

# A Humidity-Sensitive Arabidopsis Copine Mutant Exhibits Precocious Cell Death and Increased Disease Resistance

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The copines are a newly identified class of calcium-dependent, phospholipid binding proteins that are present in a wide range of organisms, including *Paramecium*, plants, *Caenorhabditis elegans*, mouse, and human. However, the biological functions of the copines are unknown. Here, we describe a humidity-sensitive copine mutant in *Arabidopsis*. Under nonpermissive, low-humidity conditions, the *cpn1-1* mutant displayed aberrant regulation of cell death that included a lesion mimic phenotype and an accelerated hypersensitive response (HR). However, the HR in *cpn1-1* showed no increase in sensitivity to low pathogen titers. Low-humidity-grown *cpn1-1* mutants also exhibited morphological abnormalities, increased resistance to virulent strains of *Pseudomonas syringae* and *Peronospora parasitica*, and constitutive expression of pathogenesis-related (PR) genes. Growth of *cpn1-1* under permissive, high-humidity conditions abolished the increased disease resistance, lesion mimic, and morphological mutant phenotypes but only partially alleviated the accelerated HR and constitutive PR gene expression phenotypes. The disease resistance phenotype of *cpn1-1* suggests that the CPN1 gene regulates defense responses. Alternatively, the primary function of CPN1 may be the regulation of plant responses to low humidity, and the effect of the *cpn1-1* mutation on disease resistance may be indirect.

## INTRODUCTION

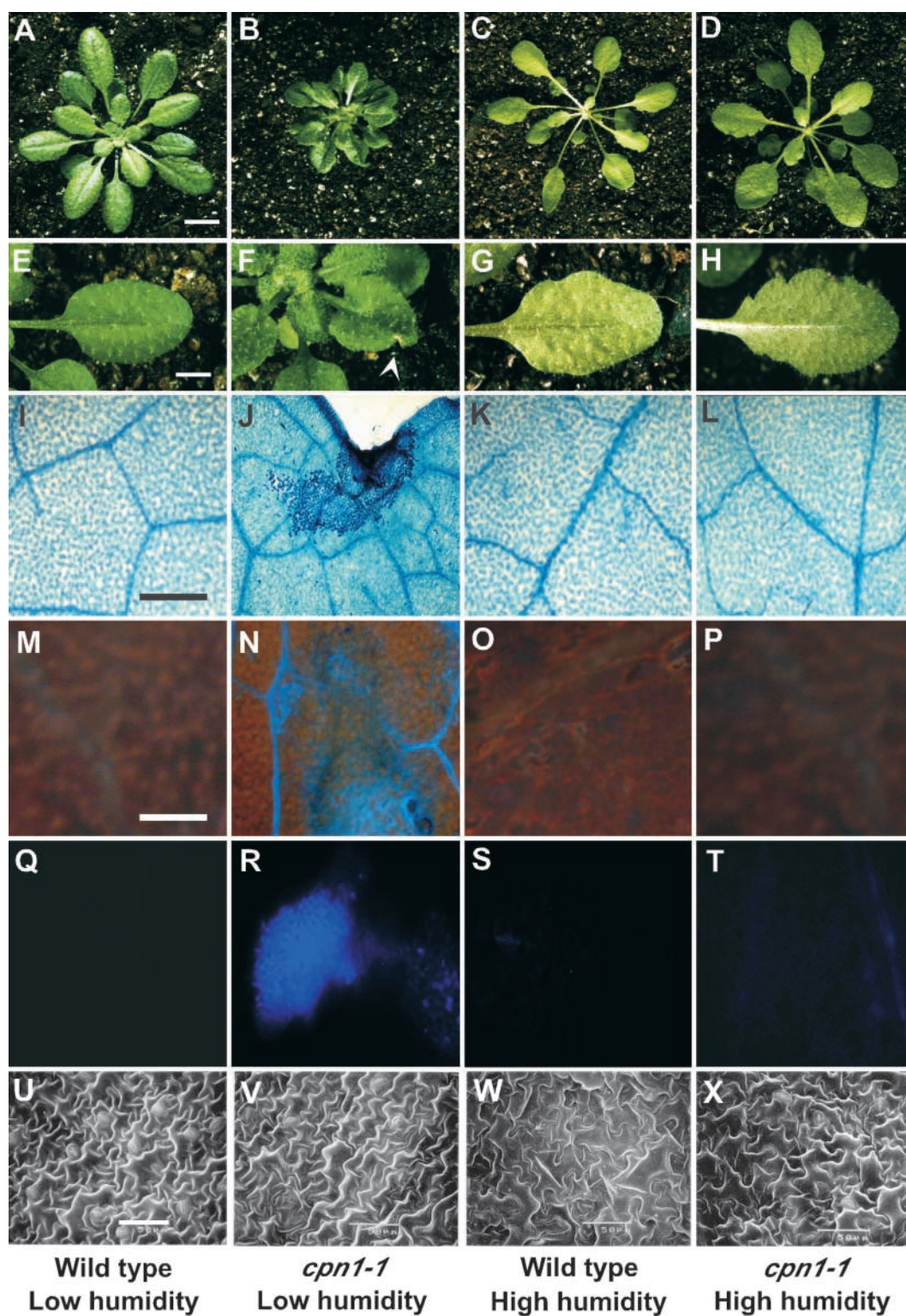
The copines are a highly conserved class of proteins found in a wide variety of organisms, including ciliates, plants, *Caenorhabditis elegans*, mouse, and human (Creutz et al., 1998). The copines are defined as a protein class by their unique combination of two C2 domains in the N-terminal region and an A domain in the C-terminal region. C2 domains are present in a large number of proteins, including protein kinase C (Azzi et al., 1992), phospholipase C (Essen et al., 1996), synaptotagmin (Brose et al., 1995), Doc2 (Duncan et al., 2000), rabphilin (Wang et al., 2000), and Munc13 (Brose et al., 1995). The C2 domain appears to confer Ca<sup>2+</sup>-dependent phospholipid binding activity to proteins, and mammalian copine I has Ca<sup>2+</sup>-dependent phospholipid binding activity (Creutz et al., 1998). The A domain may be involved in extracellular protein–protein interactions, as in the case of the von Willebrand factor (Williams et al., 1999). However, available biochemical evidence indicates that copines are intracellular proteins (Creutz et al., 1998; Tomsig and Creutz, 2000). Finally, biochemical analysis of human copine III indicates that it has an intrinsic kinase activity that may be

novel, because no homology with known kinase catalytic domains exists in copine III (Caudell et al., 2000). Together, these biochemical data raise the intriguing possibility that copines represent a new group of Ca<sup>2+</sup>-dependent signal transduction proteins that might be involved in membrane trafficking. However, the biological functions of the copines are unknown at present.

Calcium is an important second messenger involved in a variety of plant responses to stimuli, including pathogens, drought, light, hormones, touch, cold, heat, and oxidative stress (Reddy, 2001). In plant disease resistance responses, Ca<sup>2+</sup> acts as a signaling molecule, communicating primary recognition events to multiple downstream responses (Grant et al., 2000) such as phytoalexin production, induction of defense genes (Zimmermann et al., 1997), and the hypersensitive response (HR) (Xu and Heath, 1998). In parsley cells, an elicitor-responsive calcium channel has been identified and characterized (Zimmermann et al., 1997), and a transient influx of Ca<sup>2+</sup> and H<sup>+</sup> and an efflux of K<sup>+</sup> and Cl<sup>-</sup> occur within minutes of fungal elicitor addition (Hahlbrock et al., 1995; Yang et al., 1997). Pharmacological studies with tobacco By2 cells revealed that the HR requires serine proteases, calcium, and protein kinases (Sasabe et al., 2000).

The HR is a programmed cell death response: it is an active process involving transcription and translation, is under genetic control, and has some similarities to mammalian

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**Figure 1.** Humidity-Dependent Developmental and Lesion Mimic Phenotypes of *cpn1-1*.

The four columns of images, from left to right, represent Col-0 wild type grown in LH, *cpn1-1* grown in LH, Col-0 wild type grown in HH, and *cpn1-1* grown in HH, respectively. All plants were grown in SD light conditions.

apoptosis (Mittler et al., 1997; del Pozo and Lam, 1998; Maleck and Lawton, 1998). Because of the strong association of the HR with plant disease resistance, the HR is postulated to play an important role in resistance to at least some kinds of pathogens and may serve to localize a pathogen (Goodman and Novacky, 1994). A number of mutants have been identified in Arabidopsis that exhibit precocious cell death, such as the lesion mimic mutants *lsd1-lsd7*, *acd2*, *acd6*, and *cpr6* (Greenberg and Ausubel, 1993; Dietrich et al., 1994, 1997; Greenberg et al., 1994; Weymann et al., 1995; Rate et al., 1999). Many lesion mimic mutants are dependent on certain environmental conditions such as light, daylength, and RH for lesion development (Dietrich et al., 1994; Chamnongpol et al., 1996; Morel and Dangl, 1999; Yoshioka et al., 2001). Interestingly, the *DND1* gene, which eliminates the HR and can cause a microscopic lesion phenotype when mutated, encodes a putative  $\text{Ca}^{2+}$  channel, providing another potential link between  $\text{Ca}^{2+}$  and cell death (Clough et al., 2000).

Many lesion mimic mutants exhibit a state of increased disease resistance called systemic acquired resistance (SAR) and show high, constitutive levels of pathogenesis-related (*PR*) gene expression (Dietrich et al., 1994). It is possible that lesion mimic mutants represent genes that regulate the HR. However, disruptions of cellular physiology apparently unrelated to disease defense also can trigger cell death and SAR (Mock et al., 1999; Molina et al., 1999). This fact can make it difficult to determine whether or not genes defined by lesion mimic mutants play a direct role in cell death signaling and control. In addition, many Arabidopsis mutants (*cpr1*, *cpr5*, *edr1*, *mpk4*) have been identified that show constitutive expression of defense-related genes in the absence of cell death (Bowling et al., 1994, 1997; Clarke et al., 1998; Frye et al., 2001; Petersen et al., 2000). This indicates that although the HR can trigger systemic *PR* gene expression and SAR, cell death is not always required for *PR* gene expression and SAR to occur.

Here, we describe a humidity-sensitive Arabidopsis mutant that exhibited precocious cell death, increased resistance to virulent bacterial and oomyceteous pathogens, and morphological abnormalities. On the basis of molecular analyses and complementation studies, we demonstrate that the phenotype of the mutant is caused by a mutation in a copine gene, *CPN1*.

## RESULTS

### *cpn1-1* Is a Humidity-Sensitive Lesion Mimic Mutant

The *cpn1-1* mutant was identified from a population of Arabidopsis mutagenized by T-DNA insertion (Weigel et al., 2000). The *cpn1-1* mutant exhibited clear humidity-dependent developmental and growth abnormalities, as shown in Figure 1. *cpn1-1* plants grown in low-humidity (LH; 35 to 45% RH), short-day (SD; 8-hr-light/16-hr-dark) growth conditions were smaller than their wild-type Columbia (Col-0) counterparts and had a more compact growth habit (Figures 1A and 1B). The leaves of LH-grown *cpn1-1* mutants were curled, thicker than wild-type leaves, somewhat irregular in shape with short petioles, and occasionally had indentations at the margins. When *cpn1-1* mutants were grown in high-humidity (HH; 75 to 85% RH), SD conditions, these developmental defects were absent and the *cpn1-1* plants were morphologically indistinguishable from wild-type plants (Figures 1C and 1D). The phenotype was dependent on the age of the plants: *cpn1-1* seedlings were indistinguishable from wild-type seedlings for the first 1 to 2 weeks of growth in LH, SD conditions; the developmental defects manifested after that time. When grown under LH, long-day (LD; 16-hr-light/8-hr-dark) conditions, *cpn1-1* mutants were much smaller than Col-0 wild-type plants and had dramatically reduced floral apical dominance, shorter bolts, fewer flowers, and reduced seed yield (data not shown). Growth in HH, LD conditions rescued the *cpn1-1* mutant, although the mutant was still slightly smaller than the wild type under these conditions (data not shown).

The leaves of LH-grown *cpn1-1* plants displayed small necrotic lesions (Figure 1F). The largest of these lesions usually occurred at or near the margins of the leaf. No lesions were observed in HH-grown *cpn1-1* plants (Figure 1H). Lesions began to appear on the third set of true leaves after 2 weeks of growth in LH, SD conditions. These lesions occurred in *cpn1-1* plants that had not been inoculated with a pathogen, and they occurred consistently in all *cpn1-1* plants grown in LH. However, it was not clear whether or not these lesions were truly spontaneous, because it was not feasible to grow *cpn1-1* mutants under axenic, LH conditions and monitor lesion development.

**Figure 1.** (continued).

(A) to (D) Four-week-old whole plants. Bar in (A) = 1 cm for (A) to (D).

(E) to (H) Close-up images of leaves. Arrowhead in (F) denotes a lesion. Bar in (E) = 0.5 cm for (E) to (H).

(I) to (L) Sectors of leaves stained with the vital stain trypan blue. Bar in (I) = 200  $\mu\text{m}$  for (I) to (L).

(M) to (P) UV autofluorescence images of leaf sectors. Bar in (M) = 100  $\mu\text{m}$  for (M) to (P).

(Q) to (T) Leaf sectors stained with aniline blue to detect callose accumulation. The scale is the same as in (M) to (P).

(U) to (X) Scanning electron micrographs of the abaxial surfaces of leaves. Bar in (U) = 50  $\mu\text{m}$  for (U) to (X).

The *cpn1-1* lesions stained with trypan blue, indicating the presence of dead cells (Figure 1J). In addition, trypan blue staining revealed numerous microscopic lesions consisting of clusters of a few dead cells throughout LH-grown *cpn1-1* leaves (Figure 1J and data not shown). No lesions or dead cells were observed in LH- or HH-grown Col-0 wild-type leaves or in HH-grown *cpn1-1* leaves (Figures 1I, 1K, and 1L).

The leaves of LH-grown *cpn1-1* plants showed accumulation of autofluorescent phenolic compounds in the areas near the lesions and in scattered patches throughout the leaf (Figure 1N). Autofluorescent phenolic compounds were not observed in LH- or HH-grown Col-0 plant leaves or in HH-grown *cpn1-1* plant leaves (Figures 1M, 1O, and 1P). LH-grown *cpn1-1* leaves also showed substantial accumulation of callose, as determined by aniline blue staining (Figure 1R), whereas LH- and HH-grown Col-0 wild-type plant leaves and HH-grown *cpn1-1* plant leaves did not (Figures 1Q, 1S, and 1T). The accumulation of autofluorescent phenolic compounds and callose has been observed in a number of lesion mimic mutants and is a hallmark of plant stress, including pathogen stress (Bowling et al., 1994; Dietrich et al., 1994).

Because the *cpn1-1* mutant was smaller than the Col-0 wild type when grown in LH, we speculated that the cell size might be reduced in *cpn1-1*. Light microscopic analysis of LH-grown *cpn1-1* mutant leaves indicated that the cell size was reduced (Figure 1J and data not shown). In contrast, however, scanning electron microscopy of LH-grown *cpn1-1* and Col-0 wild-type abaxial leaf epidermal cells showed no dramatic difference in cell size (Figures 1U and 1V). These observations imply that the mesophyll cells may be smaller in LH-grown *cpn1-1* than in LH-grown wild type, whereas the abaxial epidermal cells in LH-grown *cpn1-1* may be normal in size but fewer in number. The borders separating cells appeared somewhat thicker in LH-grown *cpn1-1* leaves compared with LH-grown Col-0 wild-type leaves, possibly indicating the presence of thicker cell walls in LH-grown *cpn1-1* plants. Abaxial epidermal cell size in HH-grown Col-0 wild-type and *cpn1-1* mutant plant leaves was similar and larger than that observed in LH-grown wild-type and *cpn1-1* plants (Figures 1W and 1X).

### ***cpn1-1* Has an Accelerated Hypersensitive Cell Death Response**

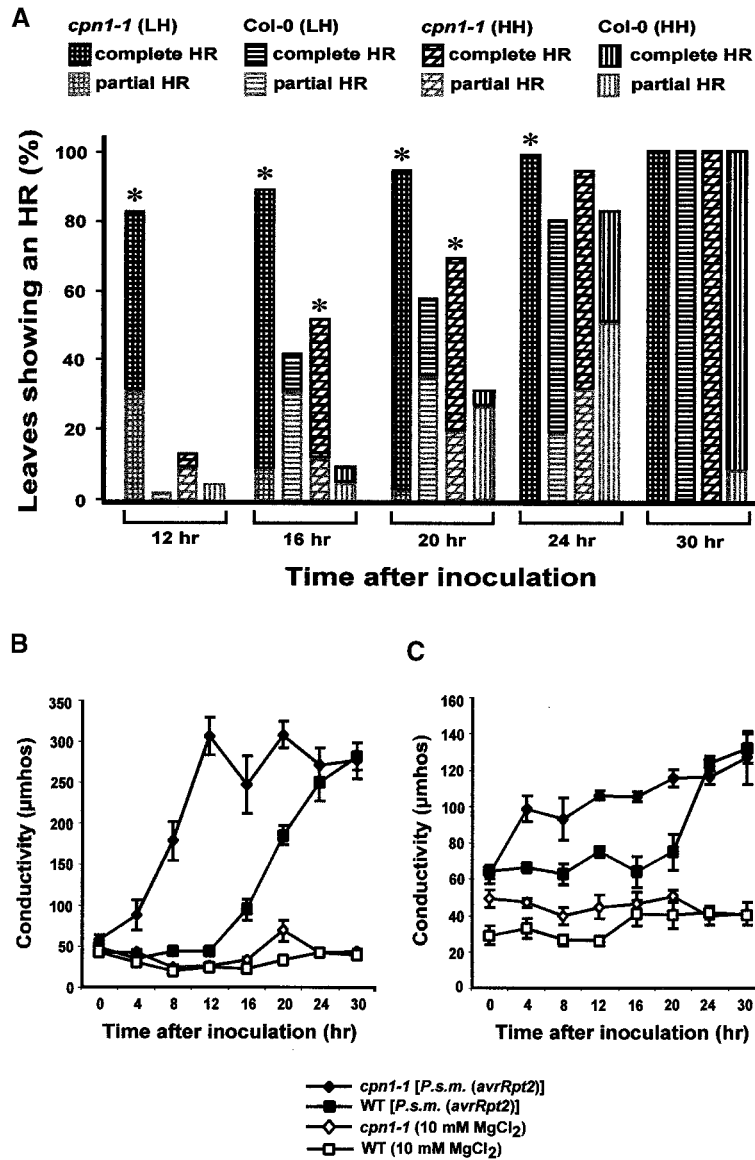
The development of necrotic lesions in *cpn1-1* plants implies that the *cpn1-1* mutation causes aberrant regulation of cell death. To investigate this possibility further, the time of HR onset after inoculation with avirulent *Pseudomonas syringae* pv *maculicola* (*avrRpt2*) (*P.s.m.* [*avrRpt2*]) was compared in *cpn1-1* and Col-0 wild-type plants. The macroscopic HR can be induced by infiltrating a high concentration ( $10^8$  colony-forming units [cfu]/mL) of avirulent bacteria into the intercellular spaces of the mesophyll of

resistant plant leaves (Klement et al., 1963). The HR is observed as a massive collapse of infiltrated tissue within ~12 to 36 hr after infiltration and involves the loss of membrane integrity and electrolyte leakage (Goodman, 1968, 1972).

Col-0 wild-type plants possess the *RPS2* disease resistance gene and are able to recognize and resist *P. syringae* bacteria bearing the corresponding *avrRpt2* avirulence gene through gene-for-gene-mediated disease resistance (Kunkel et al., 1993). *cpn1-1* was isolated from the Col-0 genetic background and appeared to have a functional *RPS2* gene (see below). *cpn1-1* and Col-0 wild-type leaves were infiltrated with high concentrations of *P.s.m.* (*avrRpt2*) bacteria, and the appearance of the HR was monitored over time. Interestingly, both LH- and HH-grown *cpn1-1* plants exhibited a faster HR than Col-0 wild-type plants grown under the same conditions, as measured by both the onset of macroscopic leaf collapse and electrolyte leakage, as shown in Figure 2.

LH-grown *cpn1-1* plants had significantly more HR responses than LH-grown Col-0 wild-type plants at 12, 16, 20, and 24 hr after inoculation, whereas HH-grown *cpn1-1* plants had significantly more HR responses than HH-grown Col-0 wild-type plants at 16 and 20 hr after infiltration (Figure 2A). By 30 hr after infiltration, all *cpn1-1* and Col-0 wild-type leaves had collapsed. Significant electrolyte leakage could be detected from LH-grown *cpn1-1* tissues infiltrated with *P.s.m.* (*avrRpt2*) within 4 to 8 hr after inoculation, whereas similar leakage from LH-grown Col-0 wild-type tissues did not occur until 16 to 20 hr after inoculation (Figure 2B). The peak of electrolyte leakage occurred at 12 hr after inoculation in LH-grown *cpn1-1*, whereas in LH-grown Col-0, the peak was observed at 24 hr after inoculation. In HH-grown *cpn1-1*, an increase in ion leakage was observed at 4 hr after inoculation, whereas a similar increase in electrolyte leakage in Col-0 wild type was not observed until 24 hr after inoculation. Some of the electrolyte leakage from HH-grown Col-0 wild-type and *cpn1-1* leaves may have been attributable to tissue damage from the infiltration, because the HH-grown leaf tissues were very tender and easily damaged. Together, these results indicated that not all aspects of the *cpn1-1* mutant phenotype were humidity sensitive.

Some lesion mimic mutants have a "hair trigger" HR, meaning that they will develop a macroscopic HR at lower bacterial inoculum concentrations than do wild-type plants (Dietrich et al., 1994). *cpn1-1* was tested for a hair trigger HR phenotype by determining the threshold concentration of *P.s.m.* (*avrRpt2*) that could trigger a macroscopic HR in *cpn1-1* compared with Col-0 wild-type plants. The minimum concentration of *P.s.m.* (*avrRpt2*) that was sufficient to evoke a macroscopic HR was  $5 \times 10^7$  cfu/mL in LH- and HH-grown Col-0 wild-type and *cpn1-1* plants (data not shown). Therefore, although *cpn1-1* plants had a faster HR than Col-0 wild-type plants, they did not appear to have increased sensitivity to low titers of *P.s.m.* (*avrRpt2*).



**Figure 2.** Accelerated HR in *cpn1-1*.

The macroscopic HR was first observed at ~12 hr after inoculation. Fifty to 100 leaves of each type were inoculated for each experiment. For details, see Methods.

**(A)** Proportions of inoculated Col-0 wild-type and *cpn1-1* leaves exhibiting partial and complete collapse over time after inoculation. Asterisks indicate percentages of leaves with partial or complete HR collapse in *cpn1-1* plants that are significantly higher than Col-0 wild-type plants grown under the same humidity conditions, at the same time during the experiment, using Student's *t* test. A leaf was scored as showing complete HR when the leaf had collapsed totally; a leaf was scored as showing partial HR when from 10 to 90% of the infiltrated leaf had collapsed. Leaves showing <10% collapse were not scored as having an HR.

**(B)** Electrolyte leakage from inoculated LH-grown Col-0 wild-type and *cpn1-1* leaves over time after infiltration with *P.s.m.* (*avrRpt2*) or 10 mM MgCl<sub>2</sub> (control).

**(C)** Electrolyte leakage from inoculated HH-grown Col-0 wild-type and *cpn1-1* leaves over time after infiltration with *P.s.m.* (*avrRpt2*) or 10 mM MgCl<sub>2</sub> (control).

WT, wild type.

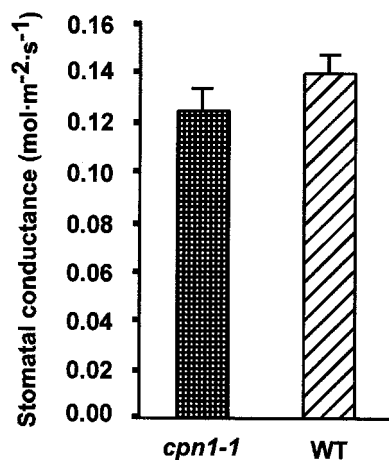
Bars in **(B)** and **(C)** indicate standard errors.

### *cpn1-1* Has Normal Stomatal Conductance

The sensitivity of *cpn1-1* to low RH led us to speculate that *cpn1-1* might be defective in responding to low humidity. For example, if *cpn1-1* were defective in controlling water loss in low humidity, this might trigger stress, cell death, and other physiological problems. However, the stomatal conductance of LH-grown *cpn1-1* and Col-0 wild-type plants was similar, as shown in Figure 3. This finding suggests that *cpn1-1* is not defective in the regulation of stomatal conductance. Also, *cpn1-1* did not appear to be particularly sensitive to drought stress, because it did not have a wilted phenotype or a lower tolerance than Col-0 wild type for low soil water potential (data not shown).

### *cpn1-1* Plants Exhibit Strong, Humidity-Dependent Resistance to Virulent *P. syringae*

Many lesion mimic mutants have increased resistance to virulent pathogens. LH-grown *cpn1-1* had dramatically increased resistance to virulent *P. syringae* pv *tomato* (*P.s.t.*) bacteria compared with Col-0 wild-type plants, as shown in Figure 4. Four days after dip inoculation with  $5 \times 10^8$  cfu/mL *P.s.t.*, wild-type plants showed severe disease symptoms typical of bacterial speck disease, including massive tissue collapse, formation of water-soaked lesions, and chlorosis (Figure 4A). In contrast, the *cpn1-1* mutant showed virtually no symptoms of the disease after inoculation (Figure 4B). The leaf tissues of inoculated *cpn1-1* mutant plants remained green and healthy, although symptoms occasion-



**Figure 3.** Stomatal Conductance in Col-0 Wild-Type and *cpn1-1* Leaves.

Data represent the average stomatal conductance for LH-grown *cpn1-1* and Col-0 wild-type (WT) plants. Stomatal conductance represents the rate of passage of water vapor and carbon dioxide through the stomata of the plant. Bars indicate standard errors.



**Figure 4.** Humidity-Dependent Resistance of *cpn1-1* to Virulent *P.s.t.*

(A) LH-grown Col-0 wild-type plants 4 days after inoculation with virulent *P.s.t.*

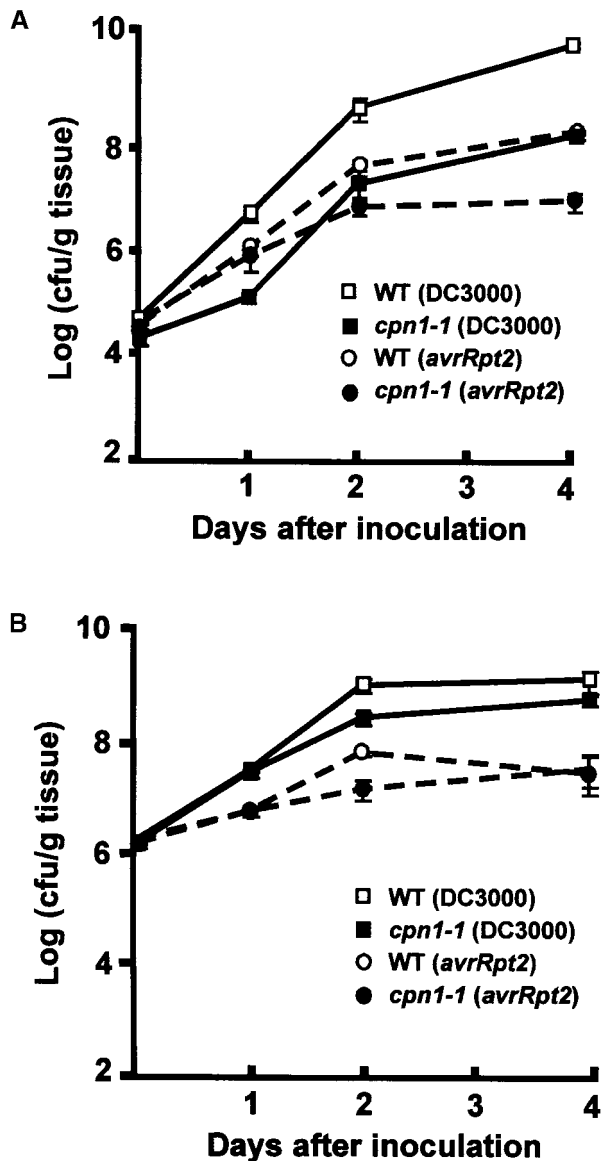
(B) LH-grown *cpn1-1* plants 4 days after inoculation with virulent *P.s.t.*

(C) HH-grown Col-0 wild-type plants 4 days after inoculation with virulent *P.s.t.*

(D) HH-grown *cpn1-1* plants 4 days after inoculation with virulent *P.s.t.*

ally occurred on older leaves, especially when the bacterial inoculum was  $>10^8$  cfu/mL (Figure 4B). HH-grown *cpn1-1* plants did not have increased resistance to *P.s.t.* compared with Col-0 wild-type plants, as measured by visible symptoms (Figures 4C and 4D).

The ability of virulent *P. syringae* bacteria to grow within LH-grown *cpn1-1* mutant leaf tissue was sharply restricted compared with that observed in LH-grown Col-0 wild-type tissue, as shown in Figure 5A. When Col-0 wild-type leaves were infiltrated with  $10^5$  cfu/mL virulent *P.s.t.*, the bacterial population increased by several orders of magnitude over the course of 4 days. In contrast, when the leaves of LH-grown *cpn1-1* mutant plants were infiltrated with virulent *P.s.t.*, the bacteria were able to multiply only to a much lower level, approximately two or three orders of magnitude less than that observed in wild-type plants. The restriction of virulent *P.s.t.* multiplication in *cpn1-1* mutant leaf tissues was consistent with the dramatic reduction of bacterial speck disease symptoms observed in inoculated LH-grown *cpn1-1* mutant plants. Virulent *P.s.t.* grew to similar levels in HH-grown Col-0 wild-type and *cpn1-1* mutant plant leaves after the plants were dip inoculated with a *P.s.t.* bacterial suspension of  $2 \times 10^7$  cfu/mL (Figure 5B). These results were consistent with the strong bacterial speck disease symptoms observed in inoculated HH-grown Col-0 wild-type and *cpn1-1* mutant plants and indicated that the increased resistance of *cpn1-1* to virulent *P.s.t.* was a humidity-sensi-



**Figure 5.** Humidity-Dependent Inhibition of Bacterial Growth in *cpn1-1*.

**(A)** Growth of virulent *P.s.t.* (DC3000) and avirulent *P.s.t.* (*avrRpt2*) bacteria in LH-grown Col-0 wild-type (WT) and *cpn1-1* plants. Plants were inoculated with the bacterial strains ( $10^5$  cfu/mL) by syringe infiltration of the leaf mesophyll. For details, see Methods.

**(B)** Growth of virulent *P.s.t.* (DC3000) and avirulent *P.s.t.* (*avrRpt2*) bacteria in HH-grown Col-0 wild-type (WT) and *cpn1-1* plants. Plants were inoculated with the bacterial strains ( $2 \times 10^7$  cfu/mL) by dipping, as described in Methods. This accounts for the higher initial bacterial count in **(B)** compared with that in **(A)**.

Bars in **(A)** and **(B)** indicate standard errors.

tive phenotype. Growth curves determined by infiltration inoculation or dip inoculation produced similar results (data not shown).

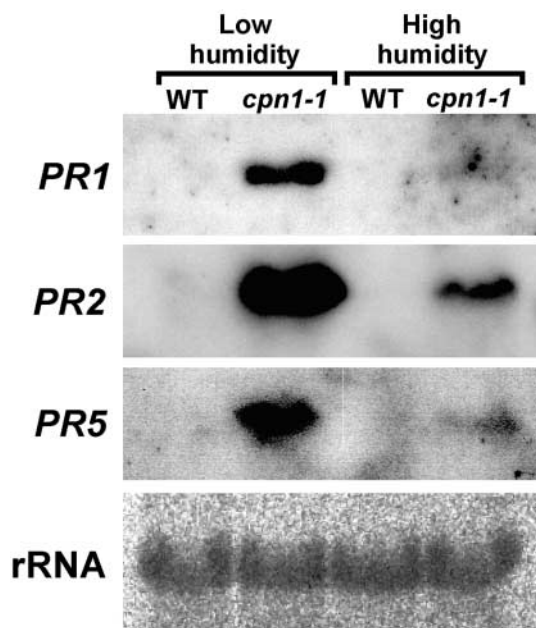
To gauge the relative level of resistance of LH-grown *cpn1-1* plants to virulent *P.s.t.*, the degree of restriction of virulent *P.s.t.* growth in *cpn1-1* was compared with the restriction of avirulent *P.s.t.* growth in Col-0 wild-type plants (Figure 5A). Growth of avirulent *P.s.t.* (*avrRpt2*) was sharply restricted in both Col-0 and *cpn1-1* plants grown under LH conditions. The degree of restriction of virulent *P.s.t.* growth in LH-grown *cpn1-1* plants was similar to the restriction of avirulent *P.s.t.* (*avrRpt2*) bacterial growth in LH-grown Col-0 wild-type plants. This indicates that the level of resistance of LH-grown *cpn1-1* plants to virulent *P.s.t.* was similar to the level of resistance of LH-grown Col-0 wild-type plants to avirulent *P.s.t.* (*avrRpt2*) bacteria. Interestingly, the growth of avirulent *P.s.t.* (*avrRpt2*) bacteria in LH-grown *cpn1-1* was moderately but significantly lower than the growth of virulent *P.s.t.* in LH-grown *cpn1-1* plants, suggesting that there may be a slight additive effect of the *cpn1-1* mutation and *RPS2*-mediated gene-for-gene resistance responses. The growth of avirulent *P.s.t.* (*avrRpt2*) bacteria was restricted in both Col-0 and *cpn1-1* plants grown under HH conditions (Figure 5B). This indicates that *RPS2*-mediated gene-for-gene recognition and resistance was intact in *cpn1-1* and that disease resistance mechanisms functioned normally in plants grown under high RH.

#### *cpn1-1* Plants Constitutively Express Defense-Related Genes

Because *cpn1-1* mutant plants had increased resistance to *P.s.t.* and also had a lesion mimic phenotype, we reasoned that LH-grown *cpn1-1* plants might be exhibiting SAR. Therefore, the expression of defense-related genes was investigated in *cpn1-1* plants. LH-grown *cpn1-1* plants showed high-level accumulation of *PR1*, *PR2*, and *PR5* transcripts, whereas these transcripts were not detectable in LH-grown Col-0 wild-type plants, as shown in Figure 6 (Uknes et al., 1992). This result suggests that the *cpn1-1* mutant was undergoing constitutive SAR in LH. Interestingly, there was moderate accumulation of *PR2* transcript and very low but detectable accumulation of *PR5* transcript in HH-grown *cpn1-1* plants. In contrast, transcripts were not detected for any of these genes in HH-grown Col-0 wild-type. This finding indicates that growth in HH did not completely abolish the effects of the *cpn1-1* mutation on *PR* gene expression.

#### *cpn1-1* Has Enhanced Resistance to an Oomyceteous Pathogen

The increased resistance to virulent *P.s.t.* and the constitutive *PR* gene expression observed in *cpn1-1* implied that



**Figure 6.** PR Gene Transcript Accumulation in *cpn1-1* and Col-0 Wild Type.

RNA gel blot analyses of the transcript levels of *PR1*, *PR2*, and *PR5* in LH- and HH-grown Col-0 wild-type (WT) and *cpn1-1* leaf tissues. Twenty micrograms of leaf total RNA were loaded in each lane. rRNA, 28S rRNA stained with methylene blue to show equal RNA loading in each lane.

*cpn1-1* was undergoing salicylic acid-dependent SAR in LH. This suggested that the mutant also might be resistant to other pathogens that are controlled by salicylic acid-dependent SAR, such as the oomyceteous pathogen *Peronospora parasitica* (Bowling et al., 1994; Nawrath and Métraux, 1999). Two-week-old LH-grown *cpn1-1* plants had dramatically increased resistance to virulent *P. parasitica* (*Ahco 2*) compared with LH-grown Col-0 wild-type plants, as shown in Figure 7. Virulent *P. parasitica* was able to sporulate on LH-grown Col-0 wild-type plants, but virtually no sporulation was observed on LH-grown *cpn1-1* plants. This finding indicated that LH-grown *cpn1-1* had improved resistance to both virulent *P. syringae* and virulent *P. parasitica*. This supports the interpretation that LH-grown *cpn1-1* was undergoing SAR.

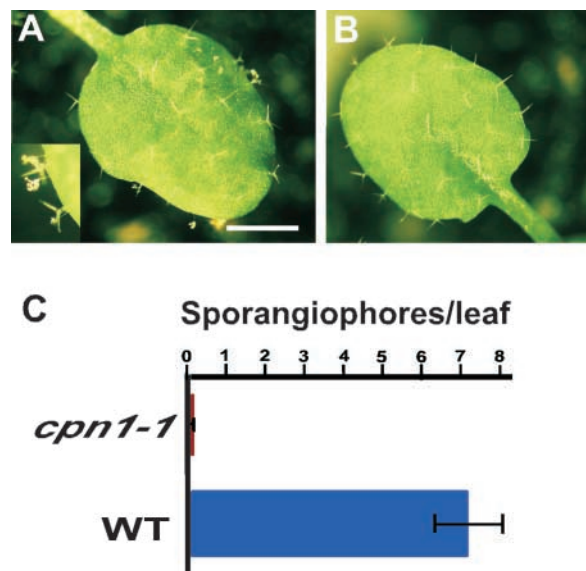
#### Genetic Analysis of *cpn1-1*

The *cpn1-1* mutation was isolated in the homozygous state. The original homozygous *cpn1-1/cpn1-1* mutant was outcrossed to Col-0 wild type (*CPN1/CPN1*). The *CPN1/cpn1-1* F1 plants were allowed to self-pollinate, and the segregation

of the *cpn1-1* phenotype was monitored in the F2 generation. The *cpn1-1* phenotype segregated in a mendelian fashion as a single, recessive locus, as indicated in Table 1. The *cpn1-1* mutant phenotype was assessed in the F2 generation based on plant morphology, lesion development, and resistance to virulent *P.s.t.* The mutant phenotypes associated with *cpn1-1* were observed to cosegregate in all 103 F2 *cpn1-1/cpn1-1* individuals examined. This result suggested that the *cpn1-1* mutant phenotype was most likely to be caused by a mutation in one gene or possibly by mutations in two or more very tightly linked genes.

#### Identification of the *CPN1* Gene

Genetic analysis indicated that the *cpn1-1* mutation likely was T-DNA tagged. The T-DNA contained the *bar* gene conferring resistance to the herbicide ammonium glufosinate as a dominant trait (D'Halluin et al., 1992; Weigel et al., 2000). Glufosinate resistance in the F2 generation of the *CPN1/CPN1* × *cpn1-1/cpn1-1* cross-segregated in a 3:1 ratio of herbicide-resistant to herbicide-sensitive plants in a group of more than 500 F2 individuals (data not shown). This finding indicated the presence of a single T-DNA integration locus



**Figure 7.** Resistance of *cpn1-1* to *P. parasitica*.

(A) LH-grown Col-0 wild-type leaf 7 days after inoculation with *P. parasitica*. Bar = 2 mm. Inset, close-up of sporangiophore at double magnification.

(B) LH-grown *cpn1-1* leaf 7 days after inoculation with *P. parasitica*. The scale is the same as in (A).

(C) Sporangiophore counts on LH-grown Col-0 wild-type (WT) and *cpn1-1* leaves 7 days after inoculation. Bars represent standard errors.



**Table 1.** Genetic Segregation Analysis of the *cpn1-1* Mutation

Female × Male	Generation	No. of Plants	Strong Mutant Phenotype	None	$\chi^2$ (P)
Col-0 × <i>cpn1-1/cpn1-1</i>	F1	4		4	
Col-0 × <i>cpn1-1/cpn1-1</i>	F2	394	103 <sup>a</sup>	291	0.27 (0.7)

<sup>a</sup>The lesion mimic, disease resistance, and leaf curling phenotypes segregated together as a recessive trait.

in the *cpn1-1* mutant line. To determine if the T-DNA was linked to the *cpn1-1* mutation, 88 of the 103 *cpn1-1/cpn1-1* F2 plants from the *CPN1/CPN1* × *cpn1-1/cpn1-1* cross were selected and tested for the presence of the T-DNA using a polymerase chain reaction (PCR) assay to amplify the *bar* gene. All 88 plants tested had the T-DNA, indicating that the T-DNA cosegregated with the *cpn1-1* mutant phenotype and that the *cpn1-1* mutation was potentially T-DNA tagged (data not shown).

DNA gel blot analysis was used to confirm that the *cpn1-1* mutant line had a single T-DNA insertion locus. The locus consisted of two copies of T-DNA in a “head-to-head” configuration, with the right T-DNA borders together and the left T-DNA borders flanking the plant DNA, as shown in Figure 8A. Plasmid rescue of the two left T-DNA borders was performed (Dilkes, 1998). The sequence of the plant DNA flanking the two left T-DNA borders was determined and used in a BLAST search of the Arabidopsis genome database. This revealed that the T-DNA was inserted into the seventh exon of a copine-like gene, *CPN1* (Figure 8A). *CPN1* is located at the bottom of chromosome 5 distal to *EIN2* and proximal to *HY5*. The Transformation-competent artificial chromosome (TAC) clone containing *CPN1* has GenBank accession number AB022212 (Liu et al., 1999).

The complete cDNA sequence of the *CPN1* gene was determined by 5′ and 3′ rapid amplification of cDNA ends (RACE) PCR. The 5′ and 3′ RACE PCR amplified single bands corresponding to the *CPN1* gene transcript (Figure 8B). This indicated that the *CPN1* gene produced a single transcript species. The 5′ and 3′ RACE PCR products were cloned and sequenced to obtain the full-length *CPN1* cDNA sequence (Figure 8C). The splicing of the *CPN1* cDNA was exactly as predicted by the annotation of the Arabidopsis genome database. The transcription start site was 82 bp upstream of the ATG, and the cDNA had a 192-bp 3′ untranslated region after the TAG stop codon. The *CPN1* gene was expressed in leaf tissue of both LH- and HH-grown Col-0 wild-type plants (Figure 8D). As expected, no complete *CPN1* transcript was detected in the *cpn1-1* mutant leaves (Figure 8D).

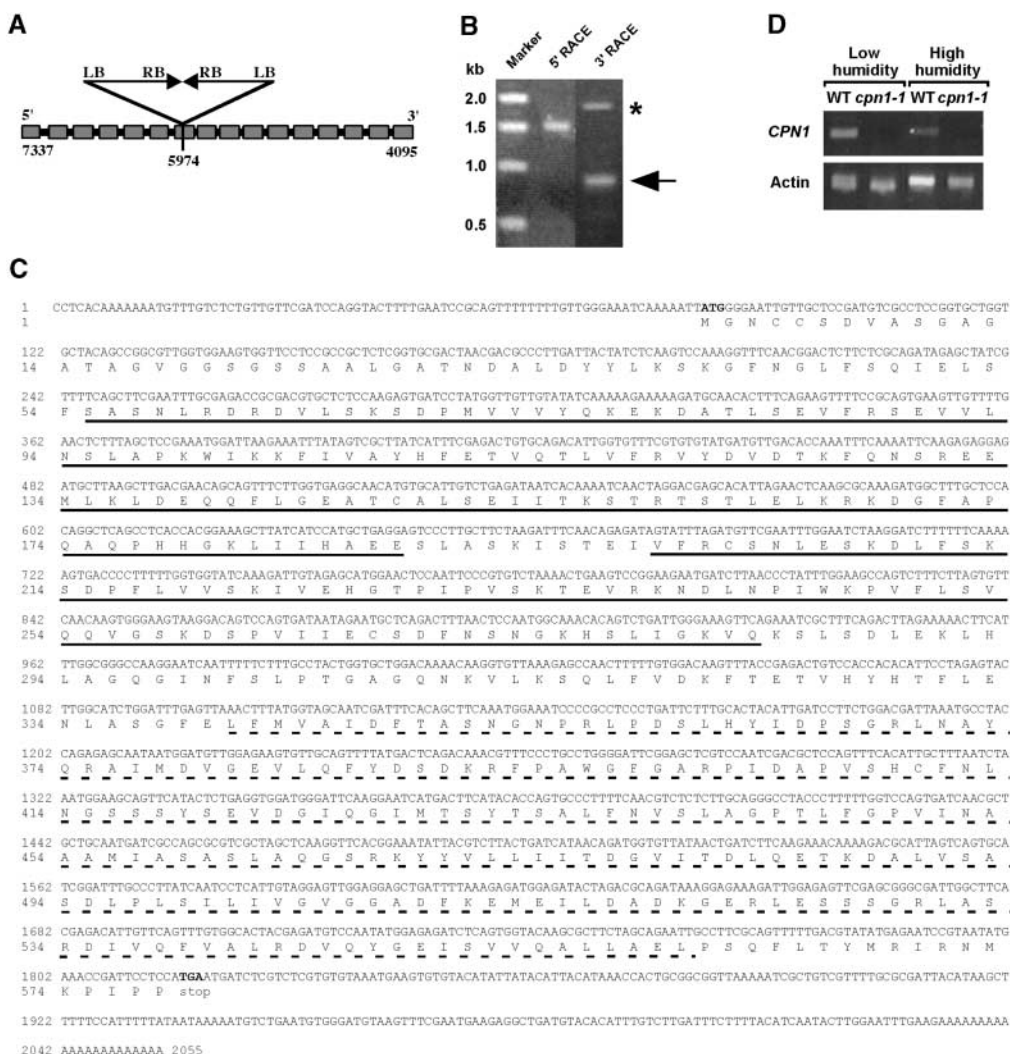
The predicted protein product of the *CPN1* gene is composed of 578 amino acids, has a molecular mass of 63.1 kD, and has homology with copine proteins (Creutz et al., 1998). The putative CPN1 protein, like other copines, has two C2 domains in the N-terminal portion of the protein and an A domain in the C-terminal region of the protein. In CPN1, the

two C2 domains are composed of amino acids 55 to 188 and 199 to 283; the A domain is composed of amino acids 341 to 560 (Figure 8C). The CPN1 protein has a high degree of homology with other copines in Arabidopsis and other species, as indicated in Table 2.

To prove that the disruption of the *CPN1* gene was responsible for the mutant phenotype in *cpn1-1* mutant plants, the *cpn1-1* mutant was complemented with a wild-type copy of the *CPN1* gene. An 8-kb genomic DNA fragment containing the *CPN1* gene coding regions plus 3.4 kb of upstream and 1.3 kb of downstream DNA was introduced into the genome of *cpn1-1/cpn1-1* mutant plants by Agrobacterium-mediated dip transformation (Clough and Bent, 1998). The 8-kb *CPN1* genomic fragment extended partly into the neighboring open reading frames on either side of *CPN1*. *cpn1-1/cpn1-1* plants transformed with the 8-kb *CPN1* genomic fragment had a wild-type phenotype, indicating that the T-DNA insertion into the *CPN1* gene is responsible for the mutant phenotype of *cpn1-1* (data not shown).

## DISCUSSION

The biological functions of genes defined by lesion mimic mutations are difficult to determine with certainty. It is likely that at least some lesion mimic mutants represent defects in genes that directly regulate the HR defense response. However, it is also likely that many lesion mimic mutants represent perturbations of plant cell physiology unrelated to disease defense responses. For example, physiological alterations caused by overexpression of genes such as pyruvate decarboxylase in potato and a bacterial proton pump in tobacco can trigger lesion formation and increased disease resistance, although the overexpressed genes do not appear to have any direct role in disease resistance responses (Mittler et al., 1995; Tadege et al., 1998). In addition, salicylic acid appears to potentiate cell death triggered by free radicals, implying that plants with increased salicylic acid might be more sensitive to the generation of free radicals by stresses (Mazel and Levine, 2001). The complexity of cell death signaling in plants makes it difficult to determine the specific function of the CPN1 protein. Nevertheless, speculation on the function of CPN1 can be made based on the results presented here and biochemical analyses of related copines, assuming that CPN1 has similar biochemical



**Figure 8.** Molecular Identification of the *CPN1* Gene.

**(A)** Scheme of the *CPN1* gene with the 16 exons depicted as boxes. In *cpn1-1*, two copies of the T-DNA are inserted in a head-to-head configuration into exon 7 of *CPN1*. The numbers indicate the base pair positions in Transformation-competent artificial chromosome (TAC) clone K22G18 for the start and stop codons of *CPN1* and the exact site of the T-DNA insertion in *cpn1-1*. RB, right T-DNA border; LB, left T-DNA border. **(B)** 5' and 3' RACE PCR products resolved on a 1% agarose gel and visualized with ethidium bromide. The 5' RACE reaction produced a single 1.5-kb band corresponding to the 5' end of the *CPN1* gene transcript. The 3' RACE reaction produced two bands. The 750-bp band, denoted by the arrow, represents the 3' end of the *CPN1* gene transcript. The 1.8-kb band, denoted by the asterisk, was determined by DNA sequencing to represent a portion of one of the other copine homologs in the Arabidopsis genome (accession number AL163912). **(C)** Complete *CPN1* cDNA sequence and predicted amino acid translation. The C2 domains are solid underlined. The A domain is dashed underlined. **(D)** Reverse transcriptase-mediated PCR analysis of *CPN1* transcript accumulation in LH- and HH-grown Col-0 wild-type (WT) and *cpn1-1* leaf tissue. *CPN1*, a 467-bp portion of the *CPN1* transcript; Actin, a 900-bp portion of an actin gene.

characteristics. One possibility is that CPN1 plays a direct role in defense cell death signaling and has indirect effects on LH adaptation. Alternatively, CPN1 may have a direct function in LH adaptation and may have indirect effects on plant defense signaling.

### CPN1 as a Regulator of Defense and Cell Death

The improved disease resistance and accelerated HR phenotypes of *cpn1-1* imply that CPN1 may play a direct role in regulating defense responses, including the HR. Because

the *cpn1-1* mutation is recessive and probably eliminates *CPN1* gene function, CPN1 could function as a repressor of cell death. An interesting possibility is that a pathogen-triggered  $\text{Ca}^{2+}$  flux may activate the repressive function of CPN1, possibly by triggering the localization of CPN1 to a membrane where it could exert a repressive influence on cell death signal transduction. This might seem counterintuitive at first, but genetic evidence suggests that the HR is under negative control as well as positive control to prevent uncontrolled cell death and lesion expansion after HR initiation (Dietrich et al., 1994). The accelerated HR in *cpn1-1* plants supports a role for CPN1 as a negative regulator of the avirulent pathogen-induced HR defense response.

However, our data indicate that CPN1 also is required for the repression of cell death and *PR* gene expression in the absence of a pathogen infection. In particular, CPN1 is essential for the repression of cell death in LH growth conditions. It is possible that some aspect of LH signaling or LH adaptation, such as a  $\text{Ca}^{2+}$  flux, has the potential to trigger the HR. *CPN1* may have a genetically nonredundant role in preventing LH-potentiated cell death from occurring. In plants, increases in intracellular calcium have been observed in response to a wide range of biotic and abiotic stimuli, including osmotic stress, cold, touch, and drought (for reviews, see Knight, 2000; Reddy, 2001). Although the calcium signatures differ for each stimulus, the calcium molecule appears to be a key shared component of signal transduction pathways controlling responses to various environmental stimuli (for reviews, see McAinsh and Hetherington, 1998; Bowler and Fluhr, 2000). CPN1 may play a role in defining  $\text{Ca}^{2+}$  signaling specificity by preventing cell death and other defense responses from being triggered by  $\text{Ca}^{2+}$  fluxes that are not induced by pathogens. According to this model, cell death occurs in the *cpn1-1* mutant in LH because there is no CPN1 protein present to prevent LH signaling events from triggering cell death.

*CPN1* also may play a direct role in suppressing *PR* gene expression in the absence of pathogen attack. Low levels of *PR* gene expression were observed in HH-grown *cpn1-1*, indicating a partial derepression or activation of *PR* gene expression in *cpn1-1* mutants even in the absence of cell death and SAR. Thus, *PR* gene expression in *cpn1-1* may occur by different mechanisms in LH and HH: in LH-grown *cpn1-1*, cell death occurs, triggering SAR and high-level *PR* gene expression; in HH-grown *cpn1-1*, low-level *PR* gene expression occurs because the repressive influence of *CPN1* on *PR* gene expression is absent.

### CPN1 as a Regulator of LH Adaptation

A low air moisture level coupled with adequate soil water potential results in high rates of water uptake in the roots and transpiration in the leaves (Mohr and Schopfer, 1995). Under these conditions, the plant must regulate cellular ion balance by reducing transpiration, controlling ion uptake,

**Table 2.** Homology of CPN1 with Other Copines

Source	Protein Accession Number	Amino Acid Identity (%)
Arabidopsis	CAB87919	80.7
	AAB70417	53.2
	CAB87781	34.7
Human	AAD46074	41.4
	AAC15920	40.4
Mouse	NP_034077	38.1
<i>C. elegans</i>	AAC71103	29.8
<i>Paramecium</i>	AAC15919	31.4

extruding ions, and sequestering ions in subcellular compartments such as the vacuole. In *cpn1-1*, the humidity-sensitive phenotype is unlikely to be attributable simply to increased or uncontrolled water loss, because stomatal conductance in *cpn1-1* was normal compared with that in Col-0 wild-type plants. Instead, CPN1 may be a key player in LH adaptation.

It is possible that *cpn1-1* is defective in controlling ion balance in LH or is defective in responding to altered ion balance in LH. As a  $\text{Ca}^{2+}$ -dependent phospholipid binding protein, CPN1 could regulate ion transport at a membrane in response to  $\text{Ca}^{2+}$  fluxes, for instance. Defective ionic homeostasis in *cpn1-1* could trigger cell death by a number of mechanisms in LH, including hyperaccumulation of essential metals or oxidative stress (Wang et al., 1996; Briat and Lebrun, 1999; Hagemeyer, 1999). For example, manganese toxicity causes leaf crinkle and lesion formation in cotton, possibly as a result of increased production of reactive oxygen species (Foy et al., 1978). Ionic imbalances in LH-grown *cpn1-1* could trigger cell death and lesion formation, which in turn could trigger SAR. Thus, the increased disease resistance phenotype of *cpn1-1* might be an indirect effect of the mutation. It is interesting, however, that HH-grown *cpn1-1* plants also exhibited an accelerated HR phenotype and very low-level expression of *PR* genes. This suggests that CPN1 might be necessary to control cellular homeostasis even in HH conditions, although the defect of the *cpn1-1* mutant is much more pronounced in LH.

### Conclusion

Plants must integrate signals from a wide range of biotic and abiotic sources to optimize their developmental pattern and physiology for their environment. The growing interconnectedness of plant signaling pathways reflects the integrated nature of plant responses to the environment. The presence of common regulatory components during signal transduction responses to abiotic and biotic stresses suggests that cellular signaling is a web, with proteins receiving

inputs from multiple partners and pathways (Bent, 2001). Slight perturbations in a particular signaling pathway could easily deregulate other signaling events in the plant as a result of interconnections between them. The phenomenon of cross-tolerance may reflect this kind of signaling interconnection (for review, see Bowler and Fluhr, 2000). Our results suggest that the *CPN1* gene may represent an important component of the plant's ability to monitor and respond to its environment. The CPN1 protein may act as a  $\text{Ca}^{2+}$ -sensitive modulator of plant responses to environmental signals, including humidity and pathogens.

## METHODS

### Plant Growth Conditions and Mutant Screening

*Arabidopsis thaliana* plants were grown in soil (Scotts Redi-Earth Plug and Seedling Mix [E.C. Geiger, Inc., Harleysville, PA]) and watered with distilled water. For low-humidity (LH) growth, plants were maintained at 24°C under either short-day (SD) or long-day (LD) conditions with 75 to 100  $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{sec}^{-1}$  light intensity and 35 to 45% relative humidity. For high-humidity (HH) growth, plants were maintained at 24°C under either SD or LD conditions with 55 to 65  $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{sec}^{-1}$  light intensity and 75 to 85% RH.

Pools of activation-tagged *Arabidopsis* seed (Columbia [Col-0] ecotype) (Weigel et al., 2000) were obtained from the *Arabidopsis* Biological Resource Center (Ohio State University, Columbus). We screened this population for mutants with improved resistance to the virulent *Pseudomonas syringae* pv *tomato* (*P.s.t.*) strain DC3000 (Cuppels, 1986). The *cpn1-1* mutant was identified from among ~10,000 independent T-DNA lines screened for mutants with increased resistance to *P.s.t.* For mutant screening, 4-week-old plants grown under LH, SD conditions were screened for increased resistance to virulent *P.s.t.* by dipping the plants into a bacterial suspension of  $5 \times 10^8$  colony-forming units [cfu]/mL in 10 mM  $\text{MgCl}_2$  with 0.00025% (v/v) Silwet L-77 surfactant (Lehle Seeds, Round Rock, TX). After inoculation, plants were maintained at 24°C under SD conditions with 55 to 65  $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{sec}^{-1}$  light intensity and 75 to 95% RH for the development of bacterial speck disease symptoms. Bacterial speck disease symptoms were assessed 4 days after inoculation.

### Bacterial Strains and Growth Conditions

For mutant screening, virulent *P.s.t.* strain DC3000 bacteria were grown on Pseudomonas Agar F (PA) (Sigma) supplemented with 100  $\mu\text{g}/\text{mL}$  rifampicin (Sigma) at 28°C. For growth curves, *P.s.t.* strain DC3000 bacteria were used carrying either the plasmid pVSP61 (empty vector) or pV288 (bearing *avrRpt2*) and were grown on PA supplemented with 100  $\mu\text{g}/\text{mL}$  rifampicin and 25  $\mu\text{g}/\text{mL}$  kanamycin (Sigma) (Whalen et al., 1991). For hypersensitive response (HR) assays, *P. syringae* pv *maculicola* (*P.s.m.*) (strain 4326) bacteria were used bearing either plasmid pVSP61 or pV288 and were grown on PA supplemented with 100  $\mu\text{g}/\text{mL}$  rifampicin and 25  $\mu\text{g}/\text{mL}$  kanamycin. The *P.s.m.* (4326) strain was used for the HR assays because it is less virulent on *Arabidopsis* than *P.s.t.* and allows for a clearer distinction between the HR and disease symptoms.

### Bacterial Growth Curves and *Peronospora parasitica* Inoculations

In planta bacterial growth assays were performed by inoculating 5-week-old plants by either infiltration or dipping. For infiltration inoculations, leaves were inoculated with a suspension of  $10^5$  cfu/mL bacteria in 10 mM  $\text{MgCl}_2$ ; for dip inoculations, plants were inoculated with a suspension of  $2 \times 10^7$  cfu/mL bacteria in 10 mM  $\text{MgCl}_2$  plus 0.00025% (v/v) Silwet L-77. Plants were maintained in SD, 55 to 65  $\mu\text{mol}\cdot\text{m}^{-2}\cdot\text{sec}^{-1}$  light intensity, 75 to 90% RH conditions for the duration of the experiment after inoculation. Bacterial populations were assessed at 0, 1, 2, and 4 days after inoculation. Three samples were taken for each data point. For each sample, three whole leaves were harvested, weighed, and ground in 1 mL of 10 mM  $\text{MgCl}_2$ . Bacteria per unit tissue weight was determined by serial dilution plating essentially as described by Whalen et al. (1991), except that bacteria were plated on PA medium supplemented with 100  $\mu\text{g}/\text{mL}$  rifampicin and 25  $\mu\text{g}/\text{mL}$  kanamycin. Each growth curve experiment was replicated three times. Statistical analyses were performed using Student's *t* test of the differences between means of log-transformed data.

Resistance to *P. parasitica* (*Ahco 2*) was tested in 18-day-old plants. The infected plants were scored after 7 days. More than 100 leaves from 20 plants were chosen at random, and the number of conidiophores per leaf was counted. Statistical significance was measured by Student's *t* test.

### Histochemistry and Microscopy

Leaf samples were taken from 3-week-old plants and stained for dead cells using trypan blue (125  $\mu\text{g}/\text{mL}$ ) (Vogel and Somerville, 2000). Samples for autofluorescence and callose examination were prepared as described by Dietrich et al. (1994). For scanning electron microscopy, leaves were fixed overnight at 4°C in 2.5% glutaraldehyde in 0.1 M cacodylate buffer, pH 7.4, rinsed in 0.1 M cacodylate buffer, pH 7.4, and then postfixed with  $\text{OsO}_4$ . Samples were dehydrated by a series of increasing concentrations of ethanol, and critical point drying was performed using  $\text{CO}_2$ . Samples were then sputter coated with a gold-palladium mixture and viewed in a scanning electron microscope at a 15° angle.

### HR Tests

Plants grown under LH and HH conditions were inoculated with the *P.s.m.* strains described above at a concentration of  $10^8$  cfu/mL in 10 mM  $\text{MgCl}_2$ . Fifty to 100 leaves were infiltrated for each plant-bacterial combination per experiment. Infiltrated leaves were scored for HR development as 0 for no HR, 1 for partial HR, and 2 for complete HR at 4, 8, 12, 16, 20, 24, and 30 hr after inoculation. Statistical analysis was performed using Student's *t* test. This experiment was repeated three times with similar results.

### Electrolyte Leakage Assay

Electrolyte leakage was measured essentially as described by Goodman (1968). LH- and HH-grown *cpn1-1* and Col-0 wild-type plants were inoculated with *P.s.m.* strains as described for the HR tests, and control plants of both genotypes were inoculated with 10 mM  $\text{MgCl}_2$ . Measurements were performed at 0, 4, 8, 12, 16, 20, 24, and 30 hr after inoculation. For each sample at each time point, five 0.07-cm<sup>2</sup> leaf

discs were excised from infiltrated leaves using a number 1 cork borer and immersed in 3 mL of distilled water and agitated gently in a shaker for 8 min at 28°C. The conductivity of the solution was measured using a Traceable Conductivity Resistivity TDS Salinity Concentration meter (Control Company, Friendswood, TX). Three independent samples were taken at each time point for each plant-inoculum combination.

#### Stomatal Conductance and Drought Tolerance Measurements

Leaves of 30-day-old LH-grown *cpn1-1* and wild-type plants were sealed in the gas exchange cuvette of a Li-Cor 6400 photosynthesis meter. The following parameters were maintained during the measurements: leaf temperature,  $24 \pm 0.5^\circ\text{C}$ ;  $\text{CO}_2$  concentration,  $350 \mu\text{L L}^{-1}$ ; air flow,  $0.1 \text{ L min}^{-1}$ ; white light illumination,  $400 \mu\text{mol m}^{-2} \text{ sec}^{-1}$ ; RH, 34 to 35%. The plants had been exposed to normal growth light for 3 to 4 hr before the beginning of the experiment. One leaf was measured at a time. Measurements were taken 5 min after leaves were sealed in the cuvette under the conditions described above, at which time the leaves had attained a steady state. The *cpn1-1* leaves were too small to fill the leaf chamber completely. Therefore, the area of each *cpn1-1* leaf used in the experiment was determined to obtain the conductance per unit area. Leaf area was determined by excising the leaf and measuring its area using NIH Image 1.62f software. The data represent the average stomatal conductance for 15 randomly selected leaves of each genotype. To test plant responses to low soil water potential, the soil was simply allowed to dry out, and the time at which 4-week-old LH- and HH-grown *cpn1-1* and Col-0 wild-type plants began to show a visible decline in turgor pressure was observed.

#### RNA Analyses

Leaf tissue samples were collected from 4-week-old plants grown under LH, SD or HH, SD conditions. Samples were flash frozen in liquid nitrogen, and RNA was extracted using the RNeasy Maxi kit (Qiagen, Valencia, CA). Twenty micrograms of total RNA was separated in a formaldehyde-agarose gel (Ambion Northern Max kit) and transferred to a Bright Star Plus membrane (Ambion, Austin, TX). After RNA transfer, the membranes were stained with methylene blue to verify RNA transfer and compare the relative levels of rRNA in each lane. Plasmids for the *PR1*, *PR2*, and *PR5* probes were obtained from Dr. Ramesh Raina (Pennsylvania State University). Probe labeling with  $^{32}\text{P}$ , hybridization, blot washing, and exposure were performed as described previously (McNellis et al., 1998).

#### Genetic Analyses

Backcrosses were performed with Col-0 wild type using *cpn1-1* mutant as both pollen donor and receiver. F1 and F2 generation plants were tested for the *cpn1-1* mutant phenotype by growing them in LH, SD conditions and inoculating with virulent *P. syringae*. F2 plants with lesions and increased resistance to *P. syringae* were tested for the presence of the T-DNA insert by polymerase chain reaction (PCR) analysis. The primers used were designed to amplify the *bar* gene (BARF, 5'-CTCTAGGATCGATCCCCCGGG-3'; BARR, 5'-GAA-TTCCTCGAGTATAAGAGC-3'; annealing at 60°C) and produce a 650-bp product.

#### DNA Gel Blotting and Plasmid Rescue

Genomic DNA was isolated using a modification of the method of Shure et al. (1983) as described previously (McNellis et al., 1998). DNA gel blot analysis was performed as described previously (McNellis et al., 1998). DNA gel blot analyses of genomic DNA of the *cpn1-1* mutant were performed using enzymes that cut the pSKI015 T-DNA once, including BamHI, SpeI, EcoRI, and HindIII (New England Biolabs, Beverly, MA). The pBluescript KS+ (Stratagene) plasmid was used as a probe for the pSKI015 T-DNA. Probe labeling and detection were performed using the Phototope-Star Chemiluminescent Detection Kit (New England Biolabs) according to the manufacturer's instructions. DNA gel blotting of *cpn1-1* DNA cut with BamHI revealed 8- and 10-kb fragments hybridizing to pBluescript. Plasmid rescue was done by digestion of 5  $\mu\text{g}$  of *cpn1-1* genomic DNA with 40 units of BamHI, self-ligation using T4 DNA polymerase (New England Biolabs), and transformation into Epicurian Coli XL-2 Blue *Escherichia coli* cells (Stratagene) according to the manufacturer's instructions, except that Luria-Bertani medium supplemented with 100  $\mu\text{g}/\text{mL}$  ampicillin (Sigma) was used to select transformants. Both the 8- and 10-kb fragments were rescued as plasmids. A partial sequence of the plant DNA flanking the T-DNA left borders in the rescued clones was obtained by cycle sequencing at the Nucleic Acid Facility at Pennsylvania State University using the T7 primer. The DNA sequence flanked by the T-DNA insert was identified by BLAST search. The exact location of the T-DNA insert was determined by sequencing of the rescued plasmids using the primers 5'-GGA-ACCTCAATCCCGTGTGC-3' (forward) and 5'-ACCTAAAATGAC-CATAAATGG-3' (reverse).

#### Reverse Transcriptase-Mediated PCR

First-strand cDNA synthesis was performed using 5  $\mu\text{g}$  of total RNA and the RETROscript kit (Ambion). As a positive control, PCR was performed to amplify the cDNA of a constitutively active actin gene using primers 5'-GTTGGGATGAACCAGAAGGA-3' (forward) and 5'-GAACCACCGATCCAGACACT-3' (reverse) (annealing at 58°C). cDNA synthesized from the total RNA of *cpn1* and Col-0 wild-type plants was tested for copine expression by PCR using the primers 5'-TCT-AGATCATGGAGGAATCGGTTTCAT-3' (forward) and 5'-ATCATGACT-TCATACACCGAGTGC-3' (reverse) (annealing at 58°C), which produce a 467-bp product.

#### Genetic Complementation of *cpn1-1*

An 8-kb BamHI fragment from Transformation-competent artificial chromosome (TAC) clone K22G18 (Liu et al., 1999) consisting of the copine gene coding regions plus upstream and downstream sequences was cloned into the BamHI site of the pCLD04541 binary vector to create pCLD04541-*CPN1* (Bent et al., 1994). pCLD04541-*CPN1* was introduced into *Agrobacterium tumefaciens* strain GV3101 by heat shock transformation, and transformants were selected on Luria-Bertani medium supplemented with 50  $\mu\text{g}/\text{mL}$  gentamycin, 3  $\mu\text{g}/\text{mL}$  tetracycline, and 25  $\mu\text{g}/\text{mL}$  kanamycin (Sigma). *cpn1-1* mutant plants were transformed with pCLD04541-*CPN1* using the floral dip method (Clough and Bent, 1998). Transformants were selected on GM medium (1  $\times$  Murashige and Skoog [1962] salts, pH 5.7 [Gibco BRL], 1  $\times$  B5 vitamins, solidified with 0.8% phytagar [Gibco BRL]) supplemented with 50  $\mu\text{g}/\text{mL}$  kanamycin. Transformants were

grown in LH, SD conditions and observed to have a wild-type phenotype.

### CPN1 Sequence Analysis

The complete sequence of the *CPN1* cDNA, including the 5' and 3' untranslated regions, was determined by performing 5' and 3' rapid amplification of cDNA ends (RACE) reactions using the SMART RACE cDNA kit (Clontech, Palo Alto, CA) and cloning the resulting PCR products using the Original TA Cloning Kit (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions. The 5' RACE gene-specific primer was 5'-GCGACGCGCTGGCGATCATTGCAG-CAGCG-3'. The 3' RACE gene-specific primer was 5'-TCCCTG-CCTGGGATTGGAGCTCGTCC-3'. The domains of the CPN1 protein were identified using Pfam and Prosite Profile Scan databases ([http://hits.isb-sib.ch/cgi-bin/hits\\_motifscan](http://hits.isb-sib.ch/cgi-bin/hits_motifscan)). Pairwise alignment of copine sequences was performed using the SIM Alignment Tool ([www.expasy.ch/tools/sim-prot.html](http://www.expasy.ch/tools/sim-prot.html)).

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# A Humidity-Sensitive Arabidopsis Copine Mutant Exhibits Precocious Cell Death and Increased Disease Resistance

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