A chloroplast-targeted DnaJ protein contributes to maintenance of photosystem II under chilling stress

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Abstract

DnaJ proteins act as essential molecular chaperones in protein homeostasis and protein complex stabilization under stress conditions. The roles of a tomato (Lycopersicon esculentum) chloroplast-targeted DnaJ protein (LeCDJ1), whose expression was upregulated by treatment at 4 and 42 °C, and with high light, NaCl, polyethylene glycol, and H2O2, were investigated here using sense and antisense transgenic tomatoes. The sense plants exhibited not only higher chlorophyll content, fresh weight and net photosynthetic rate, but also lower accumulation of reactive oxygen species and membrane damage under chilling stress. Moreover, the maximal photochemistry efficiency of photosystem II (PSII) (Fv/Fm) and D1 protein content were higher in the sense plants and lower in the antisense plants, and the photoinhibitory quenching was lower in the sense plants and higher in the antisense plants, suggesting that the inhibition of PSII was less severe in the sense plants and more severe in the antisense plants compared with the wild type. Furthermore, the PSII protein complexes were also more stable in the sense plants. Interestingly, the sense plants treated with streptomycin (SM), an inhibitor of organellar translation, still showed higher Fv/Fm, D1 protein content and PSII stability than the SM-untreated antisense plants. This finding suggested that the protective effect of LeCDJ1 on PSII was, at least partially, independent of D1 protein synthesis. Furthermore, chloroplast heat-shock protein 70 was identified as the partner of LeCDJ1. These results indicate that LeCDJ1 has essential functions in maintaining PSII under chilling stress.

Key words: Chilling stress, D1 protein, DnaJ protein, Hsp70, photosystem II, tomato.

Introduction

Photosystem II (PSII) is a large pigment–protein complex in the thylakoid membrane that performs the key reactions of photosynthesis (Shi et al., 2012). It exists mainly in dimeric form with the monomer containing at least 27–28 subunits (Rochaix, 2011). A distinct feature of PSII is that it is particularly prone to photo-oxidative damage under abiotic stresses (Aro et al., 1993). Chilling stress is one of the most significant environmental stresses on agricultural plants. Numerous studies have shown that chilling stress inhibits PSII activity (Li et al., 2010; Duan et al., 2012). Plants exposed to chilling stress show reduced metabolic rates (Partelli et al., 2009). This often leads to photoinhibition, which is referred to as inhibition of the activity of PSII, and is due to an imbalance between the rate of PSII damage and repair. The main target of PSII damage is the PSII reaction centre D1 protein, and the damaged D1 protein must be repaired by de novo D1 protein synthesis (Nishiyama et al., 2006).

In the long-term evolution of plants, many protective mechanisms were formed to quickly and effectively repair photodamaged PSII (Aro et al., 2005; Takahashi and Badger, 2011). Several auxiliary factors, such as kinases, phosphatases, proteases, and DnaJ proteins, are involved in this repair cycle.
DnaJ proteins are key components contributing to cellular protein homeostasis under stress conditions (Wang et al., 2004). They function as molecular chaperones, alone or in association with heat-shock protein 70 (Hsp70) partners, and are involved in various essential cellular processes, including protein folding, degradation, and refolding (Hennessy et al., 2005; Craig et al., 2006). Since their discovery in *Escherichia coli* as 41 kDa heat-shock proteins, DnaJ proteins have been found ubiquitously in all kingdoms of life (Georgopoulos et al., 1980; Bukau and Horwich, 1998; Craig et al., 2006). In general, DnaJ proteins contain one to four canonical domains (Silver and Way, 1993). Only the J-domain is completely conserved in all prokaryotic and eukaryotic groups. Through this domain, DnaJ proteins can interact with the ATPase domain of Hsp70 and hydrolyse ATP to ADP, facilitating client capture (Kampinga and Craig, 2010). Based on domain composition, the DnaJ family can be classified into three subtypes, with type III containing only a J-domain (Cheetham and Caplan, 1998). These subtypes are present in all major eukaryotic cell compartments, including the cytosol (Thomas and Baneyx, 1996), mitochondria (Voos and Rottgers, 2002), endoplasmic reticulum (Nicoll et al., 2006), and chloroplasts (Orme et al., 2001).

Chloroplasts are the structures in which photosynthesis mainly occurs. Previous studies have revealed that chloroplast-targeted DnaJ proteins participate in many processes that take place in the chloroplast, such as chloroplast development (Vitha et al., 2003; Liu et al., 2005, 2007; Albrecht et al., 2008), phototropin-mediated chloroplast movement (Suetsugu et al., 2005), and protein import and translocation (Chiu et al., 2010). Three Arabidopsis chloroplast DnaJ proteins, A1J8, A1J11 and A1J20, were found to be involved in optimization of photosynthetic reactions and stabilization of PSII complexes under high light stress (Chen et al. 2010). DnaJ proteins function alone or in association with Hsp70 partners. Hsp70B may protect PSII against damages through two distinct mechanisms: one that requires de novo D1 protein synthesis and one that does not (Schroda et al., 2001). Whether the small DnaJ proteins function together with their Hsp70 partner, and whether the protective effect of these proteins on PSII requires de novo D1 protein synthesis, are interesting issues that need to be addressed.

Accordingly, we isolated a chloroplast-targeted DnaJ protein (LeCDJ1) from tomato and submitted the sequence to GenBank under accession number GU925907. This protein belongs to the simplest group of DnaJ proteins (type III) characterized specifically by the J-domain. The expression of LeCDJ1 was induced by chilling stress. Overexpression of LeCDJ1 in tomato alleviated chilling stress-induced photoinhibition, whereas its suppression increased chilling sensitivity.

**Materials and methods**

*Plant materials, growth, and treatments*

Three sense T1 transgenic tomato lines (S3, S7, and S14), wild tomato cultivar (*Solanum lycopersicum* cv. Zhongshu 6), and three antisense T1 transgenic lines (A5, A11, and A13) were used as plant materials.

Seeds were sterilized, sown on Murashige and Skoog medium and incubated at 25 °C (light 16 h/dark 8 h) for 10 d. Some of the young seedlings were then exposed to 4 °C for 10 d and the growth performance was observed. The rest of the young seedlings were planted in plastic pots filled with sterilized soil and grown at 25/20 °C (day/night) with a 16 h photoperiod, a photosynthetic photon flux density (PPFD) range of 500-600 μmol m⁻² s⁻¹, and a relative humidity range of 50-60% in a greenhouse. The plants were irrigated with Hoagland nutrient solution once a week. When the sixth leaf was fully expanded (approximately 2 months old), the plants were adapted in an illuminated incubation chamber (GXZ-260C) for 2 d before treatment. For chilling treatment in light, the plants were exposed to 4 °C in the illuminated incubation chamber with a PPFD of approximately 200 μmol m⁻² s⁻¹ for 0, 3, 6, 12, and 24 h, and then recovered at 25 °C with the same PPFD for 2 and 4 h. For chilling stress in darkness, the plants were grown at 25 °C in the same illuminated incubation chamber with all lights turned off for 18 h and then exposed to 4 °C, still in darkness. For heat treatment, whole plants in pots were put in the illuminated incubation chamber at 42 °C for 0, 3, 6, 12, and 24 h. For high light treatment, 2000 μmol m⁻² s⁻¹ light from daylight-type microwave sulfur lamps (MSL-1000; Youhe, Ningbo, China) was administered by irrigating the seedlings with 20% polyethylene glycol (PEG) 6000. For oxidative treatment, the plant leaves were sprayed with 20 mM H₂O₂. The treated plant leaves were harvested at an appropriate time, frozen in liquid nitrogen, and stored at −80 °C.

In another experiment, leaf discs were taken from plants exposed to the control growth condition. Half were immediately soaked in 3 mM streptomycin (SM) solution in the dark, whereas the others were soaked in water to serve as the control. After approximately 3 h, all discs were placed on the water surface at 4 °C. The water temperature was controlled by an RTE-211 water circulator (Thermo Fisher Scientific, Worcester, MA, USA).

*Isolation and sequencing of LeCDJ1*

Total RNA was isolated using a total RNA isolation system (Tiangen, Beijing, China). For reverse transcription, 2 μg of total RNA was denatured at 70 °C for 5 min. Next, 1 μl of avian myeloblastosis virus reverse transcriptase (Fermentas) was added, mixed briefly, and incubated at 42 °C for 1 h. The reaction was terminated at 70 °C for 10 min.

For cloning of LeCDJ1 from tomato, a pair of primers (JF and JR; Supplementary Table S1 at JXB online) was designed based on the *LeCDJ1* sequence (GenBank accession no. AK323422.1). PCR amplification was performed as follows: initial denaturation at 94 °C for 5 min; 35 cycles of 94 °C for 50 s, 53 °C for 50 s, and 72 °C for 1 min; final extension at 72 °C for 10 min; and reaction termination at 4 °C. The PCR amplification products were cloned into the pMD-18T vector and then sequenced. All primers were synthesized by BioSune Biotechnology Ltd Co. (Jinan, China).

*RNA gel blot analysis*

The total RNA (20 μg) of tomato was separated in a 1.2% agarose formaldehyde gel, transferred to a nylon membrane, and fixed on the membrane by cross-linking with UV light. Pre-hybridization was performed at 65 °C for 12 h. The 3′ partial cDNA of *LeCDJ1* labelled with [α-32P]dATP by random primed labelling (Prime-a-Gene Labeling System; Promega) was used as the gene-specific probe. After 24 h of hybridization, filters were washed subsequently in 2× SSC (1× SSC: 0.15 M NaCl, 0.015 M sodium citrate, pH 7) with 0.2% SDS and 0.2× SSC with 0.2% SDS at 42 °C. Autoradiography was performed at 80 °C.

*Quantitative real-time PCR (qRT-PCR) analysis*

Total RNA extraction and reverse transcription were performed as previously above. qRT-PCR was performed on a Bio-Rad CFX96TM...
Real-time PCR System using SYBR Real Master Mix (Tiangen) with the following PCR thermal cycle conditions: denaturation at 95 °C for 30 s; and 40 cycles of 95 °C for 5 s, 58 °C for 10 s, and 68 °C for 10 s. EF-1α (GenBank accession no. X144491) was used as the actin. Template-free, negative, and single primer controls were established before the examination. The results are represented by three biological replicates (each with three technical replicates) for each sample, and a standard curve method was used for statistical analysis.

Antibody preparation, total protein extraction, and western blot analysis

The coding region of LeCDJ1 was subcloned into the pET-30a (+) vector between the BamHI and SacI sites. Expression and purification of the recombinant LeCDJ1 protein were carried out using a Ni-NTA agarose system according to the manufacturer’s instructions (Qiagen, Hilden, Germany). The purified recombinant protein was used to immunize the white mice and the obtained antisera was then purified according to the Amershams Biosciences (Piscataway, NJ, USA) antibody purification protocol. The secondary antibody was a horseradish peroixidase-conjugated goat anti-mouse IgG (Santa Cruz Biotechnology, Santa Cruz, CA, USA). The primary antibody was used at a dilution of 1:5000, whereas the secondary antibody was used at 1:5000.

Proteins were extracted from the leaves with extraction buffer (100 mM HEPES, pH 7.5, 5 mM EDTA, 5 mM EGTA, 10 mM dithiothreitol, 10 mM NaVO₄, 10 mM NaF, 1 mM phenylmethylsulfonyl fluoride, 5 mg ml⁻¹ of leupeptin, 5 mg ml⁻¹ of aprotinin, 5% glycerol, 50 mM β-glycerophosphate). After centrifugation at 15 000 g for 30 min at 4 °C, the supernatants were transferred into clean tubes, immediately frozen with liquid nitrogen, and then stored at −80 °C.

For western blotting, 20 μg of total plant proteins separated by SDS-PAGE were electrophoretically transferred to polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA) and detected with antibody preparations. The protein content was determined by a dye-binding assay. Quantitative image analysis of protein levels was performed with a Tanon Digital Gel Imaging Analysis System (Tanon-4100; Shanghai Tanon Science and Technology Co., Shanghai, China).

Subcellular localization of LeCDJ1

The N terminus of LeCDJ1 was cloned and two DNA constructs (p35S-GFP and p35S-LeCDJ1-GFP) were prepared. Isolated Arabidopsis mesophyll protoplasts were transfected with the above two constructs as described by Shah et al. (2002) and examined by dual-channel confocal microscopy (LSM510 META; Zeiss, Germany). The GFP fluorescence, red chloroplast autofluorescence, and the bright-field image of the protoplast were recorded simultaneously and compared. The potential co-localization of GFP fluorescence and red chloroplast autofluorescence was analysed further by checking for the presence of yellow signals in the superimposed images.

Plant transformation and transgenic tomato plants identification

Full-length LeCDJ1 cDNA was subcloned into the expression vector pBI121 downstream of the 35S cauliflower mosaic virus promoter. The constructs were then introduced into Agrobacterium tumefaciens LBA4404 by the freezing transformation method and verified by PCR and sequencing. Kanamycin-resistant transgenic tomato plants were generated using an A. tumefaciens-mediated leaf disk method. The DNA of the sense and antisense transgenic plants was extracted and used to amplify the target gene by PCR.

Measurement of chlorophyll content

Ten-day-old young seedlings were incubated for 10 d at 4 °C. The whole plants were homogenized in 5 ml of 80% acetone for 3 d and the homogenate was centrifuged at 3500g for 5 min. The supernatant was retained and the absorbance was recorded at 663 and 646 nm by UV spectrophotometry.

Measurement of net photosynthetic rate (Pn)

Pn was measured with a portable photosynthetic system (CIRAS-2; PP Systems, Herts, UK) under ambient CO₂ concentrations (360 μl l⁻¹), a PFD of 800 μmol m⁻² s⁻¹, and a relative humidity of 80%. The illumination source was produced by light-emitting diodes. Before Pn measurement, the plants were kept for approximately 30 min at 25 °C and at a PFD of 100 μmol m⁻² s⁻¹ to induce stomatal opening and then illuminated for approximately 15 min at a PFD of 800 μmol m⁻² s⁻¹.

Histochemical staining

Superoxide radical (O₂⁻) was detected visually by treating leaves with nitro blue tetrazolium (NBT) as described by Rao and Davis (1999). H₂O₂ was stained with 3,3’-diaminobenzidine (DAB) according to the method of Giacomelli et al. (2007). Trypan blue staining was performed as described by Choi et al. (2007).

Measurement of O₂⁻* and H₂O₂ concentration

To measure the concentration of O₂⁻*, leaves (0.5 g) were ground with liquid nitrogen in a mortar and then transferred to a centrifugal tube. Next, 3 ml of cold phosphate buffer (50 mM, pH 7.8) was added, and the homogenate was centrifuged at 5000g for 10 min at 4 °C. The supernatant with phosphate buffer (pH 7.8) and 10 mM hydroxyammonium chloride was incubated at 25 °C for 20 min, followed by the addition of 17 mM p-aminobenzene sulfonic acid and 7 mM α-naphthylamine. The mixture was then incubated at 25 °C for 20 min and then centrifuged at 1500g for 5 min. Finally, ethyl ether was added to the mixture. The water phase was used to determine the absorbance at 530 nm.

For measurement of H₂O₂ concentration, leaves (0.5 g) were ground with liquid nitrogen in a mortar and then transferred to a centrifugal tube. Subsequently, 3 ml cold phosphate buffer (50 mM, pH 6.8) was added. After centrifugation at 6000g for 15 min, 3 ml of supernatant and 1 ml of 0.1% titanium sulfate in 20% (v/v) H₂SO₄ were added into a new tube, mixed, and then centrifuged again. The absorbance was determined at 410 nm.

Measurement of malondialdehyde (MDA) and the relative electric conductivity (REC)

For measurement of MDA content, leaves (0.5 g) were ground in a cold mortar containing 10 ml of 10% trichloroacetic acid. After centrifugation at 10 000g for 10 min at 4 °C, 2 ml of supernatant with 2 ml of 0.6% thiorbarbituric acid reagent [0.6% (m/v) thiorbarbituric acid dissolved in 10% (m/v) trichloroacetic acid] was mixed, heated at 100 °C for 15 min, and then quickly cooled and centrifuged at 5000g for 10 min. The control contained 2 ml of distilled water instead of MDA extract. Absorbance was determined at 450, 532, and 600 nm. The MDA content was computed using a standard curve relating the MDA concentrations to absorbance.

Ten leaf discs (0.8 cm diameter) from each line were placed in 20 ml of distilled water, vacuumed for 30 min, and then surged for 3 h to measure the initial electric conductivity (S₀). The materials were boiled for 30 min and then cooled to room temperature to measure the final electric conductivity (S₁). Distilled water was used as a blank control and its electric conductivity (S₀) was measured. REC was calculated as REC=(S₁−S₀)/(S₂−S₀)×100.

Measurement of the chlorophyll a fluorescence transient

The chlorophyll a fluorescence transient was measured using a Handy Plant Efficiency Analyzer (Hansatech Instruments, Norfolk, UK) with dark-adapted leaves under ambient CO₂ conditions.
The transient was induced by a red light of approximately 3000 μmol m⁻² s⁻¹ provided by an array of four light-emitting diodes (peak 650 nm). The measurement protocol was as described by Zhang et al. (2012). The maximal photochemistry efficiency of photosystem II (PSII), \( F_v/F_m \), was calculated as follows: \( F_v/F_m = 1 - (F_o/F_m) \), where \( F_o \) is variable fluorescence, \( F_m \) is maximum fluorescence and \( F_o \) is minimum fluorescence (when all PSII centres are in the open state).

Measurement of chlorophyll fluorescence

A FMS-2 pulse-activated modulation fluorometer (Hansatech, Cambridge, UK) was used to measure chlorophyll fluorescence. For the assay of photoinhibitory quenching (qI) and energy-dependent quenching (qE), plants were dark adapted for 12 h before pre-stress \( F_m \) was measured. Leaves were then illuminated for 40 min under actinic light (200 μmol m⁻² s⁻¹). This length of illumination was always found to be sufficient to reach a steady-state fluorescence yield and a maximum photosynthetic rate. A pulse of saturating light (3000 μmol m⁻² s⁻¹ for 700 ms) was applied to determine the maximal fluorescence under actinic light (\( F_m' \)). After this, the actinic light was switched off, and 40 min later, the dark relaxation of fluorescence (\( F_m'' \)) was measured by applying saturating light pulses. qI and qE were calculated as follows: \( qI = F_m''/F_m - 1 \), and \( qE = F_m''/F_m' - F_m''/F_m \). Pre-stress \( F_m \) was used for the qI and qE calculation.

Thylakoid membrane preparation and SDS-PAGE

For thylakoid membrane preparation, tomato leaves were homogenized in an ice-cold isolation buffer (400 mM sucrose, 50 mM HEPES, pH 7.8, 10 mM NaCl, 2 mM EDTA, and 2 mM MgCl₂) and filtered through three layers of pledget. The filtrate was centrifuged at 5000 g for 10 min. The thylakoid pellets were washed with isolation buffer, recentrifuged, and finally suspended in an isolation buffer. The thylakoid membrane proteins were then denatured and separated using a 15% polyacrylamide gradient gel containing 6 M urea.

Blue native PAGE (BN-PAGE) and western blotting

BN-PAGE was performed as described by Sun et al. (2007). After electrophoresis, the protein complexes were denatured with 15% methanol for 15 min. The denatured protein complexes were then electroblotted onto PVDF membranes, probed with D1 antibody, and then visualized by the enhanced chemiluminescence method. X-ray films were scanned using an AlphaImager 2200 documentation system (Alpha Innotech).

Yeast two-hybrid assays

Yeast two-hybrid assays were performed according to the Matchmaker™ Gold Yeast Two-Hybrid System User Manual. The cpHsp70 coding region was amplified with cDNA from tomato seedlings and ligated to BamHI/Xhol-digested pGADT7 (Clontech). The LeCDJ1 coding region was amplified from a plasmid and then ligated to EcoRI/Sall-digested pGBK7 (Clontech). The expression vector pGADT7-cpHsp70 was co-transformed into yeast strain YI187 (Clontech) with pGBK7-LeCDJ1 using the lithium acetate transformation method. Cells were plated onto selective medium without Leu and Trp (DDO). Putative transformants were transferred to selective medium without Leu, Trp, His, and adenine and supplemented with X-α-Gal and aureobasidin (QDO/X/A). The interactions between the p35 and T proteins, as well as between the Lam and T proteins, were used as positive and negative controls, respectively. Autoactivation was analysed by a growth experiment when the detected gene was co-transformed with the pGADT7 or pGBK7 empty vector.

Statistical analysis

Data points represent the mean ± standard deviation (SD) of three replications. Statistical significance of differences in the measured parameters between the wild type (WT) and transgenic plants was tested using the software in Excel. Significant differences in comparison with the control are indicated by *P<0.05 and **P<0.01.

Results

Expression patterns of LeCDJ1 in tomato

LeCDJ1 was isolated from a tomato cDNA library representing the expression patterns of genes affected by chilling. We first studied the expression of LeCDJ1 in response to chilling using RNA gel blotting, qRT-PCR and western blotting. LeCDJ1 was clearly induced by chilling treatment (Fig. 1A, B).

![Fig. 1](image-url)

Fig. 1. Responses of LeCDJ1 to chilling stress in tomato. (A) RNA gel blot analysis of the expression of LeCDJ1 in leaves treated at 4 °C in light (200 μmol m⁻² s⁻¹) for 0, 3, 6, 12, and 24 h and then recovered at 25 °C for 2 and 4 h. Ethidium bromide-stained rRNA is shown as a loading control. (B) qRT-PCR analysis of the response of LeCDJ1 to chilling stress in light. The transcript level of LeCDJ1 was normalized to EF-1α expression. Error bars represent the SD of triplicate reactions. (C) Western blot analysis of the LeCDJ1 protein levels in the leaves treated at 4 °C in light. LC, loading control (part of the Coomassie-stained total protein SDS-PAGE gel). (D) Quantitative image analysis of the protein content in (C) by Tanon Digital Gel Imaging Analysis System. The relative protein level of LeCDJ1 was normalized to the level at 0 h.
The transcript reached a maximum level at 6 h, decreased after 12 h, and then recovered slightly to the original state after recovery at 25 °C for 4 h. Moreover, chilling stress in darkness still induced LeCDJ1 expression, although not as strongly as in light (Supplementary Fig. S1 at JXB online), suggesting that the response of LeCDJ1 to chilling did not depend on the light. Western blot analysis showed that the protein signal became gradually stronger after treatment at 4 °C, but decreased slightly during the recovery period (Fig. 1C). The quantitative image analysis of protein content in Fig. 1C showed a similar profile (Fig. 1D).

By qRT-PCR analysis, we detected the expression level of LeCDJ1 under heat (42 °C), high light (2000 μmol m⁻² s⁻¹), salt (200 mM NaCl), osmotic (20% PEG-6000) and oxidative (20 mM H₂O₂) stresses at different time points (Fig. 2). All the tested stresses induced the expression of LeCDJ1 to different extents. Heat stress induced the expression of LeCDJ1 gradually (Fig. 2A). High light stress induced the expression during the first 12 h, and the transcripts decreased at 24 h (Fig. 2B). Treated with 200 mM NaCl, the transcripts increased within 1 d, and remained high over a 3 d period (Fig. 2C). For 20% PEG treatment, transcripts reached a maximum level at 6 h and then decreased (Fig. 2D). H₂O₂ treatments also led to a rapidly accumulation of the transcripts (Fig. 2E). These results suggested that LeCDJ1 is involved in the response to various stresses.

Fig. 2. qRT-PCR analysis of the expression profiles of LeCDJ1 in tomato. (A–E) Responses of LeCDJ1 to 42 °C (A), high light (2000 μmol m⁻² s⁻¹ (B), 200 mM NaCl (C), 20% PEG-6000 (D), and 20 mM H₂O₂ (E). (F) Expression of LeCDJ1 in the roots, stems, leaves, flowers, and fruits. The transcript level of LeCDJ1 was normalized to EF-1α expression. Error bars represent the SD of triplicate reactions.
We then examined the expression patterns of *LeCDJ1* in various organs. As shown in Fig. 2F, *LeCDJ1* was constitutively expressed in all organs examined and the transcript level in the leaves was approximately 10 times higher than that in the roots. Thus, the gene is expressed preferentially in chlorophyllous tissues.

**Chloroplast targeting of LeCDJ1**

The programs ChloroP 1.1 (http://www.cbs.dtu.dk/services/ChloroP/) and TargetP 1.1 (http://www.cbs.dtu.dk/services/TargetP/) predicted that the first 45 aa of LeCDJ1 constituted the plastid target signals and that LeCDJ1 was probably a chloroplast protein (Supplementary Tables S2 and S3 at *JXB* online). To obtain direct experimental evidence, we performed targeting experiments *in vivo* in Arabidopsis protoplasts derived from leaf tissue. As shown in Fig. 3, green fluorescence in the individual protoplasts transfected with the control construct p35S-GFP (expressing the GFP coding sequence alone) was detected in the cytoplasm surrounding the chloroplasts and was not co-localized with the red autofluorescence of the chloroplasts. By contrast, when fused with LeCDJ1, the GFP signal was predominantly confined to the chloroplasts and co-localized with the red autofluorescence of the chloroplasts, suggesting that LeCDJ1 is a chloroplast protein.

In addition, amino acid sequence alignment between LeCDJ1 and DnaJ8 proteins from other plants showed that LeCDJ1 shared a high degree of sequence similarity to Arabidopsis AtDnaJ8 and pea PsJ8b (Supplementary Fig. S2 at *JXB* online). Both have been suggested to be a soluble stromal protein with a small portion peripherally associated with membranes where many key steps of PSII repair and reassembly processes take place (Chiu et al., 2010).

**Identification of transgenic plants**

A total of 31 (16 sense lines, 15 antisense lines) individual kanamycin-resistant tomato transgenic lines (T0) were harvested from tissue culture. The progeny obtained from T0 were named T1. Six sense (S1, S3, S5, S7, S8, and S14) and five antisense (A5, A7, A11, A12, and A13) T1 lines were selected for qRT-PCR. Compared with WT, the relative *LeCDJ1* mRNA level in the examined sense lines increased by 5.3-, 19.7-, 13.3-, 14.1-, 10.5-, and 14.3-fold, whereas that in the antisense lines decreased by 0.30-, 0.43-, 0.14-, 0.34- and 0.31-fold, respectively (Fig. 4A). Among these lines, S3, S7, S14, A5, A11, and A13 were selected for the following physiological measurements.

**LeCDJ1 overexpression enhanced chilling stress tolerance**

The chilling stress tolerance of the plants was determined by observing the growth performance of the young seedlings (approximately 10 d old) and grown plants (approximately 2 months old). Under normal conditions, both grew well and showed no significant difference in phenotype and physiological traits (Fig. 5). After treatment at 4 °C (young seedlings for 10 d and grown plants for 24 h), the growth of all plants was more or less suppressed. The grown plants showed little difference between sense, WT, and antisense plants (Fig. 5B). However, suppression of the growth of the young seedlings was less serious in the sense plants and more serious in the antisense plants, compared with WT. The leaves of the antisense young seedlings showed photobleaching, whereas most leaves of the sense lines remained green (Fig. 5A). Accordingly, the sense plants showed higher chlorophyll contents (approximately 83.6% that of the untreated WT plants, on average), whereas the antisense plants had a larger drop (approximately 30.3%, on average), compared with WT (approximately 76.3%) (Fig. 5C). Similarly, the fresh weight of WT plants was 0.111 ± 0.0041 g, which was lighter than S3 (0.1274 ± 0.0026 g), S7 (0.1254 ± 0.0010 g),

![Fig. 3. Intracellular targeting of LeCDJ1 in Arabidopsis protoplasts.](image-url)

*Fig. 3. Intracellular targeting of LeCDJ1 in Arabidopsis protoplasts.* (A, B) Green fluorescence of GFP (A) and the LeCDJ1–GFP fusion protein (B). (C, D) Red autofluorescence of chloroplasts. (E, F) Bright-field images of the protoplasts. (G, H) Merged images of (A), (C) and (E), and of (B), (D), and (F), respectively. Protoplasts were examined using dual-channel confocal microscopy (LSM510 META; Zeiss, Germany).
and S14 (0.1245 ± 0.00114 g) plants, but heavier than A5 (0.0741 ± 0.0041 g), A11 (0.0793 ± 0.00105 g), and A13 (0.0782 ± 0.00149 g) plants (Fig. 5D).

After chilling treatment, the \( P_e \) of both WT and transgenic grown plants markedly decreased. However, the decrease in the antisense plants was also more serious than that in the sense plants (Fig. 5E). These results indicated that the transgenic plants with higher \( \text{LeCDJ1} \) expression levels had higher chilling tolerance, whereas those with lower \( \text{LeCDJ1} \) expression levels had higher chilling sensitivity.

**LeCDJ1 overexpression alleviates reactive oxygen species (ROS) accumulation under chilling stress**

Chilling stress accelerates ROS generation, so we analysed intracellular levels of \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) by NBT and DAB staining, respectively. Prior to treatment, both \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) accumulation was low, with no significant difference between the WT and transgenic plants. After chilling treatment for 12 h, NBT staining for \( \text{O}_2^- \) showed that the blue polymerization product due to \( \text{O}_2^- \) accumulation increased, especially in WT and antisense plants (Fig. 6A). The colour was darkest in the antisense plants. Similarly, DAB staining showed that \( \text{H}_2\text{O}_2 \) accumulation also increased after chilling stress, and the accumulation was less in S3, S7, and S14 plants, but more in A5, A11, and A13 plants, than in WT (Fig. 6B). Quantitative analysis of \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) showed a similar result (Fig. 6C, D). These results suggested that \( \text{LeCDJ1} \) overexpression alleviates the accumulation of \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \).

**LeCDJ1 overexpression alleviates membrane damage under chilling stress**

As the primary target of chilling stress, membranes are particularly susceptible to ROS-initiated lipid peroxidation reactions. Trypan blue staining showed that all plants had a similar level of blue marks under normal growth condition. However, after 12 h of chilling treatment, the antisense lines exhibited darker blue marks than the WT and sense plants (Fig. 7A). For confirmation, MDA accumulation and REC, which have been reported as indicators for the membrane damage, were determined. As shown in Fig. 7B and C, both MDA accumulation and REC increased after chilling stress. In addition, the increases were more obvious in the antisense lines and less obvious in the sense lines. These results suggested that, compared with WT, membrane damage was less serious in the sense plants and more serious in the antisense plants.

**LeCDJ1 overexpression alleviates photoinhibition of PSII under chilling stress**

\( F_v/F_m \) was used to measure PSII photoinhibition. Only a slight difference was noted between the WT and transgenic lines before treatment. After chilling stress, \( F_v/F_m \) decreased slightly in the sense plants, but decreased markedly in the antisense plants, compared with WT, suggesting that \( \text{LeCDJ1} \) overexpression decreases chilling stress-induced PSII photoinhibition (Fig. 8A).

The decrease in \( F_v/F_m \) during chilling stress has various explanations: for instance, a change in \( q_I \) and \( q_E \). During and after chilling treatment, \( q_I \), part of which is due to D1 damage, was lower in the sense plants and higher in the antisense plants, compared with WT (Fig. 8B). The substantial increases in \( q_I \) indicated that the antisense plants experienced more photoinhibition than the WT and sense plants under chilling stress (Dodd *et al.*, 1998). Interestingly, \( q_E \) values among the different tested lines were not significantly different, indicating that the lower \( F_v/F_m \) in the antisense plants was most possibly caused by the higher \( q_I \) (Fig. 8C).

![Fig. 4. Identification of transgenic plants by qRT-PCR and western blotting. (A) Transcript levels of \( \text{LeCDJ1} \) in the WT and different transgenic lines. The transcript level of \( \text{LeCDJ1} \) was normalized to \( EF-1\alpha \) expression. Error bars represent the SD of triplicate reactions. (B) \( \text{LeCDJ1} \) protein levels in the WT and transgenic lines. LC, loading control (part of the Coomassie-stained total protein SDS-PAGE gel). (C) Quantitative image analysis of protein content in (B) using a Tanon Digital Gel Imaging Analysis System. The relative protein level of \( \text{LeCDJ1} \) in the transgenic lines was normalized to that in the WT. Two-month-old plants were used.](image-url)
LeCDJ1 overexpression alleviates photoinhibition of PSII whether in the presence of organelle protein synthesis or not

SM, an inhibitor of organelar translation that can inhibit de novo D1 protein synthesis, was used to investigate whether the protective effect of LeCDJ1 on PSII was dependent on D1 protein synthesis. \( F_v/F_m \) in leaf discs treated with water or SM at 25 °C decreased slightly and showed no obvious difference between the tested lines (data not shown). After chilling stress, \( F_v/F_m \) in the plants decreased markedly. It was also higher in S3 and lower in A11 leaves, compared with WT, especially in the presence of SM (Fig. 9A). For example, after chilling treatment for 9 h, \( F_v/F_m \) in S3, WT, and A11 leaves decreased to approximately 60, 53, and 48% of the original value, respectively. In the presence of SM, an approximately 48.1% decrease in \( F_v/F_m \) was observed in S3 leaves, whereas 59.9% was found in A11 leaves. Interestingly, \( F_v/F_m \) in the SM-treated S3 leaves was higher than that in the SM-untreated A11 leaves during chilling treatment. Given that this higher portion was not due to de novo D1 protein synthesis, the protective effect of LeCDJ1 on PSII, at least partially, was not dependent on D1 protein synthesis.

To investigate whether the impaired PSII function was associated with the changes in D1 protein levels, we performed western blot analysis of thylakoid membrane preparations with equal amounts of chlorophyll. As shown in Fig. 9B, D1 protein levels showed no obvious difference between the tested lines under normal condition. However, after 12 h of chilling treatment, D1 protein showed a stronger signal in the sense plants, and a weaker signal in the antisense plants, compared with WT. As a chloroplast translation inhibitor, SM further decreased the D1 protein content in all strains. However, this decrease was more obvious in the antisense plants than in the sense plants. The quantitative image analysis of protein content in Fig. 9B further proved our previous findings that the D1 protein content was more in the sense plants and less in the antisense plants, whether in the presence of SM or not (Fig. 9C). These results indicated that LeCDJ1 has a protective effect on D1 protein and this effect is at least partially not dependent on D1 protein de novo synthesis.
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Fig. 6. $O_2^{-}$ and $H_2O_2$ analysis in the WT and transgenic plants. (A) NBT staining for $O_2^{-}$. (B) DAB staining for $H_2O_2$. The top panel represents plants grown at 25 °C and the bottom panel represents plants treated at 4 °C for 12 h. (C) $O_2^{-}$ content in tomato plants. (D) $H_2O_2$ content in tomato plants. The experiment was repeated three times with similar results. The data are represented as means ±SD of three biological replicates (*$P<0.05$). Two-month-old grown plants were used. (This figure is available in colour at JXB online.)

Fig. 7. Membrane damage analysis in the WT and transgenic plants. (A) Trypan blue staining. The top panel represents plants grown at 25 °C and the bottom panel represents plants treated at 4 °C for 12 h. (B) MDA content in the WT and transgenic plants. (C) REC in the WT and transgenic plants. Three independent experiments are shown as means ±SD. Asterisks indicate significant differences from WT of the same treatment (*$P<0.05$). Two-month-old grown plants were used. (This figure is available in colour at JXB online.)
LeCDJ1 overexpression stabilizes the thylakoid protein complexes

Altered LeCDJ1 expression affected PSII activity (Fig. 8) and D1 protein levels (Fig. 9). Hence, we evaluated putative alterations in thylakoid protein complexes and performed BN-PAGE analysis. Seven major bands labelled I to VII were resolved, apparently representing PSII–LHCCI supercomplexes (band I), monomeric PSI and dimeric PSII (band II), monomeric PSII (band III), CP43-free PSII (band IV), trimeric LHCCI/PSII reaction centre (band V), monomeric LHCCI (band VI), and unassembled proteins (band VII). Few differences were apparent in the BN gel, except that the amount of PSII–LHCCI supercomplexes (band I) was a little more in S3 than in A11 plants, after chilling stress in the presence of SM (Fig. 10A). Immunoblotting of the BN gel with D1 antibody showed more clearly the decrease in PSII–LHCCI supercomplexes (band I) in the tested lines after chilling stress (Fig. 10B). S3 plants showed a smaller decrease, while A11 plants showed a larger decrease, especially in the presence of SM. The PSII–LHCCI supercomplexes completely disappeared from A11 plants in the presence of SM. Moreover, the amount of PSII dimers and monomers (band II) was also decreased, especially in WT and A11 plants, after chilling treatment. The monomeric PSII (band III) showed few changes among the tested lines after treatment. The CP43-free PSII (band IV) increased slightly after chilling stress, indicating that PSII repair was occurring. The quantitative image analysis of protein content in Fig. 10B showed the same trend (Fig. 10C). These findings indicated that LeCDJ1 provides a stabilizing force for these protein complexes.

In the present study, determination of an interaction between LeCDJ1 and these protein complexes was also attempted. However, western blotting of the BN gel with LeCDJ1 antibody showed no positive signal (data not shown). This finding may be attributed to the fact that their interaction was not strong enough to bear the solubilization of n-dodecyl β-d-maltoside.

Interaction between LeCDJ1 and a chloroplast Hsp70

To date, DnaJ proteins are known to act as important chaperones for Hsp70. Hence, we postulated that LeCDJ1 interacts with a specific Hsp70(s). According to its chloroplast localization (Fig. 3), the partner of LeCDJ1 was anticipated to be a tomato chloroplast-located protein. We then searched the NCBI database (http://www.ncbi.nlm.nih.gov) for ‘Solanum lycopersicum chloroplast Hsp70’ and identified a cDNA (GenBank accession no EU195057.1) encoding a chloroplast Hsp70. The programs ChloroP 1.1 and TargetP 1.1 also predicted that the isolated Hsp70 was probably a chloroplast protein (Supplementary Tables S2 and S3). A yeast two-hybrid assay was then performed, and the capacity of the yeast to grow on QDO/X/A was used as an interaction marker. All yeast transformants grew normally on DDO. The yeast co-transformed with LeCDJ1 and pGADT7 or cpHsp70 and pGBK T7 empty vectors did not grow on

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**Fig. 8.** Changes in \( \frac{F_v}{F_m} \) (A), qI (B), and qE (C) under chilling treatment in tomato plants. Plants grown at 25 °C in a greenhouse were exposed to 4 °C in the chambers for 0, 3, 6, 12, and 24 h in the light (200 μmol m⁻² s⁻¹). Plants were dark adapted for 30 min before measurement. The values are represented as means ±SD of three independent experiments (*P<0.05). Two-month-old grown plants were used.
QDO/X/A (Fig. 11A), indicating that none of the detected genes was autoactivated. However, when LeCDJ1 was co-transformed with cpHsp70, blue colonies growing on QDO/X/A were found, indicating an interaction between LeCDJ1 and cpHsp70 (Fig. 11B).

**Discussion**

DnaJ proteins act as molecular chaperones re-establishing normal protein conformation and cellular protein homeostasis under stress conditions (Wang et al., 2004). They are often referred to as cellular stress sensors and their expression is induced by heat, high light, methyl viologen and cold (Piippo et al., 2006; Scarpeci et al., 2008; Rajan and D’Silva, 2009). In this study, the expression of LeCDJ1 was induced by chilling, whether in light or darkness, suggesting that the response of LeCDJ1 to chilling did not depend on light (Fig. 1, Supplementary Fig. S1). We also established that LeCDJ1 was involved in the response to heat, high light, NaCl, PEG and \( \text{H}_2\text{O}_2 \) (Fig. 2). This elevated expression may be attributed to the enhanced protection of plants against the damaging effect of these environmental stresses. In fact, the roles of DnaJ proteins in response to environmental stresses have already been established by several studies (Li et al., 2007; Yang et al., 2009; Chen et al., 2010; Zhou et al., 2012; Bekh-Ochir et al., 2013). However, direct physiological evidence that supports the involvement of these proteins in chilling tolerance is still lacking. The data of the present study revealed the role of LeCDJ1, a type III chloroplast-targeted DnaJ protein (Fig. 3), in the maintenance of PSII activity under chilling stress.

Chilling stress inhibits PSII activity in several ways, for example lowering the rate of photosynthesis, blocking the PSII photosynthetic electron transport, and accelerating the generation of ROS, as well as damaging the membrane lipids and photosynthetic pigment complex systems (Suzuki et al., 2008). Several lines of evidence present in this study indicated that LeCDJ1 was involved in the maintenance of PSII activity under chilling stress. After chilling treatment, the sense
plants exhibited not only higher chlorophyll content, fresh weight, and \( P_n \), but also lower membrane damage (Figs 5 and 7). DnaJ proteins are involved in ROS level regulation in the chloroplast (Chen et al., 2010) and mitochondria (Zhou et al., 2012). In the present experiment, compared with WT, the S3, S7, and S14 plants maintained lower \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) levels, whereas the A5, A11, and A13 plants accumulated more \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \), suggesting that LeCDJ1 could help reduce the ROS level (Fig. 6). Moreover, the photochemical efficiency of PSII, the \( F_v/F_m \) ratio, in the sense lines decreased less markedly than that in the WT and antisense plants, suggesting that LeCDJ1 overexpression decreased PSII photoinhibition (Fig. 8A). Accordingly, the lower increases in \( q_I \) also indicated that the sense plants experienced less photoinhibition...
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| than WT and antisense plants under chilling stress (Fig. 8C). Thus, as discussed above, LeCDJ1 is involved in plant chilling tolerance and its overexpression alleviates PSII photoinhibition under chilling stress.

Additional lines of evidence supporting the roles of LeCDJ1 in PSII maintenance included the increased amounts of PSII reaction centre D1 protein and the increased stability of the PSII protein complexes in the sense plants (Figs 9B and 10). PSII is particularly prone to photo-oxidative damage as the water splitting reaction catalysed by this complex inevitably leads to the generation of ROS that damage the complex. Among the PSII proteins, the PSII reaction centre D1 protein is the main target of damage. Plants compromised in PSII photosynthetic activity have low levels of D1 protein (Suorsa et al., 2004). After chilling treatment for 12 h, the relative abundances of D1 protein were slightly higher in the sense plants and lower in the antisense plants, suggesting that PSII photosynthetic activity was compromised in the antisense plants (Fig. 9B, C). In addition, the PSII–LHCII supercomplexes and PSII dimers and monomers were also increased in the sense plants and reduced in the antisense plants at this time point, suggesting that LeCDJ1 may provide a stabilizing force for these protein complexes (Fig. 10B, C). Thus, our preliminary results suggest the importance of LeCDJ1 in the maintenance of PSII activity under chilling stress.

The process of PSII maintenance that might involve LeCDJ1 was also determined in this study. It is known that the extent of PSII photoinhibition is associated with a balance between the rate of PSII damage and repair (Takahashi and Murata, 2006). The main target of PSII damage is the PSII reaction centre D1 protein, whereas PSII repair requires D1 protein degradation, synthesis, and then reassembly into the PSII complex (Nishiyama et al., 2006). To distinguish between the two processes, SM, an inhibitor of organellar translation (inhibitor of de novo D1 protein synthesis), was used. The sense plants displayed a higher F_v/F_m, D1 protein content, and PSII protein complex stability whether in the presence of SM or not (Figs 9 and 10). Interestingly, after chilling stress, F_v/F_m, D1 protein content, and PSII protein complex stability in the SM-treated sense lines were higher than those in the SM-untreated antisense plants (Figs 9 and 10). Given that this higher proportion was not caused by de novo D1 protein synthesis, the protective effect of LeCDJ1 on PSII was, at least partially, not dependent on D1 protein synthesis. The contribution of plant chaperone proteins, such as heat-shock proteins, to the stability of proteins has been shown by studies in vivo (Low et al., 2000; Basha et al., 2010) and in vitro (Lee and Vierling, 2000; Basha et al., 2004). Considering the general function of DnaJ proteins as chaperone proteins (Walsh et al., 2004), an effect of LeCDJ1 on PSII that does not depend on de novo D1 protein synthesis could be explained by the possible role of LeCDJ1 in D1 protein conformation protection.

DnaJ proteins act as co-chaperones of the Hsp70 machinery and are important in the stimulation of Hsp70s ATPase activity, thereby stabilizing its interaction with client proteins (Mayer and Bukau, 2005; Kampinga and Craig, 2010). Subsequently, members of the DnaJ proteins have been found to function as molecular chaperones, alone or in association with Hsp70 partners. A chloroplast Hsp70 (cpHsp70) in

| Fig. 11. Interaction between LeCDJ1 and cpHsp70. (A) Test for autoactivation. (B) Yeast two-hybrid analysis of the interaction between LeCDJ1 and cpHsp70. An interaction between the p53 and T proteins was used as positive control (middle panel) and interaction between the Lam and T proteins (bottom panel) was used as negative control. Yeast strains harbouring the indicated plasmids were grown on selective medium without Leu and Trp (DDO; left panel) or selective medium without Leu, Trp, His, and adenine and with X-α-Gal and aureobasidin (QDO/X/A; right panel).
tomato, which could interact with LeCDJ1 directly, was identified through a yeast two-hybrid assay in this study (Fig. 11). Therefore, the LeCDJ1/cpHsp70 machinery may work together in PSII maintenance. In fact, the involvement of Hsp70B as a stabilizing force of the thylakoid membrane protein complexes is not without precedent (Schroda et al., 1999; Yokthongwattana et al., 2001). In general, DnaJ proteins drive the functional diversity of Hsp70s through two mechanisms. First, they interact with Hsp70 directly and stimulate the ATPase activity necessary for the stable binding of Hsp70 to their protein substrate (Bukau and Horwich, 1998). Secondly, DnaJ proteins localized to a particular site in a cellular compartment could be maintained at a high local concentration, thus targeting Hsp70 to particular client proteins at these sites (Nakatsukasa et al., 2008). The present data are insufficient to depict the definite role of LeCDJ1 in the Hsp70/DnaJ protein machinery. The effect of LeCDJ1 on the ATPase-affecting conformational changes in cpHsp70 should be determined to differentiate ultimately the two possible ways.

In conclusion, overexpression of LeCDJ1 in vivo resulted in increased tolerance of PSII against chilling stress, whereas suppression of LeCDJ1 increased PSII chilling sensitivity. Based on our data and previous knowledge, we postulate that the protective effect of LeCDJ1 on PSII is, at least partially, not dependent on de novo D1 protein synthesis and perhaps occurs by conformation protection of the D1 protein, together with cpHsp70.

Supplementary data

Supplementary data are available at JXB online.

Supplementary Fig. S1. Response of LeCDJ1 to chilling stress in darkness.

Supplementary Fig. S2. Amino acid sequence alignment between LeCDJ1 and DnaJ8.

Supplementary Table S1. Specific primers used in this study.

Supplementary Table S2. Prediction of subcellular localization of LeCDJ1 and cpHsp70 by the software program ChloroP 1.1.

Supplementary Table S3. Prediction of subcellular localization of LeCDJ1 and cpHsp70 using the software program TargetP 1.1.

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