



Circadian regulation of abiotic stress tolerance in plants

Jack Grundy, Claire Stoker and Isabelle A. Carré*

School of Life Sciences, University of Warwick, Coventry, UK

Extremes of temperatures, drought and salinity cause widespread crop losses throughout the world and impose severe limitations on the amount of land that can be used for agricultural purposes. Hence, there is an urgent need to develop crops that perform better under such abiotic stress conditions. Here, we discuss intriguing, recent evidence that circadian clock contributes to plants' ability to tolerate different types of environmental stress, and to acclimate to them. The clock controls expression of a large fraction of abiotic stress-responsive genes, as well as biosynthesis and signaling downstream of stress response hormones. Conversely, abiotic stress results in altered expression and differential splicing of the clock genes, leading to altered oscillations of downstream stress-response pathways. We propose a range of mechanisms by which this intimate coupling between the circadian clock and environmental stress-response pathways may contribute to plant growth and survival under abiotic stress.

Keywords: circadian clock, rhythmic gene expression, abiotic stress, plant hormones

Introduction

The world-wide population is predicted to increase from 7.2 billion to between 9.6 and 12.3 billion by 2100, increasing the demand on global food productivity (Gerland et al., 2014). However, nutrient deficiency, mineral toxicity and salinity severely limit the amount of land that is available for food production. In addition, world-wide crop losses due to extreme temperatures, drought, and flooding are estimated to rise about 50% by 2050 (Boyer, 1982). Water deficit was estimated to have caused €100 billion worth of damage in Europe between 1977 and 2007 (European Commission, 2007). The severe drought of 2012 affected 80% of the crop-growing territory in the USA (Hoerling et al., 2013), and such droughts are predicted to become more frequent as a result of global warming (Hansen et al., 2010; Lawrimore et al., 2011; Lobell and Gourdji, 2012; Cook et al., 2015).

There is an urgent need for development of crop varieties with improved tolerance to abiotic stress, but the development of such improved varieties will require a thorough understanding of the mechanisms by which plants are affected by these conditions, and how they can tolerate them. Abiotic stress can affect cellular homeostasis via a variety of mechanisms. Freezing temperatures promote intra- and extra-cellular ice crystal formation, leading to cellular damage, dehydration and disruption of the plasma membrane (Steponkus and Webb, 1992; Pearce, 2001; Yamazaki et al., 2008, 2009). High temperatures damage the plasma membrane by increasing its fluidity, leading to ion leakage. Heat stress can also result in inhibition of enzymatic activity and photosynthesis (Salvucci and Crafts-Brandner, 2004; Sharkey, 2005; Wahid et al., 2007; Allakhverdiev et al., 2008). High temperatures, salinity and drought stress are often accompanied by production of reactive oxygen species (ROS) which can lead to denaturation of functional and structural proteins (Gill and Tuteja, 2010; Suzuki et al., 2012). Gradual exposure to these different stresses causes plants to become more tolerant, however, and acclimated plants can survive harsher conditions than non-acclimated plants. The acclimation process includes the expression of new transcripts and proteins, and the accumulation

OPEN ACCESS

Edited by:

Seth J. Davis,
Max Planck Institute for Plant
Breeding Research, Germany

Reviewed by:

Elsebeth Kolmos,
BioConsortia, Inc., USA
Heather Knight,
Durham University, UK

*Correspondence:

Isabelle A. Carré,
School of Life Sciences, University
of Warwick, Gibbet Hill Road,
Coventry CV4 7AL, UK
isabelle.carre@warwick.ac.uk

Specialty section:

This article was submitted to
Plant Physiology,
a section of the journal
Frontiers in Plant Science

Received: 11 April 2015

Accepted: 04 August 2015

Published: 27 August 2015

Citation:

Grundy J, Stoker C and Carré IA
(2015) Circadian regulation of abiotic
stress tolerance in plants.
Front. Plant Sci. 6:648.
doi: 10.3389/fpls.2015.00648

of metabolites that result in a new homeostatic state that is more compatible with the change in environmental conditions. This allows stabilization of proteins and membranes, osmotic adjustment and re-establishment of the redox balance (Krasensky and Jonak, 2012; Mittler and Blumwald, 2015).

A number of transgenic lines with improved tolerance to environmental stress have been generated in a variety of crop species, and a few of these have been approved for commercialization (Lawlor, 2012; Cabello et al., 2014; Waltz, 2014). These transgenic lines exploit a variety of transgenes that act to maintain osmotic balance under abiotic stress conditions. For example, Monsanto's DroughtGard maize (MON87460) was transformed with the cold-shock protein CspB from *Bacillus subtilis* to improve water efficiency under drought stress (Nemali et al., 2014). The *betA* gene from the bacterium *Rhizobium meliloti* was introduced into Indonesian sugar cane in order to increase production of the osmoprotectant glycine betaine, resulting in enhanced resistance to drought (Marshall, 2014). An osmoregulator protein (osmotin) from tobacco was also used to improve the salinity tolerance of chili pepper, cotton, tomato, and strawberry plants (Husaini and Abdin, 2008; Parkhi et al., 2009; Goel et al., 2010; Subramanyam et al., 2010; Patade et al., 2013).

While these approaches have led to significant improvements, the genetic manipulation of master regulators of stress responses would potentially be more powerful because of the broader spectrum of resistance mechanisms that can be triggered by such modifications. However, as natural stress responses often lead to growth arrests, such modifications often cause detrimental pleiotropic effects, leading to reduced yields under normal conditions (Hussain et al., 2011). These problems can be alleviated by the use of stress inducible promoters, such that the genetic modification is only expressed at times when it is beneficial. Alternatively, the judicious choice of transcription factors that control specific pathways with minimum unwanted side effects can minimize such issues (Hussain et al., 2011). Nevertheless, an ideal approach to improving crop performance under abiotic stress conditions would be to exploit a genetic pathway that (i) contributes to tolerance to a broad range of environmental stresses and (ii) can naturally adapt in to enhance stress tolerance mechanisms when plants are exposed to deleterious conditions. Here, we will discuss recent evidence that the plant circadian clock may represent such a mechanism.

Circadian clocks are 24-h biological oscillators, which generally enable organisms to anticipate predictable, daily changes that are linked to the rotation of the earth. Such clocks have been found in a wide range of organisms, including cyanobacteria, fungi, insects, and mammals (Young and Kay, 2001). Having a clock is not essential for plant viability, and arrhythmic mutants are still capable of developing (Green et al., 2002). However, there is convincing evidence that having a clock that is well tuned to diurnal cycles of light and darkness makes an important contribution to plant fitness. For example, plants whose clocks were a good match to their rhythmic environment performed best in competition experiments (Dodd et al., 2005). This may be explained, at least in part, by the role of the clock to ensure the optimum use of starch at night while avoiding starvation before dawn (Graf et al., 2010; Scialdone et al., 2013; Mugford et al., 2014;

Webb and Satake, 2015). However, recent data suggest that the circadian clock may also contribute to plant fitness by enhancing their ability to tolerate abiotic stress.

Abnormal Function of the Circadian Clock Results in Altered Abiotic Stress Tolerance

The molecular mechanism of the plant circadian clock was investigated using the genetic model system *Arabidopsis thaliana*, and was found to be composed of a transcriptional regulatory network. The mechanism by which this network oscillates has been reviewed in detail elsewhere (see, for example Carré and Veflingstad, 2013; Hsu and Harmer, 2014; Seo and Mas, 2014), and we will only describe it minimally here, to introduce the relevant genetic components. Briefly, the day starts with expression of two closely related transcription factors, LATE ELONGATED HYPOCOTYL (LHY) and CIRCADIAN CLOCK ASSOCIATED 1 (CCA1), which peak in the early morning (Schaffer et al., 1998; Wang and Tobin, 1998). Transcription of the *LHY* and *CCA1* genes is down-regulated in the afternoon and through the night by a set of sequentially expressed PSEUDO-RESPONSE REGULATOR proteins known as PRR 9, 7, 5, and 1, where PRR1 is also known as TIMING OF CAB or TOC1 (Matsushika et al., 2000; Strayer et al., 2000; Alabadi et al., 2001; Nakamichi et al., 2010). The inhibition of *LHY/CCA1* transcription by the PRR proteins is released late at night through the action of an evening complex (EC) composed of three proteins, EARLY FLOWERING (ELF) 3, ELF4, and LUX ARRHYTHMO (LUX). The EC shuts down the expression of the *PRR* genes, which allows expression of *LHY* and *CCA1* to rise again in the following morning (Hicks et al., 2001; Doyle et al., 2002; Hazen et al., 2005; Onai and Ishiura, 2005; Nusinow et al., 2011; Pokhilko et al., 2012). *LHY* and *CCA1* subsequently bring down expression of *TOC1* and the cycle restarts (Alabadi et al., 2001). The F-box proteins ZEITLUPE (ZTL) and LOV KELCH PROTEIN 2 (LKP2), function as a complex with GIGANTEA (GI) to control the light-dependent proteolysis of the *TOC1* protein, and this acts to increase to amplitude of oscillations of the gene network under diurnal light–dark cycles (Mas et al., 2003; Kim et al., 2007). The function of the central oscillator is synchronized to environmental light dark cycles via input pathways mediated by phytochrome and cryptochrome photoreceptors (Somers et al., 1998). The circadian clock in turn controls downstream physiological processes via a number of mechanisms, termed output pathways.

Mutations in components of the circadian clock not only impacts on the function of the central oscillator but also alter plants' ability to both tolerate environmental stress conditions, and acclimate to them. For example, plants with mutations in *CCA1*, *LHY*, *ELF3*, *ELF4*, *LUX*, *PRR5*, *PRR9*, and *PRR7* are hypersensitive to ROS-generating agents, whereas *CCA1*-overexpressing plants (*CCA1-ox*) are more tolerant than wild-type plants (Lai et al., 2012). Overexpression of *LKP2* enhances drought tolerance (Miyazaki et al., 2015). The *prr5,7,9* triple mutant is more tolerant to high salinity and drought stresses (Nakamichi et al., 2009). *GI-ox* plants show enhanced salt sensitivity, whereas *gi* mutants show enhanced salt tolerance and enhanced survival

under drought stress and oxidative stress (Kim et al., 2013; Fornara et al., 2015). Reduced expression of *TOC1* in *toc1-RNAi* plants results in improved drought tolerance, whereas overexpression of *TOC1* causes increased water loss and reduced survival under drought conditions (Legnaioli et al., 2009). The *prr5,7,9* and *toc1* mutants show significantly increased freezing survival (Nakamichi et al., 2009; Keily et al., 2013), but *gi*, *lux*, *lhy*, *cca1*, and *lhy cca1* mutants have reduced constitutive and acclimated freezing tolerance (Cao et al., 2005; Espinoza et al., 2010; Dong et al., 2011; Chow et al., 2014).

These findings demonstrate that the circadian clock contributes to plants' ability to tolerate environmental stress. While the effects observed may seem relatively small (only a 1–2°C difference in freezing tolerance for *lhy cca1* double mutants, for example), this may prove valuable to allow recovery after a frost or after extreme drought for example (Lawlor, 2012). Interestingly, there is no correlation between the effect of mutations on the function of the clock and their effects on abiotic stress tolerance. For example, mutations which cause a short-period phenotype (*cca1*, *lhy*, *prr5*), a long-period phenotype (*prr9*) or complete arrhythmia (*elf3*, *elf4*, *lux*) cause similar hypersensitivity to ROS-generating agents (Green and Tobin, 1999; Mizoguchi et al., 2002; Ito et al., 2003; Yamamoto et al., 2003; Lai et al., 2012). Furthermore, plants that show similar arrhythmic phenotypes can have opposite phenotypes with regard to ROS sensitivity (*elf3*, *elf4*, and *lux* mutants vs *CCA1-ox*), drought tolerance (*LKP2-ox* vs *prr5,7,9* mutants) or freezing tolerance (*lux* vs *prr5,7,9* mutants; Hicks et al., 1996; Wang and Tobin, 1998; Schultz et al., 2001; Doyle et al., 2002; Hazen et al., 2005; Nakamichi et al., 2005, 2009; Chow et al., 2014; Miyazaki et al., 2015). This suggests that the effect of these mutations on circadian period or on the ability of the clock to oscillate is not the primary cause of the altered stress tolerance phenotype. A more likely hypothesis is that the altered expression level of the clock genes in the mutants has a direct impact to either up- or down-regulate abiotic stress responses, as discussed in the following sections.

Circadian Regulation of Stress-Responsive Genes

The effects of the clock on plant physiology and development are largely mediated through the regulation of gene expression. Approximately 80% of transcripts cycle in *Arabidopsis* plants under diurnal light–dark or temperature cycles, and 30–40% of these oscillations persist upon transfer to constant conditions (Harmer et al., 2000; Edwards et al., 2006; Covington et al., 2008; Michael et al., 2008). Similar observations have been made in a number of crop plants and tree species, including rice, poplar, maize, tomato, and soybean (Facella et al., 2008; Khan et al., 2010; Filichkin et al., 2011; Marcolino-Gomes et al., 2014). Remarkably, the phasing of expression for the majority of genes was found to be conserved across monocots and dicots (Filichkin et al., 2011). One particularly interesting outcome from transcriptomic analyses was that many genes that are responsive to abiotic stress were found to be under the control of the circadian clock. Thus, in *Arabidopsis*, approximately 50% of genes that are responsive to heat, osmoticum, salinity or water deprivation were found to

be expressed rhythmically in constant light, and 40% of cold-regulated genes (Covington et al., 2008). Rhythmic expression of abiotic stress-responsive genes was also reported for soybean and barley (Habte et al., 2014; Marcolino-Gomes et al., 2014).

This raises the question of how the daily timing of expression of these genes (i.e., their phase) relates to rhythmic changes in environmental conditions. This was examined in **Figure 1**, based on phase data for *Arabidopsis* genes retrieved from the Diurnal database (Blasing et al., 2005; Mockler et al., 2007; Covington et al., 2008) and on lists of cold, heat, drought and mannitol-responsive genes derived from existing datasets (Lee et al., 2005; Kilian et al., 2007; Wilkins et al., 2010; Barah et al., 2013). This showed that although individual genes within abiotic stress-response pathways were expressed across a broad range of phases, peaks of expression were clustered around specific times of the day. This was true for responses to environmental conditions that vary in a diurnal manner, such as cold or heat, but also for responses to conditions that progress over a number of days, such as drought, or conditions that are constant, such as high osmoticum. This suggests that circadian regulation of abiotic stress responses may confer an advantage to the plant even under conditions that do not change in a diurnal fashion.

The majority of cold-inducible genes peaked in the afternoon, a few hours ahead of cooler temperatures at night (**Figure 2**). This included the cold-responsive transcription factors *CRE/DRE-BINDING FACTORS* 1, 2, and 3, which function as master regulators of the cold acclimation response (*CBF1,2,3*, also known as *Dehydration-Responsive Element Binding* or *DREB1–B, C, and A*, respectively), and the circadian and cold-regulated genes *CCR1*, *CCR2*, and *RD29A* (Carpenter et al., 1994; Harmer et al., 2000; Dodd et al., 2006). On the other hand, genes that are down-regulated by cold peaked around dawn (**Figure 2**). Thus, the timing of expression of cold-responsive genes was consistent with a role in anticipation of predictable, diurnal changes in temperature. In cotton and soybean plants, maximum low-temperature resistance was observed at the beginning of the night, corresponding to the time when most cold-responsive genes are expressed in unstressed conditions (Couderchet and Koukkari, 1987; Rikin et al., 1993). This suggests that rhythmic transcription of cold-responsive genes may form the basis of rhythmic changes in chilling tolerance.

In *Arabidopsis*, the majority of heat-inducible genes were expressed during the day, and heat-repressed genes were broadly expressed at night (**Figure 2**). Similarly, in spinach, five genes encoding heat shock proteins (HSP70s) were found to exhibit maximum expression in the middle of the day. This was associated with a diurnal variation in thermo-tolerance. Plants exhibited minimum heat-induced electrolyte leakage following treatments in the afternoon and early evening, which was consistent with the hypothesis, that the rhythmic expression of HSP70s confers heat tolerance at the end of the day (Li and Guy, 2001). The production of ROS-scavenging enzymes also peaks mid-day in *Arabidopsis* and may contribute to heat tolerance by minimizing of heat-induced oxidative stress (Lai et al., 2012).

The majority of drought-inducible genes peaked around dawn (**Figure 2**). Plants experience a water deficit in the daytime, due to the prolonged opening of stomata for photosynthetic gas exchange

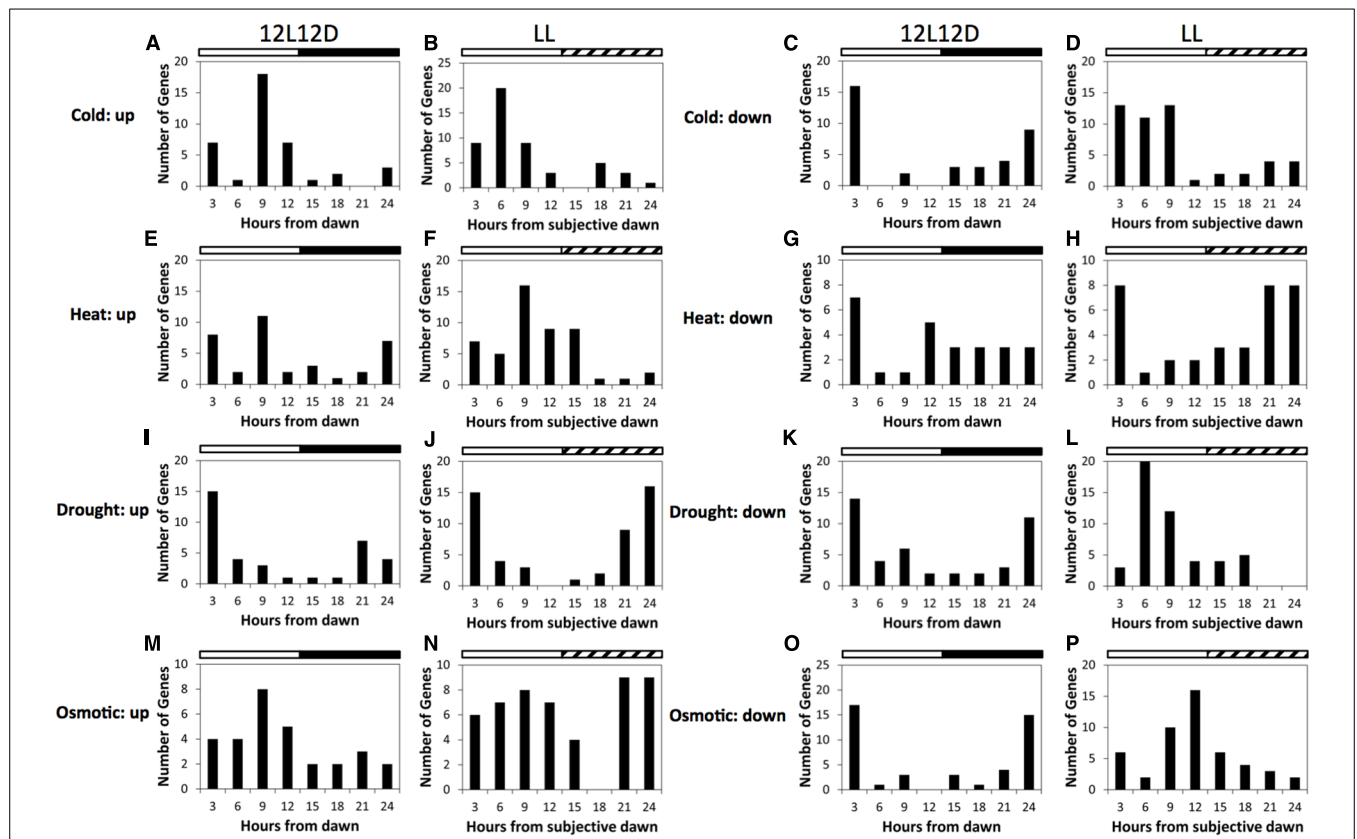


FIGURE 1 | Circadian regulation of (A-D) cold, (E-H) heat, (I-L) drought and (M-P) osmoticum-responsive genes in *Arabidopsis*. Histograms show the timing of peak mRNA levels in 12L12D or following transfer to constant light, for the 50 most significantly stress-induced (up) or repressed (down) genes that also exhibit circadian regulation. Lists of stress-responsive genes were based on the following datasets: 24 h at 0°C (Lee et al., 2005), 3 h of at 38°C (Barah

et al., 2013), water limitation for 4 days (Wilkins et al., 2010), and growth on 300 mM mannitol (Kilian et al., 2007). Phase data for *Arabidopsis* genes were obtained from experiments LDHH-Stitt and LL12 in the Diurnal database (Blasing et al., 2005; Mockler et al., 2007; Covington et al., 2008). Solid white and black bars indicate intervals of light and darkness, respectively, whereas hatched bars indicate subjective nights.

(Jarvis, 1976), and the rhythmic expression of drought-inducible genes may allow plants to minimize water loss at this crucial time. However, many drought-repressible genes were also found to peak during the day, when they would have been expected to peak at night. The physiological significance of this is unclear. In summary, while the timing of expression of stress-responsive genes is mostly consistent with the timing of daily environmental changes, this is not always the case. One possible explanation is that, genes which play a role in tolerance to a given type of abiotic stress may also contribute to other pathways, and that there may be a greater selective pressure for optimal timing of their expression relative to these other processes.

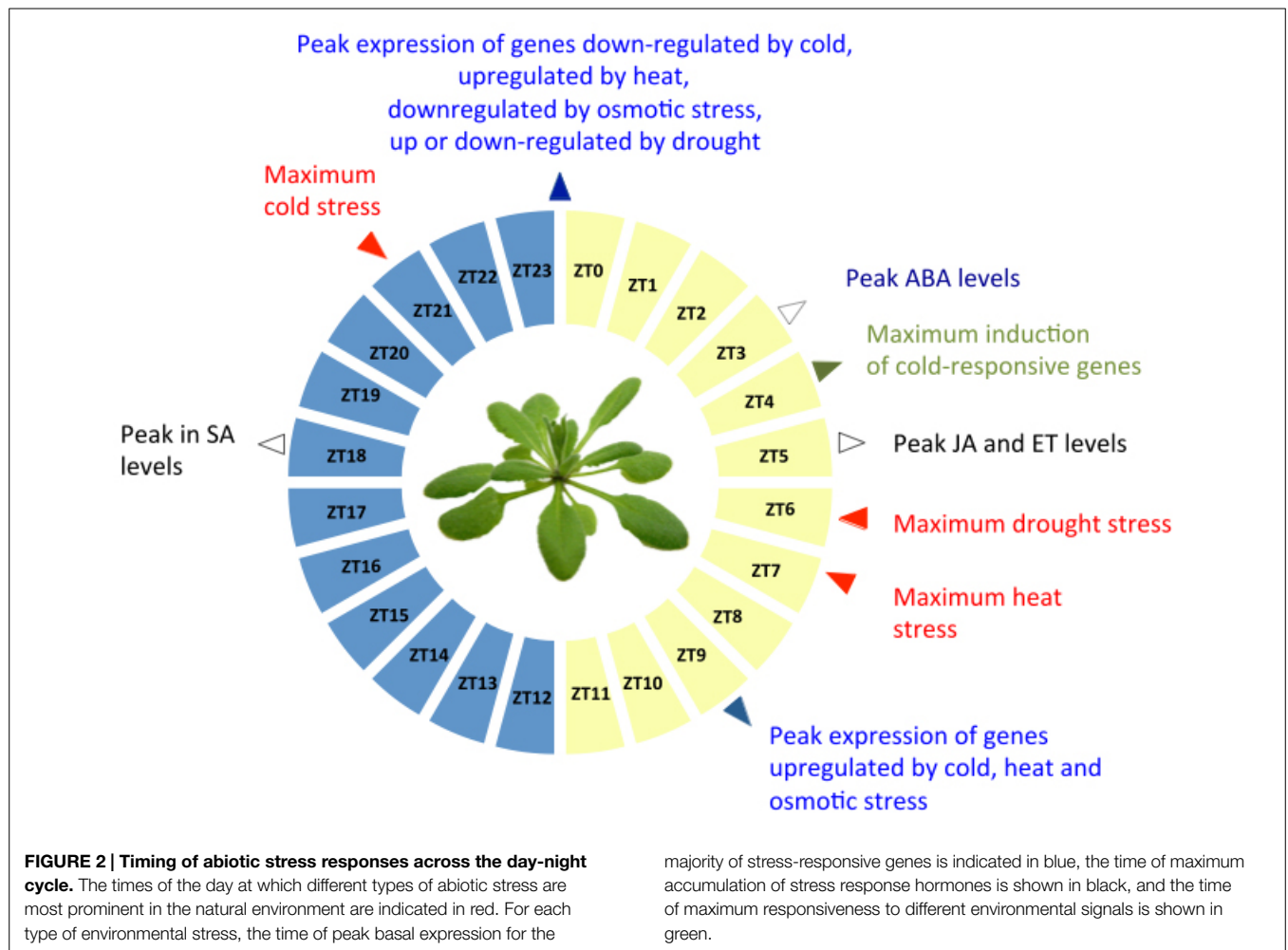
Circadian Gating of Abiotic Stress Responses

In addition to driving the rhythmic, basal expression of stress-responsive genes, the clock controls the amplitude of their responses to environmental signals, in a process called gating. For example, cold treatments at different times of the day result in differential induction of *CBF1,2,3* transcription, with

maximal induction observed in the morning (Fowler et al., 2005; Thomashow, 2010). Consistent with the role of CBF transcription factors as master regulators of the chilling response, about 75% of cold-responsive genes exhibit similar gating of cold responses with maximum responsiveness in the morning. A similar timing of maximal cold-responses was observed in peach trees. Out of five *CBF* homologs, four showed maximal cold-induction in the morning, and this was associated with induction of downstream targets including *DEHYDRIN (DHN) 1* and 3 (Artlip et al., 2013).

The conserved gating of low temperature responses across distant plant species strongly suggests that the circadian gating of chilling responses confers a selective advantage. Little is known, however, about how the gating of temperature responses contributes to plant fitness. One hypothesis is that, while the rhythmic expression of cold-responsive genes allows anticipation of predictable cold temperatures at night, gated responses may be important to allow rapid changes in gene expression in response to unexpectedly cold mornings.

Responses to water deficit are also gated by the circadian clock, and maximal drought-induced changes in gene expression are observed at dusk (Wilkins et al., 2010; Kielbowicz-Matuk et al., 2014). Interestingly, different sets of genes are induced by identical



cold or drought treatments given at different time of the day (Bieniawska et al., 2008; Wilkins et al., 2010). Similarly, significant differences in heat stress responsive pathways were observed in grapevine, depending on whether the treatment was applied during the day or during the night (Rienth et al., 2014), and the physiological implications of these time of the day-differences remain to be investigated.

Regulation of Abiotic Stress Responses by Core Oscillator Components

A number of components of the circadian clock encode transcription factors, which have a dual role, both to ensure the function of the central oscillator and to control the rhythmic expression of downstream genes.

Thus, a number of abiotic stress-responsive genes are under the control of core components of the circadian oscillator. For example, the LHY and CCA1 transcription factors play a key role to regulate low temperature responses by controlling expression of the *CBF* genes. The promoters of *CBF1*, 2, and 3 contain LHY and CCA1 binding sites, also known as Evening Elements or EEs (AAATATCT). A CCA1:GFP fusion protein was found to bind the *CBF1-3* locus in chromatin immunoprecipitation

experiments (Harmer et al., 2000; Dong et al., 2011), and loss of LHY and CCA1 function in the *lhy cca1* double mutant resulted in arrhythmic basal expression of *CBF1*, 2, and 3. This provided strong evidence that LHY and CCA1 play an essential role in the rhythmic expression of the *CBF* genes and of the cold responses that they regulate. The TOC1 and LUX proteins were also found to bind the promoter of *CBF3*, and mathematical modeling showed that the regulation of *CBF3* transcription by TOC1 and LUX was important for the correct gating of *CBF3* responses (Keily et al., 2013). Accurate gating patterns could only be reproduced in simulations when (i) expression of *CBF3* was down-regulated by TOC1 and the EC; (ii) it was up-regulated by LHY/CCA1, and (iii) cold was assumed to increase the level of LHY/CCA1 protein. In addition to controlling master regulators of the low temperature response, LHY and CCA1 control expression of cold-responsive genes in a direct manner. Thus, cold-responsiveness of the cold-inducible genes *CONSTANS-LIKE 1 (COL1)* and *COLD REGULATED GENE 27 (COR27)* was shown to result from a synergy between EEs and ABA-responsive elements (ABREs, CACGTG) in their promoters (Mikkelsen and Thomashow, 2009).

LKP2 was recently suggested to play a role to control responses to drought. LKP2-overexpressing plants exhibited a smaller

TABLE 1 | Direct regulation of stress hormone biosynthesis and signaling by core components of the *Arabidopsis* circadian clock.

Hormone	Gene ID	Gene name	Role	TOC1 target	PRR5 target	PRR7 target
JA	AT1G17420	LOX3	JA biosynthesis	Y	N	N
JA	AT1G72520	LOX4	JA biosynthesis	Y	N	N
JA	AT3G45140	LOX2	JA biosynthesis	Y	N	N
JA	AT1G19180	JAZ1	Repressor of JA-responsive genes	Y	N	N
JA	AT1G70700	JAZ9	Repressor of JA-responsive genes	Y	N	N
JA	AT5G11270	OCP3	JA signaling	Y	N	Y
JA	AT1G25540	PFT1	Regulator of JA responses	N	Y	N
JA	AT1G80840	WRKY40	JA-responsive transcription factor	N	Y	N
JA	AT3G06490	MYB108	JA-responsive transcription factor	N	Y	N
JA	AT1G52890	ANAC019	JA-responsive transcription factor	N	Y	N
ET	AT4G11280	ACS6	ET biosynthesis	Y	N	N
ET	AT5G03280	EIN2	ET signaling	Y	N	N
ET	AT5G47230	ERF5	ET-responsive transcription factor	Y	N	N
ET	AT1G53170	ERF8	ET-responsive transcription factor	Y	N	N
ET	AT4G39403	PLS	Negative regulator of ET signaling	N	Y	N
ABA	AT5G13630	ABAR	Putative ABA receptor	Y	Y	N
ABA	AT5G59220	PP2C:HA1	Catalytic subunit of ABA receptor	N	Y	N
ABA	AT3G11410	PP2CA/AHG3	Catalytic subunit of ABA receptor	Y	Y	N
SA	AT2G41010	CAMB25	Regulation of SA biosynthesis	Y	N	N
SA	AT1G20020	LFNR2	Regulation of SA biosynthesis	N	Y	N
SA	AT3G56710	SIB1	Regulation of SA biosynthesis	N	Y	N

Evidence for direct regulation of genes in the JA, SA, ET, or ABA pathways was obtained from genome-wide analyses of binding sites for the TOC1, PRR5, and PRR7 proteins (Huang et al., 2012; Nakamichi et al., 2012; Liu et al., 2013).

stomatal opening and showed elevated expression of several drought-responsive genes (Miyazaki et al., 2015). LKP2's protein partner, GI was also found to regulate the expression of genes involved in drought and cold stress responses, including a number of *EARLY RESPONSIVE TO DEHYDRATION* genes (*ERD10*, *ERD7*, and *AT4G04340*) and cold-regulated genes (*COR78*, *KIN2*, *COR15a*, *COR15b*, *COR314*, and *COR413*; Fornara et al., 2015). This effect of GI was dependent on CYCLING DOF FACTORS (CDFs), a family of rhythmically expressed transcriptional repressors known for their roles in the photoperiodic regulation of flowering time (Imaizumi et al., 2005). As LKP2 and its homolog FKF1 interact with GI to control the light-dependent degradation of CDF proteins, these observations provide independent evidence for a role of the LKP2/GI/CDF regulatory module in drought tolerance.

Circadian Control of Hormone Biosynthesis and Signaling

Many plant responses to abiotic stress are mediated by phytohormones. For example, abscisic acid (ABA) accumulates rapidly upon exposure to water stress or salinity and controls many stress-adaptation responses (Agarwal and Jha, 2010). ABA levels are under circadian control. Peak levels were observed during the day in the angiosperms *Senecio minimus*, *Arbutus unedo*, and *Arabidopsis thaliana*, and in the gymnosperm *Callitris rhomboidea* (Burschka et al., 1983; Lee et al., 2006; McAdam et al., 2011). More complex temporal patterns have been observed in *Nicotiana tabacum*, where ABA levels showed a first peak immediately after dawn, a second in the afternoon and a third at night (Novakova et al., 2005). Circadian regulation takes place at the level of ABA biosynthesis. In *Arabidopsis*, the

enzymes responsible for biosynthesis of carotenoid precursors and the rate-limiting enzyme *NINE-CIS-EPOXYCAROTENOID DIOXYGENASE* (NCED) peak in the morning (Thompson et al., 2000; Covington et al., 2008). In maize, the rhythmic expression of ABA biosynthetic enzymes was also consistent with ABA synthesis taking place in the early morning (Khan et al., 2010). In sugar cane, the temporal pattern of ABA biosynthesis may be more complex as one gene involved in ABA synthesis peaked in the early morning, and two others peaked during the night (Hotta et al., 2013). The rhythmic production of ABA was proposed to be controlled by the PRR5, 7, and 9 proteins, because analysis of a triple mutant (*prr5,7,9*) revealed increased ABA levels (Fukushima et al., 2009). However, this effect is likely to be indirect because ChIP-seq analyses of genome-wide binding sites for PRR5, PRR7, and TOC1 did not reveal binding of these proteins to the promoters of any component of ABA biosynthesis pathways (Table 1; Nakamichi et al., 2010; Huang et al., 2012; Liu et al., 2013).

Plants' ability to respond to ABA also varies with the time of the day. For example, stomatal closure is more sensitive to ABA in the afternoon, allowing for a more sensitive control of stomatal aperture (Correia et al., 1995). *TOC1*-overexpressing plants had widely open stomata throughout the diurnal cycle and exhibited increased sensitivity to drought. In contrast, plants with reduced expression of *TOC1* had the opposite phenotype, suggesting that *TOC1* may act to limit stomatal aperture (Legnaioli et al., 2009). This effect of *TOC1* was proposed to be mediated through the ABA-binding protein ABAR (also known as CHLH or GUN5) as expression of the *ABAR* gene was directly controlled by *TOC1* binding to its promoter (Legnaioli et al., 2009; Huang et al., 2012). However, the function of ABAR in ABA signaling remains controversial as the mechanisms by which it regulates ABA responses are not known (Hubbard et al., 2010). *TOC1* may

nevertheless regulate ABA responses through a different signaling pathway, as ChIP-seq data indicate that it binds to the promoters of two class A phosphatases 2C, HAI1 and PP2CA/AHG3, that function as catalytic subunits of another ABA receptor [regulatory components of ABA receptor (RCAR), also known as pyrabactin resistance 1 (PYR) or PYR1-like (PYL); Huang et al., 2012].

The clock also regulates the expression of ABA-responsive genes in a direct manner. In *Arabidopsis*, 40% of ABA responsive genes were found to also be under circadian control (Covington et al., 2008). Analysis of motifs in the promoters of these genes revealed over-representation for LHY and CCA1 binding sites, suggesting that LHY and CCA1 control expression of large fraction of ABA-responsive genes by binding directly to their promoters (Huang et al., 2007).

While ABA is the main hormone involved in responses to drought and to osmotic stress, salicylic acid (SA), jasmonic acid (JA), ethylene (ET) also play a role in responses to other types of abiotic stress (Yang et al., 2004; Cheng et al., 2013; Wasternack and Hause, 2013; Kazan, 2015), and reduction of GA levels and signaling contributes to the resulting growth arrests (Colebrook et al., 2014).

Ethylene and JA act cooperatively in the regulation of responses to cold, salinity, drought and heat. The production of ET peaks during the subjective day in *Arabidopsis* and sorghum, and at the end of the day in potato (Finlayson et al., 1999; Thain et al., 2004; Chincinska et al., 2013). In *Arabidopsis*, rhythmic expression of the rate-limiting enzyme *ACC SYNTHASE 8 (ACS8)* is key to the rhythmicity of ET biosynthesis, and a knock-out mutation abolished circadian ET accumulation (Thain et al., 2004). Another *ACS* gene (*ACS6*) may also be under circadian control, as *TOC1* was found to bind its promoter (Huang et al., 2012; **Table 1**). Two putative *ACC OXIDASE* enzymes may also play a role since they exhibit rhythmic expression with a similar phase as *ACS8* (Covington et al., 2008). Sucrose transport from source to sink tissue may contribute to the rhythmic production of ET, as mutation of the rhythmically expressed sucrose transporter *SUT4* abolished diurnal changes in ET levels in potatoes (Chincinska et al., 2013). The clock may also influence signaling downstream of ET, since the ethylene-responsive transcription factor (ERF) *ETHYLENE-INSENSITIVE 3 (EIN3)* is expressed rhythmically in *Arabidopsis* under the control of the clock-associated *XAP5 CIRCADIAN TIMEKEEPER (XCT)*; Ellison et al., 2011). Furthermore, ChIP-seq analyses revealed the binding of *TOC1* and *PRR5* proteins to genes encoding *ERFs* 8 and 5, and to regulators of ET signaling such as *EIN2* and *POLARIS (PLS)*; Huang et al., 2012; Nakamichi et al., 2012; **Table 1**).

Jasmonic acid levels also peak in the middle of the day (Goodspeed et al., 2012; Zhang et al., 2013). Rhythmic changes in JA levels may be controlled by the *TOC1* protein, as *TOC1* was found at the promoters of several genes encoding 13-lipoxygenase enzymes (*13-LOX*) which initiate the biosynthesis of JA (**Table 1**). Responses to JA may also be regulated by *TOC1*, *PRR5*, and *PRR7*, as ChIP-seq analyses revealed binding of these proteins to the promoters of several genes encoding regulators of JA responses. This included two negative regulators of the JA signaling pathway (JASMONATE ZIM DOMAIN

or JAZ proteins) and several JA-induced transcription factors (Huang et al., 2012; Nakamichi et al., 2012; Liu et al., 2013; **Table 1**). However, the gating of JA responses seems to be primarily mediated by the clock-associated protein TIME FOR COFFEE (TIC; Shin et al., 2012). TIC interacts with a positive regulator of JA signaling, *MYC2*, leading to its proteasomal degradation and to the down-regulation of JA responses (Kazan and Manners, 2013). *MYC2* mRNA accumulation is also under circadian control, providing another layer of rhythmic regulation of JA signaling. While *tic* mutants retain some level of gating of JA responses, this is completely abolished in *myc2* mutants (Shin et al., 2012).

Salicylic acid is increased in response to most biotic stresses and was proposed to have an acclimation-like effect, due primarily to enhanced anti-oxidative capacity (Horváth et al., 2007; Miura and Tada, 2014). SA is generally thought to act antagonistically with JA, at least with regard to defenses against pathogens (Spoel et al., 2007). As might be expected, SA levels peaks at night in *Arabidopsis*, 180° out of phase with JA levels (Goodspeed et al., 2012, 2013; Zhang et al., 2013). *PRR5* and *TOC1* may play a role to up-regulate the production of SA at night, as they bind to the promoters of several genes encoding regulators of SA biosynthesis (Huang et al., 2012; Nakamichi et al., 2012; Liu et al., 2013; **Table 1**). SA accumulation is negatively influenced by a rhythmically expressed phosphate transporter gene (*PHT4;1*) and this has also been proposed to contribute to oscillations in SA levels (Wang et al., 2014).

Altogether, these studies reveal a pervasive effect of the clock on stress-responsive hormonal pathways. Remarkably, circadian regulation occurs at multiple levels, including hormone biosynthesis, signaling and rhythmic expression of hormone-responsive genes. ChIP-seq analyses have revealed the direct regulation of key components of hormone signaling pathways by *PRR5*, *PRR7*, and *TOC1* (**Table 1**) and it is likely that further genome-wide analyses will reveal additional roles for other components of the clock. Interestingly, data available so far reveal a division of labor between the *PRR* proteins. For example *TOC1* was found at the promoter of several JA biosynthesis genes (*LOX2,3,4*) and of two repressors of JA responses (*JAZ1,9*) whereas *PRR5* was found at a distinct set of genes encoding regulators of JA responses and JA-responsive transcription factors (*PFT1*, *WRKY40*, *MYB108: ANAC019*; see **Table 1**) and *PRR7* only regulated one JA signaling component (*OCP3*). This may reflect a need for sequential expression of these genes, or alternatively, for independent control of their expression in response to changes in the environment. Understanding how regulation by the different clock proteins is integrated to regulate flux through the different signaling pathways will be a major challenge, which can only be addressed through mathematical modeling.

Effects of Abiotic Stress on the Transcription of Clock Genes

While the clock has a clear impact on how plants respond to environmental stress, there is also evidence that the function of

the clock is altered in response to abiotic stress conditions. For example, high salinity resulted in lengthening of the circadian period in wheat, and osmotic stress increased the expression level of barley clock genes and advanced their phase of expression (Erdei et al., 1998; Habte et al., 2014). Similarly, drought stress reduced the expression of evening-specific components of the clock in soybean (*TOC1*, *LUX*, *ELF4*, and *PRR*-like genes), and this led to disruption of the circadian system (Marcolino-Gomes et al., 2014). Although the period of the clock varies by less than 1 h in *Arabidopsis* over the physiological range of temperatures (Gould et al., 2006; Salome et al., 2010), the expression level of oscillator components is altered. Peak expression levels of the *TOC1* and *LUX* transcripts increase as temperatures get warmer, whereas *CCA1* and *LHY* RNA rhythms increase in amplitude and peak expression level as temperatures get colder, from 17 to 12°C (Gould et al., 2006; Mizuno et al., 2014). On the other hand, heat stress was reported to shorten the circadian period (Kusakina et al., 2014), whereas cold stress leads to a loss of rhythmicity. At 4°C, most clock-associated genes oscillate with reduced amplitudes under day-night cycles and become arrhythmic upon transfer to constant light. Only a small proportion of the genes that cycle at 20°C maintain clear diurnal oscillations at 4°C (Martino-Catt and Ort, 1992; Bieniawska et al., 2008; Espinoza et al., 2010).

These effects of abiotic stress on the function of the clock may be mediated by changes in the level of stress response hormones. In *Arabidopsis*, SA application did not influence circadian period length, but application of the ET precursor ACC (1-aminocyclopropane-1-carboxylic acid) shortened the period length (Hanano et al., 2006). In contrast, ABA treatment lengthened the period of the *Arabidopsis* clock (Hanano et al., 2006). This was associated with a reduction in *CCA1* mRNA levels, which is surprising because loss of function of *CCA1* causes a short period phenotype (Green and Tobin, 1999). However, ABA may affect the oscillator mechanism by altering expression of other clock genes, which would result in lengthening of period. Evolutionary conserved ABREs are present in the promoters of *TOC1*, *LHY*, and *CCA1*, and may represent multiple points of entry for ABA signals into the clock (Bieniawska et al., 2008; Spensley et al., 2009; Picot et al., 2010; Habte et al., 2014).

Several core clock components are regulated by stress-related transcription factors. For example, the heat shock transcription factor HsfB2b represses transcription from the *PRR7* promoter at high temperatures and in response to drought (Kolmos et al., 2014), and binding of CBF1 to the promoter of the *LUX* gene plays a role to maintain robust transcriptional oscillations at low temperatures (Chow and Kay, 2013). In addition, the transcriptional repressor FLOWERING BASIC HELIX-LOOP-HELIX 1 (FBH1) binds to the *CCA1* promoter. FBH1 expression is up-regulated by heat and down-regulated by cold, and its overexpression in transgenic plants leads to reduced expression of *CCA1* and shortening of circadian period by approximately 1 h. This suggests that, in wild-type plants, *FBH1* plays a role to down-regulate *CCA1* expression at warm temperatures (Nagel et al., 2014).

Differential Splicing of Clock Genes in Response to Environmental Stress

The plant clock is also regulated by alternative splicing, and mutation of a spliceosomal component (*SPICEOSOMAL TIMEKEEPER LOCUS1* or *STIPL1*) produced a long-period phenotype in *Arabidopsis* (Jones et al., 2012). Many clock-associated genes including *CCA1*, *LHY*, *PRR7*, *PRR9*, *TOC1*, *ELF3*, and *ZTL* undergo extensive alternative splicing through diverse mechanisms including intron retention, exon skipping and selection of alternative 5' splice sites (Filichkin et al., 2010; Filichkin and Mockler, 2012; James et al., 2012; Seo et al., 2012; Kwon et al., 2014). Many of these alternative splicing events occur in response to abiotic stress conditions (Filichkin et al., 2010, 2015; Filichkin and Mockler, 2012; James et al., 2012; Seo et al., 2012; Kwon et al., 2014). For example, low temperatures suppress alternative splicing of *ELF3* but induce that of *TOC1*. In contrast, heat induces the alternative splicing of *CCA1*, *PRR7*, *TOC1*, and *ELF3* genes, and high salinity suppresses that of *ELF3* (Kwon et al., 2014).

Alternatively spliced isoforms often contain a premature termination codon that can give rise to a shorter but still functional protein, or to a non-functional protein that will be degraded by the non-sense-mediated decay pathway. For example, the *CCA1* and *LHY* transcripts are both alternatively spliced to form either the full-length protein or a truncated protein lacking the MYB DNA-binding region (James et al., 2012; Seo et al., 2012). Both truncated proteins, produced through intron retention, suppress the formation of functional homo- and heterodimers (*CCA1-CCA1*, *CCA1-LHY*, and *LHY-LHY*) through competitive inhibition. This impairs the ability of *LHY* and *CCA1* proteins to bind DNA and results in dampened rhythmicity. Interestingly, the production of the truncated isoform of *CCA1* (*CCA1-β*) decreases at low temperature, while that of truncated isoform of *LHY* increases. This results in a relatively constant level of functional homo- and heterodimers and enables the clock to maintain a constant period length across the physiological range of temperatures (James et al., 2012). Overexpression of the full-length form of *CCA1* (35S::*CCA1-α*) enhanced freezing tolerance whereas overexpression of the truncated variant (35S::*CCA1-β*) decreased it (Seo et al., 2012). This was taken to suggest that the regulation of *CCA1* alternative splicing by cold temperatures is important for freezing tolerance. However, the effect of these transgenes is not an accurate reflection of what happens in response to temperature in wild-type plants, as any change in the splicing of *CCA1* is normally compensated for by an opposite change in the splicing of *LHY*. An alternative and perhaps simpler interpretation for the effects of *CCA1-α* and *CCA1-β* overexpression is that reduced function of both *LHY* and *CCA1* proteins in 35S::*CCA1-β* plants may result in a cold-sensitive phenotype, whereas increased function of *CCA1* in 35S::*CCA1-α* may result in enhanced chilling resistance. More work is required to fully understand the importance of *CCA1* alternative splicing for plant tolerance to cold temperatures.

The molecular mechanisms linking the plant circadian clock to the control of RNA processing are largely unknown, but

PROTEIN ARGININE METHYLTRANSFERASE 5 (PRMT5) is thought to play a key role in this process. This rhythmically expressed protein acts to transfer methyl groups to arginine residues present in histones and Sm spliceosomal proteins, and contributes to the regulation of many pre-messenger-RNA splicing events (Sanchez et al., 2010). In particular, *prmt5* mutant plants produce an alternatively spliced form of the core clock component *PRR9*, resulting in a long period phenotype (Hong et al., 2010; Sanchez et al., 2010; Hernando et al., 2015). The rhythmic expression of PRMT5 was responsive to both light and temperature cues, but interestingly the transcript accumulated at different phases in different environmental conditions. Expression of PRMT5 peaked at dusk under diurnal light–dark cycles, but peaked just before dawn under thermocycles (Hong et al., 2010). This altered phase in response to thermocycles was also seen when plants were exposed to photocycles suggesting that the PRMT5 rhythm is preferentially synchronized to temperature cycles. Thus, PRMT5 may play an important role to ensure changes in splicing of the clock genes in response to diurnal changes in temperature. Interestingly, *prmt5* mutants also exhibit reduced tolerance to salinity (Hernando et al., 2015), but it is not clear at this point whether this is related to the altered function of their circadian clock, or whether this is due to direct effects on abiotic stress response pathways.

Alternative splicing is known to mediate clock responses to environmental changes in other organisms. In *Neurospora*, temperature-dependent alternative splicing of the *FREQUENCY* transcript allows adaptation of the circadian clock to seasonal variations of photoperiod and temperature (Diernfellner et al., 2005, 2007). In *Drosophila*, thermosensitive splicing in the 3′-UTR of the clock gene *period* (*per*) leads to phase advanced accumulation of *per* RNA on cold days, which is crucial for seasonal adaptation of the circadian locomotor activity rhythm (Majercak et al., 1999). Interestingly, the *dart5-1* mutant, which is affected in the *Drosophila* PRMT5 homolog, also exhibited an abnormal period of locomotor activity (Sanchez et al., 2010). This suggested that PRMT5 may also play a role to regulate the alternative splicing of one or more clock genes in flies, and may identify a conserved mechanism for alternative splicing of clock-associated genes in response to environmental signals.

Abiotic stress may also affect the abundance of clock-associated proteins at the post-translational level. For example, salt stress induces the proteasomal degradation of the GI protein, thus reducing the amplitude of its oscillations (Kim et al., 2013). Reduced GI protein levels contribute to salt acclimation by freeing the SOS2 kinase to activate the Na⁺/H⁺ antiporter SOS1. It is likely that further examples of such post-translational regulation will emerge in the near future.

These findings are interesting because they show that the function of the circadian clock is altered in response to abiotic stress. Furthermore, the presence of evolutionarily conserved stress-responsive promoter elements in the promoters of clock genes from distantly related species suggests that these effects of abiotic stress on the clock mechanism may provide a selective advantage.

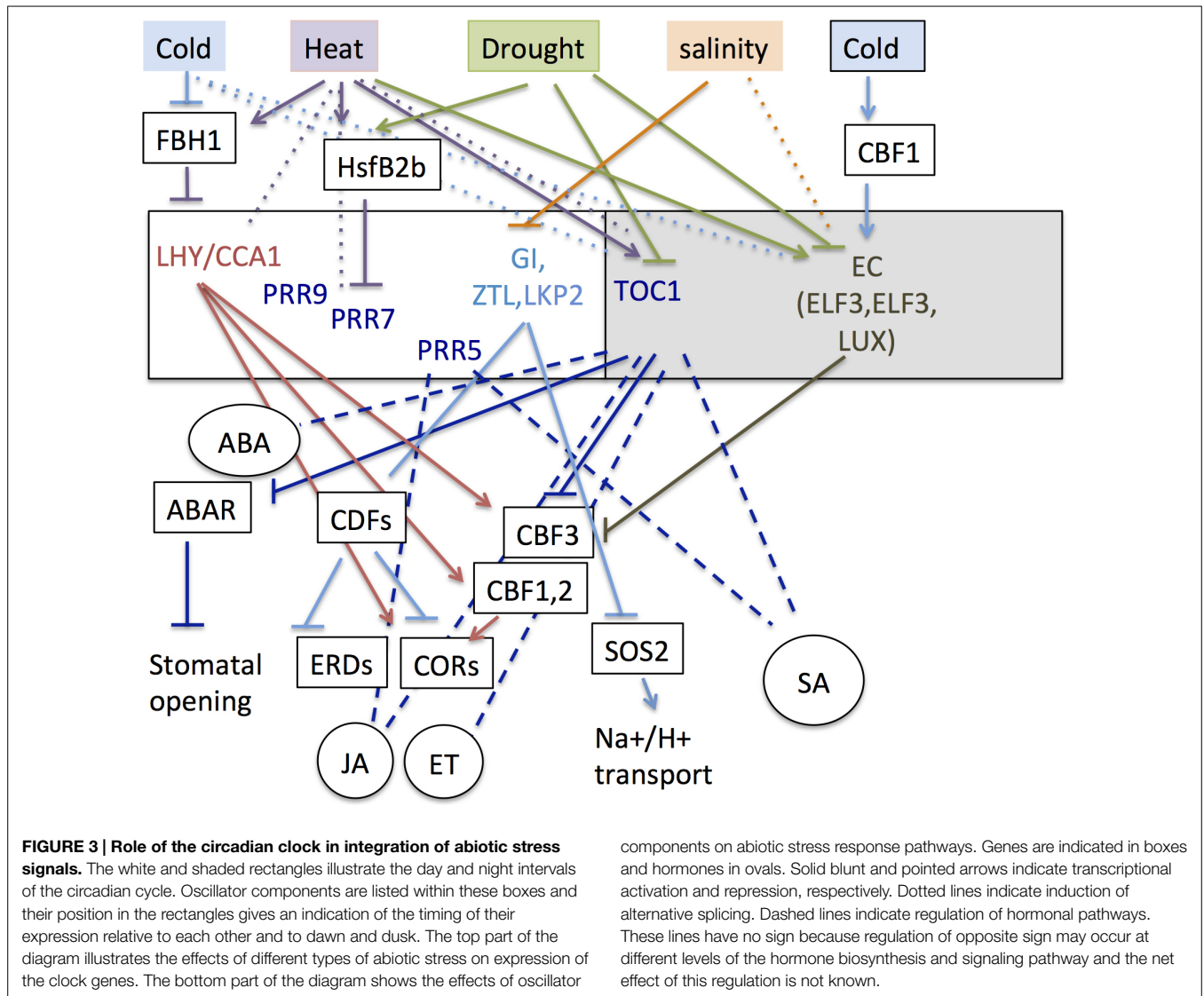
Conclusions and Perspectives

The past few years of research have revealed a web of regulatory connections between abiotic stress response pathways and the plant circadian clock. The current knowledge of these interactions is summarized in **Figure 3**, but the picture is unlikely to be complete at this stage. Many aspects of abiotic stress responses are regulated directly by components of the central oscillator. The clock modulates the activity of stress response pathways at multiple levels and through a range of mechanisms, including the transcriptional regulation of key regulators of the stress response, the direct regulation of stress-responsive genes and signaling by stress response hormones. Most intriguingly, abiotic stress causes altered transcription and/or splicing of the clock genes, suggesting that abiotic stress response pathways feedback onto the oscillator to regulate its function.

How does circadian regulation of abiotic stress responses provide an advantage to plants, when their constitutive expression would provide a high level of stress tolerance at all times? As stress responses in plants are frequently associated with a growth arrest, we suggest that circadian regulation may provide an advantage by ensuring that plants can grow during part of the diurnal cycle, while only expressing stress tolerance mechanisms at times of the day when they are required most critically. However, this does not explain the contribution of the clock to salinity and osmotic stress tolerance, as these stress conditions do not vary in a diurnal fashion. Therefore, the clock may also contribute to abiotic stress tolerance via some other mechanisms.

As the clock allows plants to perceive changes in season (Carré, 2001; Yanovsky and Kay, 2002), it may act to modulate abiotic stress responses depending on the time of the year. This would ensure that a strong response is only triggered at those times of the year when a stress is likely to be pronounced and persistent. Thus, plant responses to cold were recently found to be regulated by seasonal changes in photoperiod (Lee and Thomashow, 2012). Under warm temperatures, the *CBF2* gene oscillated with higher amplitude under short day conditions (8L16D) than under long day conditions (16L8D). This resulted in up-regulation of the CBF pathway and increased freezing tolerance in the winter. Another study suggested that CDFs, a family of transcription factors known for their roles in the photoperiodic regulation of flowering, may mediate improved cold tolerance under short-day photoperiods. Short days lead to the stabilization of CDF proteins, which act to up-regulate expression of cold-responsive genes, resulting in enhanced cold tolerance in the winter (Fornara et al., 2015). Future work may reveal similar seasonal regulation of other types of abiotic stress responses in plants.

As the function of the clock is altered in response to abiotic stress, circadian regulation may also provide a form of memory of the signal, which may contribute to the stress acclimation process. Just like perturbing the oscillation of a pendulum results in lasting changes in its swinging pattern, acute up- or down-regulation of the clock genes in response to abiotic stress would be expected to result in persistent changes in the phase or amplitude of their oscillation. This may, in



turn, result in long-term changes in the expression of the stress response pathways that they regulate. For example, water deficit causes an increase in the amplitude of the rhythmic expression of plasma membrane aquaporins, and this was proposed to benefit the plant by decreasing transpiration and by avoiding excessive dehydration of the rhizosphere that would dramatically decrease its hydraulic conductivity (Caldeira et al., 2014).

Figure 3 suggests a number of mechanisms by which the clock might ensure that stress responses are sustained following initial exposure. For example, cold temperatures increase the expression of *LHY* and *CCA1*, and this would in turn act to enhance expression of cold-responsive genes (*CBFs* and *CORs*) over multiple circadian cycles. Heat stress, on the other hand, down-regulates *LHY* and *CCA1* expression while increasing that of *TOC1*, and this would be expected to reduce the peak expression of *CBF* genes. Drought stress represses expression of *TOC1*, resulting in reduced stomatal opening, while salinity

represses expression of *GI*, which promotes the activity of the Na^+/H^+ transporter *SOS1*. However, as plants in their natural environment are often exposed to multiple types of abiotic stresses simultaneously, the level and temporal pattern of expression of the clock genes is more likely to reflect the integration of multiple signals. For example, drought and heat stress, which are often experienced in combination, have opposite effects on *TOC1* and *LUX* transcription, but act synergistically to enhance *PRR7* expression by promoting that of *HSFB2b*. Therefore the predicted combined effect would be elevated expression of *PRR7* and intermediate levels of *TOC1* and *LUX* transcription. On the other hand, cold promotes the expression of *LUX* and *CCA1* but has no known effect on *TOC1*. Therefore, the predicted combined effect of cold and drought would again be intermediate levels of *LUX* expression, but this would be associated with low levels of *TOC1* and high levels of *CCA1*. Thus, different combinations of environmental stresses may lead to distinct profiles of expression of the clock genes and to distinct effects on downstream stress

responses. Further work will be required to test whether such integration of abiotic stress signals by the clock does take place, and whether it plays a role to maximize plant survival or productivity.

In summary, circadian regulation may provide a sophisticated, tunable mechanism to ensure an optimum compromise between plant growth and abiotic stress tolerance. Adjustments in clock function in response to changes in the environment may allow plants to re-adjust this balance and to ensure a sustained increase in abiotic stress tolerance when needed. If these hypotheses are correct, such mechanisms may in the future be exploited for improving abiotic stress tolerance in crops without reducing yield.

References

- Agarwal, P. K., and Jha, B. (2010). Transcription factors in plants and ABA dependent and independent abiotic stress signalling. *Biol. Plant.* 54, 201–212. doi: 10.1007/s10535-010-0038-7
- Alabadi, D., Oyama, T., Yanovsky, M. J., Harmon, F. G., Mas, P., and Kay, S. A. (2001). Reciprocal regulation between TOC1 and LHY/CCA1 within the *Arabidopsis* circadian clock. *Science* 293, 880–883. doi: 10.1126/science.1061320
- Allakhverdiev, S., Kreslavski, V., Klimov, V., Los, D., Carpentier, R., and Mohanty, P. (2008). Heat stress: an overview of molecular responses in photosynthesis. *Photosyn. Res.* 98, 541–550. doi: 10.1007/s11120-008-9331-0
- Artlip, T. S., Wisniewski, M. E., Bassett, C. L., and Norelli, J. L. (2013). CBF gene expression in peach leaf and bark tissues is gated by a circadian clock. *Tree Physiol.* 33, 866–877. doi: 10.1093/treephys/tpt056
- Barah, P., Jayavelu, N. D., Mundy, J., and Bones, A. M. (2013). Genome scale transcriptional response diversity among ten ecotypes of *Arabidopsis thaliana* during heat stress. *Front. Plant Sci.* 4:532. doi: 10.3389/fpls.2013.00532
- Bieniaszka, Z., Espinoza, C., Schlereth, A., Sulpice, R., Hinch, D. K., and Hannah, M. A. (2008). Disruption of the *Arabidopsis* circadian clock is responsible for extensive variation in the cold-responsive transcriptome. *Plant Physiol.* 147, 263–279. doi: 10.1104/pp.108.118059
- Blasing, O. E., Gibon, Y., Gunther, M., Hohne, M., Morcuende, R., Osuna, D., et al. (2005). Sugars and circadian regulation make major contributions to the global regulation of diurnal gene expression in *Arabidopsis*. *Plant Cell* 17, 3257–3281. doi: 10.1105/tpc.105.035261
- Boyer, J. S. (1982). Plant productivity and environment. *Science* 218, 443–448. doi: 10.1126/science.218.4571.443
- Burschka, C., Tenhunen, J. D., and Hartung, W. (1983). Diurnal variations in abscisic acid content and stomatal response to applied abscisic acid in leaves of irrigated and non-irrigated *Arbutus unedo* plants under naturally fluctuating environmental conditions. *Oecologia* 58, 128–131. doi: 10.1007/BF00384552
- Cabello, J. V., Lodeyro, A. F., and Zurbriggen, M. D. (2014). Novel perspectives for the engineering of abiotic stress tolerance in plants. *Curr. Opin. Biotechnol.* 26, 62–70. doi: 10.1016/j.copbio.2013.09.011
- Caldeira, C. F., Jeanguenin, L., Chaumont, F., and Tardieu, F. (2014). Circadian rhythms of hydraulic conductance and growth are enhanced by drought and improve plant performance. *Nat. Commun.* 5:5365. doi: 10.1038/ncomms6365
- Cao, S., Ye, M., and Jiang, S. (2005). Involvement of GIGANTEA gene in the regulation of the cold stress response in *Arabidopsis*. *Plant Cell Rep.* 24, 683–690. doi: 10.1007/s00299-005-0061-x
- Carpenter, C. D., Kreps, J. A., and Simon, A. E. (1994). Genes encoding glycine-rich *Arabidopsis thaliana* proteins with RNA-binding motifs are influenced by cold treatment and an endogenous circadian rhythm. *Plant Physiol.* 104, 1015–1025. doi: 10.1104/pp.104.3.1015
- Carré, I. A. (2001). Day-length perception and the photoperiodic regulation of flowering in *Arabidopsis*. *J. Biol. Rhythms* 16, 415–423. doi: 10.1177/074873001129002006
- Carré, I. A., and Veflingstad, S. R. (2013). Emerging design principles in the *Arabidopsis* circadian clock. *Semin. Cell Dev. Biol.* 24, 393–398. doi: 10.1016/j.semcdb.2013.03.011
- Cheng, M.-C., Liao, P.-M., Kuo, W.-W., and Lin, T.-P. (2013). The *Arabidopsis* ETHYLENE RESPONSE FACTOR1 regulates abiotic stress-responsive gene expression by binding to different *cis*-acting elements in response to different stress signals. *Plant Physiol.* 162, 1566–1582. doi: 10.1104/pp.113.221911
- Chincinska, I., Gier, K., Krügel, U., Liesche, J., He, H., Grimm, B., et al. (2013). Photoperiodic regulation of the sucrose transporter StSUT4 affects the expression of circadian-regulated genes and ethylene production. *Front. Plant Sci.* 4:26. doi: 10.3389/fpls.2013.00026
- Chow, B. Y., and Kay, S. A. (2013). Global approaches for telling time: omics and the *Arabidopsis* circadian clock. *Semin. Cell Dev. Biol.* 24, 383–392. doi: 10.1016/j.semcdb.2013.02.005
- Chow, B. Y., Sanchez, S. E., Breton, G., Prunedo-Paz, J. L., Krogan, N. T., and Kay, S. A. (2014). Transcriptional Regulation of LUX by CBF1 mediates cold input to the circadian clock in *Arabidopsis*. *Curr. Biol.* 24, 1518–1524. doi: 10.1016/j.cub.2014.05.029
- Colebrook, E. H., Thomas, S. G., Phillips, A. L., and Hedden, P. (2014). The role of gibberellin signalling in plant responses to abiotic stress. *J. Exp. Biol.* 217, 67–75. doi: 10.1242/jeb.089938
- Cook, B. I., Ault, T. R., and Smerdon, J. E. (2015). Unprecedented 21st century drought risk in the American Southwest and Central Plains. *Sci. Adv.* 1, e1400082. doi: 10.1126/sciadv.1400082
- Correia, M. J., Pereira, J. S., Chaves, M. M., Rodrigues, M. L., and Pacheco, C. A. (1995). ABA xylem concentrations determine maximum daily leaf conductance of field-grown *Vitis vinifera* L. plants. *Plant Cell Environ.* 18, 511–521. doi: 10.1111/j.1365-3040.1995.tb00551.x
- Couderchet, M., and Koukkari, W. (1987). Cold sensitivity oscillations of young soybean plants. *Prog. Clin. Biol. Res.* 227A, 59–65.
- Covington, M. F., Maloof, J. N., Straume, M., Kay, S. A., and Harmer, S. L. (2008). Global transcriptome analysis reveals circadian regulation of key pathways in plant growth and development. *Genome Biol.* 9, R130. doi: 10.1186/gb-2008-9-8-r130
- Diernfellner, A., Colot, H. V., Dintsis, O., Loros, J. J., Dunlap, J. C., and Brunner, M. (2007). Long and short isoforms of *Neurospora* clock protein FRQ support temperature compensated circadian rhythms. *FEBS Lett.* 581, 5759–5764. doi: 10.1016/j.febslet.2007.11.043
- Diernfellner, A. C. R., Schafmeier, T., Merrow, M. W., and Brunner, M. (2005). Molecular mechanism of temperature sensing by the circadian clock of *Neurospora crassa*. *Genes Dev.* 19, 1968–1973. doi: 10.1101/gad.345905
- Dodd, A. N., Jakobsen, M. K., Baker, A. J., Telzerow, A., Hou, S.-W., Laplaze, L., et al. (2006). Time of day modulates low-temperature Ca²⁺ signals in *Arabidopsis*. *Plant J.* 48, 962–973. doi: 10.1111/j.1365-313X.2006.02933.x
- Dodd, A. N., Salathia, N., Hall, A., Kevei, E., Toth, R., Nagy, F., et al. (2005). Plant circadian clocks increase photosynthesis, growth, survival, and competitive advantage. *Science* 309, 630–633. doi: 10.1126/science.1115581
- Dong, M. A., Farré, E. M., and Thomashow, M. F. (2011). CIRCADIAN CLOCK-ASSOCIATED 1 and LATE ELONGATED HYPOCOTYL regulate expression of the C-REPEAT BINDING FACTOR (CBF) pathway in *Arabidopsis*. *Proc. Natl. Acad. Sci. U.S.A.* 108, 7241–7246. doi: 10.1073/pnas.1103741108
- Doyle, M. R., Davis, S. J., Bastow, R. M., Mcwatters, H. G., Kozma-Bognar, L., Nagy, F., et al. (2002). The ELF4 gene controls circadian rhythms and flowering time in *Arabidopsis thaliana*. *Nature* 419, 74–77. doi: 10.1038/nature00954

Author Contributions

JG and CS carried out the initial research and wrote the first draft of the manuscript. JG prepared **Figure 1**, and CS **Figure 2** and **Table 1**. IC edited the work, wrote the conclusions and perspectives section, prepared **Figure 3** and finalized the manuscript.

Acknowledgments

JG was funded by a BBSRC-KTN CASE studentship and CS by the BBSRC MIBTP training programme.

- Edwards, K., Anderson, P., Hall, A., Salathia, N., Locke, J., Lynn, J., et al. (2006). FLOWERING LOCUS C mediates natural variation in the high-temperature response of the *Arabidopsis* circadian clock. *Plant Cell* 18, 639–650. doi: 10.1105/tpc.105.038315
- Ellison, C. T., Vandenbussche, F., Van Der Straeten, D., and Harmer, S. L. (2011). XAP5 CIRCADIAN TIMEKEEPER regulates ethylene responses in aerial tissues of *Arabidopsis*. *Plant Physiol.* 155, 988–999. doi: 10.1104/pp.110.164277
- Erdei, L., Szegletes, Z., Barabás, K. N., Pesténác, A., Fülöp, K., Kalmár, L., et al. (1998). Environmental stress and the biological clock in plants: changes of rhythmic behavior of carbohydrates, antioxidant enzymes and stomatal resistance by salinity. *J. Plant Physiol.* 152, 265–271. doi: 10.1016/S0176-1617(98)80141-7
- Espinoza, C., Degenkolbe, T., Caldana, C., Zuther, E., Leisse, A., Willmitzer, L., et al. (2010). Interaction with diurnal and circadian regulation results in dynamic metabolic and transcriptional changes during cold acclimation in *Arabidopsis*. *PLoS ONE* 5:e14101. doi: 10.1371/journal.pone.0014101
- European Commission. (2007). Addressing the challenge of water scarcity and droughts in the European Union. *Communication from the Commission to the European Parliament and the Council*, COM(2007) 414, Brussels.
- Facella, P., Lopez, L., Carbone, F., Galbraith, D. W., Giuliano, G., and Perrotta, G. (2008). Diurnal and circadian rhythms in the tomato transcriptome and their modulation by cryptochrome photoreceptors. *PLoS ONE* 3:e2798. doi: 10.1371/journal.pone.0002798
- Filichkin, S. A., Breton, G., Priest, H. D., Dharmawardhana, P., Jaiswal, P., Fox, S. E., et al. (2011). Global profiling of rice and poplar transcriptomes highlights key conserved circadian-controlled pathways and cis-regulatory modules. *PLoS ONE* 6:e16907. doi: 10.1371/journal.pone.0016907
- Filichkin, S. A., Cumbie, J. S., Dharmawardhana, P., Jaiswal, P., Chang, J. H., Palusa, S. G., et al. (2015). Environmental stresses modulate abundance and timing of alternatively spliced circadian transcripts in *Arabidopsis*. *Mol. Plant* 8, 207–227. doi: 10.1016/j.molp.2014.10.011
- Filichkin, S. A., and Mockler, T. C. (2012). Unproductive alternative splicing and nonsense mRNAs: a widespread phenomenon among plant circadian clock genes. *Biol. Dir.* 7, 20–20. doi: 10.1186/1745-6150-7-20
- Filichkin, S. A., Priest, H. D., Givan, S. A., Shen, R., Bryant, D. W., Fox, S. E., et al. (2010). Genome-wide mapping of alternative splicing in *Arabidopsis thaliana*. *Genome Res.* 20, 45–58. doi: 10.1101/gr.093302.109
- Finlayson, S. A., Lee, I.-J., Mullet, J. E., and Morgan, P. W. (1999). The mechanism of rhythmic ethylene production in sorghum. The role of phytochrome B and simulated shading. *Plant Physiol.* 119, 1083–1090. doi: 10.1104/pp.119.3.1083
- Fornara, F., De Montaigu, A., Sánchez-Villarreal, A., Takahashi, Y., Ver Loren Van Themaat, E., Huettel, B., et al. (2015). The GI-CDF module of *Arabidopsis* affects freezing tolerance and growth as well as flowering. *Plant J.* 81, 695–706. doi: 10.1111/tpl.12759
- Fowler, S. G., Cook, D., and Thomashow, M. F. (2005). Low temperature induction of *Arabidopsis* CBF1, 2, and 3 is gated by the circadian clock. *Plant Physiol.* 137, 961–968. doi: 10.1104/pp.104.058354
- Fukushima, A., Kusano, M., Nakamichi, N., Kobayashi, M., Hayashi, N., Mizuno, T., et al. (2009). Impact of clock-associated *Arabidopsis* pseudo-response regulators in metabolic coordination. *Proc. Natl. Acad. Sci. U.S.A.* 106, 7251–7256. doi: 10.1073/pnas.0900952106
- Gerland, P., Raftery, A. E., Ševčíková, H., Li, N., Gu, D., Spoorenberg, T., et al. (2014). World population stabilization unlikely this century. *Science* 346, 234–237. doi: 10.1126/science.1257469
- Gill, S. S., and Tuteja, N. (2010). Reactive oxygen species and antioxidant machinery in abiotic stress tolerance in crop plants. *Plant Physiol. Biochem.* 48, 909–930. doi: 10.1016/j.plaphy.2010.08.016
- Goel, D., Singh, A. K., Yadav, V., Babbar, S. B., and Bansal, K. C. (2010). Overexpression of osmotin gene confers tolerance to salt and drought stresses in transgenic tomato (*Solanum lycopersicum* L.). *Protoplasma* 245, 133–141. doi: 10.1007/s00709-010-0158-0
- Goodspeed, D., Chehab, E. W., Min-Venditti, A., Braam, J., and Covington, M. F. (2012). *Arabidopsis* synchronizes jasmonate-mediated defense with insect circadian behavior. *Proc. Natl. Acad. Sci. U.S.A.* 109, 4674–4677. doi: 10.1073/pnas.1116368109
- Goodspeed, D., Liu, J. D., Chehab, E. W., Sheng, Z., Francisco, M., Kliebenstein, D. J., et al. (2013). Postharvest circadian entrainment enhances crop pest resistance and phytochemical cycling. *Curr. Biol.* 23, 1235–1241. doi: 10.1016/j.cub.2013.05.034
- Gould, P., Locke, J., Larue, C., Southern, M., Davis, S., Hanano, S., et al. (2006). The molecular basis of temperature compensation in the *Arabidopsis* circadian clock. *Plant Cell* 18, 1177–1187. doi: 10.1105/tpc.105.039990
- Graf, A., Schlereth, A., Stitt, M., and Smith, A. M. (2010). Circadian control of carbohydrate availability for growth in *Arabidopsis* plants at night. *Proc. Natl. Acad. Sci. U.S.A.* 107, 9458–9463. doi: 10.1073/pnas.0914299107
- Green, R. M., Tingay, S., Wang, Z.-Y., and Tobin, E. M. (2002). Circadian rhythms confer a higher level of fitness to *Arabidopsis* plants. *Plant Physiol.* 129, 576–584. doi: 10.1104/pp.004374
- Green, R. M., and Tobin, E. M. (1999). Loss of the CIRCADIAN CLOCK-ASSOCIATED protein 1 in *Arabidopsis* results in altered clock-regulated gene expression. *Proc. Natl. Acad. Sci. U.S.A.* 96, 4176–4179. doi: 10.1073/pnas.96.7.4176
- Habte, E., Muller, L. M., Shtaya, M., Davis, S. J., and Von Korff, M. (2014). Osmotic stress at the barley root affects expression of circadian clock genes in the shoot. *Plant Cell Environ.* 37, 1321–1327. doi: 10.1111/pce.12242
- Hanano, S., Domagalska, M. A., Nagy, F., and Davis, S. J. (2006). Multiple phytohormones influence distinct parameters of the plant circadian clock. *Genes Cells* 11, 1381–1392. doi: 10.1111/j.1365-2443.2006.01026.x
- Hansen, J., Ruedy, R., Sato, M., and Lo, K. (2010). GLOBAL SURFACE TEMPERATURE CHANGE. *Rev. Geophys.* 48, RG4004. doi: 10.1029/2010RG000345
- Harmer, S. L., Hogenesch, J. B., Straume, M., Chang, H. S., Han, B., Zhu, T., et al. (2000). Orchestrated transcription of key pathways in *Arabidopsis* by the circadian clock. *Science* 290, 2110–2113. doi: 10.1126/science.290.5499.2110
- Hazen, S. P., Schultz, T. F., Prunedo-Paz, J. L., Borevitz, J. O., Ecker, J. R., and Kay, S. A. (2005). LUX ARRHYTHMO encodes a Myb domain protein essential for circadian rhythms. *Proc. Natl. Acad. Sci. U.S.A.* 102, 10387–10392. doi: 10.1073/pnas.0503029102
- Hernando, C. E., Sanchez, S. E., Mancini, E., and Yanovsky, M. J. (2015). Genome wide comparative analysis of the effects of PRMT5 and PRMT4/CARM1 arginine methyltransferases on the *Arabidopsis thaliana* transcriptome. *BMC Genomics* 16:192. doi: 10.1186/s12864-015-1399-2
- Hicks, K. A., Albertson, T. M., and Wagner, D. R. (2001). EARLY FLOWERING 3 encodes a novel protein that regulates circadian clock function and flowering in *Arabidopsis*. *Plant Cell* 13, 1281–1292. doi: 10.1016/S1360-1385(01)02184-7
- Hicks, K. A., Millar, A. J., Carre, I. A., Somers, D. E., Straume, M., Meeks-Wagner, D. R., et al. (1996). Conditional circadian dysfunction of the *Arabidopsis* early-flowering 3 mutant. *Science* 274, 790–792. doi: 10.1126/science.274.5288.790
- Hoerling, M., Schubert, S., Mo, K., Aghakouchak, A., Berbery, H., Dong, J., et al. (2013). An interpretation of the origins of the 2012 central great plains drought. *Assessment Report from the NOAA Drought Task Force Narrative Team*.
- Hong, S., Song, H.-R., Lutz, K., Kerstetter, R. A., Michael, T. P., and McClung, C. R. (2010). Type II protein arginine methyltransferase 5 (PRMT5) is required for circadian period determination in *Arabidopsis thaliana*. *Proc. Natl. Acad. Sci. U.S.A.* 107, 21211–21216. doi: 10.1073/pnas.1011987107
- Horváth, E., Szalai, G., and Janda, T. (2007). Induction of abiotic stress tolerance by salicylic acid signaling. *J. Plant Growth Regul.* 26, 290–300. doi: 10.1007/s00344-007-9017-4
- Hotta, C. T., Nishiyama, M. Y. Jr., and Souza, G. M. (2013). Circadian rhythms of sense and antisense transcription in sugarcane, a highly polyploid crop. *PLoS ONE* 8:e71847. doi: 10.1371/journal.pone.0071847
- Hsu, P. Y., and Harmer, S. L. (2014). Wheels within wheels: the plant circadian system. *Trends Plant Sci.* 19, 240–249. doi: 10.1016/j.tplants.2013.11.007
- Huang, D., Jaradat, M. R., Wu, W., Ambrose, S. J., Ross, A. R., Abrams, S. R., et al. (2007). Structural analogs of ABA reveal novel features of ABA perception and signaling in *Arabidopsis*. *Plant J. Cell Mol. Biol.* 50, 414–428. doi: 10.1111/j.1365-313X.2007.03056.x
- Huang, W., Pérez-García, P., Pokhilko, A., Millar, A. J., Antoshechkin, I., Riechmann, J. L., et al. (2012). Mapping the core of the *Arabidopsis* circadian clock defines the network structure of the oscillator. *Science* 336, 75–79. doi: 10.1126/science.1219075
- Hubbard, K. E., Nishimura, N., Hitomi, K., Getzoff, E. D., and Schroeder, J. I. (2010). Early abscisic acid signal transduction mechanisms: newly discovered components and newly emerging questions. *Genes Dev.* 24, 1695–1708. doi: 10.1101/gad.1953910
- Husaini, A. M., and Abidin, M. Z. (2008). Development of transgenic strawberry (*Fragaria × ananassa* Duch.) plants tolerant to salt stress. *Plant Sci.* 174, 446–455. doi: 10.1016/j.plantsci.2008.01.007

- Hussain, S. S., Kayani, M. A., and Amjad, M. (2011). Transcription factors as tools to engineer enhanced drought stress tolerance in plants. *Biotechnol. Prog.* 27, 297–306. doi: 10.1002/btpr.514
- Imaizumi, T., Schultz, T. F., Harmon, F. G., Ho, L. A., and Kay, S. A. (2005). FKF1 F-box protein mediates cyclic degradation of a repressor of CONSTANS in *Arabidopsis*. *Science* 309, 293–297. doi: 10.1126/science.1110586
- Ito, S., Matsushika, A., Yamada, H., Sato, S., Kato, T., Tabata, S., et al. (2003). Characterization of the APRR9 pseudo-response regulator belonging to the APRR1/TOC1 quintet in *Arabidopsis thaliana*. *Plant Cell Physiol.* 44, 1237–1245. doi: 10.1093/pcp/pcg136
- James, A. B., Syed, N. H., Bordage, S., Marshall, J., Nimmo, G. A., Jenkins, G. I., et al. (2012). Alternative splicing mediates responses of the *Arabidopsis* circadian clock to temperature changes. *Plant Cell* 24, 961–981. doi: 10.1105/tpc.111.093948
- Jarvis, P. G. (1976). The interpretation of the variations in leaf water potential and stomatal conductance found in canopies in the field. *Philos. Trans. R. Soc. B* 273, 593–610. doi: 10.1098/rstb.1976.0035
- Jones, M. A., Williams, B. A., Mcnicol, J., Simpson, C. G., Brown, J. W. S., and Harmer, S. L. (2012). Mutation of *Arabidopsis* spliceosomal timekeeper locus1 causes circadian clock defects. *Plant Cell* 24, 4066–4082. doi: 10.1105/tpc.112.104828
- Kazan, K. (2015). Diverse roles of jasmonates and ethylene in abiotic stress tolerance. *Trends Plant Sci.* 20, 219–229. doi: 10.1016/j.tplants.2015.02.001
- Kazan, K., and Manners, J. M. (2013). MYC2: the master in action. *Mol. Plant* 6, 686–703. doi: 10.1093/mp/sss128
- Kelly, J., Macgregor, D. R., Smith, R. W., Millar, A. J., Halliday, K. J., and Penfield, S. (2013). Model selection reveals control of cold signalling by evening-phased components of the plant circadian clock. *Plant J.* 76, 247–257. doi: 10.1111/tpj.12303
- Khan, S., Rowe, S., and Harmon, F. (2010). Coordination of the maize transcriptome by a conserved circadian clock. *BMC Plant Biol.* 10:126. doi: 10.1186/1471-2229-10-126
- Kielbowicz-Matuk, A., Rey, P., and Rorat, T. (2014). Interplay between circadian rhythm, time of the day and osmotic stress constraints in the regulation of the expression of a *Solanum* Double B-box gene. *Ann. Bot.* doi: 10.1093/aob/mct303
- Kilian, J., Whitehead, D., Horak, J., Wanke, D., Weinl, S., Batistic, O., et al. (2007). The AtGenExpress global stress expression data set: protocols, evaluation and model data analysis of UV-B light, drought and cold stress responses. *Plant J.* 50, 347–363. doi: 10.1111/j.1365-313X.2007.03052.x
- Kim, W.-Y., Ali, Z., Park, H. J., Park, S. J., Cha, J.-Y., Perez-Hormaeche, J., et al. (2013). Release of SOS2 kinase from sequestration with GIGANTEA determines salt tolerance in *Arabidopsis*. *Nat. Commun.* 4, 1352. doi: 10.1038/ncomms2357
- Kim, W. Y., Fujiwara, S., Suh, S.-S., Kim, J., Kim, Y., Han, L., et al. (2007). ZEITLUPE is a circadian photoreceptor stabilized by GIGANTEA in blue light. *Nature* 449, 356–360. doi: 10.1038/nature06132
- Kolmos, E., Chow, B. Y., Pruneda-Paz, J. L., and Kay, S. A. (2014). HsfB2b-mediated repression of PRR7 directs abiotic stress responses of the circadian clock. *Proc. Natl. Acad. Sci. U.S.A.* 111, 16172–16177. doi: 10.1073/pnas.1418483111
- Krasensky, J., and Jonak, C. (2012). Drought, salt, and temperature stress-induced metabolic rearrangements and regulatory networks. *J. Exp. Bot.* 63, 1593–1608. doi: 10.1093/jxb/err460
- Kusakina, J., Gould, P. D., and Hall, A. (2014). A fast circadian clock at high temperatures is a conserved feature across *Arabidopsis* accessions and likely to be important for vegetative yield. *Plant Cell Environ.* 37, 327–340. doi: 10.1111/pce.12152
- Kwon, Y.-J., Park, M.-J., Kim, S.-G., Baldwin, I., and Park, C.-M. (2014). Alternative splicing and nonsense-mediated decay of circadian clock genes under environmental stress conditions in *Arabidopsis*. *BMC Plant Biol.* 14:136. doi: 10.1186/1471-2229-14-136
- Lai, A. G., Doherty, C. J., Mueller-Roeber, B., Kay, S. A., Schippers, J. H. M., and Dijkwel, P. P. (2012). CIRCADIANT CLOCK-ASSOCIATED 1 regulates ROS homeostasis and oxidative stress responses. *Proc. Natl. Acad. Sci. U.S.A.* 109, 17129–17134. doi: 10.1073/pnas.1209148109
- Lawlor, D. W. (2012). Genetic engineering to improve plant performance under drought: physiological evaluation of achievements, limitations, and possibilities. *J. Exp. Bot.* 64, 83–108. doi: 10.1093/jxb/ers326
- Lawrimore, J. H., Menne, M. J., Gleason, B. E., Williams, C. N., Wuertz, D. B., Vose, R. S., et al. (2011). An overview of the Global Historical Climatology Network monthly mean temperature data set, version 3. *J. Geophys. Res. Atmos.* 116, D19121. doi: 10.1029/2011JD016187
- Lee, B.-H., Henderson, D. A., and Zhu, J.-K. (2005). The *Arabidopsis* cold-responsive transcriptome and its regulation by ICE1. *Plant Cell* 17, 3155–3175. doi: 10.1105/tpc.105.035568
- Lee, C.-M., and Thomashow, M. F. (2012). Photoperiodic regulation of the C-repeat binding factor (CBF) cold acclimation pathway and freezing tolerance in *Arabidopsis thaliana*. *Proc. Natl. Acad. Sci. U.S.A.* 109, 15054–15059. doi: 10.1073/pnas.1211295109
- Lee, K. H., Piao, H. L., Kim, H.-Y., Choi, S. M., Jiang, F., Hartung, W., et al. (2006). Activation of glucosidase via stress-induced polymerization rapidly increases active pools of abscisic acid. *Cell* 126, 1109–1120. doi: 10.1016/j.cell.2006.07.034
- Legnaioli, T., Cuevas, J., and Mas, P. (2009). TOC1 functions as a molecular switch connecting the circadian clock with plant responses to drought. *EMBO J.* 28, 3745–3757. doi: 10.1038/emboj.2009.297
- Li, Q.-B., and Guy, C. L. (2001). Evidence for non-circadian light/dark-regulated expression of Hsp70s in spinach leaves. *Plant Physiol.* 125, 1633–1642. doi: 10.1104/pp.125.4.1633
- Liu, T., Carlsson, J., Takeuchi, T., Newton, L., and Farré, E. M. (2013). Direct regulation of abiotic responses by the *Arabidopsis* circadian clock component PRR7. *Plant J.* 76, 101–114. doi: 10.1111/tpj.12276
- Lobell, D. B., and Gourdj, S. M. (2012). The influence of climate change on global crop productivity. *Plant Physiol.* 160, 1686–1697. doi: 10.1104/pp.112.208298
- Majercak, J., Sidote, D., and Hardin, P. E. (1999). How a circadian clock adapts to seasonal decreases in temperature and day length. *Neuron* 24, 219–230. doi: 10.1016/S0896-6273(00)80834-X
- Marcolino-Gomes, J., Rodrigues, F. A., Fuganti-Pagliarini, R., Bendix, C., Nakayama, T. J., Celaya, B., et al. (2014). Diurnal oscillations of soybean circadian clock and drought responsive genes. *PLoS ONE* 9:e86402. doi: 10.1371/journal.pone.0086402
- Marshall, A. (2014). Drought-tolerant varieties begin global march. *Nat. Biotech.* 32, 308–308. doi: 10.1038/nbt.2875
- Martino-Catt, S., and Ort, D. R. (1992). Low temperature interrupts circadian regulation of transcriptional activity in chilling-sensitive plants. *Proc. Natl. Acad. Sci. U.S.A.* 89, 3731–3735. doi: 10.1073/pnas.89.9.3731
- Mas, P., Kim, W. Y., Somers, D. E., and Kay, S. A. (2003). Targeted degradation of TOC1 by ZTL modulates circadian function in *Arabidopsis thaliana*. *Nature* 426, 567–570. doi: 10.1038/nature02163
- Matsushika, A., Makino, S., Kojima, M., and Mizuno, T. (2000). Circadian waves of expression of the APRR1/TOC1 family of pseudo-response regulators in *Arabidopsis thaliana*: insight into the plant circadian clock. *Plant Cell Physiol.* 41, 1002–1012. doi: 10.1093/pcp/pcd043
- McAdam, S. a. M., Brodribb, T. J., Ross, J. J., and Jordan, G. J. (2011). Augmentation of abscisic acid (ABA) levels by drought does not induce short-term stomatal sensitivity to CO₂ in two divergent conifer species. *J. Exp. Bot.* 62, 195–203. doi: 10.1093/jxb/erq260
- Michael, T. P., Mockler, T. C., Breton, G., Mcentee, C., Byer, A., Trout, J. D., et al. (2008). Network discovery pipeline elucidates conserved time-of-day-specific cis-regulatory modules. *PLoS Genet.* 4:e14. doi: 10.1371/journal.pgen.0040014
- Mikkelsen, M., D., and Thomashow, M. F. (2009). A role for circadian evening elements in cold-regulated gene expression in *Arabidopsis*. *Plant J.* 60, 328–339. doi: 10.1111/j.1365-313X.2009.03957.x
- Mittler, R., and Blumwald, E. (2015). The roles of ROS and ABA in systemic acquired acclimation. *Plant Cell* 27, 64–70. doi: 10.1105/tpc.114.133090
- Miura, K., and Tada, Y. (2014). Regulation of water, salinity, and cold stress responses by salicylic acid. *Front. Plant Sci.* 5:04. doi: 10.3389/fpls.2014.00004
- Miyazaki, Y., Abe, H., Takase, T., Kobayashi, M., and Kiyosue, T. (2015). Overexpression of LOV KELCH PROTEIN 2 confers dehydration tolerance and is associated with enhanced expression of dehydration-inducible genes in *Arabidopsis thaliana*. *Plant Cell Rep.* 34, 843–852. doi: 10.1007/s00299-015-1746-4
- Mizoguchi, T., Wheatley, K., Hanzawa, Y., Wright, L., Mizoguchi, M., Song, H. R., et al. (2002). LHY and CCA1 are partially redundant genes required to maintain circadian rhythms in *Arabidopsis*. *Dev. Cell* 2, 629–641. doi: 10.1016/S1534-5807(02)00170-3
- Mizuno, T., Nomoto, Y., Oka, H., Kitayama, M., Takeuchi, A., Tsubouchi, M., et al. (2014). Ambient temperature signal feeds into the circadian clock transcriptional circuitry through the EC night-time repressor in *Arabidopsis thaliana*. *Plant Cell Physiol.* 55, 958–976. doi: 10.1093/pcp/pcu030

- Mockler, T.C., Michael, T.P., Priest, H.D., Shen, R., Sullivan, C.M., Givan, S.A., et al. (2007). The Diurnal project: diurnal and circadian expression profiling, model-based pattern matching, and promoter analysis. *Cold Spring Harbor Symposia on Quantitative Biology: Clocks and Rhythms*. 72. doi: 10.1101/sqb.2007.72.006
- Mugford, S. T., Fernandez, O., Brinton, J., Flis, A., Krohn, N., Encke, B., et al. (2014). Regulatory properties of ADP glucose pyrophosphorylase are required for adjustment of leaf starch synthesis in different photoperiods. *Plant Physiol.* 166, 1733–1747. doi: 10.1104/pp.114.247759
- Nagel, D. H., Pruneda-Paz, J. L., and Kay, S. A. (2014). FBH1 affects warm temperature responses in the *Arabidopsis* circadian clock. *Proc. Natl. Acad. Sci. U.S.A.* 111, 14595–14600. doi: 10.1073/pnas.1416666111
- Nakamichi, N., Kiba, T., Henriques, R., Mizuno, T., Chua, N. H., and Sakakibara, H. (2010). PSEUDO-RESPONSE REGULATORS 9,7 and 5 are transcriptional repressors in the *Arabidopsis* circadian clock. *Plant Cell* 22, 594–605. doi: 10.1105/tpc.109.072892
- Nakamichi, N., Kiba, T., Kamioka, M., Suzuki, T., Yamashino, T., Higashiyama, T., et al. (2012). Transcriptional repressor PRR5 directly regulates clock-output pathways. *Proc. Natl. Acad. Sci. U.S.A.* 109, 17123–17128. doi: 10.1073/pnas.1205156109
- Nakamichi, N., Kita, M., Ito, S., Yamashino, T., and Mizuno, T. (2005). PSEUDO-RESPONSE REGULATORS, PRR9, PRR7 and PRR5, together play essential roles close to the circadian clock of *Arabidopsis thaliana*. *Plant Cell Physiol.* 46, 686–698. doi: 10.1093/pcp/pci086
- Nakamichi, N., Kusano, M., Fukushima, A., Kita, M., Ito, S., Yamashino, T., et al. (2009). Transcript profiling of an *Arabidopsis* PSEUDO RESPONSE REGULATOR arrhythmic triple mutant reveals a role for the circadian clock in cold stress response. *Plant Cell Physiol.* 50, 447–462. doi: 10.1093/pcp/pcp004
- Nemali, K. S., Bonin, C., Dohleman, F. G., Stephens, M., Reeves, W. R., Nelson, D. E., et al. (2014). Physiological responses related to increased grain yield under drought in the first biotechnology-derived drought-tolerant maize. *Plant Cell Environ.* 38, 1866–1880. doi: 10.1111/pce.12446
- Novakova, M., Motyka, V., Dobrev, P. I. M., Malbeck, J., Gaudinova, A., and Vankova, R. (2005). Diurnal variation of cytokinin, auxin and abscisic acid levels in tobacco leaves. *J. Exp. Bot.* 56, 2877–2883. doi: 10.1093/jxb/eri282
- Nusinow, D. A., Helfer, A., Hamilton, E. E., King, J. J., Imaizumi, T., Schultz, T. F., et al. (2011). The ELF4-ELF3-LUX complex links the circadian clock to diurnal control of hypocotyl growth. *Nature* 475, 398–402. doi: 10.1038/nature10182
- Onai, K., and Ishiura, M. (2005). PHYTOCLOCK 1 encoding a novel GARP protein essential for the *Arabidopsis* circadian clock. *Genes Cells* 10, 963–972. doi: 10.1111/j.1365-2443.2005.00892.x
- Parkhi, V., Kumar, V., Sunilkumar, G., Campbell, L., Singh, N., and Rathore, K. (2009). Expression of apoplastically secreted tobacco osmotin in cotton confers drought tolerance. *Mol. Breed.* 23, 625–639. doi: 10.1007/s11032-009-9261-3
- Patade, V. Y., Khatri, D., Kumari, M., Grover, A., Mohan Gupta, S., and Ahmed, Z. (2013). Cold tolerance in Osmotin transgenic tomato (*Solanum lycopersicum* L.) is associated with modulation in transcript abundance of stress responsive genes. *Springerplus* 2, 117. doi: 10.1186/2193-1801-2-117
- Pearce, R. S. (2001). Plant freezing and damage. *Ann. Bot.* 87, 417–424. doi: 10.1006/anbo.2000.1352
- Picot, E., Krusche, P., Tiskin, A., Carré, I., and Ott, S. (2010). Evolutionary analysis of regulatory sequences (EARS) in plants. *Plant J.* 64, 165–176. doi: 10.1111/j.1365-313X.2010.04314.x
- Pokhilko, A., Fernandez, A. P., Edwards, K. D., Southern, M. M., Halliday, K. J., and Millar, A. J. (2012). The clock gene circuit in *Arabidopsis* includes a repressilator with additional feedback loops. *Mol. Syst. Biol.* 8, 574. doi: 10.1038/msb.2012.6
- Rienth, M., Torregrosa, L., Luchaire, N., Chatbanyong, R., Lecourieux, D., Kelly, M., et al. (2014). Day and night heat stress trigger different transcriptomic responses in green and ripening grapevine (*Vitis vinifera*) fruit. *BMC Plant Biol.* 14:108. doi: 10.1186/1471-2229-14-108
- Rikin, A., Dillwith, J. W., and Bergman, D. K. (1993). Correlation between the circadian rhythm of resistance to extreme temperatures and changes in fatty acid composition in cotton seedlings. *Plant Physiol.* 101, 31–36.
- Salome, P. A., Weigel, D., and McClung, C. R. (2010). The role of the *Arabidopsis* morning loop components CCA1, LHY, PRR7, and PRR9 in temperature compensation. *Plant Cell* 22, 3650–3661. doi: 10.1105/tpc.110.079087
- Salvucci, M. E., and Crafts-Brandner, S. J. (2004). Inhibition of photosynthesis by heat stress: the activation state of Rubisco as a limiting factor in photosynthesis. *Physiol. Plant.* 120, 179–186. doi: 10.1111/j.0031-9317.2004.0173.x
- Sanchez, S. E., Petrillo, E., Beckwith, E. J., Zhang, X., Rugnone, M. L., Hernando, C. E., et al. (2010). A methyl transferase links the circadian clock to the regulation of alternative splicing. *Nature* 468, 112–116. doi: 10.1038/nature09470
- Schaffer, R., Ramsay, N., Samach, A., Corden, S., Putterill, J., Carré, I. A., et al. (1998). The late elongated hypocotyl mutation of *Arabidopsis* disrupts circadian rhythms and the photoperiodic control of flowering. *Cell* 93, 1219–1229. doi: 10.1016/S0092-8674(00)81465-8
- Schultz, T. F., Kiyosue, T., Yanovsky, M., Wada, M., and Kay, S. A. (2001). A role for LKP2 in the circadian clock of *Arabidopsis*. *Plant Cell* 13, 2659–2670. doi: 10.1105/tpc.010332
- Scialdone, A., Mugford, S. T., Feike, D., Skeffington, A., Borrill, P., Graf, A., et al. (2013). *Arabidopsis* plants perform arithmetic division to prevent starvation at night. *Elife* 2, e00669. doi: 10.7554/eLife.00669
- Seo, P. J., and Mas, P. (2014). Multiple layers of posttranslational regulation refine circadian clock activity in *Arabidopsis*. *Plant Cell* 26, 79–87. doi: 10.1105/tpc.113.119842
- Seo, P. J., Park, M.-J., Lim, M.-H., Kim, S.-G., Lee, M., Baldwin, I. T., et al. (2012). A self-regulatory circuit of CIRCADIAN CLOCK-ASSOCIATED1 underlies the circadian clock regulation of temperature responses in *Arabidopsis*. *Plant Cell* 24, 2427–2442. doi: 10.1105/tpc.112.098723
- Sharkey, T. D. (2005). Effects of moderate heat stress on photosynthesis: importance of thylakoid reactions, rubisco deactivation, reactive oxygen species, and thermotolerance provided by isoprene. *Plant Cell Environ.* 28, 269–277. doi: 10.1111/j.1365-3040.2005.01324.x
- Shin, J., Heidrich, K., Sanchez-Villarreal, A., Parker, J. E., and Davis, S. J. (2012). TIME FOR COFFEE represses accumulation of the MYC2 transcription factor to provide time-of-day regulation of jasmonate signaling in *Arabidopsis*. *Plant Cell* 24, 2470–2482. doi: 10.1105/tpc.111.095430
- Somers, D. E., Devlin, P. F., and Kay, S. A. (1998). Phytochromes and cryptochromes in the entrainment of the *Arabidopsis* circadian clock. *Science* 282, 1488–1490. doi: 10.1126/science.282.5393.1488
- Spensley, M., Kim, J.-Y., Picot, E., Reid, J., Ott, S., Helliwell, C., et al. (2009). Evolutionarily conserved regulatory motifs in the promoter of the *Arabidopsis* clock gene LATE ELONGATED HYPOCOTYL. *Plant Cell* 21, 2606–2623. doi: 10.1105/tpc.109.069898
- Spoel, S. H., Johnson, J. S., and Dong, X. (2007). Regulation of tradeoffs between plant defenses against pathogens with different lifestyles. *Proc. Natl. Acad. Sci. U.S.A.* 104, 18842–18847. doi: 10.1073/pnas.0708139104
- Steponkus, P. L., and Webb, M. S. (1992). “Freeze-induced dehydration and membrane destabilization in plants,” in *Water and Life*, eds G. Somero, C. Osmond, and C. Bolis (Berlin: Springer), 338–362.
- Strayer, C., Oyama, T., Schultz, T. F., Raman, R., Somers, D. E., Mas, P., et al. (2000). Cloning of the *Arabidopsis* clock gene TOC1, an autoregulatory response regulator homolog. *Science* 289, 768–771. doi: 10.1126/science.289.5480.768
- Subramanyam, K., Sailaja, K. V., Subramanyam, K., Rao, D. M., and Lakshmidivi, K. (2010). Ectopic expression of an osmotin gene leads to enhanced salt tolerance in transgenic chilli pepper (*Capsicum annum* L.). *Plant Cell Tissue Organ Cult.* 105, 181–192. doi: 10.1007/s11240-010-9850-1
- Suzuki, N., Koussevitzky, S., Mittler, R. O. N., and Miller, G. A. D. (2012). ROS and redox signalling in the response of plants to abiotic stress. *Plant Cell Environ.* 35, 259–270. doi: 10.1111/j.1365-3040.2011.02336.x
- Thain, S. C., Vandenbussche, F., Laarhoven, L. J., Dowson-Day, M. J., Wang, Z. Y., Tobin, E. M., et al. (2004). Circadian rhythms of ethylene emission in *Arabidopsis*. *Plant Physiol.* 136, 3751–3761. doi: 10.1104/pp.104.042523
- Thomashow, M. F. (2010). Molecular basis of plant cold acclimation: insights gained from studying the CBF cold response pathway. *Plant Physiol.* 154, 571–577. doi: 10.1104/pp.110.161794
- Thompson, A., Jackson, A., Parker, R., Morpeth, D., Burbidge, A., and Taylor, I. (2000). Abscisic acid biosynthesis in tomato: regulation of zeaxanthin epoxidase and 9-cis-epoxycarotenoid dioxygenase mRNAs by light/dark cycles, water stress and abscisic acid. *Plant Mol. Biol.* 42, 833–845. doi: 10.1023/A:1006448428401
- Wahid, A., Gelani, S., Ashraf, M., and Foolad, M. R. (2007). Heat tolerance in plants: an overview. *Environ. Exp. Bot.* 61, 199–223. doi: 10.1016/j.envexpbot.2007.05.011
- Waltz, E. (2014). Beating the heat. *Nat. Biotechnol.* 32, 610–613. doi: 10.1038/nbt.2948
- Wang, G., Zhang, C., Battle, S. L., and Lu, H. (2014). The phosphate transporter PHT4;1 is a salicylic acid regulator likely controlled by the circadian clock protein CCA1. *Front. Plant Sci.* 5:701. doi: 10.3389/fpls.2014.00701

- Wang, Z.-Y., and Tobin, E. M. (1998). Constitutive expression of the CIRCADIAN CLOCK ASSOCIATED (CCA1) gene disrupts circadian rhythms and suppresses its own expression. *Cell* 93, 1207–1217. doi: 10.1016/S0092-8674(00)81464-6
- Wasternack, C., and Hause, B. (2013). Jasmonates: biosynthesis, perception, signal transduction and action in plant stress response, growth and development. An update to the 2007 review in *Annals of Botany*. *Ann. Bot.* 111, 1021–1058. doi: 10.1093/aob/mct067
- Webb, A. A. R., and Satake, A. (2015). Understanding circadian regulation of carbohydrate metabolism in *Arabidopsis* using mathematical models. *Plant Cell Physiol.* 56, 586–593. doi: 10.1093/pcp/pcv033
- Wilkins, O., Brautigam, K., and Campbell, M. M. (2010). Time of day shapes *Arabidopsis* drought transcriptomes. *Plant J.* 63, 715–727. doi: 10.1111/j.1365-313X.2010.04274.x
- Yamamoto, Y., Sato, E., Shimizu, T., Nakamichi, N., Sato, S., Kato, T., et al. (2003). Comparative genetic studies on the APRR5 and APRR7 genes belonging to the APRR1/TOC1 quintet implicated in circadian rhythm, control of flowering time, and early photomorphogenesis. *Plant Cell Physiol.* 44, 1119–1130. doi: 10.1093/pcp/pcg148
- Yamazaki, T., Kawamura, Y., and Uemura, M. (2008). Cryobehavior of the plasma membrane in protoplasts isolated from cold-acclimated *Arabidopsis* leaves is related to surface area regulation. *Plant Cell Physiol.* 49, 944–957. doi: 10.1093/pcp/pcn068
- Yamazaki, T., Kawamura, Y., and Uemura, M. (2009). Extracellular freezing-induced mechanical stress and surface area regulation on the plasma membrane in cold-acclimated plant cells. *Plant Signal. Behav.* 4, 231–233. doi: 10.4161/psb.4.3.7911
- Yang, Y., Qi, M., and Mei, C. (2004). Endogenous salicylic acid protects rice plants from oxidative damage caused by aging as well as biotic and abiotic stress. *Plant J.* 40, 909–919. doi: 10.1111/j.1365-313X.2004.02267.x
- Yanovsky, M. J., and Kay, S. A. (2002). Molecular basis of seasonal time measurement in *Arabidopsis*. *Nature* 419, 308–312. doi: 10.1038/nature00996
- Young, M. W., and Kay, S. A. (2001). Time Zones: a comparative genetics of circadian clocks. *Nat. Rev. Gen.* 2, 702–715. doi: 10.1038/35088576
- Zhang, C., Xie, Q., Anderson, R. G., Ng, G., Seitz, N. C., Peterson, T., et al. (2013). Crosstalk between the circadian clock and innate immunity in *Arabidopsis*. *PLoS Pathog.* 9:e1003370. doi: 10.1371/journal.ppat.1003370

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2015 Grundy, Stoker and Carré. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) or licensor are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.