

A circadian oscillator in the fungus *Botrytis cinerea* regulates virulence when infecting *Arabidopsis thaliana*

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The circadian clock of the plant model *Arabidopsis thaliana* modulates defense mechanisms impacting plant–pathogen interactions. Nevertheless, the effect of clock regulation on pathogenic traits has not been explored in detail. Moreover, molecular description of clocks in pathogenic fungi—or fungi in general other than the model ascomycete *Neurospora crassa*—has been neglected, leaving this type of question largely unaddressed. We sought to characterize, therefore, the circadian system of the plant pathogen *Botrytis cinerea* to assess if such oscillatory machinery can modulate its virulence potential. Herein, we show the existence of a functional clock in *B. cinerea*, which shares similar components and circuitry with the *Neurospora* circadian system, although we found that its core negative clock element FREQUENCY (BcFRQ1) serves additional roles, suggesting extracircadian functions for this protein. We observe that the lesions produced by this necrotrophic fungus on *Arabidopsis* leaves are smaller when the interaction between these two organisms occurs at dawn. Remarkably, this effect does not depend solely on the plant clock, but instead largely relies on the pathogen circadian system. Genetic disruption of the *B. cinerea* oscillator by mutation, overexpression of BcFRQ1, or by suppression of its rhythmicity by constant light, abrogates circadian regulation of fungal virulence. By conducting experiments with out-of-phase light:dark cycles, we confirm that indeed, it is the fungal clock that plays the main role in defining the outcome of the *Arabidopsis*–*Botrytis* interaction, providing to our knowledge the first evidence of a microbial clock modulating pathogenic traits at specific times of the day.

circadian clock | virulence | plant–pathogen interaction | *Arabidopsis thaliana* | *Botrytis cinerea*

Circadian clocks are autonomous oscillators based on transcription–translation negative feedback loops (TTFLs), which temporally coordinate a series of processes, from gene expression to metabolism (1, 2). From an evolutionary perspective, these biological clocks have independently emerged at least three times throughout evolution and its molecular description is well documented in photosynthetic organisms like cyanobacteria, algae, and plants, as well as in insects, mammals, birds, and fungi (2). Recent evidence indicates that these machineries provide organisms with adaptive advantages (3–7), whereas the relevance of circadian regulation on the control of immune and defense responses in mammals (8, 9), flies (10, 11), and plants (12–16) has also been documented. Therefore, the time of the day at which a host–pathogen interaction first occurs has measurable consequences in the overall development of the pathogenesis process, at least in all cases where this paradigm has been assessed.

The idea that a circadian clock could modulate the outcome of a plant–pathogen interaction was postulated in recent years (17), and since then, its importance has been experimentally addressed in the plant (18). Nevertheless, the focus has been mostly centered on the circadian system of the host, partially due to the lack of molecular description of oscillatory machineries

in the pathogens under study. Thus, no molecular evidence of a circadian clock modulating a pathogen's behavior has been reported, with the exception of phenotypic evidence observed for insects (16) and malaria parasites (19).

The nonpathogenic fungus *Neurospora crassa* has been a premier model for chronobiology studies (1, 20, 21) but surprisingly, even after 25 y of the characterization of the central component of the *N. crassa* clock—the protein FREQUENCY (FRQ) (22)—no other fungal circadian system has been molecularly described. Thus, the work in *Neurospora* has unveiled that in its core oscillator the transcription factor (TF)/photoreceptor White Collar-1 (WC-1) associates with its partner White Collar-2 (WC-2) forming the White Collar Complex (WCC) to activate the expression of the *frequency* (*frq*) gene. FRQ is produced, associates with the FRQ-interacting RNA helicase (FRH), and acts as the negative element of the TTFL shutting down its own expression by inhibiting the WCC (21). As FRQ is progressively phosphorylated, it decreases its negative effect on WCC, and it is finally degraded around the time that a new cycle of *frq* expression is initiated (23). Thus, this process can be visualized as daily oscillations in *frq* mRNA and protein levels.

To unravel the role of a circadian clock on pathogenic traits, herein we characterize and describe the circadian oscillator of the plant pathogen *Botrytis cinerea*. This necrotrophic ascomycete is responsible for the gray mold disease on more than 200 different dicotyledonous plants (24) and is considered the second most important phytopathogenic fungus based on its scientific and economic importance (25). By generating arrhythmic

Significance

Circadian clocks are molecular machineries that allow organisms to anticipate daily cyclic challenges and to temporally modulate different processes. Thus, plant defense mechanisms against pathogens have been reported to vary daily in *Arabidopsis thaliana*. Although the plant–pathogen interaction is a two-sided story, nothing is known regarding circadian regulation of pathogenic traits. Herein we characterize a functional circadian clock in the necrotrophic fungal plant pathogen *Botrytis cinerea*, postulating additional roles for BcFRQ1, the *Botrytis* ortholog of the core clock component FREQUENCY of *Neurospora crassa*. By using different plant and *Botrytis* clock-null mutants, we demonstrate that the interaction between this pathogen and its host varies with the time of day, being the *B. cinerea* circadian clock key in regulating this outcome.

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of the leaves surface, respectively). Nevertheless, from these results we cannot rule out if the differences are explained by a cyclical behavior—driven by the environment—in one of the organisms or in both, or if instead it is being endogenously (circadianly) controlled. Consequently, we proceeded to define the contribution of each circadian system in this time-of-day effect on the pathogenic outcome.

An Endogenous Circadian Clock Controls the Infection Process of *B. cinerea* on *A. thaliana*. To confirm that the observed effects were not just driven responses to environment cycles (LD), additional analyses were conducted under constant (circadian) conditions. First, to assess the role of the plant circadian clock we used both WT and arrhythmic *A. thaliana* ecotypes [Columbia 0 (Col-0) and constitutive expression of the *CIRCADIAN CLOCK ASSOCIATED 1* (CCA1ox), respectively], keeping them for 72 h under DD after their inoculation at dawn or dusk. The results show that, under FRCs, significant differences in the percentage of the leaves infected areas were once again observed when B05.10 WT *B. cinerea* infected Col-0 plants, with the smaller lesions being produced when inoculation occurred at dawn (14.8% and 19.6% at dawn and dusk, respectively) (Fig. 3A and Fig. S9). Remarkably, when CCA1ox arrhythmic plants were challenged with the B05.10 strain, significant differences were still detected, with a bigger percentage of lesion area at dusk (20.7% and 26.8% at dawn and dusk, respectively; Fig. 3A and Fig. S9). Importantly, the basal level of infection was higher in the CCA1ox genetic background, indicative of an increased susceptibility. The fact that there is a time-of-day difference even when an *A. thaliana* clock-deficient mutant is used, strongly suggests that the fungal clock is playing a key role in the outcome of this plant–pathogen interaction. In agreement with this observation, the arrhythmic $\Delta bcwcl1$ strain (Fig. 1F) (28) is unable to produce a statistically significant different lesion area when inoculated at dusk compared with dawn, in the Col-0 plant (Fig. 3A and Fig. S9). To confirm this result, arrhythmic CCA1ox plants were challenged with the $\Delta bcwcl1$ mutant, revealing the same behavior (Fig. 3A and Fig. S9).

To exclude the possibility that the loss of temporal variation seen for the infections with $\Delta bcwcl1$ in DD resulted from defects in the blue light signaling pathway (28) rather than by the absence of a functional circadian clock, the arrhythmic OE::bcfrq1 mutant (Fig. S4) was used. Notably, the OE::bcfrq1 strain also lacks the ability to achieve a better infection at night when infecting Col-0 or CCA1ox plants (Fig. 3B and Fig. S9). Surprisingly, this strain not only has lost the circadian modulation of virulence, but moreover it displays smaller lesions and shows the inability to form secondary spreading lesions (Fig. S9, third row).

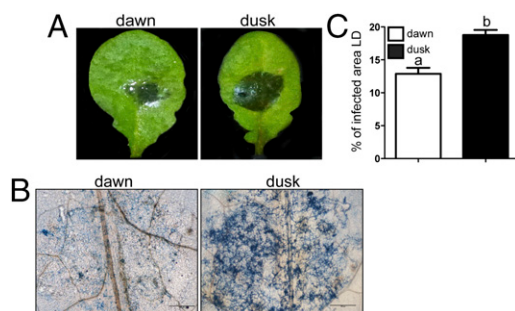


Fig. 2. The outcome of the *B. cinerea*–*A. thaliana* interaction differs with the time of day. (A) Lesion spreading of B05.10 strain on *A. thaliana* Col-0 plants. *Arabidopsis* leaves were inoculated at dawn or dusk, and then plants were incubated in LD for 72 h. Representative pictures are shown. (B) Trypan blue staining showing fungal growth after 3 dpi. (Scale bars, 500 μ m.) (C) Quantification of lesion spreading of three independent virulence assays. Bars represent mean \pm SEM (different letters indicate statistical differences).

Finally, using the macroconidia from $\Delta bcfrq1$ we further confirmed the influence of the fungal clock in the plant–pathogen circadian phenotype under study. Consistent with the previous observations, infection with this clock-deficient strain resulted in no significant difference in the percentage of infected leaf area when inoculation was performed at dawn or at dusk (Fig. 3C and Fig. S9). In aggregate, these results (Fig. 3) indicate that the outcome of the *B. cinerea*–*A. thaliana* interaction is strongly influenced by the fungal circadian clock leading the pathogen to achieve differential infection levels at night versus at day.

Light:Dark Cycles Confirm the Importance of the *B. cinerea* Circadian Clock in Regulating Virulence Under Environmentally Relevant Conditions. To obtain further data defining the importance of fungal versus plant clocks in mediating the described interaction, additional infection assays were performed under LD cycling conditions. Fig. 4A and B depict that, whereas the *B. cinerea* WT strain presents differences in its infection capability between inoculations at dawn and dusk on both Col-0 and CCA1ox plants, neither of the *B. cinerea*'s clock mutants (OE::bcfrq1 or $\Delta bcfrq1$) displayed time-differential infection capabilities. This finding reflects that the differences in virulence observed between dawn and dusk for the *B. cinerea* WT strain under both DD and LD conditions are indeed due to the control of its circadian clock. Thus, these results are consistent with a scenario in which the fungal clock, rather than an environmentally driven behavior or the plant clock, is largely responsible for the outcome of the interaction.

Under Constant Light the *A. thaliana* Clock Acquires a Dominant Role. Additional assays were performed under constant light (LL) because this setting allows the plant clock to “free run” (31), whereas the *B. cinerea* clock (as the one in *Neurospora*) fails to do so (Fig. 1G). Thus, inoculation of Col-0 plants at dusk with the B05.10 strain showed larger lesions, but notably, no significant differences were observed between dawn and dusk when the CCA1ox plants were challenged with the same fungal strain (Fig. 4C), demonstrating that under LL conditions—and in contrast to what is seen in LD and DD—the *A. thaliana* circadian clock acquires a preponderant role.

Additional Plant Arrhythmic Mutants and Inverted-Cycle Infection Further Confirm the Relevance of the Fungal Clock Under FRCs. To provide additional lines of evidence supporting the importance of the fungal versus the plant clock under FRCs (DD), we tested the behavior of WT *B. cinerea* on the triple mutant PSEUDO-RESPONSE REGULATORS *prp9 prp7 prp5* (*d975*) arrhythmic *Arabidopsis* mutant (32). Once again, we observed that despite the absence of a functional clock in the plant, the WT *B. cinerea* strain is capable of producing differential lesions with larger ones at night, and that moreover, this arrhythmic triple *Arabidopsis* mutant is—as also observed for CCA1ox—more sensitive to this pathogen (Fig. 4D).

Finally, we proceeded to work with inverted cycles to determine in a different manner the relevance of both clocks. Thus, the fungus was entrained in an inverted light:dark cycle (12 h out of phase) to the one under which the plants were grown. Then we proceeded to infect the plants at dawn or dusk with *B. cinerea* that had the same internal time or that had the opposite time of the plants, maintaining them thereafter in FRCs for 72 h. Plants inoculated with the fungus that had been under the same LD entrainment recapitulated the type of results already shown (Fig. 3). Notably, in the inverted cycle experiment, we observed that plants challenged at dawn showed larger lesions as the fungus that was inoculated on them had dusk time, whereas plants inoculated at dusk with fungus that had dawn time displayed smaller lesions (Fig. 4E). In aggregate, these experiments provide additional and compelling evidence supporting the importance of the *B. cinerea* clock in controlling its pathogenic potential.

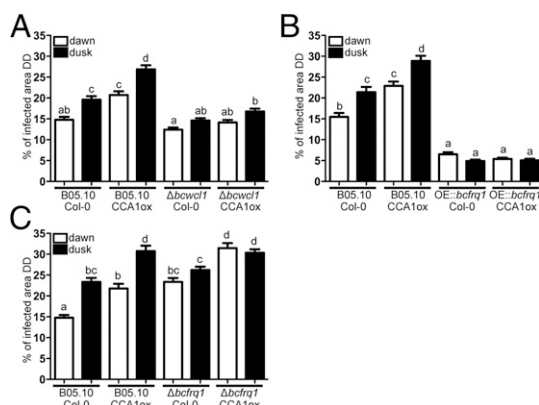


Fig. 3. A fungal circadian clock controls the infection process of *B. cinerea* on *A. thaliana*. (A) Measurements of lesion spreading of B05.10 and $\Delta bcwcl1$ strains. (B) Comparison of lesion spreading between B05.10 and OE::*bcfrq1* strains. (C) Lesion spreading of B05.10 and $\Delta bcfrq1$ strains. Conidia were inoculated at dawn or dusk on *A. thaliana* Col-0 and CCA1ox plants, as indicated. Lesions were quantified 3 dpi under DD. Bars represent mean \pm SEM (different letters indicate statistical differences).

Discussion

In *B. cinerea* B05.10, rhythms in macroconidiation can be observed under light:dark cycles, but not under constant darkness (28). This result differs from what is known in *Neurospora*, where conidiation rhythms can also be detected in the latter condition, but only in strains containing a mutant allele of the *ras-1* gene (*ras-1^{bd}*), which leads to enhanced circadian output (33). The absence of overt rhythms in *B. cinerea* could be due to a similar reason, to the divergence of output pathways, or to the absence of a functional clock. Therefore, motivated by the interest in understanding the impact of circadian regulation on fungal virulence, we were able to establish the existence of an endogenous circadian clock in this ascomycete by using several molecular assays. *bcfrq1* mRNA levels oscillate in LD as well as under free running conditions, whereas rhythms are lost under LL stimulation, or upon deletion of *bcwcl1* (Fig. 1 A, C, F, and G). These results are consistent with the notion that in *N. crassa* *frq* transcript levels oscillate in a cyclical environment (1) or in constant darkness (34). Likewise, in *Neurospora* a dysfunctional circadian clock is evidenced under LL—due to high *frq* expression levels (35)—and also in a $\Delta wc-1$ strain (36). The latter observations are also consistent with the fast light responsiveness of *bcfrq1* mRNA upon light stimulation, which depends on *bcwcl1* (28).

In addition, by overexpressing an extra copy of *bcfrq1* under a constitutive promoter, we demonstrated that *bcfrq1* expression levels at the endogenous locus become low and arrhythmic (Fig. S4), recapitulating the iconic experiment performed by Aronson et al. (34) in *N. crassa*, which provided solid evidence for the central role of negative feedback loops in circadian core mechanisms. Additionally, we showed that BcFRQ1 protein levels not only oscillate under DD culture conditions, but also anticipate changes in temperature cycles and can be entrained by cycles of different T (Fig. 1 B, D, and E). Together, these data demonstrate the presence of a functional circadian oscillator in *B. cinerea*, closing the gap in knowledge regarding clock mechanisms in fungi other than *Neurospora*. Whereas circadian rhythms (output) have been described in *Aspergillus* (37) and yeast (38), and more recently in bioluminescence in a mushroom (7), the molecular bases of the underlying oscillators remain obscure.

Remarkably, the absence of *bcfrq1* has a strong impact on sexual/asexual reproduction and virulence, which is further modulated by modifying media culture conditions (Fig. S6). Such a widespread effect of *bcfrq1* in *B. cinerea* biology clearly differs with what has been reported for the *frq* knockout strain (*frq¹⁰*) of *N. crassa*, which shows no reported phenotypic alterations related to the production of conidia (39). This result suggests novel

extra circadian functions for BcFRQ1, because the developmental phenotype of $\Delta bcfrq1$ is observed even under conditions where the fungal clock is already disrupted by a constant environmental cue (LL; Fig. S10), and also because in the absence of BcWCL1 (where the clock is also broken) multinucleated conidia capable of infection are easily produced (28). With respect to the latter, we noticed that the phenotypes of $\Delta bcwcl1$ and $\Delta bcfrq1$ strains are quite different, even though one of the molecular consequences of BcWCL1 absence is low level of *bcfrq1* transcripts (28). Therefore, although part of the phenotypes observed in the $\Delta bcwcl1$ strain should overlap with the ones seen in $\Delta bcfrq1$, this overlap does not occur. A plausible explanation resides in the known ability of FRQ to modify WCC activity. Thus, in *Neurospora*, the absence of FRQ leads to hyperactivity of WC-1 and its destabilization due to transcription-associated turnover (40). So, defects in signaling connected with BcWCL1 and other associated proteins involved in light signaling and development (i.e., BcVEL1) (41) could be impacted by the absence of BcFRQ1. Importantly, media composition can revert the sexually related phenotype of *B. cinerea*, suggesting that nutritional cues can partially override some of the developmental alterations observed in this mutant. The mechanisms behind these extracircadian functions of BcFRQ1 will be the subject of a different study.

Previous studies have highlighted the ways by which circadian clocks regulate plant defense mechanisms against bacteria, insects, and oomycetes (18). However, this paradigm had not been tested for fungi in general and for necrotrophic fungi in particular. Based on a typical type of assay conducted for these experiments (infection of WT and clock-deficient *Arabidopsis* plants at different circadian times followed by the development of infection under constant light conditions) the conclusion would have been that the plant clock solely regulates this interaction. Indeed, as we demonstrated herein, in LL, a condition known to reduce *B. cinerea* basal infection levels (28), the environmental disruption of the fungal clock does not dramatically

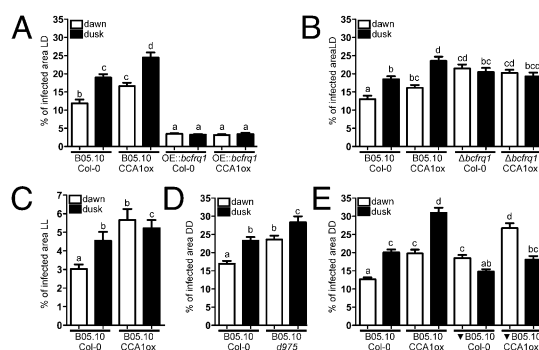


Fig. 4. The *B. cinerea* circadian clock influences the outcome of the interaction under 12:12 photoperiods as well as under constant darkness conditions. (A) Disruption of the clock by *bcfrq1* overexpression (OE::*bcfrq1*) affects temporal variation of virulence in LD. (B) There are no differences in the infection behavior observed between dawn and dusk for the $\Delta bcfrq1$ strain in LD. (C) Under LL, the *Arabidopsis* clock—but not the *Botrytis* clock—provides a time-of-day effect on susceptibility. (D) The B05.10 strain has the ability to produce bigger lesions at dusk in both Col-0 and *d975* arrhythmic mutant plants. (E) The relative magnitude of lesions achieved at dawn or dusk follows the time sensed by the fungus and not the plant. WT B05.10 strain was grown under the same LD cycle or 12 h out of phase (represented as a black inverted triangle) than the plants. *Arabidopsis* were inoculated at dawn or dusk with *B. cinerea* grown in the same phase or having opposite times. In all graphs, after macroconidia inoculations, WT (Col-0) and mutant plants were kept under the specified LD, LL, or DD regimes. White and black bars indicate conidia inoculation at dawn and dusk, respectively. Lesion area measurements were performed 3 dpi. Bars represent mean \pm SEM (different letters indicate statistical differences).

change the time-dependent outcome of the interaction, whereas in return the plant clock seems to adopt an important role as revealed by the data in the CCA1ox mutant (Fig. 4C), results that are aligned to what has been previously described for bacterial and oomycete pathogens (12, 13, 15). Nevertheless, by generating and evaluating *B. cinerea* clock mutants, we have demonstrated that in this fungus, a functional clock modulates its virulence under DD (Figs. 3 and 4D) and also under conditions that better reflect the real world (LD cycling conditions) (Fig. 4A and B). In addition, the experiments conducted with inverted cycles (Fig. 4E) are key in ruling out potentially strange explanations of the results, further supporting the importance of the fungal clock in the outcome of the *Botrytis*–*Arabidopsis* interaction.

Intriguingly, once we compensated $\Delta bcfq1$'s failure to produce microconidia—by manipulating media composition—we observed that its macroconidia have a bigger infection capability than the B05.10 WT strain. However, the temporal difference to infect better at night is lost (Figs. 3C and 4B). It is intriguing that when removing a central clock component, such as BcFRQ1 (but not BcWCL1), the infection is significantly enhanced. Nonetheless, similar results have been described for *A. thaliana*: Zhang et al. (15) reported that arrhythmic CCA1ox plants have enhanced resistance to the oomycete pathogen *Hpa*, which is in agreement with the results published by Wang et al. (13). Nevertheless, Bhardwaj et al. (12) and Zhang et al. (15) showed that the CCA1ox plants have enhanced susceptibility and loss of temporal regulation in defense against the virulent bacteria *Pseudomonas syringae*. Thus, although one would intuitively predict that an arrhythmic plant is always more prone to infection (due to the loss of the adaptive advantage of the clock), some of the results in *A. thaliana* reveal the opposite for the CCA1ox strain. Thus, as it was described for the plant clock protein CCA1, BcFRQ1 could be acting on other output pathways not regulated by the circadian clock (42), further complicating the molecular dissection of clock regulation on virulence mechanisms.

In this fencing contest between these two organisms, the dominance of each clock is judge and the results presented here indicate that under DD and LD conditions, the *B. cinerea* clock plays a fundamental role in the interaction, thus explaining the major part of the temporal variation of the infection process. In this regard, the most important plant defense mechanism against necrotrophic infection is the jasmonic acid (JA) pathway (43). Interestingly, a circadian gating effect allows higher JA-signaling activity response during the morning (14, 16). Although it is reasonable to hypothesize that the circadian clock of *B. cinerea* allows the fungus to anticipate the JA plant defense response, optimizing and generating its maximal virulence potential at dusk, the experiments using out-of-phase light:dark cycles (Fig. 4E) argue against this possible explanation because they show that the fungus can achieve maximum virulence even at dawn, as long as its internal clock indicates dusk time. A likely explanation, therefore, is that fungal virulence potential can override plant defense mechanisms. Further studies will help us to elucidate the mechanisms by which the circadian clock is modulating the pathogenic potential of this hostile organism at different times of the day.

Materials and Methods

***B. cinerea* Strains and Culture Conditions.** Strain B05.10 (26) of *B. cinerea* Pers.: Fr. [*Botryotinia fuckeliana* (de Bary) Whetzell], originally isolated from *Vitis vinifera* (Germany), was used as the recipient strain for genetic modifications. *B. cinerea* strains were cultivated in Petri dishes containing synthetic CM (44), PDA (Applichem), or Gamborg B5 (Duchefa Biochemie) supplemented with 2% (wt/vol) glucose. The strains were grown at 20 °C using Percival incubators, equipped with cool white light fluorescent tubes (light intensity up to 100 $\mu\text{M}/\text{m}^2/\text{s}$; wavelength 400–720 nm) in a 12:12-h LD regime.

Time Course Experiments, RNA Extraction, and Real-Time Quantitative PCR. All experiments were conducted in Percival incubators kept at 20 °C. For circadian experiments under DD conditions, B05.10, $\Delta bcwcl1$, and OE::*bcfrq1* strains were grown on PDA plates covered with cellophane under LL conditions for 24 h and then transferred to DD every 4 h. For LL circadian

experiments, B05.10 was grown on PDA plates covered with cellophane under DD for 24 h and then transferred to LL every 4 h. After 48 h, mycelia samples (obtained from independent PDA plates) were harvested. For the LD time course experiment, the B05.10 strain was inoculated on cellophane-covered PDA Petri dishes under LD culture conditions, during 96 h. Thereafter, cultures were harvested every 2 h, during 32 h. RNA isolation and real-time quantitative PCR (RT-qPCR) procedures were conducted as previously described (28). Expression values refer to the culture grown 4 h in the dark (1 = DD4) for Fig. 1A, C, and F (1 = DD16) and for Fig. 1G expression levels refer to the culture obtained in the dark (1 = LL0). The plot for Fig. 1A represents four biological with three technical replicates each and for Fig. 1C, F, and G two biological with two technical replicates each. In the case of Fig. S4A, transcript levels refer to the culture grown 4 h in the dark determined for the B05.10 strain (1 = DD4) and all of the plots represent three biological replicates. Each point, in black, represents mean values \pm SEM, whereas a trend line is depicted in orange.

Cloning of Replacements Cassettes. All of the replacement cassettes were assembled using a previously described strategy (28). The fully codon optimized (*oluc*) *N. crassa* luciferase gene (45) was used to generate a BcFRQ1-LUC reporter, as described for *Neurospora frq* (46).

Transformation of *B. cinerea*. *B. cinerea* transformations were carried out using protoplast-mediated transformation as previously described (27).

Southern Blot. PCR-verified mutants for $\Delta bcfq1$ were checked by Southern blot hybridization using the DIG Easy Hyb Hybridization solution (Roche) and the PCR DIG Probe Synthesis Kit (Roche) following manufacturer's instructions.

Assessment of Rhythmic Luminescence. The luciferase reporter BcFRQ1-LUC (two biological replicates) was analyzed using a PIXIS-CCD camera (Princeton Instruments) at 20 °C, using Petri plates filled with Gamborg B5, as described (45). Before being transferred to DD, plates were entrained for 3 d in LD. The reporter strain was also monitored under temperature entrainment conditions using steps from 22 °C to 27 °C under DD conditions (12:12) and by using 22 °C to 27 °C temperature cycles of varying period length (16, 20, and 24 T). Bioluminescence traces were acquired using the WinView software and analyzed in more detail using a custom-made ImageJ macro. For Fig. 1B, D, and E, the plots represent the mean of two biological with two technical replicates each.

Fluorescence Microscopy. Conidia from 2-wk-old culture were collected by glass wool filtration, diluted in water to 10^8 conidia per milliliter and then pelleted by centrifugation ($1,000 \times g$ for 20 min). Thereafter, conidia were stained with a dilution of 1 mg/mL of Hoechst 33342 (Life Technologies) for 15 min in darkness and then rinsed twice with 1 \times phosphate-buffered saline. Conidia were mounted in 30% (vol/vol) glycerol. Fluorescence microscopy was performed with Cytation 3 Cell Imaging Multi-Mode Reader (Biotek). Photos were obtained using Gen5 Data Analysis Software (Biotek).

***A. thaliana* Growth.** Accession Col-0, constitutive expression of the *CIRCADIAN CLOCK ASSOCIATED 1* (CCA1ox), and the triple mutant of PRR9, PRR7, and PRR5 (*d975*) were grown at 22 °C for 14 d under LL and then for another 2 wk at 20 °C under LD using Percival incubators. CCA1ox and *d975* plants are arrhythmic with respect to circadian outputs in LL, DD, and LD (15, 32, 47, 48).

Virulence Assays. Before infection, *A. thaliana* and *B. cinerea* were grown under LD conditions for 2 wk at 20 °C. Thereafter, leaves of approximately 1-mo-old plants were inoculated with *B. cinerea* conidia, which were obtained essentially as described previously (28). Briefly, spores were harvested from the same *B. cinerea* plate at dawn or dusk, and 7 μL of conidial suspensions of 2×10^5 spores per milliliter were used to inoculate unwounded *A. thaliana* leaves. Infections were done at dawn or dusk, according to the photoperiod of the plant growth chamber. All inoculated plants were kept inside plastic boxes at 20 °C under a humid environment, in LL, LD, or DD conditions for 72 h and then immediately analyzed (3 d postinoculation, dpi). Lesions on *A. thaliana* leaves—at least 180 infected leaves per *B. cinerea* strain and condition—were measured semiautomatically with ImageJ software using an external calibration scale. Because the different *A. thaliana* ecotypes have different leaf sizes, the lesion area was calculated by measuring the total and infected area of the leaf, being the total area of the leaf 100%. Therefore, measurements in the graphs are expressed as percentage. For Fig. 2B, the quantification was obtained from three independent virulence assays; the Mann-Whitney test was performed and *P* values < 0.05 were used as cutoff values for showing significance. For Figs. 3 and 4, the results were obtained from four and three independent virulence assays, respectively. For Figs. 3 and 4A, B,

and *E*, three-way analysis of variance and Tukey's HSD ($P < 0.05$) were used. The statistical analysis for Fig. 4 *C* and *D* was done by using two-way analysis of variance and Tukey's honest significant difference test ($P < 0.05$). For Figs. S7 and S8, two independent virulence assays were done; the Kruskal-Wallis test resulted in $P < 0.05$.

Trypan Blue Staining. Detection of *B. cinerea* hyphae in infected plant tissues was done as previously described (28).

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