND-1000). To remove contaminating genomic DNA in RNA preparations, 10 µg of total RNA was treated with 4 units of RNase-free DNase I (Takara) at 37°C for 30 min. The removal of DNA was verified under the same conditions as those used for the RT-PCR, except that the 30-min cDNA synthesis step at 37°C was omitted. First-strand cDNA was synthesized using Moloney murine leukemia virus reverse transcriptase (RNase free) and oligo(dT)18 primer (Invitrogen). Quantitative RT-PCR was performed in 20-µL reactions including 20 ng of cDNA, 0.2 µM gene-specific primer or reference actin gene (Supplemental Table S1), 10 µL of SYBR Premix Ex Taq (Takara), and 6.8 µL of deionized water. PCR was performed on an ABI PRISM 7300 Fast Real-Time PCR System (Applied Biosystems) under the following conditions: 95°C for 5 s, 40 cycles of 95°C for 5 s and 60°C for 31 s to calculate cycle threshold values, followed by a dissociation program of 95°C for 15 s, 60°C for 1 min, and 95°C for 15 s to obtain melt curves. The 7300 System Sequence Detection Software (version 1.2; SDS) was used to obtain relative expression levels of each sample.

P. sojae Strains, Manipulation, and Inoculation Assays

P. sojae reference strain P6497 (race 2; Forster et al., 1994) was routinely grown and maintained on V8 agar (Erwin and Riberio, 1996).
transformation was carried out as described (Dou et al., 2008a). Putative P. sojae transformants were screened for PsCRN63 transgenes by amplifying genomic DNA from each line as described previously (Dou et al., 2008a). For screening transformants for silencing of the PsCRN63 gene, total RNA was extracted from mycelia with the above methods. The silenced transformants were identified by quantitative real-time RT-PCR assay following the same steps as above.

The pathogenicity phenotypes of selected transformants were determined by inoculation of etiolated soybean (Glycine max) seedlings (Dou et al., 2009) and by hypocotyl inoculation (Tyler et al., 1995). For inoculation of etiolated soybean seedlings, cv Williams (rps) was used. The lesion diameter was measured as lesion length for quantitative virulence of P. sojae transformants. There were three replicates of five to 10 plants each within each experiment. For light microscopy, after inoculation of P. sojae, the epidermis of hypocotyls of etiolated soybean seedlings was detached using an Olympus 1X71 inverted microscope to visualize hyphae in infected tissue. For colostage staining, after the inoculation, the hypocotyls of etiolated soybean seedlings was stained with anilin blue for 30 min, and then the epidermal cells were observed with an Olympus IX71 inverted fluorescent microscope. The average number of colostage deposits per microscopic field of 1 mm² was calculated from all the tested hypocotyls using the ImageJ software (http://www.uhnresearch.ca/wefi).

For hypocotyl inoculation, the following differential cultivars were used: Harlon (Rps1a), L77-1863 (Rps1b), Williams79 (Rps1c), PI103091(Rps1d), Williams82 (Rps1k), L76-1988 (Rps2), Chapman (Rps3a), PRX146-36 (Rps3b), PRX145-48 (Rps5), L85-2352 (Rps6), L85-3509 (Rps9), Harosovy62ccx (Rps6), and Williams (rps). Seedlings of each soybean cultivar were grown in the greenhouse as described (Dou et al., 2008b) for 7 d. Each virulence determination was repeated at least three times. A strain was considered avirulent if more than 60% of the inoculated seedlings survived.

**Agrobacterium tumefaciens Infiltration Assays**

The *A. tumefaciens* infiltration assays were performed as described by Dou et al. (2008a), except that *A. tumefaciens* strain GV3101 (Hellens et al., 2000) was used. For infiltration, recombinant strains were cultured in Luria-Bertani medium supplemented with 50 mg mL⁻¹ kanamycin in a test tube at 28°C to 30°C and 220 rpm for 48 h. The cells were collected by centrifugation (3,000g, 5 min), washed three times in 10 mM MgCl₂, and then resuspended in 10 mM MgCl₂ to an optical density at 600 nm of 0.4 to 0.6. Infiltration experiments were performed on 7- to 8-week-old *Nicotiana benthamiana* plants. Plants were grown and maintained throughout the experiments in a greenhouse with an ambient temperature of 22°C to 25°C and high light intensity under a 16-h/8-h light/dark photoperiod. For cell death induction experiments, *A. tumefaciens* cell solutions carrying the respective constructs were infiltrated into *N. benthamiana* leaves by pressure infiltration: a small nick was placed in each leaf with a needle, and then 30 to 50 μL of cell suspension was infiltrated through the nick using a syringe without a needle. For cell death suppression experiments, *A. tumefaciens* cells carrying the cell death-inducing genes (*PsCRN63, PsqNIP*, *Bax*, and the combination of *Avr3a* and *R3a*) were infiltrated into the same site 12 h later. Symptoms development was monitored from 4 to 8 d after infiltration, and photographs were taken after 5 d. The experiments were repeated at least three times. Although it is likely that PVX replication occurred in the transformed plant cells, resulting in amplified expression of the genes in the PVX vector, no attempt was made to quantitate PVX replication (Dou et al., 2008a).

**Particle Bombardment Assays**

Particle bombardment assays were performed using a double-barreled extension of the Bio-Rad He/1000 particle delivery system (Dou et al., 2008a, 2008b; Kale et al., 2010). Analyzing the bombardment data as a ratio between the test and control shots improves the reproducibility of the measurements greatly (Dou et al., 2008a, 2008b; Kale et al., 2010). The cell death-induction activity of *PsCRN63* and *PsCRN115* constructs was measured as the reduction in the number of blue spots comparing the *PsCRN63* or *PsCRN115* + GUS bombardment with GUS + GFP bombardment. The cell death suppression activity of *PsCRN115* was measured in two ways: (1) the number of blue spots comparing the *PsCRN115* + cell death inducer (PsqNIP or *PsCRN63*) + GUS bombardment with the cell death inducer (PsqNIP or PsCRN63) + GUS bombardment (direct assay); and (2) the number of blue spots comparing the ratio of *PsCRN115* or *PsqNIP/ PsCRN63* + GUS bombardment with the GFP + GUS bombardment control with the ratio of the *PsqNIP/ PsCRN63* + GUS bombardment with the GFP + GUS bombardment control (indirect assay). For each paired shot, the logarithm of the ratio of the spot numbers of *PsCRN63* or *PsCRN115* to that of the control was calculated. P values was calculated using the Wilcoxon rank sum test for the direct assay results and the Wilcoxon signed rank test for the direct assay results.

**Supplemental Data**

The following materials are available in the online version of this article.

**Supplemental Figure S1.** Text file of unmasked alignment corresponding to the phylogenetic tree in Figure 1.

**Supplemental Figure S2.** *PsCRN63* triggers cell death when expressed from plant expression vector pCHF3.

**Supplemental Table S1.** Oligonucleotides used for PCR and plasmid construction.

**Supplemental Table S2.** Description of plasmids used.

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Functional Analysis of *Phytophthora* CRN Effectors