Co-regulation role of endogenous hormones and transcriptomics profiling under cold stress in *Tetrastigma hemsleyanum*

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Abstract

Background: *Tetragonia hemsleyana* Diels et Gilg is a valuable medicinal herb, Chilling sensitivity is the dominant environmental factor limiting the artificial domestication of the plant. Hormone-related gene regulation and hormone signaling pathways in response to cold stress in *T. hemsleyana* remain unknown.

Results: Some key genes involved in hormones biosynthesis, such as ZEP and NCED genes of ABA biosynthesis, GA2ox, GA3ox, and GA20ox genes of GA biosynthesis, ACO genes of ET biosynthesis pathway were screened to be crucial in cold response. Consistently, the response of ABA and ABA/GA1+3 to cold stress was prior to that of GA1+3, ZR, ABA/IAA, and ABA/ZR. The increasing changes in ABA/GA1+3 turned to a steep decline with the extension of stress time, which might be one factor contributing to cold-sensitivity of *T. hemsleyana* under prolonged stress. The cold tolerance of *T. hemsleyana* would be enhanced by ABA but repressed by GA3 when each phytohormone was applied alone. The ABA-mediated promotion and GA-mediated repression of cold tolerance could both be attenuated by the co-application of the two phytohormones within 6h. When the biosynthesis of endogenous ABA and GA were inhibited by FLU and PAC, respectively, the effects of GA and ABA treatment were reversed partially.

Conclusions: In summary, we presented the first study of global expression patterns of hormone-regulated transcripts in *T. hemsleyana*. This study suggested that GA and ABA could work antagonistically to balance the responses to cold in *T. hemsleyana*. PAC, a GA biogenesis inhibitor, as well as exogenous ABA, might be potential plant growth regulators that can promote cold tolerance of *T. hemsleyana*. The study also provided valuable hints in revealing a new theoretical basis and potential candidate genes that govern cold tolerance of *T. hemsleyana*.

Background

Tetragonia hemsleyana Diels et Gilg is a perennial herb, and its underground root contains abundant flavonoids and polysaccharides that are beneficial to human health [1, 2]. Due to over exploitation, wild resources are on the verge of extinction in recent years. However, it is considered to be a cold-sensitive species, with the best growing temperatures ranging from 20°C to 30°C. Low temperature susceptibility has become the dominant environmental factor limiting the artificial domestication of *T. hemsleyana*. So the development of cold stress tolerance in *T. hemsleyana* is important for improving production but it is hindered by lack of more precise physiological knowledge. A better understanding of the gene expression profile of *T. hemsleyana* under cold stress is imperative, which could be an important theoretical basis to study cold tolerance mechanisms for improving the cold resistance.

Studies have been reported to provide evidence of global changes in gene expression in response to cold stress [3, 4], which highlighted the significance of transcriptional regulation in plant adaptation to low temperature. Phytohormone are generally considered to participate in the regulation of organ differentiation and development. Recently, the development of stress tolerance has been reported in some plants to be accompanied by changes in hormonal-related genes expression and subsequently physiochemical modifications [5]. Plants could survive from stress by utilizing hormones to regulate and coordinate growth. These hormones-related genes could act in the upstream of stress response and trigger hormone signal pathway.

Absciscic acid (ABA) has been shown to mediate plant responses to various kinds of environmental stresses like cold [6], drought [7], and so on. Exogenous ABA application before cold stress improved resistance of sugarcane seedlings by increasing contents of proline and soluble sugar [8]. Recent studies showed the existence of potential co-regulation among the ABA and other hormones in plants. For example, auxin controlled seed dormancy by activating the ABA response [9]. Changes in ABA status can regulate cytokinin (CK) concentrations via altering cytokinin oxidase activity [10]. Moreover, the balance between ABA and gibberellic acid (GA) has been proposed to be correlated with stress responses [11]. Both the elevated content of endogenous ABA and the higher proportion of ABA/IAA and ABA/GA played an important role in kiwifruit resistance to cold [12].

However, hormone-related gene regulation and hormone signaling pathways in response to cold stress in *T. hemsleyanum* remain unknown. Our aim was therefore to elucidate the effects, as well as the transcriptomic adaptive mechanisms of cold stress on endogenous plant hormones in *T. hemsleyanum*. The results will facilitate the identification of potential candidate genes involved in cold stress tolerance in *T. hemsleyanum*, which should contribute to improving cold tolerance in future breeding programs, and should contribute to our understanding of the hormone responsive regulatory mechanisms of plant in response to environmental stress.

Results

In previous research, we measured the phenotypes and some physiological and biochemical parameters [13]. Our results showed that the 0 °C exposure was the optimum treatment temperature. The physiological and biochemical parameters remained essentially stable during 0 h and 8 h, significantly changed from 12 h to 48 h, and remained stable after 48 h. So RNA-seq was employed to investigate the changes in gene expression under 0 °C exposure for four time points (0, 12, 24, and 48 h).

Overview Of Transcriptome Sequencing

Four *T. hemsleyanum* leaves cDNA libraries were constructed and sequenced. A total of 216,043,064 raw reads were generated. After removing low-quality reads and adapter sequences, 181,412,182 clean reads were obtained, with 45,346,334; 46,460,402; 44,804,246; 44,801,200 clean reads generated for the 0, 12, 24, and 48 h treatment group libraries, respectively.

We obtained 106,275 unigenes annotated in Nr, NT, BLASTX, and BLASTP databases with an average length of 676 bp and N50 of 1121 bp (Fig. 1). Among these unigenes, 18082 unigenes had hits in all four public databases with informative annotations. 53,511 unigenes had hits that significantly matched against the Nt database (50.35% of all unigenes). According to Nt annotations, 41,743 unigenes were matched to *Vitis vinifera*, followed by *Zea mays*. The E-value distribution of the Nr blast results indicated that 81.91% of the matched sequences (43,831) had strong homologies (smaller than $1.0e^{-60}$), while the other 18.09% had homologies that ranged between $1.0e^{-5}$ to $1.0e^{-60}$. To produce more definitive annotations, the Nr, BLASTX, and BLASTP database were also used to align all of the unigenes. 43,827; 28,775 and 19,598 high-score annotate unigenes were matched, respectively.

Go Enrichment Analysis Of Degs

To characterize the functional classifications of differentially expressed genes, GO and KEGG analyses were performed to access the distributions of functional categories. In the 12, 24, and 48 h cold treatment samples,

8205(72.4%), 14400(83.5%), and 15402(85.1%) unigenes were annotated in GO, respectively. In the 12 h treatment group (Fig. 2A), 'membrane' (1194; 77.5% of 1541), 'membrane part' (986; 63.4% of 1541), and 'intrinsic component of membrane' (943; 63.4% of 1541) were the terms that dominated in the cellular component category. Response to stimulus(972; 48.1% of 2022) 'response to stress(652; 32.3% of 2022)' 'defense response(329; 16.3% of 2022)' 'hormone-mediated signaling (164; 8.11% of 2022)' were the most representative terms in the biological process category. 'oxidoreductase activity' (397; 25.88% of 1534), 'nucleic acid binding transcription factor activity' (247; 16.0% of 1534) and 'sequence-specific DNA transcription factor binding transcription factor activity' (247; 16.0% of 1534) were the most abundant terms in the molecular function category. During the 24 h (Fig. 2B) and 48 h (Fig. 2C) cold stress treatment, 'response to stimulus', 'response to stress' remained relatively stable in the biological process category, while 'response to hormone' and 'response to endogenous stimulus' skyrocketed in 24 h cold treatment samples. 'membrane', 'membrane part', 'intrinsic component of membrane' remained relatively stable in the cellular component category, while 'integral component of membrane' was suddenly increased in 24 h cold treatment samples. 'oxidoreductase activity' remained predominating during the whole cold stress. In the 12 h cold treatment samples, various of transcription factor activity were the most abundant terms in the molecular function category, such as 'nucleic acid binding transcription factor activity', 'sequence-specific DNA transcription factor binding transcription factor activity'. But with the increase of cold stress time, 'kinase activity' and 'catalytic activity' dramatically predominated in the molecular function category.

In general, 'response to stimulus', 'response to stress', 'membrane', 'membrane part', 'intrinsic component of membrane', 'oxidoreductase activity' remained predominating during the whole cold stress, while some biological process categories related to endogenous hormone response skyrocketed in the early stress stage, such as 'hormone-mediated signaling' 'response to hormone' and 'response to endogenous stimulus' in 12 h and 24 h. 'single organism process', 'integral component of membrane', and 'kinase activity' were suddenly increased in the intermediate stress stage, 'catalytic activity' dramatically predominated in the late stress stage.

Kegg Pathways Enrichment Analysis Of Degs

Out of total annotated unigenes, 12,936 unigenes were significantly matched to KEGG database and were assigned to 272 KEGG pathways. Pathways displaying significant changes ($Q \text{ value} \leq 0.05$) in response to the cold treatment were identified in each comparison group. The numbers of up-regulated genes and down-regulated genes in enriched pathways for each comparison group were listed in Fig. 3. In the 12 h (Fig. 3A), 24 h (Fig. 3B), and 48 h (Fig. 3C) cold treatment samples, 12, 6, and 11 KEGG pathways were significantly enriched, respectively. The top 5 enriched pathways of the 12 h cold treatment sample were 'metabolic pathways', 'biosynthesis of secondary metabolites', 'starch and sucrose metabolism', 'phenylpropanoid biosynthesis', and 'phenylalanine metabolism'. The top 5 enriched pathways of the 24 h cold treatment sample were 'metabolic pathways', 'biosynthesis of secondary metabolites', 'phenylpropanoid biosynthesis', 'steroid biosynthesis', and 'diterpenoid biosynthesis'. In the 48 h cold treatment sample, 'metabolic pathways', 'biosynthesis of secondary metabolites', 'phenylpropanoid biosynthesis' were also the top three enriched pathways, but it is noteworthy that various degradation metabolisms started to appear in enriched pathways, such as 'bisphenol degradation', 'polycyclic aromatic hydrocarbon degradation', 'limonene and pinene degradation'.

In general, 'metabolic pathways', 'biosynthesis of secondary metabolites', and 'phenylpropanoid biosynthesis' were common regulated during the whole cold stress stage. The cold responses were characterized by protective response through stimulating secondary metabolism and inhibiting primary metabolism to modulate cellular

metabolic homeostasis. In the early stress stage, primary metabolism still remained vigorous, such as 'starch and glucose metabolism' and 'phenylalanine metabolism', while the related secondary metabolism enzyme activities were gradually activated. It would lead to the stimulation of secondary metabolisms, such as 'steroid biosynthesis', 'diterpenoid biosynthesis' and so on. With the increase of cold stress time and chilling injury, many degradation pathways were strengthened, such as 'bisphenol degradation', 'polycyclic aromatic hydrocarbon degradation', 'limonene and pinene degradation', and ultimately led to cell apoptosis and necrosis.

Responses Of Hormone-related Genes

Based on the above overview of gene ontology enrichment and KEGG pathways present during cold treatment, we speculated that endogenous hormone-mediated signaling process would play an important role in early-time responses of *T. hemsleyanum* under cold exposure. So here we mainly focused on groups of differentially expressed genes involved in different plant hormone biosynthesis pathways, such as Auxin, abscisic acid (ABA), jasmonic acid (JA), gibberellic acid (GA), brassinosteroid (BR), cytokinin (CK) and ethylene (ET) pathways. Some representative hormone-related genes that were differentially expressed during the cold treatment are shown in Supplementary file 1–7. The heat map of hormones-related transcripts of *T. hemsleyanum* during 0 to 48 h of cold stress are shown in Fig. 4.

Absciscic acid (ABA) is an important phytohormone in plant stress response. Its metabolic changes are one of the determinants of endogenous ABA levels and control diverse stress response. Among the mapped enzymes, we identified four 9-cis-epoxycarotenoid dioxygenase (NCED) homologs of the ABA biosynthesis pathway in annotated transcriptome unigene dataset, and all of them were highly expressed with RPKM values > 100. Two of them were significantly up-regulated by 2.0-2.5 folds at 24 h. Zeaxanthin epoxidase (ZEP) is another key enzyme in ABA biosynthesis. Three highly expressed ZEP genes were identified, with c95330_g1 and c81504_g1 being the most abundantly and the remarkable differentially expressed. c35330_g2 was the most particularly noteworthy ZEP gene, which was almost undetectable in control and was significantly up-regulated by 9.8 folds and 21 folds under 24 h and 48 h of cold treatment, with an increased RPKM value from 0.024 to 0.237 and 0.511, respectively.

Four indole-3- acetaldehyde oxidase(AAO1) genes, two cellulose synthase A catalytic(CESA)genes and three abscisic-aldehyde oxidase(AAO3) genes were all down-regulated significantly under cold stress. So in general, in the ABA biosynthesis pathway, most of ZEP genes were strongly induced significantly during early cold stress, most of NCED genes were up-regulated significantly during the medium stress stage, while other genes were down-regulated significantly. But with the increase of cold stress time and chilling injury, all the above genes were down-regulated significantly at 48 h.

Four Gibberellin 20 oxidase(GA20ox) genes were identified in our dataset, and the fact of c98675_g1 and c93517_g2 were the most highly expressed suggested they may be crucial for gibberellic acid (GA) biosynthesis, with a decreased RPKM value from 214 to 21 and from 430 to 20 under cold stress, respectively. Six Gibberellin 2-beta-dioxygenase(GA2ox) genes were identified in our dataset. c92581_g1 was the most highly expressed GA2ox gene, with an increased RPKM value from 1051 to 3426 under cold stress. c93844_g1 was the most remarkable differentially expressed GA2ox gene, with a 7.6 folds significant up-regulation under cold stress, the RPKM value increased from 0.339 to 2.586. Four GA3ox were identified, among which c78816_g1 was the most highly expressed, with a 6.2 folds significant up-regulation under cold stress.

All differentially expressed genes in the auxin biosynthesis pathway were down-regulated, including three indole-3-acetaldehyde oxidase (AAO1) genes, four indole-3-pyruvate monooxygenase (YUCCA) genes, and three abscisic-aldehyde oxidase (AAO3) genes.

Five CK riboside 5'-monophosphate phosphoribohydrolase (CRMP_LOG) genes were differentially expressed in cytokinin (CK) biosynthesis pathway. Among them, LOG1, 5, 7 were up-regulated, while LOG3, 8 were down-regulated significantly. and the fact of c72906_g1 was the most abundantly and the most remarkable differentially expressed suggested it might be crucial for CK biosynthesis, with an increased RPKM value from 624 to 2292 after 24 h of cold treatment.

Eight 1-aminocyclopropane-1-carboxylate oxidase (ACO) genes of ethylene (ET) biosynthesis pathway were differentially expressed under cold treatment. Most of them were up-regulated in early stress stage and then maintained a high level, among which c90679_g3 was the most highly and the most remarkably up-regulated ACO gene, with an increased RPKM value from 12152 to 68974 after 24 h of cold treatment. However, three 1-aminocyclopropane-1-carboxylate synthase (ACS) genes of ET biosynthesis pathway were identified to be differentially expressed during cold acclimation. Two of them were down-regulated significantly, with a decreased RPKM value from 112 to 3.2 and from 12 to 1 after 24 h of cold treatment, respectively.

One 3-ketoacyl-CoA thiolase (KCT) gene, three 4-coumarate-CoA ligase (4CL) genes and one allene oxide cyclase (AOC) genes in jasmonic acid (JA) biosynthesis pathways were identified in our dataset. KCT and 4CL genes were highly expressed with RPKM values > 100 and were significantly down-regulated after 24 h of cold treatment. AOC gene, known as a key enzyme in JA biosynthesis, was significantly up-regulated by 10.3 and 9.2 folds at 24 h and 48 h of cold treatment, with an increased RPKM value from 0.024 to 0.251 and 0.223, respectively..

Validation of the differentially expressed genes (DEGs) via RT-qPCR.

To verify the reliability and accuracy of this Illumina RNA-seq data, quantitative real-time reverse transcription-PCR (qRT-PCR) was used to assess the relative expression levels for 14 selected key genes of six endogenous hormone synthesis pathways in every treatment group. GAPDH was selected for internal control reference gene. The correlation between qRT-PCR and RNA-seq was measured by scatter plotting log₂-fold changes, which showed a positive correlation coefficient in both techniques (Fig. 5). This independent evaluation revealed the reliability of the RNA-seq data.

Influence of cold stress on endogenous hormone contents and hormone ratios

The effect of cold stress on the endogenous hormone contents were examined. The dynamic changes in ABA, GA₁₊₃, IAA and ZR in three stages under 25°C or 0°C were shown in Fig. 6. The result indicated that there were no significant differences in all the four endogenous hormone contents between 0–48 h under 25°C. During 0°C cold stress, the ABA content strongly increased by 1.75- and 1.83-fold ($P < 0.01$) at 12 and 24 h, compared with the control, respectively. And then gradually decreased to 25% higher than the control. GA₁₊₃ levels increased to peak at 24 h by 161% and then maintained the higher level. The ratio of ABA to GA first skyrocketed to peak at 12 h by

88.3% and then decreased to a very low level. IAA content basically maintained constant. The ratio of ABA to IAA increased by 151.8%, 238.5% and 63.7% at 12 h, 24 h and 48 h, respectively. On the contrary, ZR levels decreased significantly at 24 h by 42.8% ($P < 0.05$) and then maintained the low level. Accordingly, the elevated ratio of ABA to ZR ranged from 121.5–395.8% under cold stress. The most significant increase was observed at 24 h by 395.8% as compared to the control.

In general, ABA content was strongly induced by cold stress in the early stages, $GA_1 + 3$ content was significantly higher in the middle-late stages, while ZR content was significantly lower, especially in the middle-late stages. The changes of ABA/GA, ABA/IAA, and ABA/ZR ratio under cold stress showed an increasing trend in the early-middle stages, but then gradually decreased as the stress time prolonged.

Effect of exogenous phytohormones and biosynthesis inhibitors on cold stress response

A systematic pilot experiments were conducted by pot experiments to determine the effects of multiple plant growth regulators on cold stress response of *T. hemsleyanum*. The results showed that GA_3 and ABA were the most important phytohormones affecting the stress related indicators.

After exogenous hormonal applications, the stress related indicators, including MDA content, soluble protein content and electrolyte leakage were simultaneously analyzed. As shown in Fig. 7A, ABA-treated seedlings had significantly more soluble protein content than control. ABA spraying dramatically decreased the MDA content and electrolyte leakage. The MDA content plummeted from $7.1 \pm 1.4 \mu\text{mol/g}$ (CK) to $5.1 \pm 1.1 \mu\text{mol/g}$ (ABA). The electrolyte leakage plummeted from $73.5 \pm 8.4\%$ (CK) to $50.0 \pm 11.2\%$ (ABA). FLU-treatment drastically increased electrolyte leakage and MDA content to $94.1 \pm 13.2\%$ and $11.2 \pm 1.3 \mu\text{mol/g}$, respectively. By contrast, the combined use of ABA and FLU had relatively stable MDA content, electrolyte leakage, and soluble protein content. On the other hand, GA_3 -treatment had significant less soluble protein content than control. GA_3 spraying dramatically increased the MDA content and electrolyte leakage. The MDA content soared from $7.1 \pm 1.4 \mu\text{mol/g}$ (CK) to $11.8 \pm 1.3 \mu\text{mol/g}$ (GA_3). The electrolyte leakage soared from $73.5 \pm 8.4\%$ (CK) to $92.3 \pm 16.2\%$ (GA_3). Compared with GA_3 -treated seedlings, the electrolyte leakage and MDA content were greatly decreased to $59.9 \pm 8.1\%$ and $6.6 \pm 1.4 \mu\text{mol/g}$ in the presence of PAC, by 35.5% and 44.1%, respectively. The combined use of GA_3 and PAC had relatively stable MDA content and electrolyte leakage.

We found that treatment with ABA or GA can suppress or enhance chilling damage, respectively. However, we thought therefore that ABA-mediated suppression and GA-mediated enhancement of chilling damage could be reversed by the application of exogenous FLU and PAC, respectively.

The combined effects of GA and ABA on cold stress response were further tested (Fig. 7B C). GA-mediated promotion and ABA-mediated repression of chilling damage could both be attenuated by the co-application of ABA and GA_3 , respectively, suggesting that the effects of GA_3 and ABA are antagonistic. Interestingly, the promotion of chilling damage by GA_3 was only significantly attenuated when ABA was applied within 6 h of the initial GA_3 treatment. No significant attenuation was seen if ABA was added at 12 or 24 h (Fig. 7B). Moreover, ABA-mediated alleviation of chilling damage was aggravated by supplementation with GA_3 at 2 or 6 h after the initial application of ABA. No significant attenuation was seen if GA_3 was added at 12 or 24 h (Fig. 7C).

Discussion

In our previous study [13], we found that 0–48 h is the critical period of the chilling damage in *T. hemsleyanum*. After that time, serious injury or even death are inevitable. In the present study, the transcriptomes of *T. hemsleyanum* during 0–48 h of cold-treatment duration were compared by using the Illumina sequencing platform. Then GO and KEGG enrichment analysis of the DEGs were used to indicate the biological processes involved in response to cold stress. The results indicated that some biological process categories related to endogenous hormone response skyrocketed in the early stress stage, such as ‘hormone-mediated signaling’ ‘response to hormone’ and ‘response to endogenous stimulus’ in 12 h and 24 h. Therefore, we speculated that phytohormones regulatory network was a significant mechanism in response to cold stress of *T. hemsleyanum*.

Metabolic changes of endogenous hormones in response to cold stress

Phytohormones, such as ABA, IAA, GA, and so on are known to play critical roles in the complex signaling cascades and also in adaptive responses to various biotic and abiotic stresses. The cooperation and antagonism among the various hormones are recognized to be existent objectively in the growth and development of plants, such as dormancy release and germination. However, it remains poorly understood whether such an antagonistic relationship exists in the regulation of plant stress response.

ABA is perhaps the most well characterized phytohormone of various stress signals. ABA biosynthesis is one of the fastest responses of plants to abiotic stress, triggering downstream genes expression and causing stomatal closure, thereby enhancing cell water retention and the content of osmotic adjustment substances, such as proline, soluble sugar, and soluble protein. This could ultimately lead to restricting cellular growth. There was a significant positive correlation between the ABA accumulation and the drought and cold tolerance in rice [14]. Exogenous ABA could improve the cold tolerance of plants [15]. In higher plants, some major genes for the enzymes in ABA biosynthetic pathway have been identified. The first step in the ABA biosynthesis pathway is catalyzed by a zeaxanthine epoxidase (ZEP), whose molecular identity was first revealed in tobacco [16]. The 9-cis epoxycarotenoid dioxygenase (NCED) has been suggested to be a key rate-limiting enzyme during ABA biosynthesis [17]. It was originally identified from maize viviparous 14 mutants, which cleaves the ABA precursor C40-cis-epoxycarotenoids, 9-cis-neoxanthin, and so on to produce xanthoxin, the direct precursor of ABA [18]. NCED is a rate-limiting enzyme in ABA synthesis and various studies reported transcriptional regulation of NCED following abiotic stress [19]. In *T. hemsleyanum*, three ZEP genes were up-regulated by approximately 2.5–9.8 folds after 12 h of cold treatment, and two NCED genes were up-regulated by approximately 2.0–2.5 folds after 24 h of cold treatment. Up-regulation of these genes implied that synthesis of ABA might be elevated under cold, which is consistent with results reported previously in *Zea mays* ssp. *mexicana* L [20] and rice [21]. It is notable that most of these ABA biosynthesis-related genes were suddenly down-regulated after 24 h or 48 h of cold treatment. Accordingly, the ABA content was also first strongly increased, continuously came up to a "peak value" at 24 h, and then gradually decreased to a comparatively low level. This result is consistent with studies of *Poncirus trifoliata* [22] and *P. euphratica* [23]. It is suggested that the increased levels of expression of ABA-responsive genes and an increased ABA content would be beneficial for plants under environmental stress [24]. Thus, a series of related genes involved in ABA biosynthesis and signaling pathway of *T. hemsleyanum* were expressed at low levels after 48 h of cold treatment, which might be one factor contributing to cold-sensitivity of *T. hemsleyanum*.

GAs, as a group of tetracyclic diterpene carboxylic acids, regulate many critical biological events in plants, such as stem elongation, seed germination, and so on. Among the different GAs identified from plants, a few act as endogenous plant growth regulators including GA₁, GA₃, and so on. GA₃ is considered to be the earliest known cold-related phytohormone. Plant growth could be restrained partly by the accumulation of DELLA proteins, a critical component in GA signaling [25, 26], suggesting the plants could balance their growth and the resistance to abiotic stress though down-regulating GA levels [27].

Bioactive GAs are synthesized from geranyl geranyl diphosphate (GGDP), a common C₂₀ precursor for diterpenoids [28]. The final stage of bioactive GA synthesis is catalyzed by GA20ox and GA3ox, respectively. GA₁ content in rice would be reduced by the suppression of GA3ox2, or the activation of GA2ox1 and GA2ox3 [29]. GA20ox is a rate-limiting enzyme in GA synthesis and regulation pathway. Over-expression of GA20ox often leads to plant growth acceleration and internode elongation by promoting GA synthesis [30]. Researches indicated that it would be beneficial for plants to survive in adversity at the cost of growth loss. The inhibition of plant endogenous hormones, particularly auxin and GA, would dramatically increase the surviving ability of plant under environmental stress. This could be a proactive self-protection mechanism of plant [31]. Niu et al[32] reported that low temperature delayed the growth and development of tobacco adventitious root by inhibiting GA signal transduction and dramatically reducing GA content. Similarly, the GA20ox genes in *Zea mays* were also down-regulated by cold treatment [33]. In our study, two GA20ox genes were down-regulated significantly under cold stress, which is in accord with this theory that growth retardation is a typical evolution characteristic of plant adaptation to environmental stress. But there were still some opposite reports, GA20ox genes of citrus were stimulated by chilling treatment [34]. The enzymes involved in GA synthesis pathway of *Thlaspi arvensis* were induced by low temperature [35].

GA2ox could transform bioactive GA₁ and GA₃ to inactive GA₈ and GA₃₄ by catalyzing hydroxylation reactions at C₂. Over-expression of GA2ox would inhibit GA synthesis, and thus it would lead to plant growth retardation. The up-regulation of GA2ox genes was found to be concerned with the high tolerance to drought [36]. Similarly, in our study, six putative members of the GA2ox gene family were identified in *T. hemsleyanum*. Among them, four genes exhibited a similar expression profile and all were significantly up-regulated by approximately 3–7 folds following cold treatment, with a peak at 12 h after exposure. But then their expression level declined rapidly with the extension of stress time.

On the other hand, GA3ox was a positively regulatory factor of GA synthesis. An A20/AN1 zinc finger protein negatively regulated plant height in rice by reducing expression of GA3ox, enhancing expression of GA2ox1 and GA2ox3, and decreasing the level of GA₁ [37]. In our study, a significantly up-regulated GA3ox was annotated, which was up-regulated by approximately 6 folds following cold treatment, with a peak at 24 h after exposure. Accordingly, GA content increased to peak at 24 h by 161% and then maintained at the higher level. Previous studies on *Zea mays* ssp. *maize* L. transcript profile under cold, salt, and drought stresses indicated that the abundance of bioactive GAs was reduced, suggesting a negative effects of GA in stress tolerance[20]. This is in disagreement with what we observed in cold-stressed *T. hemsleyanum*. The up-regulation of GA3ox (c78816_g1, C8451_g1 and c57017_g2) and the down-regulation of GA20ox (c98675_g1, c54888_g1, c93517_g2, c85773_g1) were induced by cold, suggesting an inconsistent role of GA in this response. Thus, we speculated that there was a more complicated genes network of GA biosynthesis involving in cold-induced responses, besides these conventional rate-limiting enzymes. Future more comparative study will provide better understanding of the molecular basis of the GA associated response to cold in *T. hemsleyanum*.

Indole-3-acetic acid (IAA), the major auxin which has been recognized for involving in plant cell division, differentiation and so on. Recent reports suggested that auxin is also involved in stress responses. Transcript profiling of rice revealed that the expression of many auxin-related genes were regulated in response to dehydration stress [38]. With the increase of cold stress intensity, the content of IAA decreased significantly in roots of Sea buckthorn, which was a cold-tolerance plant species [39]. So far, the main IAA biosynthesis and regulation pathways were still largely unknown. Mashiguchi et al suggested that IAA is produced from IPA by the YUC family of flavin-containing monooxygenase (FMO)[40]. The OsGH3-2 overexpression rice showed increased resistance to cold stress due to the combined effects of reduced free IAA content, decreased membrane penetrability, and so on[14]. Our study was similar to those results, two FOM unigenes were identified in *T. hemsleyanum* transcriptome, and both them were down-regulated after 48 h of cold stress. But as for the content of IAA, although a slight increase was noted at 12 h, the difference was not statistically significant, and then it remained basically stable. This could be because the expression levels of the two FOM genes were comparatively low, with RPKM values < 60. So the effects of them on IAA metabolism should be diminished.

Jasmonates(JAs) are the important signal molecules that regulate and enhance resistance in plants to abiotic stress, such as cold stress, and biotic stress. The application of exogenous methyl jasmonate (MeJA) in *Camellia japonica*[41] and banana fruit [42] significantly enhanced cold tolerance by increasing endogenous JA accumulation during cold storage. A recent study showed that the JA signaling pathway plays an important role in the early cold response by modulating the CBF signaling pathway. In plants, Allene oxide synthases (AOCs) are known as key enzymes in JA biosynthesis. JA is synthesized from α -linolenic acid. AOS and allene oxide cyclase activity subsequently results in *cis*-(+)-12-oxo-phytodienoic acid (OPDA). AOC is required for formation of *cis*-(+)-enantiomer (9S, 13S) of OPDA, which is thought to be the precursor of JA. OPDA, JA, and their derivatives can stimulate AOC expression. Suppressing AOC1 expression in *Medicago truncatula* roots could lower JA levels significantly [43]. Biotic or abiotic stresses that lead to endogenous increases in JAs are usually accompanied by upregulation of AOC and AOS transcription [44]. Our results also showed that the expression of AOC genes encoding the main JA biosynthetic enzymes were up-regulated significantly, indicating a high level of JA in *T. hemsleyanum* during cold stress stage.

Ethylene (ET) is a growth regulator involved in leaf abscission, fruit ripening, growth of adventitious roots, and so on. Moreover, it is synthesized in response to stresses such as wounding, chilling, and freezing [45]. ET biosynthesis is originated from the amino acid methionine generated by the Yang cycle, in which the precursor S-adenosylmethionine (AdoMet) is converted into 1-aminocyclopropane-1-carboxylic acid (ACC) by the enzyme 1-aminocyclopropane – 1-carboxylase synthase (ACS)[46]. ACO (1-aminocyclopropane-1-carboxylate oxidase), termed as ET synthase, can directly catalyze the conversion of ACC to ET. Also, the ACO would directly control the content of ET because of its effect in the final step of the pathway. An increase in levels of free ACC content was observed to coincide with an increase in ET production of the drought-tolerant soybean cultivar[47]. In the present study, the expression levels of ACO genes were increased dramatically under cold stress, yet simultaneously, a significant decrease in the transcript levels of ACS genes occurred in *T. hemsleyanum*. It is interesting to note that the expression trend of ACO and ACS genes were divergent. It can be deduced from the above information that the ET biosynthesis was terribly sensitive to cold stress. Both the ACO and ACS were members of the gene family which is termed as a responsive gene family to cold stress, characteristic with increase and decrease at the early stage, respectively. We supposed that when a short interval of cold stress was applied, ET biosynthesis was promoted briefly because of the overproduction of ACO, but when the stress duration was increased, ET biosynthesis would be increasingly suppressed because of the inactivation of ACS. Previous studies had showed

partial similarity to our works. Sun et al [45] reported that ET positively regulated cold tolerance in grapevine, the cold tolerance was reduced by the application of the ET biosynthesis inhibitor aminoethoxyvinylglycine but increased by the treatment of the ET precursor ACC. But on the contrary, Shi et al [48] reported that ET negatively regulated plant responses to freezing stress in *Arabidopsis thaliana*. Freezing tolerance was decreased by the application of the ET precursor ACC but increased by the addition of the ET biosynthesis inhibitor, and the ET-insensitive mutants displayed enhanced freezing tolerance. In general, these results suggested that the role of ET in cold tolerance varied in different species.

The proportion change and interaction of various endogenous hormones in response to cold stress

It is generally perceived that ABA/GA ratio could affect seed germination and sex differentiation, ABA/IAA ratio could affect growing potential, and ABA/ZA could affect stomatal oscillation. However, it remains much less understood whether such an antagonistic relationship exists in the regulation of plant stress response. Some studies showed that under environment stress, a variety of phytohormones promoted and restricted mutually to reach a new balance by inducing stomatal closure, weakening transpiration, and slowing down growing rate, which is tightly related with the plant's ability to withstand stress and disturbance.

ABA and GA were thought to be hormone partners that functioned via a complicated and antagonistic network in which transcription factors DREB [49] and OsAP2-39 [50] were found to play a critical role in rice. It has been reported that restricting GA-promoted processes would in turn to promote ABA accumulation in *Arabidopsis* [25]. In contrast, it was found that the biosynthesis of both GA and ABA were inhibited by an A20/AN1-type zinc finger protein, which led to the decreased stress tolerance of rice [51]. Some evidence at the transcriptional level demonstrated that GA and ABA had different and contrasting effects in a variety of developmental processes of plant [52]. ABA levels and the ratio of ABA/GA in leaves of cold-tolerant bamboo species were both higher than those of cold-sensitive bamboo species for all cold treatments, while GA₁ and GA₄ were conversely lower [11]. Similarly, Huang et al [8] verified that exogenous ABA treatment could promote the increase of endogenous ABA content and the ratio of ABA/GA, in contrast with the decreased contents of MDA and GA₃, and thereby induced cold stress tolerance of sugarcane seedling. In accordance with the previous reports, our study indicated that ratio of ABA/GA in *T. hemsleyanum* was increased steeply by 1.9-fold at 12 h, but then declined sharply to about the level of CK.

Previous reports suggested that a crosstalk of the signaling pathway also occurred between the ABA and auxin responses. For example, ABL1, an ABA signaling component of rice could modulate auxin responses by regulating the expression level of ABRE-containing genes related to auxin metabolism [53]. Auxin response factor₂ (ARF₂) was revealed to play a role in the ABA signaling pathway [54]. IAA content was significantly declined in rice treated with the ABA biosynthesis inhibitor, and expression of auxin synthesis and metabolism-related genes were changed in the ABA-inhibited rice. Endogenous IAA levels were decreased in the carotenoid-deficient rice mutants, demonstrating that ABA and IAA could mutually affect each other at the biosynthesis level, and the increased resistance to cold stress was probably related to the deficiency in IAA [6]. In the current study, we found that endogenous IAA levels decreased slightly in the cold-treated *T. hemsleyanum* under cold stress, but the change was not significant. Ratio of ABA/IAA was increased gradually from 0 h to 24 h, then declined at 48 h, but the value was still higher than control. The increase amplitude of ABA/GA ratio was greater than that of ABA/IAA, and

as well the peak time for enhancement of ABA/GA ratio was earlier than that of ABA/IAA, which demonstrated that ABA/GA had a greater impact on the stress response than ABA/IAA.

Few studies have been reported the relationship between ABA and other phytohormones. Ratio of ABA/GA₃ was one of the important factors that could effectively increase cold stress tolerance in sugarcane plants, along with the increased ratio of ABA/IAA and the ratio of ABA/ZR, and exogenous ABA application reduced GA₃ and ZR content but increased endogenous ABA content [8]. Guo et al [55] reported that a combination of the increased ZT/IAA ratio and H₂O₂ concentration was the basis for the enhanced shoot morphogenesis in cold-treated *Saussurea involucreata* explants. In our study, ABA/ZR ratio in *T. hemsleyanum* also showed an obviously increasing trend, which was correlated with the increase of ABA content and the decrease of ZR content under prolonged cold stress. On this basis, it could be inferred that other phytohormones also could influence stress tolerance through cross-talk mechanisms.

The antagonistic relationship of ABA and GA₃ in response to cold stress

Hormones co-regulate plant growth and stress response in a critical way, and they act either synergistically or antagonistically[37], among which ABA and GA were showed to be the most effective hormones in cold stress response of *T. hemsleyanum*. It was verified that GA regulated a variety of critical biological events in plants, including petal growth and seed germination [52]. As a versatile regulator, ABA has been shown to act antagonistically to the function of GA in many developmental processes, including petal elongation [52] and seed germination [56]. However, it remains poorly understood whether such an antagonistic relationship exists in the regulation of stress response.

The higher the electrolyte leakage rate is, the more serious the damage by cold stress is. Besides, MDA content can reflect the severity of membrane lipid peroxidation and the stress tolerance of plants. So they are negatively correlated with stress tolerance ability of plants. The soluble proteins could play important roles in osmotic adjustment, and it is positively correlated with stress tolerance of plants. In this study, the electrolyte leakage rate and MDA content of the GA-treated plantlets were promoted, while the soluble protein content was reduced than control. And instead, ABA-treatment had contrary influences on these stress related indicators. Moreover, we also found that when the biosynthesis of endogenous ABA and GA were inhibited by FLU and PAC, respectively, the effects of GA and ABA treatment were reversed. The combined effects of GA and ABA on cold response were further tested. Two findings in our current study showed that GA and ABA had antagonistic effects on cold tolerance of *T. hemsleyanum*: (1) cold-endurance ability was enhanced by ABA but repressed by GA when each phytohormone was applied alone; (2) the ABA-mediated promotion and GA-mediated repression of cold tolerance could both be attenuated by the co-application of the two phytohormones, respectively. However, the antagonistic effect of GA and ABA was no longer apparent when the treatment was extended to longer than 6 h, indicating that transcriptional regulation of hormone-associated pathways was an early and transient event during stress response.

Conclusions

In summary, the transcriptome of *T. hemsleyanum* during 0–48 h of cold-treatment indicated that phytohormones regulatory network was a significant mechanism in response to cold stress. Some hormone-related genes, such as NCED, ZEP, GA3ox, FOM, AOC and so on, were crucial enzyme genes, which resulted in the significant changes of hormones content. The response of ABA and ABA/GA₁₊₃ to cold stress was prior to that of GA₁₊₃, ZR, ABA/IAA, and ABA/ZR. The increasing changes in ABA/GA₁₊₃ turned to a steep decline with the extension of stress time, which might be one factor contributing to cold-sensitivity of *T. hemsleyanum* under prolonged stress. Our data supported the hypothesis that GA and ABA could work antagonistically to balance the early responses to cold in *T. hemsleyanum*. This study also suggested that PAC, a GA biogenesis inhibitor, as well as exogenous ABA, might be used as early interventions for chilling damage of *T. hemsleyanum*. Further study in the relationship among ABA/GA homeostasis and the regulation of the stress-related genes involved in the phytohormone signaling network can provide a new clue for understanding their connection in stress response.

Methods

Plant materials and cold stress treatments

Healthy sterile test-tube plantlets of *T. hemsleyanum* were obtained and cultured as the same as our previous report [57]. Based on our preliminary work results of physiological parameters determination, the temperature and sampling time of cold stress treatment were optimized [13]. For both the control and cold stress tests, these seedlings were cultured in the same conditions except the different temperatures at 25 °C or 0 °C. After 0 h, 12 h, 24 h, and 48 h of cold stress treatment, the young leaves of 10 individual 4 week-old seedlings were collected and immersed in liquid nitrogen immediately and stored at – 80 °C for RNA extraction and quantitative real-time PCR analysis. Three biological replicates were performed at the same time. Each biological replicate consisted of a pool of leaves from 10 seedlings.

Rna Extraction, Quality Determination, And De Novo Assembly

Total RNA was extracted with Trizol reagent (Invitrogen, CA, USA), following the manufacturer's protocol. RNA purity was checked using the kaiaoK5500@Spectrophotometer (Kaiao, Beijing, China). RNA integrity and concentration was assessed using the RNA Nano 6000 Assay Kit of the Bioanalyzer 2100 system (Agilent Technologies, CA, USA) at Annoroad Gene Technology (Beijing) co., LTD. The clustering was performed on a cBot cluster generation system using HiSeq PE Cluster Kit v4-cBot-HS (Illumina), and sequenced by Illumina HiSeq2500. The software Trinity was used for de novo assembly which was developed at the Broad Institute and the Hebrew University of Jerusalem. Trinity partitions the sequence data into many individual de Bruijn graphs, each representing the transcriptional complexity at a given gene or locus. The clean data were mapped to the assembled transcripts by Bowtie 2(v2.2.3).

Coding Regions Prediction And Transcriptome Functional Annotation

The candidate coding regions within transcript sequences, such as those generated by de novo RNA-Seq transcript assembled by Trinity were identified by TransDecoder. The functional annotation of unigenes and ORFs were performed by Trinotate.

Rpkm Estimation And Deg Analysis

Reads Count for each gene in each sample was counted by HT Kilobase Million Mapped Reads each sample, the formula is shown as:

$$RPKM = 10^6 * R / (NL / 10^3)$$

R is the number of reads in a certain sample that is assigned to a certain gene, N is the total number of mapped reads in the certain sample and L is the length of the certain gene.

The differential gene expression between samples was analyzed by DESeq(v1.16). P-value could be assigned to each gene and adjusted by the Benjamini and Hochberg's approach for controlling the false discovery rate. "q ≤ 0.05 and |log2_ratio| ≥ 1" was set as the threshold to determine the differentially expressed genes (DEGs).

Functional Annotation And Classification Of Degr

The GO (Gene Ontology, <http://geneontology.org/>) enrichment of DEGs was implemented by hypergeometric test. GO terms with q < 0.05 were considered to be significantly enriched. KEGG (Kyoto Encyclopedia of Genes and Genomes <http://www.kegg.jp/>) analysis of the DEGs was performed to identify the associated biochemical and signal transduction pathways. KEGG terms with q < 0.05 were considered to be significantly enriched.

Validation Of Rna-seq Data By Qrt-pcr

Quantitative real-time RT-PCR (qPCR) was performed to examine expression patterns of the candidate unigenes using a StepmoPlus real-time PCR system (Applied Biosystems, Forster City, CA, USA). The primers for amplification are listed in Table 1. Each reaction contained 2 µl of cDNA, 10 µl of 2x Power SYBR green PCR master mix (Applied Biosystems, Forster City, CA, USA), and 2 µl of forward and reverse primers in a final volume of 20 µl. The PCR reactions were conducted by incubation at 95 °C for 3 min followed by 40 cycles of 95 °C for 15sec and 60 °C for 45sec. The GAPDH gene was used as the reference gene to normalize target gene expression. The calculation for relative gene expression was determined by the $2^{-\Delta\Delta C_t}$. Dissociation-curve analysis was carried out to confirm the amplification specificity. Three technical replicates were performed for each sample and the data are shown as means ± standard errors (SE) (n = 3).

Determination Of Phytohormones Content By Elisa

The double-antibody sandwich ELISA was performed as our previously described [54] with slight modifications for the endogenous phytohormones quantification. Briefly, 50 mg of fresh leaves from *T. hemsleyanum* seedlings were ground and then subjected to extraction (80% methanol containing 10 mg/L butylhydroxytoluene v/v) for 18 h at 4 °C, and then centrifuged at 4000 rpm for 15 min at 4 °C. The supernatant was passed through C18 solid phase extraction cartridge (Waters Corporation, Millford, MA, USA) to remove the chlorophyll. The products were dried by pure N₂ at 20 °C, and resuspended in 20 µl cold solution from the kit for phytohormones-testing (MLBIO, Shanghai El Biotechnology Co., Ltd. China). Endogenous phytohormones levels were determined according to the manufacturer's instructions, respectively. The imark microplate reader (Bio-Rad Co. Ltd., Hercules, USA) was used to measure the ABA, IAA, ZR and GA content at 450 nm of the optical density (OD).

Hormone And Inhibitor Treatments

For the evaluation of hormonal mediation in cold response, GA and/or ABA treatments were employed in the in vivo experiments. The young leaves of 10 individual seedlings were included for each treatment. In vivo treatments were performed by spraying with 5 ml 20 μ M GA3 or 100 μ M ABA at 6 h, then were sampled after 48 h. As a control, seedlings sprayed with 0.1% ethanol in deionized water were sampled in parallel. The widely used inhibitors of GA and ABA biosynthesis, paclobutrazol (PAC) and fluridone (FLU), were also sprayed in this study. In the experiment protocol described above, 10 μ M PAC or 0.1 μ M FLU were sprayed with or without the phytohormones.

To evaluate the dynamic interaction between the effects of GA and ABA, the experiment using a combination of GA and ABA was further performed, in which, after spraying with ABA for 2, 6, 12, or 24 h, GA was sprayed, with the final measurements being performed at 48 h. Conversely, where GA was the initial hormone sprayed, ABA was then added during the experiment. Three biological replicates were performed at the same time. Each biological replicate consisted of a pool of leaves from 10 seedlings.

Physiological Parameters Determination

MDA content was measured as our previous report[13]. $\text{MDA content } (\mu\text{mol/g}) = (C_{\text{MDA}} \times V_r \times 10^{-3} \times V_t) \times (W \times V_s)^{-1}$. $C_{\text{MDA}} = 6.45(A_{532} - A_{600}) - 0.56A_{450}$, $\mu\text{mol/L}$; V_r : reaction volume (mL); V_t : total volume of total extractive enzyme (mL), V_s : tested enzyme volume (mL); W : the weight of leaves (g).

Electrolyte leakage was determined as our previous report [13]. The fresh leaves (1 g) were incubated with 15 ml deionized water at 25 °C for 24 h. The electrolyte leakage was expressed as percent (EL%) = $EC_1/EC_2 \times 100\%$. Deionized water without leaves incubated in it was taken as control.

Soluble protein content was measured by Coomassie Brilliant Blue G-250 staining. Soluble protein content (mg/g) = $C \times V_t \times (V_s \times W \times 1000)^{-1}$; C : protein quantity determined from the standard curve (μg); V_s : sample solution volume (mL); V_t : total volume of the extracted solution (mL); W : the weight of samples (g).

Statistical Analysis

All data were collected from three biological replicates and three technical replicates. Values are presented as mean $\pm$ SD of three independent experiments. All statistical analyses were performed using the SPSS 21.0 software (SPSS, Chicago, IL, USA). The difference among the multiple groups was analyzed by one-way analysis of variance followed by LSD multiple comparison tests. $P < 0.05$ was considered significant.

Abbreviations

Table 1

List of primer sequences of randomly selected unigenes used for qRT-PCR analysis

GeneID	Gene Name	Tm	Primer F(5'-3')	Primer R(5'-3')
c81504_g1	ZEP_4	55.8	TACAGTTTATTTAGCCCGCC	TTTCCTACTCGTTTCCATCC
c102056_g1	AAO3_2	53.7	AACGGTGAGTTGTGGTGGTG	GAAGGGTGGATGGTGGAGAC
c66096_g3	NCED_3	56.9	CACCTTCTGCTTCCACCTC	TTTCTCCTCGCATTCTGTTA
c94964_g1	AAO1_3	54.8	TAGAGTCCAACAGCAGAGCAG	GGAAGGGAGATAGTAAAGAAGTG
c72906_g1	CRMP_LOG1	55.0	AGACAGTTGGGGAAGTGAAGG	TTATCTATGAATGAGAGGAGGGAG
c73984_g2	CRMP_LOG7	51.2	AAATACCTCCCTTCCCTCCT	GCCTACACTTCCTCCACCAT
c80750_g2	CRMP_LOG8	56.5	CTTCTTGCTTGGGTATTGAC	AGTTTATGGCGTCGGGTTC
c100822_g2	ACO_1	55.3	CTCACCAAACAATGCTCCTTC	TTTTCTTGATTTCGAACCCAG
c90679_g2	ACO_4	52.5	ACTGTTTGTGTCACCACTCTATG	GTCAGCAACTACCCTCCCT
c99260_g1	ACS_1	56.0	TTTTTTGACGGGTATGGTAGAG	GATTTGAAGGATTGGTGATGAG
c93517_g2	GA2ox_2	52.0	AATGGGGTTGTTTCAGAGTC	AAAGAGGATTGATGTCGCTG
c92581_g1	GA2ox_4	56.4	TGGGGGGAGTTTGAGAGG	GCGTGAAGCATAGGGTGTTG
c90130_g1	KCT	55.8	CTTCTTCTTCGCCTTCTCGT	ACAACCACAATGTCATCCCC
c52785_g1	AOC	56.0	CAGATCCCCGCCGTGAT	GCGCCAAGCAGAGCCAG

Abbreviation	Full name	Abbreviation	Full name
ABA	abscisic acid	NCED	9-cis-epoxycarotenoid dioxygenase
GA	gibberellic acid	TatC	sec-independent protein translocase protein
CK	cytokinin	IPT	adenylate isopentenyltransferase
ZR	zeatin riboside	CRMP	cytokinin riboside 5'-monophosphate phosphoribohydrolase
PAC	paclobutrazol	STM	homeobox protein Shoot Meristemless
ET	ethylene	YUCCA	indole-3-pyruvate monooxygenase
JA	jasmonic acid	ACC	aminocyclopropane carboxylate oxidase
BR	brassinosteroid	ACS	1-aminocyclopropane-1-carboxylate synthase
FLU	fluridone	DRBP	dehydration-responsive element-binding protein
AAO3	abscisic-aldehyde oxidase	CTR1	serine/threonine-protein kinase
CESA	cellulose synthase A	AAO1	indole-3-acetaldehyde oxidase
ZEP	zeaxanthin epoxidase	ADLT	AMP dimethylallyltransferase
EUPL	E3 ubiquitin-protein ligase	CODH	copal-8-ol diphosphate hydratase
ELP	ETO1-like protein	ECDS	ent-copalyl diphosphate synthase
LOX2S	lipoxygenase	EKAH	ent-kaurenoic acid hydroxylase
AOC	allene oxide cyclase	GA20ox	gibberellin 20-oxidase
4CL	4-coumarate-CoA ligase	GA2ox	gibberellin 2-beta-dioxygenase
EKES	Ent-kaur-16-ene synthase	GA3ox	gibberellin 3-beta-dioxygenase
ATH1	homeobox protein ATH1	ACO	1-aminocyclopropane-1-carboxylate oxidase
ACDD	acyl-CoA dehydrogenase	CRMP_LOG	Cytokinin riboside 5'-monophosphate phosphoribohydrolase LOG
KCT	3-ketoacyl-CoA thiolase		

Declarations

Ethics approval and consent to participate: Not applicable

Consent to publish: All Authors have read the manuscript and have agreed to submit it in its current form for consideration for publication.

Availability of data and materials: The datasets generated during the current study are available in Supplementary files 1-7 were unigenes expression matrix data of ABA, Auxin, GA, Ethylene, JA, Cytokinin, and Brassinosteroid synthesis pathways in *T. hemsleyanum* transcriptome, respectively.

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Author Contributions: Peng X conceived the study. Li M J performed GO and KEGG Pathways enrichment analysis; Wu H performed physiological indicators determination; Qiu W Y performed endogenous hormone determination; Chen H J performed RNA Extraction and quality determination. Peng X wrote the manuscript, designed the experiments; Zhang Z Y carried out the analysis.

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Supplementary Files Legend

Additional files

Supplementary files 1 was unigenes expression matrix data of ABA synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files 2 was unigenes expression matrix data of Auxin synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files 3 was unigenes expression matrix data of GA synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files 4 was unigenes expression matrix data of Ethylene synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files 5 was unigenes expression matrix data of JA synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files 6 was unigenes expression matrix data of Cytokinin synthesis pathways in *T. hemsleyanum* transcriptome.

Supplementary files7 was unigenes expression matrix data of Brassinosteroid synthesis pathways in *T. hemsleyanum* transcriptome.

Figures

Figure 1

Venn diagram showing distribution of similar annotations from Nr, NT, BLASTX, and BLASTP databases. The overlapping regions among the four circles contain the number of unigenes that share similar hits with respective databases.

Figure 2

GO classifications of differentially expressed unigenes of *T. hemsleyanum* in response to 12h(A), 24h(B), and 48h(C) of cold exposure

Figure 3

KEGG pathway classification of differentially expressed unigenes of *T. hemsleyanum* in response to 12h(A), 24h(B), and 48h(C) of cold exposure

Figure 5

Expression pattern validation (A) and linear dependence relation (B) between the log₂ of the key genes expression ratios obtained from RNA-seq and qRT-PCR during cold stress. The transcriptional level of genes was examined by real-time PCR with three biological replicates. RNA-Seq data were highly consistent with qRT-PCR results ($R^2 = 0.865$). The candidate genes arranged from left to right as: AAO3_2; NCED_3; ZEP_4; AAO1_3; CRMP_LOG1; CRMP_LOG7; CRMP_LOG8; ACO_1; ACO_4; ACS_1; GA20ox_2; GA20ox_4; KCT; AOC

Figure 6

Variation of endogenous phytohormones level and ratio of ABA/GA (A); ABA/IAA (B); ABA/ZR(C) under 25°C or 0°C. Values are presented as mean±SD of three independent experiments. Bars annotated by the different lowercase letters indicate the significant differences between the samples ($P < 0.05$). Curves annotated by * indicate the significant differences between the treated and control groups ($P < 0.05$).

Figure 7

Antagonistic effects of GA and ABA on cold tolerance of *T. hemsleyanum*. (A) Effect of exogenous GA or ABA used alone or in combination with biosynthesis inhibitors on cold tolerance; (B) Effects of GA on cold tolerance are attenuated by co-application of ABA at 2h, 6h, 12h, and 24h; (C) Effects of ABA on cold tolerance are attenuated by co-application of GA at 2h, 6h, 12h, and 24h. Values are presented as mean±SD of three independent experiments. Bars annotated by the different lowercase letters indicate the significant differences between the samples ($P<0.05$).

Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [Supplementaryfile1ABA2.xlsx](#)
- [Supplementaryfile6Cytokinin2.xlsx](#)
- [Supplementaryfile4Ethylene2.xlsx](#)
- [Supplementaryfile7Brassinosteroid.xlsx](#)
- [Supplementaryfile3GA2.xlsx](#)
- [Supplementaryfile2AUXIN.xlsx](#)
- [Supplementaryfile5JA2.xlsx](#)