Article
Suppression of Rice Cryptochrome 1b Decreases Both Melatonin and Expression of Brassinosteroid Biosynthetic Genes Resulting in Salt Tolerance

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Abstract: We investigated the relationship between the blue-light photoreceptor cryptochrome (CRY) and melatonin biosynthesis by generating RNA interference (RNAi) transgenic rice plants that suppress the cryptochrome 1b gene (CRY1b). The resulting CRY1b RNAi rice lines expressed less CRY1b mRNA, but not CRY1a or CRY2 mRNA, suggesting that the suppression is specific to CRY1b. The growth of CRY1b RNAi rice seedlings was enhanced under blue light compared to wild-type growth, providing phenotypic evidence for impaired CRY function. When these CRY1b RNAi rice plants were challenged with cadmium to induce melatonin, wild-type plants produced 100 ng/g fresh weight (FW) melatonin, whereas CRY1b RNAi lines produced 60 ng/g FW melatonin on average, indicating that melatonin biosynthesis requires the CRY photoreceptor. Due to possible feedback regulation, the expression of melatonin biosynthesis genes such as T5H, SNAT1, SNAT2, and COMT was elevated in the CRY1b RNAi lines compared to the wild-type plants. In addition, laminar angles decreased in the CRY1b RNAi lines via the suppression of brassinosteroid (BR) biosynthesis genes such as DWARF. The main cause of the BR decrease in the CRY1b RNAi lines seems to be the suppression of CRY rather than decreased melatonin because the melatonin decrease suppressed DWARF4 rather than DWARF.

Keywords: cadmium; blue-light photoreceptors; brassinosteroids; CRY1b; melatonin; transgenic rice

1. Introduction

Light-dependent melatonin synthesis occurs in some plant species, whereas in other plants, melatonin exhibits a nocturnal increase similar to that in animals [1–4]. Melatonin is broadly implicated in many physiological activities in plants, where it mostly serves as a potent free radical scavenger, signal molecule, or hormone via the mitogen-activated protein kinase (MAPK) signaling pathway [5–9]. Although a plant melatonin receptor was recently reported [10], its integrity remains controversial and requires further study [11].

The major known functions of melatonin are plant defense responses against both biotic and abiotic stresses [12,13], plant growth, and reproduction [7,14]. Briefly, melatonin is closely involved in conferring tolerance against endoplasmic reticulum stress [15,16], salt [17,18], pathogens [19,20], high temperatures [21,22], high light stress [23], and other stresses [24]. Melatonin also plays roles in plant growth and reproduction, including the synthesis of secondary metabolites [25,26], germination [27], seed longevity [28], hormone synthesis [29,30], fruit yield [31], and other processes [32].

The melatonin biosynthesis pathways have been well-characterized with the successful cloning of relevant biosynthesis genes [8]. It is clear that melatonin induction is dependent on light based on experiments in which rice plants were challenged with cadmium, which is the best inducer of melatonin [2]. The pivotal gene responsible for melatonin induction under light is that encoding tryptophan decarboxylase (TDC), the first committed step enzyme in melatonin biosynthesis, which converts tryptophan into...
tryptamine. The optimal light for the maximum induction of melatonin was found to be a combination of red and blue light, rather than red or blue alone; far-red light is unable to induce melatonin in response to cadmium treatment in rice, indicating that melatonin synthesis is predominantly associated with photosynthesis rather than with other photoreceptors such as phytochromes [33].

Light is indispensable for plant growth via photosynthesis, during which mainly red (720 nm) and blue (450 nm) wavelengths are absorbed via photosynthetic pigments, including chlorophyll [34]. Light is also an important signal that influences plant life and death from germination to reproduction. Plants have evolved specific photoreceptors that respond to different light wavelengths. For instance, phytochromes absorb red (600–700 nm) and far-red (700–750 nm) light as part of their biological functions, whereas phototropins and cryptochromes (CRY) sense ultraviolet A/blue light (390–500 nm). The first molecular genetic analysis of the possible involvement of photoreceptors in melatonin synthesis was recently reported, in which rice phytochrome mutants exhibited decreased melatonin synthesis compared to wild-type plants, confirming a previous report that melatonin synthesis requires light [33]. However, the possible involvement of CRY in melatonin biosynthesis in plants remains unclear [35]. Here, we generated transgenic rice plants that suppressed rice CRY1b to investigate if CRY are also involved in melatonin synthesis in rice. The rice genome contains a small CRY gene family that includes CRY1a (AB024337), CRY1b (AB098568), and CRY2 (AK065669). CRY1a exhibits greater homology with CRY1b (75% identity), whereas CRY1a and CRY1b exhibit less homology with CRY2, with amino-acid sequence identities of 46% and 47%, respectively [36]. Of the three CRY genes, CRY1b was reported to be associated with blue light-dependent phenotypes in Arabidopsis and rice [37]. Therefore, we chose rice CRY1b to develop transgenic RNAi rice plants to examine whether CRY is involved in melatonin biosynthesis (Figure 1a). Due to the high homology between CRY1a and CRY1b, we selected a C-terminal portion of CRY1b that exhibits greater sequence diversity from that in CRY1a for RNA silencing.

Figure 1. The structure of binary vector pTCK303 and the generation of CRY1b-suppressed transgenic rice plants. (a) Schematic diagram of the pTCK303:CRY1b binary vector. (b) Results of quantitative real-time polymerase chain reaction (PCR) analyses. (c) Results of reverse transcription-PCR analyses of the wild-type and transgenic (T2) lines. Rice seeds were germinated in half-strength Murashige and Skoog (MS) medium and grown for 8 days at 28 °C under a 14-h light/10-h dark cycle. Total RNA was extracted from these seedlings. CRY1b, Oryza sativa cryptochrome 1b; Ubi-P, maize ubiquitin promoter; HPT, gene encoding hygromycin phosphotransferase. Asterisks (***)) indicate significant differences from the wild type (p < 0.001). GenBank accession numbers are AB024337 (CRY1a), AB098568 (CRY1b), AK065669 (CRY2), and Os03g13170 (UBQ5). Numbers in parentheses indicate the numbers of PCR cycles.
2. Results and Discussion

2.1. Generation of Transgenic Rice Plants Suppressing Rice CRY1b

From four independent T1 lines, we selected three homozygous T2 transgenic lines (Figure 1b,c). CRY1b was significantly suppressed in the three CRY1b RNAi lines, whereas CRY1a and CRY2 mRNA levels were comparable to those in the wild type according to qRT-PCR and semiquantitative RT-PCR analyses. These data indicated the successful, specific suppression of CRY1b using RNAi. The lengths and second leaf sheath lengths of 8-day-old wild-type and CRY1b RNAi seedlings grown in half-strength Murashige and Skoog (MS) medium under a 12-h light/12-h dark regime with cool fluorescent light were similar (Figure 2a–c). However, seedling length was longer in the CRY1b RNAi lines than in the wild type when seedlings were grown under blue light, as was the length of the second leaf sheath (Figure 2d–f). Notably, wild-type seedling length decreased under blue light versus cool fluorescent light because blue light inhibits seedling growth in monocots, including rice [37,38]. Consistent with this, our transgenic CRY1b RNAi seedlings exhibited significantly reduced inhibition of seedling growth and second leaf sheath length, confirming that our CRY1b RNAi rice plants had a functional defect in the blue-light photoreceptor (CRY). In contrast to results obtained for CRY1b RNAi, the overexpression of CRY1b induced an increased responsiveness to blue light, resulting in severe seedling growth inhibition compared to the wild type [36,37].

2.2. Melatonin Levels in CRY1b RNAi Rice Seedlings

To determine whether the blue-light photoreceptor (CRY) was associated with melatonin biosynthesis, 8-day-old rice seedlings were challenged with cadmium to induce melatonin. As shown in Figure 3, melatonin levels decreased two-fold in CRY1b RNAi rice
lines 2 and 3 and by 20% in line 1 compared to those in the wild type (Figure 3c). A drastic decrease in N-acetylserotonin (NAS) levels was observed in all CRY1b RNAi lines, with NAS contents 10-fold less in RNAi lines (166 ng/g fresh weight (FW)) than in the wild type (1750 ng/g FW) (Figure 3b). Conversely, serotonin levels in the CRY1b RNAi lines were indistinguishable from those in the wild type (Figure 3a). These data indicate that the blue-light photoreceptor is involved in melatonin biosynthesis when rice seedlings are challenged with cadmium. Next, we measured the expression of melatonin biosynthesis gene mRNA in 7-day-old rice seedlings in the absence of cadmium. TDC is the first enzyme to participate in melatonin biosynthesis, followed by tryptamine 5-hydroxylase (T5H), which synthesizes serotonin from tryptamine. Serotonin is then acetylated into NAS by serotonin N-acetyltransferase (SNAT), which has two isogenes in the rice genome—SNAT1 and SNAT2. The last enzyme in melatonin synthesis is NAS O-methyltransferase (ASMT), which catalyzes the conversion of NAS into melatonin. The caffeic acid O-methyltransferase (COMT) exhibiting ASMT enzyme activity plays pivotal roles in melatonin synthesis in rice [39]. The expression of the melatonin biosynthesis genes T5H, SNAT1, SNAT2, and COMT was all elevated in the CRY1b RNAi lines, whereas that of TDC1 was not (Figure 4). The high levels of melatonin biosynthesis gene expression in the CRY1b RNAi lines suggest that feedback regulation compensates for low melatonin levels.

![Figure 3](image-url)

**Figure 3.** Serotonin, N-acetylserotonin, and melatonin contents in WT and CRY1b RNAi lines. (a) Serotonin, (b) N-acetylserotonin (NAS), and (c) melatonin levels in response to cadmium treatment under cool fluorescent light. Eight-day-old rice seedlings were challenged rhizosperically with 0.5 mM CdCl₂ for 3 days under cool fluorescent light at a rate of 60 μmol m⁻² s⁻¹. Values are presented as the means ± standard deviations (SDs) of three independent experiments. FW, fresh weight. *** p < 0.001, Tukey’s HSD post-hoc test.

![Figure 4](image-url)

**Figure 4.** Quantitative real-time PCR analyses of melatonin biosynthesis genes. TDC1, tryptophan decarboxylase 1; T5H, tryptamine 5-hydroxylase; SNAT1, serotonin N-acetyltransferase 1; SNAT2, serotonin N-acetyltransferase 2; COMT, caffeic acid O-methyltransferase. * p < 0.05 and *** p < 0.001 vs. the WT. GenBank accession numbers are AK069031 (TDC1), AK071599 (T5H), AK059369 (SNAT1), AK068156 (SNAT2), AK064768 (COMT), and Os03g13170 (UBQ5).
2.3. Decreased Leaf Angle in the CRY1b RNAi Rice Seedlings

Blue light promotes leaf bending by upregulating brassinosteroid (BR) biosynthesis genes in rice [40]. Similarly, the overexpression of CRY genes increases the leaf angle in rice, implying a positive relationship between CRY and leaf angle [36]. Additionally, melatonin is positively associated with the leaf angle in rice by regulating BR levels [29]. To investigate the role of CRY1b in determining leaf angle and BR-related gene expression, lamina angles of the second leaves of 10-day-old seedlings grown in soil pots were measured. As shown in Figure 5, the CRY1b RNAi lines had smaller leaf angles than the wild type. To examine the possible contribution of BR-related genes to the decreased leaf angles in the CRY1b RNAi lines, we quantified the expression of various BR-related genes. **DWARF, DWARF4, CPD (CONSTITUTIVE PHOTOMORPHOGENIC DWARF), and DET2 (DEETIOLATED2)** are BR biosynthesis genes. DWARF converts 6-deoxo-catasterone into catasterone via a C6 oxidation reaction, whereas DWARF4 converts 6-oxo-campestanol into cathasterone via C22 hydroxylation. **CPD and DET2** encode C3 oxidase and 5-reductase, respectively [40]. BR-responsive genes include **BZR1 (BRASSINAZOLE RESISTANT1; a BR receptor), TXR3 (XYLOGLUCAN ENDOTRANSGLYCOSYLASE RELATED), and BRI1 (BRASSINOSTEROID-INSENSITIVE 1; a transcription factor for BR biosynthesis).** TXR3 is a BR-inducible gene encoding a cell wall-loosening enzyme. All BR biosynthesis genes except for **DWARF4** were downregulated in the CRY1b RNAi lines compared to the wild type, suggestive of reduced BR synthesis. Our data implicate that CRY suppression leads to BR suppression in rice. Intriguingly, the decrease in melatonin caused by the suppression of **SNAT2** led to a decrease in BR via the suppression of DWARF4 [29]. Collectively, we attribute the BR decrease in the CRY1b RNAi lines mainly to the downregulation of CRY, rather than a decrease in melatonin caused by the suppression of CRY1b. The expression of **BZR1** also decreased in the CRY1b RNAi lines relative to the wild type, indicating that BR levels decreased in the CRY1b RNAi lines. Contrary to the results observed for BR biosynthesis genes, BR-responsive genes such as **TXR3** and **BRII** were significantly upregulated in the CRY1b RNAi lines relative to the wild type, suggesting that these genes are regulated by feedback from reduced levels of endogenous BR.

![Figure 5](image-url)

**Figure 5.** Lamina joint angles and transcript induction profiles of brassinosteroid (BR)-related genes in WT and CRY1b RNAi (1–3) rice plants. (a) Photograph of representative lamina joints in plants grown for 10 days in soil pots. (b) Lamina joint angles in WT and transgenic rice seedlings. (c) Quantitative real-time PCR analysis of the expression of various BR-related genes in WT and CRY1b RNAi plants. GenBank accession numbers are Os03g0602300 (**DWARF**), Os03g12660 (**DWARF4**), Os11g0143200 (**CPD1**), Os12g0139300 (**CPD2**), Os01g0851600 (**DET2**), Os07g39220 (**BZR1**), AP005859 (**TXR3**), AK101085 (**BRII**), and Os03g13170 (**UBQ5**). Lines 1–3 are RNAi rice plants with downregulated CRY1b genes. * p < 0.05, *** p < 0.001, Tukey’s HSD post-hoc test.
2.4. CRY1b RNAi Rice Exhibits Salt Tolerance, Possibly via a Decrease in BR

Exposure to blue light may increase BR synthesis, possibly via CRY in *Arabidopsis* and rice [40,41]. In fact, the overexpression of CRY1a conferred hypersensitivity against salt in *Arabidopsis*, possibly through ABA (abscisic acid) signaling [41]. BR levels have been shown to be negatively associated with abiotic stresses in plants [42–44]. Therefore, a decrease in endogenous BR levels or signaling can lead to abiotic stress tolerance in plants via the ABA and BR signaling pathways [29,41,44]. To test whether the CRY1b RNAi lines showing the reduced expression of BR-biosynthetic genes than the wild type confer abiotic tolerance or not, the CRY1b RNAi lines were challenged with salt, and their responses were monitored. As shown in Figure 6, CRY1b RNAi rice exhibited a salt-tolerant phenotype, reflected by reduced production of malondialdehyde (MDA) compared to the wild type. Our data strongly suggest that CRY1b RNAi rice produces less BR than the wild type, which accounts for the enhanced salt tolerance.

**Figure 6.** Effects of salt treatment on CRY1b RNAi rice plants. (a) Photograph of representative phenotypes after salt treatment. (b) Malondialdehyde (MDA) levels in WT and transgenic rice plants. (c) Proposed model of CRY-mediated salt tolerance. Values are presented as the means ± SDs of three independent experiments. FW, fresh weight; d, day. ***p < 0.001, Tukey’s HSD post-hoc test.

3. Materials and Methods

3.1. Plant Growth Conditions

Rice (*Oryza sativa* cv. Dongjin) seeds were sterilized and grown in soil or half-strength Murashige and Skoog (MS) under cool daylight fluorescent lamps (60 μmol m⁻² s⁻¹) (Philips, Amsterdam, Netherlands) in a 14-h light/10-h dark photoperiod or blue light (15 μmol m⁻² s⁻¹) (STECH LED, Gyeonggi-do, Korea) at 28 °C/24 °C (day/night). The angles of the lamina joint were measured in the second leaf of 10 d-old rice seedlings grown in soil. For tolerance assay in response to salt stress, 10-d old seedlings were exposed to 200 mM NaCl for 6 days.

3.2. Generation of CRY1b Suppression Transgenic Rice by RNA Interference (RNAi)

The pTCK303 binary vector was used to suppress rice CRY1b gene expression as previously described [45]. Briefly, a C-terminal 288 bp CRY1b cDNA fragment was amplified by PCR using the following primer set: CRY1b-F 5'-ACT AGT GAA AAT TTCCGTACCACT-3' (SptI site underlined) and CRY1b-R 5'-GAG CTC GTG GGA TAA TTG ACT CCA-3' (SacI site underlined), with the cDNA templates synthesized from the total RNA from rice seedlings. The CRY1b PCR product was first subcloned into the T&A cloning vector (T&A:CRY1b; RBC Bioscience, New Taipei City, Taiwan) for further cloning experiments. From the T&A:CRY1b plasmid, the antisense CRY1b insert was digested by SacI and SpeI, whereas the sense CRY1b insert was digested by KpnI and BamHI in the T&A:CRY1b vector. The antisense CRY1b fragment was first ligated into the pTCK303 vector. Thereafter,
the sense fragment of the CRY1b insert was sequentially ligated into the pTCK303 vector plasmid harboring the corresponding CRY1b antisense fragment (Figure 1a). The resulting pTCK303:CRY1b RNAi binary vector was transformed into A. tumefaciens LBA4404, and into rice (Oryza sativa cv. Dongjin), as previously described [46,47]. In brief, in order to generate transgenic rice plants, scutellum-derived calli from rice (O. sativa cv. Dongjin) were co-cultured with Agrobacterium harboring the pTCK303:CRY1b binary vector for 3 days, during which T (transfer)-DNA is incorporated into the rice genome. Thereafter, rice calli were screened on N6 selection medium containing 2 mg/L 2,4-D and 50 mg/L hygromycin for 30 days in the dark to generate transgenic calli, and transferred to regeneration media containing 3 mg/L BAP (6-benzylaminopurine), 1 mg/L NAA (1-naphthalene acetic acid), and 50 mg/L hygromycin for shoot and root development. The resulting rice seedlings were grown in a paddy field for obtaining T1 seeds.

3.3. Cadmium Treatment

Eight-day old seedlings were incubated in water containing 0.5 mM CdCl$_2$ (Sigma-Aldrich, St. Louis, MO) under cool daylight fluorescent lamps (60 µmol m$^{-2}$ s$^{-1}$) (Philips, Amsterdam, The Netherlands) for 3 days at 28 °C. The leaves and stems were harvested for further analyses.

3.4. Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR) Analysis

Total RNA of rice plants was isolated using a NucleoSpin RNA Plant Kit (Macherey-Nagel, Düren, Germany). First-strand cDNA was synthesized from 2 µg of total RNA using MG MMLV Reverse Transcriptase (MGmed, Inc., Seoul, Korea) and an oligo dT$_{18}$ primer at 42 °C for 1 h. qRT-PCR was performed in a Mic qPCR Cycler system (Biomolecular Systems, Queensland, VIC, Australia) with specific primers and the Luna Universal qPCR Master Mix (New England Biolabs, Ipswich, MA, USA), as described previously. The expression of genes was analyzed using Mic’s RQ software (Biomolecular Systems) and normalized to ACT1. Semi-quantitative RT-PCR was performed as described previously [2].

3.5. Quantification of Serotonin, N-acetylserotonin, and Melatonin

Frozen samples (0.1 g) were pulverized to a powder in liquid nitrogen using the TissueLyser II (Qagen, Tokyo, Japan). For serotonin and N-acetylserotonin, rice samples were extracted with 1 mL of methanol for 1 h at room temperature. The extracts were centrifuged for 10 min at 12,000 × g, and the supernatants (20 µL) were subjected to high performance liquid chromatography (HPLC) using a fluorescence detector system (Waters, Milford, MA, USA), as described previously [2]. For melatonin measurement, the samples were extracted with 1 mL of chloroform for 16 h at 4 °C. The extracts were centrifuged for 10 min at 12,000 × g, and the supernatants (20 µL) were completely evaporated and dissolved in 0.1 mL of 40% methanol, and 20-µL aliquots were subjected to HPLC using a fluorescence detector system (Waters, Milford, MA, USA), as described previously [2]. All measurements were performed in triplicate.

3.6. Malondialdehyde Measurement

The rice samples (50 mg) were ground and extracted with 1.5 mL of reaction buffer (containing 0.5% thiobarbituric acid (TBA) and 20% trichloroacetic acid (TCA)). The extracts were centrifuged for 15 min at 12,000 × g, and the resulting supernatants were collected for measurement. The supernatant was boiled for 25 min at 95 °C, and placed for 5 min in ice. MDA content was determined at 440, 532, and 600 nm wavelengths using a spectrophotometer (OPIZEN POP-BIO). MDA was quantified using a molar extinction coefficient of 156/nM/cm.

3.7. List of Primers for qRT-PCR

The following primers were utilized for qRT-PCR and semifquantitative RT-PCR. CRY1a (forward 5’-AGA TGG AAG TTG ACC GAG-3’; reverse 5’-AGG CTG GAA GAG
we found that light quality played a role in light-induced melatonin biosynthesis, and the
when plants were incubated in light [2]. From the results of more detailed experiments,
ware (IBM Corp. Armonk, NY, USA). Means with asterisks indicate significantly different

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