Jasmonate Regulates the INDUCER OF CBF EXPRESSION–C-REPEAT BINDING FACTOR/DRE BINDING FACTOR1 Cascade and Freezing Tolerance in *Arabidopsis*

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The INDUCER OF CBF EXPRESSION (ICE)–C-REPEAT BINDING FACTOR/DRE BINDING FACTOR1 (CBF/DREB1) transcriptional cascade plays a critical role in modulating cold stress responses in *Arabidopsis thaliana*. Dissecting crucial upstream regulatory signals or components of the ICE-CBF/DREB1 cascade will enhance our understanding of plant cold-tolerance mechanisms. Here, we show that jasmonate positively regulates plant responses to freezing stress in *Arabidopsis*. Exogenous application of jasmonate significantly enhanced plant freezing tolerance with or without cold acclimation. By contrast, blocking endogenous jasmonate biosynthesis and signaling rendered plants hypersensitive to freezing stress. Consistent with the positive role of jasmonate in freezing stress, production of endogenous jasmonate was triggered by cold treatment. In addition, cold induction of genes acting in the CBF/DREB1 signaling pathway was upregulated by jasmonate. Further investigation revealed that several JASMONATE ZIM-DOMAIN (JAZ) proteins, the repressors of jasmonate signaling, physically interact with ICE1 and ICE2 transcription factors. JAZ1 and JAZ4 repress the transcriptional function of ICE1, thereby attenuating the expression of its regulon. Consistent with this, overexpression of JAZ1 or JAZ4 represses freezing stress responses of *Arabidopsis*. Taken together, our study provides evidence that jasmonate functions as a critical upstream signal of the ICE-CBF/DREB1 pathway to positively regulate *Arabidopsis* freezing tolerance.

INTRODUCTION

Temperatures outside an organism’s optimal tolerance range are regarded as a major environmental stress. Extreme low temperature disrupts cellular homeostasis and severely impairs plant growth and development. To tolerate cold stress, plants have evolved sophisticated mechanisms involving altered physiological and biochemical processes. Previous studies have revealed that the INDUCER OF CBF EXPRESSION (ICE)–C-REPEAT BINDING FACTOR/DRE BINDING FACTOR1 (CBF/DREB1) transcriptional cascade plays a critical role in the cold-response pathways in *Arabidopsis thaliana* (Thomashow, 1999; Chinnusamy et al., 2007). ICE1, a basic helix-loop-helix (bHLH) transcription factor, directly binds to cis-elements (CANNTG) in the *CBF3/DREB1a* promoter (Chinnusamy et al., 2003). Expression of *CBF3/DREB1a* and its downstream target genes is downregulated in *ice1* mutants; *ice1* plants consequently display significantly reduced cold acclimation-induced freezing tolerance (Chinnusamy et al., 2003). Similarly, ICE2, the homolog of ICE1, influences the expression of CBF1/DREB1b and participates in regulating plant cold stress responses (Fursova et al., 2009). Moreover, the SMALL UBIQUITIN-RELATED MODIFIER E3 ligase SIIZ1 (SAP and Miz) and the ubiquitin E3 ligase HIGH EXPRESSION OF OSMOTICALLY RESPONSIVE GENES1 have been shown to modify ICE1 posttranslationally and function in the ICE-CBF/DREB1 signaling pathway (Dong et al., 2006; Miura et al., 2007). However, whether there are other crucial regulators acting upstream of the ICE transcription factors needs to be investigated.

In addition to the ICE-CBF/DREB1 regulatory pathway, plant hormones have been demonstrated to modulate cold stress responses. For example, low temperature induces a transient increase in levels of the endogenous stress hormone abscisic acid (ABA), and exogenous application of ABA enhances plant cold tolerance (Lång et al., 1994; Mäntylä et al., 1995). Jeon et al. (2010) showed that cytokinin receptors *Arabidopsis* HISTIDINE KINASE2 (AHK2) and AHK3 and type-A *Arabidopsis* RESPONSE REGULATOR (ARR) proteins play regulatory roles in cold stress signaling via inhibition of ABA responses. Recently, ethylene signaling was shown to negatively regulate plant freezing stress responses by repressing the expression of CBF1/DREB1 and type-A ARR genes in *Arabidopsis* (Shi et al., 2012).

The phytohormone jasmonate acts as an important regulatory signal to influence multiple plant processes. In *Arabidopsis*, jasmonate is directly involved in root growth (Staswick et al., 1992; Feyes et al., 1994; Pauwels et al., 2010), male fertility (McConn and Browse, 1996; Sanders et al., 2000; Stintzi and Browse, 2000; Cheng et al., 2009), anthocyanin accumulation.
(Franceschi and Grimes, 1991; Shan et al., 2009), senescence (Ueda and Kato, 1980; Schommer et al., 2008; Shan et al., 2011), and defense responses (Howe et al., 1996; McConn et al., 1997; Reymond and Farmer, 1998; Vijayan et al., 1998; Farmer, 2001; Farmer et al., 2003; Browse, 2009). Jasmonate is perceived by the F-box protein CORONATINE INSENSITIVE1 (COI1), which subsequently facilitates the degradation of JAZ proteins via the SCFCOI1-26S proteasome pathway (Xu et al., 2002; Chini et al., 2007; Thines et al., 2007; Yan et al., 2009; Sheard et al., 2010). JAZ proteins, harboring the Jas domain, act as repressors of jasmonate signaling via their physical interactions with a wide array of transcription factors; their degradation releases these factors and subsequently activates downstream signal cascades. The bHLH transcription factors JASMONATE INSSENSITIVE1 (JIN1/MYC2), MYC3, and MYC4 function as direct targets of JAZ proteins to modulate a subset of jasmonate-regulated responses, such as the inhibition of root elongation and defense responses (Boter et al., 2004; Lorenzo et al., 2004; Chini et al., 2007; Dombrecht et al., 2007; Fernández-Calvo et al., 2011). The essential components of WD-repeat/bHLH/MYB transcriptional complexes, including bHLH proteins GLABRA3 (GL3), ENHANCER OF GLABRA3 (EGL3), and TRANSPARENT TESTA8 and MYB domain proteins GLABRA1 (GL1) and MYB75, also act as JAZ targets to mediate jasmonate-regulated biological processes (Qi et al., 2011). In addition, two other MYB transcription factors (MYB21 and MYB24) have recently been identified as JAZ targets to regulate jasmonate-mediated anther development and filament elongation (Song et al., 2011).

Although jasmonate has been implicated in cold storage of several tropical and subtropical fruits (González-Aguilar et al., 2000, 2004; Cao et al., 2009; Mohammad et al., 2011; Zhao et al., 2013), the exact role played by jasmonate in mediating freezing responses and the molecular mechanisms of these regulatory responses remain to be elucidated. In this study, we undertook a molecular and genetic approach to investigate the role of jasmonate in plant freezing stress responses. We found that exogenous application of jasmonate significantly improved Arabidopsis freezing tolerance, while blocking jasmonate biosynthesis and signaling decreased plant freezing tolerance. Mechanism investigation revealed that several JAZ proteins physically interact with ICE1 and ICE2 and repress their transcriptional functions and that overexpression of JAZ1 or JAZ4 represses responses to freezing stress. Our results thus provide evidence that jasmonate acts as an upstream signal of the ICE-CBF/DREB1 transcriptional pathway to positively regulate freezing stress responses of Arabidopsis.

RESULTS

Exogenous Application of Jasmonate Enhances Plant Freezing Tolerance

To explore the regulatory role of jasmonate in cold stress responses, we first examined whether the freezing tolerance of wild-type plants was affected by exogenous application of jasmonate under both nonacclimated and cold-acclimated conditions. In the presence of methyl jasmonate (MeJA; 3 and 5 \(\mu\)M), the wild-type plants were slightly smaller in size and accumulated more anthocyanin compared with nontreated plants (Figure 1A). Without cold acclimation, the survival rates of MeJA-treated plants were significantly higher than those of control plants after exposure to freezing temperatures of \(-4^\circ\) and \(-5^\circ\)C (Figures 1A and 1B). Similarly, under cold-acclimated conditions (2 weeks at \(4^\circ\)C), MeJA-treated plants also exhibited enhanced tolerance of freezing stress; their survival rates after freezing treatments (\(-8^\circ\) and \(-9^\circ\)C) were dramatically higher than those of control plants (Figures 1A and 1C). To further confirm the role of exogenous jasmonate on freezing tolerance, we analyzed the performances of soil-grown wild-type plants with or without MeJA treatments in response to freezing temperatures. As shown in Figures 1D and 1E, MeJA-treated soil-grown plants also displayed substantially increased tolerance to freezing stress under both nonacclimated and cold-acclimated conditions. Consistent with this, lower levels of relative electrolyte leakage, a parameter for evaluation of cold-induced membrane injury (Lyons, 1973), were observed in MeJA-treated plants than in control plants subjected to freezing temperatures (Figure 1F). These results suggested that jasmonate may have a positive role in modulating Arabidopsis freezing tolerance with or without cold acclimation.

Blocking Jasmonate Biosynthesis and Signaling Renders Plants Hypersensitive to Freezing Stress

To further determine jasmonate’s role in mediating freezing tolerance, we tested the involvement of jasmonate biosynthesis and signaling in freezing stress responses. ALLENE OXIDE SYNTHASE (AOS) and LIPOXYGENASE2 (LOX2) are key enzymes in jasmonate synthesis, whereas JASMONATE RESISTANT1 (JAR1) carries out a jasmonate conjugation/activation reaction (Song et al., 1993; Bell et al., 1995; Staswick et al., 2002). With or without acclimation, loss-of-function mutants lox2, aos, and jar1 displayed substantially decreased tolerance of freezing temperatures compared with wild-type plants (Figure 2A); their survival rates in response to freezing temperatures (nonacclimated plants to \(-4^\circ\) and \(-5^\circ\)C and cold-acclimated plants to \(-7^\circ\), \(-8^\circ\), and \(-9^\circ\)C) were dramatically lower than those of wild-type plants (Figures 2B and 2C). Consistent with these phenotypes, the relative electrolyte leakage levels following freezing (\(-4^\circ\) and \(-5^\circ\)C) were significantly higher in lox2, aos, and jar1 mutants than in control plants (Figure 2D). The F-box protein COI1 is the jasmonate receptor and a key positive regulator in the jasmonate signaling pathway (Xie et al., 1998; Yan et al., 2009). Freezing tolerance assays showed that disruption of COI1 rendered plants hypersensitive to freezing stress under both nonacclimated and cold-acclimated conditions compared with wild-type plants (Figures 2B to 2E). To further corroborate the biological role of jasmonate in modulating freezing tolerance, we analyzed the performances of soil-grown mutants under freezing stress. As shown in Figures 2F to 2H, with or without cold acclimation, the soil-grown mutants were also more sensitive to freezing temperatures than the wild-type plants. Taken together, these observations indicate that jasmonate biosynthesis and signaling positively regulate Arabidopsis freezing tolerance under both nonacclimated and cold-acclimated conditions.

Cold Stress Induces an Increase in Endogenous Jasmonate Levels

Having ascertained that jasmonate is involved in plant freezing tolerance, we then asked whether endogenous jasmonate production...
is affected by cold stress. To test this, we measured jasmonate production in wild-type plants following cold treatment. As shown in Figure 3A, endogenous jasmonate production was induced by cold treatment. To further understand the effect of cold stress on jasmonate production, we examined the expression of several genes encoding critical enzymes for jasmonate biosynthesis. As shown in Figure 3B, expression of analyzed genes was upregulated by cold stress. These results indicate that cold stress triggers the accumulation of endogenous jasmonate in Arabidopsis.

**Cold Induction of CBF/DREB1 and Their Targets Is Upregulated by Jasmonate**

Currently, the best understood freezing tolerance pathway is the CBF/DREB1 transcriptional regulatory cascade. As jasmonate is
Figure 2. Freezing Tolerance of Mutants Involved in Jasmonate Biosynthesis and Signaling.

(A) Freezing phenotypes of MS medium–grown lox2, aos, and jar1 mutant plants with or without cold acclimation. Nonacclimated (NA) 12-d-old seedlings were treated at −4°C for 1 h, and cold-acclimated seedlings (2 weeks at 4°C) were exposed to −8°C for 1 h, followed by recovery at 22°C for 4 d. Experiments were performed three times with similar results. WT, the wild type.

(B) and (C) Survival rates of various nonacclimated (B) and cold-acclimated (C) mutant plants after exposure to indicated freezing temperatures. Error bars show SD from three replicates. CA, cold-acclimated (2 weeks at 4°C); MS, seedlings grown on MS medium; NA, nonacclimated.

(D) Ion leakage assays of mutants treated with indicated freezing temperatures. Error bars show SD from three replicates. CA, cold-acclimated (2 weeks at 4°C); MS, seedlings grown on MS medium; NA, nonacclimated.

(E) Freezing phenotypes of coi1-1 and coi1-2 mutants with or without cold acclimation. Nonacclimated 12-d-old seedlings were treated at −4°C for 1 h, and cold-acclimated (2 weeks at 4°C) seedlings were exposed to −8°C for 1 h, followed by recovery at 22°C for 4 d. Experiments were performed three times with similar results.

(F) Freezing phenotypes of various soil-grown mutants with or without cold acclimation. Nonacclimated 18-d-old soil-grown seedlings were treated at −3 or −4°C for 1.5 h, and cold-acclimated soil-grown seedlings (2 weeks at 4°C) were exposed to −7 or −8°C for 1.5 h, followed by recovery at 22°C for 7 d. Experiments were performed three times with similar results using over 100 plants per treatment.

(G) and (H) Survival rates of various nonacclimated (G) and cold-acclimated (H) soil-grown mutant plants after exposure to indicated freezing temperatures. Error bars show SD from three replicates. CA, cold-acclimated (2 weeks at 4°C); NA, nonacclimated; Soil, seedlings grown in soil. *Differences between the mutant and the wild type are significant (P < 0.05). **Differences between the mutant and the wild type are highly significant (P < 0.01).
Initially, we analyzed expression levels of jasmonate and several of their targets, including CBF/DREB1. Initially, we analyzed expression levels of jasmonate, CBF/DREB1, and their target genes, including CBF/DREB1, to understand how jasmonate affects all cold-regulated gene expression. We also analyzed the expression of several cold-responsive genes acting in ABA or cytokinin signaling pathway and those associated with the synthesis of soluble sugars (functioning as cryoprotectant molecules) in coi1-1 mutants, such as ABSCISIC ACID RESPONSIVE ELEMENT BINDING FACTOR1 (ABF1), ARR5, and SUCROSE SYNTHASE. As shown in Supplemental Figure 2 online, their transcripts were not affected in coi1-1 mutants under cold stress, compared with those in wild-type plants. Taken together, these results indicate that jasmonate positively regulates the CBF/DREB1 transcriptional regulatory pathway in Arabidopsis.

**JAZ Proteins Physically Interact with ICE1 and ICE2**

To understand how jasmonate modulates the CBF/DREB1 cascade and freezing tolerance, we used the yeast two-hybrid system to identify potential downstream transcription factors of JAZ repressors. The full-length of JAZ1 was fused to the Gal4 DNA binding domain of the bait vector (BD-JAZ1). After screening, three independent clones encoding ICE1 were identified by prototrophy for His and Ade. To confirm the interaction, the full-length coding sequence (CDS) of ICE1 was cloned and introduced into the prey vector (AD-ICE1). The bait and prey vectors were cotransformed into yeast and the protein–protein interaction was reconstructed (Figure 5A). To confirm whether JAZ1 specifically interacts with ICE1, we also analyzed its interaction with the homolog of ICE1, ICE2. As shown in Figure 5A, JAZ1 also interacts with ICE2 in the yeast two-hybrid system. We further investigated interactions of ICE1 and ICE2 with all 12 Arabidopsis JAZ proteins in the yeast two-hybrid system. Besides JAZ1, ICE1 also strongly interacted with JAZ4 and JAZ9, and slightly interacted with JAZ3 and JAZ11 (Figure 5A). Similarly, ICE2 also interacted with JAZ4 and JAZ9 in yeast (Figure 5A). As a control, we examined the expression levels of the 12 BD-JAZ fusion proteins by immunoblot analysis, finding that all BD-JAZ fusion proteins were expressed in yeast (see Supplemental Figure 3 online).

Interactions of JAZ proteins with ICE1 and ICE2 in planta were further corroborated by bimolecular fluorescence complementation (BiFC) and communoprecipitation (CoIP) assays. JAZ1 and JAZ4 were used as representatives in the BiFC and CoIP assays. For the BiFC assays, JAZ proteins were fused to the C-terminal yellow fluorescent protein (YFP) fragment (JAZ-cYFP) and ICE factors to the N-terminal YFP fragment (ICE-nYFP). When fused JAZ1-cYFP was coexpressed with ICE1-nYFP or ICE2-nYFP in leaves of tobacco (Nicotiana benthamiana), the YFP signal was detected in the nuclear compartment of transformed cells, as revealed by staining with 4',6-diamidino-2-phenylindole (Figure 5B). Similar results were also observed for interactions of JAZ4 with ICE1 and ICE2 (Figure 5B). No fluorescence was detected in negative control experiments in which either JAZ-cYFP was coexpressed with unfused nYFP or unfused
cYFP was coexpressed with ICE-nYFP (Figure 5B). In addition to the BiFC assays, JAZ–ICE interaction was verified by CoIP assays using plant total protein (Figure 5C). Taken together, these results demonstrate that ICE1 and ICE2 interact with JAZ proteins in plant cell nuclei, implying that ICE1 and ICE2 function as direct targets of JAZ proteins.

The C-Terminal Fragment of ICE1 and the Jas Domain of JAZ1 Are Responsible for the Interaction

To investigate which region of ICE1 is required for interaction with JAZ proteins, we fused five truncated ICE1 variants to the Gal4 activation domain of the prey vector. The interaction between these derivatives and JAZ proteins was then assayed using the yeast two-hybrid system. As shown in Figure 6A, deletion of the N-terminal 260 residues of ICE1 (AD-ICE1, D1–260) did not affect its interactions with JAZ1 and JAZ4. By contrast, deletion of the C-terminal of ICE1 (AD-ICE1, D261–494) completely eliminated the interactions between ICE1 and JAZ proteins (Figure 6A; see Supplemental Figure 4A online). This finding demonstrates that the C-terminal domain of ICE1 is essential for the interaction with JAZ proteins. Further mapping revealed that the 74 amino acids at the C-terminal end are specifically responsible for the interaction because a derivative with amino acids 261 to 420 deleted from the C terminus of ICE1 could still interact with JAZ proteins (Figure 6A; see Supplemental Figure 4A online).

To identify the JAZ1 region responsible for the interaction with ICE1 and ICE2, we performed additional directed yeast two-hybrid analysis. JAZ1 was divided into N-terminal (BD-JAZ1, D121–253) and C-terminal (BD-JAZ1, D1–120) portions, and the C-terminal part was further truncated into the Jas domain (BD-JAZ1-Jas). The result showed that ICE1 and ICE2 interacted with both the C-terminal portion and the Jas domain of JAZ1 in yeast (Figure 6B; see Supplemental Figure 4B online), indicating that the Jas domain of JAZ1 is required for the interaction with ICE1 and ICE2.

Figure 4. Expression of CBF/DREB1 and Their Regulons in Response to Jasmonate under Cold Stress.

Eighteen-day-old wild-type (WT) and coi1-1 mutant plants were treated at 4°C with water or 100 μM MeJA for the indicated time periods. Error bars show so from three independent RNA extractions.
JAZ Proteins Repress the Transcriptional Function of ICE1

Because JAZ proteins directly interact with ICE1 and ICE2, we hypothesized that the physical interactions might interfere with transcriptional function of these factors. To test this possibility, we generated and analyzed a CBF3 promoter-driven CBF3-green fluorescent protein (GFP) fused protein (ProCBF3-CBF3-GFP) in a transient expression assay. When the reporter construct was transformed into the leaves of N. benthamiana and kept at 22°C, no fluorescence signal was observed (Figures 7A and 7B). However, if the reporter construct was induced at 4°C, fluorescence signals were detected in the nucleus (Figures 7A and 7B). When ProCBF3-CBF3-GFP was coinfiltrated into N. benthamiana leaves along with ICE1 driven by the CaMV35S cauliflower mosaic virus promoter (CaMV35S-ICE1), much stronger fluorescence signals were observed after 4°C treatment (Figures 7A and 7B). However, coinfiltration of ProCBF3-CBF3-GFP with CaMV35S-JAZ1 or CaMV35S-JAZ4 generated dramatically lower fluorescence levels at 4°C (Figures 7A and 7B). In addition, coinfiltration of ProCBF3-CBF3-GFP with CaMV35S-JAZ1 or CaMV35S-JAZ4 and CaMV35S-ICE1 also generated much weaker fluorescence signals in comparison with coinfiltration of ProCBF3-CBF3-GFP alone or coinfiltration of ProCBF3-CBF3-GFP with CaMV35S-ICE1 after cold treatment (Figures 7A and 7B). As a control, coinfiltration of ProCBF3-CBF3-GFP with CaMV35S-GUS and CaMV35S-ICE1 was performed, but no obvious differences in fluorescence signals were observed compared with coinfiltration of ProCBF3-CBF3-GFP with CaMV35S-ICE1 (Figures 7A and 7B). Taken together, these results demonstrate that JAZ proteins repress the transcriptional function of ICE1.

To further verify the effect of JAZ proteins on the transcriptional function of ICE1, we analyzed relative expression of CBF3-GFP in
N. benthamiana leaves in response to low temperature. As shown in Figure 7C, we detected high levels of CBF3-GFP transcripts in ProCBF3-CBF3-GFP and CaMV35S-ICE1 coinfected N. benthamiana leaves after cold treatment. By contrast, coexpression of JAZ proteins with ICE1 significantly suppressed the accumulation of CBF3-GFP transcripts (Figure 7C). These results further support the notion that JAZ proteins repress the transcriptional function of ICE1.

Overexpression of JAZ1 or JAZ4 Represses Freezing Tolerance

As several JAZ repressors interact with ICE factors and modulate the transcriptional function of ICE1, we queried whether disruption or overexpression of these JAZ proteins affected Arabidopsis freezing stress responses. To test this possibility, we first analyzed the performances of mutant plants in response to freezing stress. The tested jaz4 and jaz9 single and the jaz4 jaz9 double mutants behaved similarly to the wild-type plants after treatments, showing similar survival rates and induced expression levels of cold-responsive genes (see Supplemental Figure 5 online). However, overexpression of JAZ1 or JAZ4 rendered transgenic plants (JA-Z1OE or JA-Z4OE) sensitive to freezing stress (Figures 8A and 8B). Consistent with this, transcripts of CBF/DREB1 and their downstream targets were reduced in JAZ1OE and JAZ4OE under cold stress (Figure 8C). These results indicated that overexpression of JAZ1 or JAZ4 represses the ICE-CBF/DREB1 signaling pathway and freezing stress responses in Arabidopsis.

Previous studies have shown that disruption of the jasmonate receptor COI1 protein permits JAZ repressors to accumulate in the nucleus (Chini et al., 2007; Thines et al., 2007), thereby repressing downstream transcription factors. To further corroborate the regulatory effect of JAZ proteins on the ICE-CBF/DREB1 cascade in Arabidopsis, we investigated whether overexpression of ICE1 could restore the freezing-sensitive phenotype of coi1-1 mutants. As shown in Figures 8D and 8E, transgenic expression of ICE1 was able to rescue the phenotype of coi1-1 mutants in response to freezing stress. This observation further supports the idea that JAZ proteins repress the ICE-CBF/DREB1 transcriptional cascade in Arabidopsis.

Expression of Several CBF/DREB1 Pathway-Independent Genes Is Reduced in coi1-1

Our expression analysis showed that cold induction of well-known genes acting in the CBF/DREB1 signaling pathway was...
upregulated by jasmonate (Figure 4). To further investigate whether jasmonate modulates Arabidopsis cold acclimation–induced freezing tolerance exclusively through the CBF/DREB1 pathway, we profiled the transcriptomes of coi1-1 mutant plants after 4°C treatments for 6 or 24 h by microarray analysis. As shown in Supplemental Data Set 1 online, more than 50 cold-responsive genes were downregulated in coi1-1 mutants under cold stress, compared with wild-type plants. As expected, among the jasmonate-regulated genes were members of the CBF/DREB1 signaling pathway. In addition, several cold-responsive genes (such as
**Figure 8.** Phenotypic Characterization of the JAZ1 and JAZ4 Overexpression Plants.

(A) Survival rates of the JAZ1 and JAZ4 overexpression plants (JAZ1OE and JAZ4OE) with or without cold acclimation. Nonacclimated (NA) 18-d-old soil-grown plants were treated at −4°C for 1.5 h, and cold-acclimated seedlings (CA; 2 weeks at 4°C) were exposed to −8°C for 1.5 h, followed by recovery at 22°C for 7 d. Error bars show SD from three replicates. Soil, plants grown in soil. *Differences between overexpression plants and the wild type (WT) are significant (P < 0.05). **Differences between overexpression plants and the wild type are highly significant (P < 0.01).

(B) Ion leakage assays of the JAZ1 and JAZ4 overexpression plants (JAZ1OE and JAZ4OE) with indicated freezing temperatures. Error bars show SD from three replicates. *Differences between overexpression plants and wild type are significant (P < 0.05).

(C) Expression of CBF/DREB1 and their regulons in the JAZ1 and JAZ4 overexpression plants (JAZ1OE and JAZ4OE) under cold stress. Eighteen-day-old soil-grown plants were treated at 4°C for the indicated time periods. Error bars show SD from three replicates. *Differences between overexpression plants and wild type are significant (P < 0.05).

(D) Survival rates of coi1-1/ICE1OE plants with or without cold acclimation. Nonacclimated 18-d-old soil-grown plants were treated at −4°C for 1.5 h, and cold-acclimated seedlings (2 weeks at 4°C) were exposed to −8°C for 1.5 h, followed by recovery at 22°C for 7 d. Error bars show SD from three replicates.

(E) Ion leakage assays of coi1-1/ICE1OE treated with freezing temperatures. Error bars show SD from three replicates. coi1-1/ICE1OE, overexpression of ICE1 in coi1-1 background. Soil, plants grown in soil. *Differences between mutant plants and the wild type are significant (P < 0.05). **Differences between mutant plants and the wild type are highly significant (P < 0.01).
AT4G38960, AT2G27080, and GIGANTEA) that are not parts of the CBF/DREB1 pathway were also affected by jasmonate signaling under cold stress (see Supplemental Data Set 1 online). To confirm the reliability of the microarray data, we examined the cold-induced expression of several genes by quantitative RT-PCR (qRT-PCR) analysis. Consistent with the microarray results, the induced expression levels of these selected genes were lower in coi1-1 mutants than in the wild type (Figure 9; see Supplemental Figure 6 online). These results indicated that jasmonate may also modulate plant cold acclimation-induced freezing tolerance through the CBF/DREB1-independent pathways.

In addition to the cold acclimation–induced freezing tolerance, our results revealed that jasmonate also positively regulated Arabidopsis constitutive freezing tolerance. Under nonacclimated conditions, the jasmonate biosynthesis- and signaling-related mutant plants exhibited reduced freezing tolerance compared with wild-type plants (Figures 2), indicating that the constitutive levels of jasmonate are required for normal levels of Arabidopsis constitutive freezing tolerance. However, our expression analysis showed that the basic expression levels of CBF/DREB1 and their target genes were not affected by jasmonate signaling (Figure 4). These results indicated that jasmonate may modulate Arabidopsis constitutive freezing tolerance through CBF/DREB1-independent pathways. To understand how jasmonate modulates Arabidopsis constitutive freezing tolerance, we compared the transcriptomes of wild-type and the coi1-1 mutant plants grown under normal growth conditions. By comparing the microarray results, we identified several cold-responsive but CBF/DREB1 pathway-independent genes (such as GIGANTEA, AT4G38960, and AT3G22840) that showed lower expression levels in coi1-1 mutant plants (Figure 10A; see Supplemental Data Set 1 and Supplemental

**Figure 9.** Induced Expression of Genes Responsive to Cold but Independent of the CBF/DREB1 Signaling Pathway in Soil-Grown coi1-1 under Cold Stress.

Eighteen-day-old soil-grown wild-type (WT) and coi1-1 mutant plants were treated at 4°C for the indicated time periods. Error bars show ±S from three independent RNA extractions used for qRT-PCR.
Moreover, a number of genes encoding enzymes for synthesis of secondary metabolites (such as polyamine, glutathione, and anthocyanins) were also affected by jasmonate signaling (Figure 10B; see Supplemental Data Set 1 and Supplemental Figure 7B online). These results suggested that jasmonate may regulate Arabidopsis constitutive freezing tolerance through modulating the base expression of some CBF/DREB1 pathway-independent cold-responsive genes and the synthesis of cold stress–related secondary metabolites.

DISCUSSION

The plant hormone jasmonate is ubiquitous in the plant kingdom and is required for regulation of multiple physiological processes. In this study, we investigated jasmonate’s role in freezing stress in Arabidopsis by examining the effect of exogenous application of jasmonate on plant freezing tolerance, the effect of impaired jasmonate biosynthesis and signaling on freezing stress, and the changes in endogenous jasmonate levels in response to cold. Our results demonstrate that jasmonate has a positive role in both constitutive and cold acclimation–induced freezing tolerance of Arabidopsis. Treating wild-type plants with exogenous MeJA significantly enhanced tolerance to freezing stress (Figure 1). By contrast, blocking endogenous jasmonate biosynthesis and signaling conferred decreased freezing tolerance under both nonacclimated and cold-acclimated conditions (Figure 2). Furthermore, endogenous jasmonate production was triggered by cold treatment (Figure 3). We therefore conclude that jasmonate positively regulates Arabidopsis constitutive and cold acclimation–induced freezing tolerance.

Previous studies indicated that jasmonate-induced chilling tolerance of several fruits may be associated with the increase of cryo-protective compounds (González-Aguilar et al., 2000, 2004; Cao et al., 2009; Mohammad et al., 2011; Zhao et al., 2013). However, the exact molecular mechanisms of jasmonate-mediated cold stress responses remained unclear.

**Figure 10.** Base Expression of Genes Responsive to Cold but Independent of the CBF/DREB1 Signaling Pathway and Genes for the Synthesis of Secondary Metabolites in Soil-Grown coi1-1.

(A) Base expression of genes responsive to cold but independent of the CBF/DREB1 signaling pathway in 18-d-old soil-grown coi1-1. WT, the wild type.

(B) Base expression of genes encoding enzymes for the synthesis of polyamine, anthocyanins, and glutathione in 18-d-old soil-grown coi1-1. Error bars show sd from three independent RNA extractions used for qRT-PCR.
jasmonate-regulated responses requires a profound transcriptional reprogramming of cellular genetic programs involved in the complex interplay between negative and positive regulators (e.g., JAZ repressors and downstream transcription factors). JAZ proteins repress jasmonate-mediated responses through interaction and attenuation of their downstream transcription factors. The bHLH transcription factors MYC2, MYC3, and MYC4, essential components of the WD-repeat/bHLH/MYB transcriptional complexes such as GL3, EGL3, and GL1, MYB21 and MYB24, and ETHYLENE INSENSITIVE3 (EIN3), were recently identified as direct targets of JAZ proteins mediating various jasmonate-regulated processes (Chini et al., 2007; Fernández-Calvo et al., 2011; Qi et al., 2011; Song et al., 2011; Zhu et al., 2011). In this study, we found that the ICE1 and ICE2 transcription factors also function as targets of JAZ proteins (Figures 5 and 6) and demonstrated that JAZ proteins repress the transcriptional function of ICE1 (Figure 7). Consistent with these findings, overexpression of JAZ1 or JAZ4 repressed the cold-induced expression of CBF/DREB1 and their regulons, thereby rendering transgenic plants sensitive to freezing (Figure 8). Moreover, overexpression of ICE1 was able to rescue freezing sensitive phenotype of coi1-1 mutant plants (Figure 8).

The transcription factors ICE1 and ICE2 can activate expression of downstream CBF3/DREB1a and CBF1/DREB1b genes, respectively, in response to cold stress (Chinnusamy et al., 2003; Fursova et al., 2009). Our expression analysis showed that CBF3/DREB1a, CBF1/DREB1b, and their targets were induced by MeJA treatment under cold stress (Figure 4), consistent with the enhanced freezing tolerance of MeJA-treated wild-type plants and the repressive effect of JAZ proteins on ICE1 and ICE2 transcription factors (Figures 1 and 7). CBF2/DREB1c expression in response to cold stress was also upregulated by MeJA treatment (Figure 4). This was surprising because previous studies have suggested that CBF2/DREB1c expression is not influenced by ICE1 and ICE2 (Chinnusamy et al., 2003; Fursova et al., 2009); furthermore, cold-induced expression of CBF2/DREB1c was not found to be associated with CBF3/DREB1a and CBF1/DREB1b in a previous study (Novillo et al., 2007). Further research is thus required to investigate how jasmonate signaling affects CBF2/DREB1c expression under cold stress. Recently, Zhao et al. (2013) showed that two MYC2 proteins of banana (Musa acuminata) interact with ICE1. This suggests that banana MYC2 proteins may participate in regulating the expression of cold-responsive genes. However, in our study, the expression of three CBF/DREB1 genes was not affected in the myc2-2 mutant compared with the wild type under cold stress (see Supplemental Figure 8 online). This observation suggests that MYC2 may be not involved in cold stress responses in Arabidopsis.

Recently, Shi et al. (2012) demonstrated that ethylene signaling negatively regulates freezing tolerance in Arabidopsis. Further genetic and biochemical analysis in their study revealed that EIN3 and EIL1 directly bind CBF/DREB1 promoters and negatively regulate expression of these cold-induced genes. EIN3 and EIL1 were also identified in another study as direct targets of JAZ repressors mediating jasmonate-regulated responses (Zhu et al., 2011). Consequently, jasmonate signaling appears to negatively modulate CBF/DREB1 gene expression via EIN3 and EIL1. We speculate that the dual regulations of the CBF/DREB1 signaling pathway by jasmonate may be a balancing mechanism between establishment of the appropriate stress tolerance and minimization of detrimental effects on plant growth and development. Nevertheless, the final outcome of jasmonate regulation is the enhancement of plant tolerance to freezing stress.

Under normal growth conditions, JAZ repressors interact with ICE transcription factors. The physical interactions attenuate transcriptional functions of these downstream transcription factors. Under cold conditions, production of endogenous jasmonate is induced. The COI1 receptor perceives jasmonate and targets JAZ proteins for degradation. Upon degradation of JAZ proteins, the ICE factors are subsequently released to activate CBF/DREB1 gene expression and their downstream cold-responsive genes. Jasmonate also regulates Arabidopsis constitutive and cold acclimation–induced freezing tolerance by influencing the synthesis of secondary metabolites and via some CBF/DREB1-independent pathways. The final outcome of jasmonate regulation is the enhancement of plant tolerance to freezing stress.

Figure 11. Model for Jasmonate-Regulated Cold Signaling Pathway in Arabidopsis.

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In addition to the CBF/DREB1 signaling pathway, our results revealed that jasmonate also regulates expression of several genes independent of the CBF/DREB1 signaling pathway under cold stress (see Supplemental Data Set 1 online; Figure 9). These findings indicate that jasmonate also modulates plant cold acclimation through the CBF/DREB1-independent pathways. Fowler and Thomashow (2002) showed that only 12%
of the cold-responsive genes were regulated by CBF/DREB1 transcription factors, suggesting that the CBF/DREB1-independent pathways may also play important roles in mediating plant cold stress responses. Interestingly, endogenous jasmonate has a peak of induction by low temperature at 12 h (Figure 3), which appears to be inconsistent with the results that the cold-induced expression of CBF/DREB1 genes peaks at 3 h and decreases at 12 h of cold treatment (Figure 4). It is possible that the high levels of endogenous jasmonate at 12 to 24 h may be involved in the induction of CBF/DREB1-independent genes. Consistent with this possibility, most of the selected genes independent of the CBF/DREB1 signaling pathway were strongly induced at 12 to 24 h by cold treatment (Figure 9). Further investigation of these jasmonate-regulated pathways (Figures 9 and 10; see Supplemental Data Set 1 online) revealed several common chemicals were obtained from Shanghai Sangon. Taq DNA polymerases were purchased from Takara Biotechnology, and other common chemicals were obtained from Shanghai Sangon. Arabidopsis thaliana plants were grown in an artificial growth chamber at 22°C under a 10-h-light/14-h-dark photoperiod. Wild-type and mutant plants used in this study were in the Columbia-0 genetic background. The mutants coi1-1 (Xie et al., 1998) and coi1-2 (Xu et al., 2002) were described as previously. Other mutants used in this study are listed as follows: tox2 (CS3748), aox1 (CS6149), jar1 (CS8072), and myc2-1 (CS6320). To generate the overexpression transgenic plants, the full-length cDNA of JAZ1, JAZ4, or ICE1 were cloned into the pOCA30 vector in the sense orientation behind the CaMV 3SS promoter (Hu et al., 2013). All primers used for clones are listed in Supplemental Table 1 online.

Freezing Tolerance Assays

The freezing tolerance assays were performed as described (Chinnusamy et al., 2003), with some modifications. Briefly, 12-d-old plants grown at 22°C on Murashige and Skoog (MS) agar medium with or without cold acclimation (2 weeks at 4°C) were placed in a freezing chamber set to −1°C and programmed to cool at −1°C per hour. Petri dishes of plants were removed after exposure to the desired temperatures. After the freezing treatment, plants were incubated at 4°C in the dark for 10 h and then transferred to light at 22°C. The survival rates of the seedlings were scored visually after 4 d. For soil-grown plants, 18-d-old plants with or without cold acclimation (2 weeks at 4°C) were placed in a freezing chamber set to 0°C and programmed to cool at −1°C per 1.5 h. Plants were removed after exposure to the desired temperatures. After the freezing treatment, plants were incubated at 4°C in the dark for 10 h and then transferred to light at 22°C. The survival rates of the seedlings were scored visually after 7 d. For jasmonate treatment, 3 or 5 μM MeJA was added to the MS agar medium. Seeds of the wild type were germinated on this MeJA-containing medium, and seedlings were grown at 22°C for 12 d. Soil-grown plants were sprayed with water or a 30 μM MeJA solution diluted from the stock. The water- and MeJA-treated plants were maintained in a small chamber with a transparent cover for 3 d before freezing treatment (under nonacclimated conditions). Under cold-acclimated conditions, plants were treated with MeJA at 4°C for 3 d and then were removed from the chamber and kept at 4°C for another 11 d.

Relative Electrolyte Leakage

To measure relative electrolyte leakage, 12-d-old Arabidopsis seedlings were treated at freezing temperatures. After treatment, entire seedlings were harvested and relative electrolyte leakage was measured according to Jiang et al. (2007).

Determination of Jasmonate Contents

To measure jasmonate contents, 18-d-old soil-grown wild-type plants were treated at 4°C or kept at 22°C. After treatment, entire seedlings were harvested and jasmonate was analyzed by ELISA using monoclonal antibodies of jasmonate according to Deng et al. (2008).

RNA Extraction and qRT-PCR

Total RNA was extracted from Arabidopsis seedlings using the Trizol reagent (Invitrogen). qRT-PCR was performed as described by Hu et al. (2012). Briefly, first-strand cDNA was synthesized from 1.5 μg DNase-treated RNA in a 20-μL reaction volumes using M-MuLV reverse transcriptase. MATERIALS AND PLANT GROWTH CONDITIONS

The plant hormone MeJA was purchased from Sigma-Aldrich. Taq DNA polymerases were purchased from Takara Biotechnology, and other common chemicals were obtained from Shanghai Sangon. Arabidopsis thaliana plants were grown in an artificial growth chamber at 22°C under a 10-h-light/14-h-dark photoperiod. Wild-type and mutant plants used in this study were in the Columbia-0 genetic background. The mutants coi1-1 (Xie et al., 1998) and coi1-2 (Xu et al., 2002) were described as previously. Other mutants used in this study are listed as follows: tox2 (CS3748), aox1 (CS6149), jar1 (CS8072), and myc2-1 (CS6320). To generate the overexpression transgenic plants, the full-length cDNA of JAZ1, JAZ4, or ICE1 were cloned into the pOCA30 vector in the sense orientation behind the CaMV 3SS promoter (Hu et al., 2013). All primers used for clones are listed in Supplemental Table 1 online.

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The full-length JAZ1 CDS was cloned into the bait vector pGBKT7 and

expression. Gene-specific primers used to detect transcripts are listed in Supplemental Table 2 online.

Yeast Two-Hybrid Screening and Confirmation
The full-length JAZ1 CDS was cloned into the bait vector pGBKT7 and then transformed into the yeast strain Y2HGold (Clontech). The cDNA library was obtained from Clontech (catalog number 630487). Two-hybrid screening was performed via the mating protocol described in Clontech’s Matchmaker Gold Yeast Two-Hybrid user manual. To confirm protein–protein interactions, the full-length ICE1, ICE2, or truncated ICE1 CDS were cloned into the prey vector pGADT7. Primers used for amplifying these truncated or mutated fragments are listed in Supplemental Table 1 online.

BiFC Assays
cDNA sequences of the N-terminal, 173-amino acid, enhanced YFP (nYFP) and C-terminal, 64-amino acid (cYFP) fragments were PCR amplified and cloned into pFGC5941 to generate pFGC-nYFP and pFGC-cYFP, respectively (Kim et al., 2008). The full-length JAZ1 and JAZ4 CDS was inserted into pFGC-cYFP to generate C-terminal in-frame fusions with cYFP, while ICE1 and ICE2 CDS were introduced into pFGC-nYFP to form N-terminal in-frame fusions with nYFP. The resulting plasmids were introduced into Agrobacterium tumefaciens (strain GV3101), and infiltration of Nicotiana benthamiana was performed as described previously (Hu et al., 2013). Infected tissues were analyzed 48 h after infiltration. YFP and 4′,6-diamidino-2-phenylindole fluorescence were observed under a confocal laser scanning microscope (Olympus).

CoIP Assays
The full-length CDS of JAZ1, JAZ4, ICE1, and ICE2 were individually cloned into tagging plasmids behind the MYC or FLAG tag sequence in the sense orientation behind the CaMV 35S promoter. The constructs were transformed into Agrobacterium GV3101, MYC-fused JAZ1 or JAZ4 and FLAG-fused ICE1 or ICE2 were then transiently coexpressed in N. benthamiana. Infected leaves were sectioned 48 h after infiltration. CoIP assays were performed using leaf protein extracts as described by Shang et al. (2010). Briefly, MYC-fused JAZ1 and JAZ4 were immunoprecipitated using an anti-MYC antibody and the communoprecipitated protein was then detected using an anti-FLAG rabbit antibody (Sigma-Aldrich).

Transient Expression Assays
The transient expression assays were performed in N. benthamiana leaves. The full-length CDS of CBF3 was fused with GFP reporter gene behind the native promoter of CBF3. The full-length CDS of JAZ1, JAZ4, and ICE1 were driven by the CaMV 35S promoter. These constructs were then introduced into the Agrobacterium strain GV3101. The infiltration of N. benthamiana was performed as described previously (Kim et al., 2008). Plants were kept at 22°C for 48 h before induced by cold stress (4°C for 4 h). Infected leaves were observed under a confocal laser scanning microscope (Olympus). All experiments were repeated with five independent biological replicates with similar results.

Microarray Analysis
Eighteen-day-old soil-grown wild type and coi-1-1 mutant plants were used for microarray analysis. Wild-type and coi-1-1 mutant plants grown under normal growth conditions were treated at 4°C for 6 or 24 h. Total RNA was isolated from three replicates of control or treated plants using the Trizol reagent (Invitrogen). RNA quantity was assessed using the NanoDrop ND-1000 and the integrity was assessed using standard denaturing agarose gel electrophoresis. The microarray analysis was performed by the Shanghai Kangchen Biological Technology Company. For microarray analysis, the Agilent Array platform was employed. The sample preparation and microarray hybridization were performed based on the manufacturer’s standard protocols. Briefly, total RNA from each sample was amplified and transcribed into fluorescent complementary RNA using Agilent’s Quick Amp Labeling protocol (version 5.7; Agilent Technologies). The labeled cRNAs were hybridized onto the Whole Genome Oligo Microarray (4x44K; Agilent Technologies). After the slides were washed, the arrays were scanned by the Agilent Scanner G2505C. Agilent Feature Extraction software (version 11.0.1.1) was used to analyze acquired array images. Quantile normalization and subsequent data processing were performed using the GeneSpring GX v11.5.1 software package (Agilent Technologies). Differentially expressed genes were identified through fold change filtering.

Accession Numbers
Arabidopsis Genome Initiative numbers for the genes discussed in this article are as follows: LOX1, AT1G55020; LOX2, AT3G45140; LOX3, AT1G17420; LOX4, AT1G25250; AOS, AT5G2650; AOC1, AT3G25760; AOC2, AT3G25770; AOC3, AT3G25780; AOC4, AT1G13280; JAR1, AT2G46370; CO1, AT2G39940; JAZ1, AT1G19180; JAZ2, AT1G74850; JAZ3, AT3G17860; JAZ4, AT1G48500; JAZ5, AT1G17380; JAZ6, AT1G72450; JAZ7, AT2G34600; JAZ8, AT1G30135; JAZ9, AT1G70700; JAZ10, AT5G13220; JAZ11, AT3G43440; JAZ12, AT5G20900; ICE1, AT3G26744; ICE2, AT1G12860; CBF1, AT4G5490; CBF2, AT4G25470; CBF3, AT4G25480; RD29A, AT5G2310; RD29B, AT5G2300; KIN1, AT5G15960; COR47, At1g20440; COR414, At1g29395; COR15b, At2g42530; ADC1, AT2G16500; SAMDC, AT3G02470; SPD1, AT1G23820; DFR, AT5G42800; LDOX, AT4G22880; UF37G, AT5G54060; GSH1, AT4G23100; GSTu4, AT2G39460; GSTF12, AT5G17220; GIGANTEA, AT1G22770; WRKY22, AT4G01250; and ACTIN2, AT3G18780.

Supplemental Data
The following materials are available in the online version of this article.

Supplemental Figure 1. Cold-Induced Expression of CBF/DREB1 and Their Regulons in Response to Jasmonate in MS Medium

Supplemental Figure 2. Induced Expression of Genes Responsive to Cold under Cold Stress

Supplemental Figure 3. Base Expression of Genes Responsive to Cold but Independent of the CBF/DREB1 Signaling Pathway

**Supplemental Figure 8.** Expression of CBF/DREB1 Genes in myc2-2 Mutant Plants under Cold Stress.

**Supplemental Table 1.** Primers Used for Generating Clones.

**Supplemental Table 2.** Primers Used for qRT-PCR.

**Supplemental Data Set 1.** MS Excel File with Probe Names, Fold-Change Information, AGI Numbers, and Annotation of Cold-Regulated Genes Affected by Jasmonate Signaling.

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AUTHOR CONTRIBUTIONS

Y.H. designed and performed experiments, analyzed data, and wrote the article. L.J. and F.W. performed experiments and helped to analyze data. D.Y. designed experiments and helped to interpret data and edit the article. All authors read and approved the final article.

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Jasmonate Regulates the INDUCER OF CBF EXPRESSION–C-REPEAT BINDING FACTOR/DRE BINDING FACTOR1 Cascade and Freezing Tolerance in *Arabidopsis*
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