Expression of the Sweet Potato MYB Transcription Factor IbMYB48 Confers Salt and Drought Tolerance in Arabidopsis

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Abstract: Sweet potato (Ipomoea batatas (L.) Lam) is one of the most crucial food crops widely cultivated worldwide. In plants, MYB transcription factors play crucial roles in plant growth, defense regulation, and stress resistance. However, the regulatory mechanism of MYBs in salt and drought response remains poorly studied in sweet potato. By screening a transcriptome database for differentially expressed genes between the sweet potato variety Jingshu 6 and its mutant JS6-5 with high anthocyanin and increased tolerance to salt and drought stresses, we identified a R2R3-MYB gene IbMYB48, for which expression was induced by PEG6000, NaCl, abscisic acid (ABA), methyl jasmonic acid (MeJA), salicylic acid (SA) and H2O2. Particle-mediated transient transformation of onion epidermal cells showed IbMYB48 is localized in the nucleus. Transactivation activity assay in yeast cells revealed that IbMYB48 has transactivation activity, and its active domain is located in the carboxyl (C)-terminal region. Furthermore, expression of IbMYB48 confers enhanced tolerance to salt and drought stresses in transgenic Arabidopsis. The contents of endogenous ABA, JA, and proline in transgenic lines were higher than control, and the activity of superoxide dismutase (SOD) was significantly increased under salt and drought stress conditions. By contrast, the accumulation of malondialdehyde (MDA) and H2O2 were lower. Moreover, genes encoding enzymes involved in ABA biosynthetic pathway, JA biosynthesis and signaling pathway, and reactive oxygen species (ROS) scavenging system were significantly up-regulated in transgenic Arabidopsis under salt or drought stress. Altogether, these results suggest IbMYB48 may be a candidate gene for improvement of abiotic stress tolerance.

Keywords: sweet potato; IbMYB48; MYB transcription factor; salt and drought tolerance

1. Introduction

Plants are all the products of eons of evolution from primal organisms in response to abiotic and biotic stress conditions [1]. Salt and drought stresses are major abiotic stress in plant agriculture worldwide, because these unfavorable environmental factors will negatively affect the normal growth and development of crops [2–4]. Therefore, a more detailed reaction processes are needed to understand the mechanism of the stress response [5]. Plants have evolved a series of mechanisms to respond and adapt to adverse environment [6]. The regulatory networks consist of stress sensors, a network of protein–protein reactions, transcription factors and promoters, and eventually the output proteins or metabolites [2]. For instance, ABA plays a key role in the plant’s response to abiotic stress, such as drought, salinity, cold, and hypoxia [7]. Stress responses primarily include transcriptional regulation of gene expression, which depends on the interaction of transcription factors with cis-regulatory sequences [2]. Numerous signaling proteins including transcription factors, and protein kinases, play signal transduction roles during...
plant adaptation to abiotic stress, when plants are involved in ranging from stress signal perception to stress-responsive gene expression [8]. Transcription factors (TFs) are generally the earliest response to abiotic stress, which act as significant coordinators of signal transduction [9]. Core sets of transcription factor family genes are distinctively expressed in response to increased abiotic stress, including bZIP, WRKY, AP2/ERF, MYB, bHLH, and NAC families [10].

Among the different TFs, MYBs, as one of the most widely distributed transcription factors in plants are engaged in plant development and response to stresses by binding to cis-elements in promoter regions of target genes [11]. MYB protein are characterized by a highly conserved DNA-binding domain of approximately 50 amino acids, which usually comprises up to four imperfect amino acid sequence repeats [12]. Depending on number of consecutive repeats, MYB protein can be divided into four classes: 1R-MYB, 2R3-MYB, 3R-MYB and 4R-MYB [12]. The majority of plant MYB genes encode a protein of the 2R3-MYB subfamily, by the absence of the sequences encoding the R1 repeat and following expansion of the gene family [13]. The functions of several 2R3-MYB TFs have been proven in different plants using genetic and molecular methods. MYBs play an important role in abiotic and biotic stresses [11–13]. In Arabidopsis, such as AtMYB2, AtMYB15, AtMYB44 and AtMYB60 regulate ABA and abiotic stresses [14–17]. AtMYB15 is responsible for cold-regulation of CBF genes and in freezing tolerance, mutation of AtMYB15 shows increased tolerance to freezing stress in contrast to overexpression of AtMYB15 [16]. AtMYB44, a 2R3-MYB gene of Arabidopsis, is activated under various abiotic stress, such as dehydration, low temperature and salinity [17]. Correspondingly, AtMYB44 overexpression transgenic plants obviously enhance tolerance to salt and drought stress compared to wild-type (WT) plants [17]. Overexpression of wheat TaMYB19 and TaMYB30-B improve drought stress tolerance in transgenic Arabidopsis [18,19]. Overexpression of soybean GmMYB76 or GmMYB177 confers plant to salt and freezing tolerance in Arabidopsis, while transgenic soybean plants overexpressing GmMYB84 have a superior ability to withstand drought stress [20,21].

Sweet potato counts among the most widely staple crops cultivated worldwide [22,23]. Due to its strong ecological adaptability, a high yield potential, and excellent nutritional value, sweet potato has become an indispensable food crop, especially in developing countries [24]. Although sweet potato has many unique properties, its normal growth can also be affected by adverse conditions, such as freezing, drought, and salt stress [25]. Up to now, several genes involved in drought or salinity have been isolated and characterized in sweet potato, including IbLCYB2, IbC3H18, IbGTAT24, IbWRKY22, IbFWRKY70, IbMIPS1, IbMYB116, etc. [25–31]. Although the functions of hundreds of TFs in stress response have been extensively examined in sweet potato, the role of 2R3-MYB TFs in salt tolerance and drought resistance in sweet potato remains unclear.

In this study, a novel 2R3-MYB gene IbMYB48 was cloned from the sweet potato line JS6-5. Overexpression of IbMYB48 gene in Arabidopsis enhanced salt and drought tolerance. The levels of ABA, jasmonic acid (JA), proline and superoxide dismutase (SOD) activities were significantly higher in transgenic plants under the salt and drought stress treatment, while the contents of malondialdehyde (MDA) and H2O2 were markedly lower. Overexpression of IbMYB48 up-regulated genes involved in ABA biosynthetic pathway, JA biosynthesis and signaling pathway and reactive oxygen species (ROS) scavenging system, thereby improving salt and drought resistance in transgenic Arabidopsis.

2. Materials and Methods

2.1. Plant Materials and Growth Conditions

Sweet potato line JS6-5 was used for the cloning of IbMYB48 gene, as well as JS6-5 was applied to the expression analysis of IbMYB48 gene with multiple abiotic stress in this study. Arabidopsis (Columbia-0) plants were grown in a greenhouse maintained at 22 °C under 16/8 h day/night regime or on 1/2 MS medium at 22 °C on a 16/8 h day/night system.
2.2. Cloning and Sequence Analysis of IbMYB48 Gene and Its Promoter Region

The total procedure of RNA extraction experiment followed the manufacturer’s protocol of TransZol Up Kit (TransGen Biotech, Beijing, China), and first-strand cDNA was synthesized using PrimeScript™ II 1st Strand cDNA Synthesis Kit (Takara, Dalian, China). According to the EST obtained in a previous study [32] and referring to the genomic data of Sweet potato GARDEN (http://sweetpotato-garden.kazusa.or.jp/, accessed on 13 January 2018), the cDNA sequence of IbMYB48 was obtained. The PCR conditions were as follows: denaturation, 95 °C 20 s; annealing, 55 °C 20 s; and extension, 72 °C 60 s for 35 cycles.

Genomic DNA extracted from JS6-5 was used to amplify the genomic sequence of IbMYB48 using LA Taq enzyme (Takara, Dalian, China). The PCR conditions were as follows: denaturation, 95 °C 30 s; annealing, 55 °C 30 s; and extension, 72 °C 360 s for 32 cycles. We used the Genome Walking Kit (Clontech, Palo Alto, CA, USA) to amplify the promoter sequence of the IbMYB48 gene as described previously [33].

Multiple protein sequences of IbMYB48 were aligned by the DNAMAN software (Lynnon Biosoft, Quebec, Canada). The molecular weight and theoretical isoelectric point (pI) of IbMYB48 were estimated from ExPASy (http://web.expasy.org/compute_pi/, accessed on 3 January 2018). The phylogenetic tree was constructed using MEGA 6.0 software with the neighbor-joining method. The IbMYB48 promoter region was screened for cis-acting regulatory elements with Plant CARE (http://bioinformatics.psb.ugent.be/webtools/plantcare/html/, accessed on 10 March 2020). The primers were listed in Supplementary Materials Table S1.

2.3. Expression Analysis of IbMYB48 in Sweet Potato

The transcript levels of IbMYB48 in the roots, stems, and leaves, and under different stresses of JS6-5 plants were analyzed using quantitative real-time PCR (qRT-PCR). The roots of 4-week-old in vitro-grown plants were soaked in 1/2 Hoagland solution with 200 mM NaCl, 20% PEG6000, 100 µM MeJA, 100 µM ABA, 100 µM salicylic acid (SA) and 10 mM H2O2, respectively. The whole plants were sampled 0, 0.5 h, 1 h, 3 h, 6 h, 12 h, 24 h and 48 h after treatments and then analyzed for the expression of IbMYB48.

The RNA was extracted using the TransZol method mentioned above. The qRT-PCR assay conditions were 95 °C 5 s for denaturation, 60 °C 34 s for annealing and extension, and 45 cycles. The gene relative mRNA expression levels were measured with the primers IbMYB48-qPCR-F/R, and IbActin was used as the internal control. The genes relative expression level was counted with the comparative C_T method [34]. Results are indicated as means ± SE of biological replicates (n = 3). The primers were listed in Supplementary Materials Table S1.

2.4. Subcellular Localization of IbMYB48

The coding region of IbMYB48 was amplified using PCR primers 83-IbMYB48-F/R listed in Table S1. PacI/Ascl-digested plasmids pMDC83-green fluorescent protein (GFP) and pMDC83-IbMYB48-GFP were constructed and separately transformed into fresh onion epidermal cells by particle bombardment with a Gene Gun (Bio-Rad Laboratories, Hercules, CA, USA). The onion cells were observed with fluorescent microscope Nikon TE-2000E (Nikon, Tokyo, Japan) using an excitation wavelength of 488 nm. The primers were listed in Supplementary Materials Table S1.

2.5. Transcriptional Activation Activity Analysis of IbMYB48

The open reading frame (ORF) of IbMYB48 and two truncated variants were cloned separately into the NdeI/SalI-digested pGBKTK7 vector (ClonTech, Beijing, China). The pGBKTK7-Lam and pGBKTK7-53 vectors were used as a negative and positive control, respectively. The fusion plasmids, negative control, and positive control were transformed into the yeast strain Y2HGold (ClonTech, Beijing, China), individually. The transformed yeast was subjected to selection on SD/-Trp and SD/-Trp/-His/3-AT/X-α-gal mediums.
to observe the growth of yeast at 30 °C for 3–5 days. The primers used for transcriptional activation activity analysis were listed in Supplementary Materials Table S1.

2.6. Vector Construction and Transformation of IbMYB48 into Arabidopsis

The coding sequence of IbMYB48 was amplified by using primers IbMYB48-OE-F/R and then was inserted into the KpnI/SalI-digested pCAMBIA1300-GFP expressing vector under the control of the CaMV 35S promoter. The promoter sequence of IbMYB48 was amplified via using 162-IbMYB48-F/R. This DNA fragment was then inserted into a binary vector pMDC162 and fused to a β-glucuronidase (GUS) reporter gene to create a recombinant transcription unit, IbMYB48pro::GUS. The pCAMBIA1300-IbMYB48-GFP (Figure S1) and pMDC162-IbMYB48pro::GUS vectors were introduced into Arabidopsis using Agrobacterium-mediated floral dip method, respectively [35].

Seeds were surface sterilized and grown on 1/2 MS medium containing 50 mg/L hygromycin for selection. Independent IbMYB48 overexpression lines from homozygous in T3 generation were obtained. DNA was extracted from leaves and the presence of the hygromycin gene was detected by PCR. Three transgenic lines (L6, L8 and L12) with the higher expression of IbMYB48 were selected for subsequent experiments. The sequences of the primers used were listed in Supplementary Materials Table S1.

2.7. Stress Treatments of IbMYB48 Transgenic Plants

The method of stress treatments of IbMYB48 transgenic plants according to the methods described by Du et al. [36] and Kang et al. [37]. Firstly, 30 seeds of IbMYB48 transgenic Arabidopsis and WT were germinated on 1/2 MS medium. Then, seedlings were cultured in 1/2 MS medium with 100 mM NaCl or 300 mM mannitol, respectively. After 7 days, the root length, fresh weight and germination rate of transgenic plants and WT plants were measured by the method of Du et al. [36]. Furthermore, transgenic plants and WT seedlings were grown in pots and regularly watered for 10 days. The plants were then watered with 100 mM NaCl or 300 mM mannitol every 2 days for 3 weeks. The drought treatment adopted the natural drought method for 3 weeks, and the plant growth status was observed after the re-watering treatment for 2 days. The contents of ABA, JA, proline, MDA, H2O2 and SOD were measured by the methods of Zhai et al. [28], Zhou et al. [26] and Kang et al. [37].

2.8. Histochemical GUS Staining

Indirect GUS histochemical staining of transgenic Arabidopsis plants containing IbMYB48pro::GUS fusion construct followed the previously described method [38–40]. The stained plants were photographed using a digital camera (Nikon D800).

2.9. Statistical Analysis

All the experiments were repeated three times individually. The data are presented as the mean ± SD. The results were analyzed using Student’s t-test in a two-tailed analysis.

3. Results

3.1. IbMYB48 Is a Potential Candidate Gene Involved in the Regulation of Tolerance to Abiotic Stress

We previously conducted root transcriptomes of the sweetpotato variety Jingshu 6 and its mutant JS6-5 with high anthocyanin and increased tolerance to salt and drought stresses by high-throughput RNA sequencing [32]. Among these differential genes, the expression of a transcription factor gene, IbMYB48 was obviously up-regulated in JS6-5 [32]. This trend was further supported by qRT-PCR analysis (Figure 1A). Therefore, IbMYB48 gene was selected for subsequent analysis.
Figure 1. Sequence analysis of IbMYB48. (A) Expression level analysis of IbMYB48 in storage root of Jingshu 6 and JS6-5. (B) Multiple sequence alignment of IbMYB48 and its closest orthologues from various species. The black lines represent the R2R3 domain of MYB transcription factor. (C) Phylogenetic analysis of MYB proteins from different plants. (D) An overview of IbMYB48 promoter which contains different kinds of cis-acting elements, as well as the structure of the IbMYB48 gene. Black arrows represent different cis-acting elements. Exon sequence is represented by blue boxes and introns are represented by yellow lines.

The full-length of IbMYB48 cDNA was isolated from mutant line JS6-5. The ORF of IbMYB48 consists of 801 bp, encoding a protein of 266 amino acids with a calculated molecular mass of 30.9 kDa and a calculated isoelectric point of 8.73. The IbMYB48 belongs to R2R3-MYB transcription factor subgroup (Figure 1B). Based on multiple sequence alignment, this protein shared a high sequence identity with MYB48 proteins in Ipomoea triloba (XP_031122863.1, 79.35%), Capsicum annuum (XP_016577226.2, 47.10%), Nicotiana tomentosiformis (XP_009590049.1, 46.38%), Sesamum indicum (XP_011089823.1, 45.65%), Solanum lycopersicum (XP_019069701.1, 44.57%), Lactuca sativa (XP_023748294.1, 43.12%), Arabidopsis thaliana (AT3G46130.2, 38.97%). Phylogenetic analysis showed that IbMYB48 has a close relationship with that of Ipomoea triloba (Figure 1C).

Next, the promoter region of the IbMYB48 gene was cloned from the JS6-5 genomic DNA by the Genome Walking method [41]. The size of 2293 bp promoter contained many kinds of regulatory elements, such as TCA (cis-acting element involved in salicylic acid responsiveness), TGACG (cis-acting regulatory element involved in the MeJA-responsiveness), LTR (cis-acting element involved in low-temperature responsiveness), ERE (ethylene-responsive element), MBS (MYB binding site involved in drought-inducibility) and O2-site (cis-acting regulatory element involved in zein metabolism regulation) motif, of which we mainly focused on elements associated with abiotic stress resistance (Figure 1D). Moreover, The IbMYB48 gene consists of a total of 3 exons and 2 introns through comparison of ORF and genomic sequence of IbMYB48 (Figure 1D).

3.2. Expression Level of IbMYB48 Responses to Salt or Drought Treatments

To gain more insight into the biological function of IbMYB48, the expression profiling of IbMYB48 in vitro-grown plants of JS6-5 were examined by qRT-PCR. Our results revealed that IbMYB48 was widely detected in different tissues such as stem, root, and leaf, but it was predominantly expressed in leaf and root (Figure 2A).
Figure 2. Expression analysis of \textit{IbMYB48}. (A) Tissue-specific expression of \textit{IbMYB48} in the stem, root, and leaf of JS6-5 plants. (B) Histochemical staining of GUS activity in \textit{IbMYB48pro::GUS} transgenic Arabidopsis plants subjected to salt stress. Bars = 3 mm. (C) Expression analysis of \textit{IbMYB48} in whole plants of JS6-5 in response to 100 µM ABA, 10 mM H$_2$O$_2$, 100 µM MeJA, 200 mM NaCl, 20% PEG6000 and 100 µM SA, respectively. The \textit{IbActin} was used as the reference. Error bars reflect ± SE scores (\(n = 3\)); Asterisk (*) and (**) indicate a significant difference at the level \(p < 0.05\), \(p < 0.01\), individually.

To investigate whether \textit{IbMYB48} is involved in salt or drought stress in sweet potato, the \textit{IbMYB48pro::GUS} fusion plasmid was transformed into Arabidopsis. Two weeks old transgenic Arabidopsis plants were treated with 100 mM NaCl and 300 mM mannitol for 12 h and stained before and after treatment. The results showed that promoter region from \textit{IbMYB48} was sufficient to confer GUS expression in whole plants. We observed stronger GUS staining in both roots and leaves of \textit{IbMYB48pro::GUS} lines after NaCl and mannitol treatments, indicating that the \textit{IbMYB48} promoter contains cis-regulatory elements responding to NaCl and mannitol (Figure 2B).
To further analyze its potential function, the expression of *IbMYB48* was analyzed under different stress conditions, including 100 µM ABA, 10 mM H$_2$O$_2$, 100 µM MeJA, 200 mM NaCl, 20% PEG6000, and 100 µM SA for 0, 0.5 h, 1 h, 3 h, 6 h, 12 h, 24 h and 48 h, respectively. These results showed that the expression of *IbWRKY48* was significantly induced by ABA, H$_2$O$_2$, MeJA, NaCl, PEG, and SA, and peaked at 48 h with 1.9-fold, 12 h with 3.0-fold, 24 h with 5.3-fold, 24 h with 11.0-fold, 24 h with 4.5-fold, and 24 h with 4.9-fold, respectively (Figure 2C).

### 3.3. *IbMYB48* Is Localized in the Nucleus

To explore the subcellular localization of the *IbMYB48*, we performed a transient expression experiment of *IbMYB48* in onion epidermal cells. The C terminus of *IbMYB48* was fused with GFP under the control of the CaMV 35S promoter, and the construct was introduced into onion epidermal cells using gene gun method [42]. In the cells expressing GFP alone, the signal was detected in the cytoplasm and nucleus. In contrast, the green fluorescence signal from the *IbMYB48*-GFP fusion protein was mainly detected in the nucleus, indicating that *IbMYB48* is a nucleus protein (Figure 3).

![Figure 3](image_url)

**Figure 3.** Transient expression and localization analysis of *IbMYB48*-GFP in onion epidermal cells. Specimens were observed using fluorescent microscope Nikon TE-2000E with 488 nm excitation wavelength laser. Bars = 30 µm.

### 3.4. *IbMYB48* Possesses Transcriptional Activation Activity

To analyze whether *IbMYB48* can serve as a transcriptional activator and which region of *IbMYB48* has the transcriptional activity, full-length and different truncation of *IbMYB48* construct vectors (pGBK7-*IbMYB48* $^{1-109}$, pGBK7-*IbMYB48* $^{110-266}$, pGBK7-*IbMYB48* $^{1-266}$) were constructed for transactivation analysis in yeast. The resulting plasmids, pGBK7-Lam (negative control) and pGBK7-53 (positive control) were then transformed into yeast strain Y2HGold, respectively (Figure 4A).
Figure 4. Experiment of IbMYB48 transactivation activity in yeast using full-length as well as truncated forms. (A) The pGBK7-53 (P) and pGBK7-Lam (N) were used as positive and negative control, respectively. Part I, Part II, and Part III represents the pGBK7-IbMYB48\(^{1-266}\), pGBK7-IbMYB48\(^{1-109}\) and pGBK7-IbMYB48\(^{110-266}\) were transformed into Y2HGold, individually. (B) The transformed yeast cells were grown in SD/-Trp/-His/3-AT medium with X-\(\alpha\)-gal. Yeast cells carrying any one of the five vectors grew well on SD/-Trp medium. Yeast cells containing pGBK7-IbMYB48\(^{110-266}\), pGBK7-IbMYB48\(^{1-266}\) and the positive control grew properly on SD/-Trp/-His/3-AT/X-\(\alpha\)-gal medium displaying \(\alpha\)-galactosidase activity, whereas the yeast cell containing pGBK7-IbMYB48\(^{1-109}\) or negative control did not grow (Figure 4B). These results demonstrate that IbMYB48 is a nuclear protein and C-terminal region of IbMYB48 is responsible for its transcriptional activation activity.

3.5. Expression of IbMYB48 Enhances Salt and Drought Tolerance in Arabidopsis

To further analyze the biological function of IbMYB48 in plants, we next generated transgenic plants carrying IbMYB48 under the control of the CaMV 35S promoter. A total of 8 positive T\(_3\) transgenic lines were obtained (Figure 5A). The mRNA expression level of IbMYB48 in different lines was analyzed by qRT-PCR (Figure 5B). Three independent lines (L6, L8, and L12) with the high expression level were chosen for subsequent physiology experiments.

Then, transgenic plants and WT seedlings were treated with different stress conditions. T\(_3\) transgenic lines L6, L8, L12 and WT seeds were cultured on 1/2 MS medium which contains 100 mM NaCl or 300 mM mannitol for 7 days, respectively. There were no phenotypic differences between the WT and transgenic plants under normal conditions (Figure 6A). However, the growth status of WT was inferior to the transgenic lines under NaCl or mannitol treatments. We assessed plant tolerance to abiotic stress by measuring the length and fresh weight of roots. The root length of transgenic plants was longer than WT and the fresh weight of transgenic plants was higher than WT under both salt and mannitol treatments (Figure 6B,C).
During the seed’s germination stage, the germination rate of the WT seeds was markedly decreased, while \( \text{IbMYB48} \)-overexpression transgenic plants exhibited higher germination rate compared with WT under NaCl and mannitol stresses. These results indicated that overexpression of \( \text{IbMYB48} \) could enhance the resistance to salt and drought stresses (Figure 6D).

To further verify \( \text{IbMYB48} \) could contribute to salt resistance, the above transgenic lines were grown in potting soil mixture under normal condition for 10 days; And then, these lines were treated with \( \frac{1}{2} \) Hoagland solution supplemented with 300 mM NaCl for 3 weeks. The leaves of the WT exhibited chlorosis and wilted, while the leaves of the transgenic lines remained green and the overall growth performances was better than WT under salt stress. Under drought treatment, most of the WT wilted and died, while the majority of the transgenic lines survived and showed relatively good growth (Figure 7A). Taken together, these morphological changes indicated that transgenic lines have better growth under salt and drought stresses.
Figure 6. Response of transgenic and WT seedlings under salt and drought stresses. (A–C) Root length and fresh weight of WT and transgenic lines cultivated separately on 1/2 MS medium with normal (A), 100 mM NaCl (B) or 300 mM mannitol (C) for 7 days. (D) The germination rate of transgenic and WT seeds, which were cultured on 1/2 MS medium with 100 mM NaCl or 300 mM mannitol, respectively. Asterisk (*) and (**) indicate a significant difference from the WT at values of \( p < 0.05 \) and \( p < 0.01 \), individually, by Student’s \( t \)-test.
Figure 7. Responses of transgenic lines and WT grown in pots under salt and drought stresses. (A) The transgenic lines and WT grown in 1/2 Hoagland solution with 300 mM NaCl. Performance of transgenic lines and WT treated with soil drought stress without water for 3 weeks and then recovered for 2 days. Bar = 2 cm. (B–G) ABA (B), JA (C), proline (D), MDA (E), and H$_2$O$_2$ (F) contents and SOD activity (G) in the transgenic Arabidopsis plants and WT under salt and drought stress. Values are shown as mean ± SE (n = 3); Asterisk (*) and (**) indicate that there is a significant difference at p < 0.05 and p < 0.01, individually.

To explore the potential mechanism of IbMYB48 in stress response, several physiological indexes related to stress tolerance were evaluated. The contents of ABA, JA and proline were considerably increased in transgenic lines compared to the WT, and the contents of MDA and H$_2$O$_2$ were significantly decreased in the transgenic plants after salt and drought treatments (Figure 7B–F). In addition, SOD activity was increased by 28% and 86% in transgenic plants under salt and drought treatments, respectively (Figure 7G). All these findings indicated that transgenic lines show decreased sensitivity to salt and drought stresses by regulating the plant hormone levels and osmotic balance.

3.6. Expression of the Stress-Related Genes Are Up-regulated in Transgenic Arabidopsis Plants

To further investigate the molecular mechanism underlying the IbMYB48 involved in salt and drought resistance, the expression levels of stress-responsive genes were investigated. Under salt and drought treatments, overexpression of IbMYB48 up-regulated the genes involved in ABA biosynthesis (AAO, ZEP and ABA1), JA biosynthesis and signaling
pathway (AOC1, LOX2, OPR3 and MYC2), proline biosynthesis (P5CS and P5CR), and the reactive oxygen species (ROS) scavenging (DHAR, SOD and CAT) in transgenic lines compared to WT (Figure 8).

Figure 8. The expression profile of stress-related genes in transgenic lines and WT under salt and drought treatments for 2 weeks. The aerial parts of transgenic and WT plants were collected to extract total RNA. qRT-PCR analysis of the expression of ABA biosynthetic genes (A–C), JA biosynthesis and signaling pathway genes (D–G), proline biosynthetic genes (H,I) and ROS-scavenging genes (J–L). Values are indicated as means ± SE (n = 3); Asterisk (**) indicates that there is a significant difference at p < 0.01, individually. AAO, ABA-aldehyde oxidase; ZEP, zeaxanthin epoxidase; ABA1, ABA-deficient 1; AOC1, allene oxide cyclase 1; LOX2, 13-lipoxygenase 2; OPR3, 12-oxophytodienoate reductase 3; P5CS, pyrroline-5-carboxylate synthase; P5CR, pyrroline-5-carboxylate reductase; DHAR, dehydroascorbate reductase; SOD, superoxide dismutase; CAT, catalase.

4. Discussion

TFs act as principal modulators of plant growth progresses. Hence, they are expected to be superb candidates for improving important traits in crops [43]. MYB transcription factors represent one of the largest protein family in plants [44]. A growing number of studies have shown that the R2R3-MYB presumably has various functions in the regulation of different
abiotic stress responses. For instance, it is demonstrated that AtMYB2 and AtMYB60 are engaged in drought tolerance stress in plants [14,15]. AtMYB44 and AtMYB2 respond to salt stress at the transcript level and overexpression of AtMYB44 and AtMYB2 could enhance the resistance of transgenic Arabidopsis plants to salt stress [14,17]. Heterologous expression of GmMYB81 in Arabidopsis can intensify salt and drought tolerance during seed germination [45]. Several R2R3-MYB genes, such as rice OsMYB91, apple MdSIMYB1 and wheat TaMYB33, can increase salt and drought tolerance of transgenic plants [46–48]. However, few MYB family members have been studied in sweet potato under abiotic stress conditions.

Here, we cloned a MYB member obtained from the transcriptome database. The R2R3-MYB functional domain has been identified in IbMYB48 (Figure 1B). IbMYB48 possesses transcriptional activation activity, implying that IbMYB48 may act as an activator to affect the expression of downstream genes (Figure 4B). In further experiments, IbMYB48 is proved to be induced by ABA, H2O2, MeJA, NaCl, PEG6000 and SA treatments. Overexpression of IbMYB48 confers salt and drought tolerance in Arabidopsis (Figures 2C and 7A).

In Arabidopsis, AtMYB60 transcript is primarily expressed in guard cells [49]. Mutation of this gene leads to guard-cell-specific defects without affecting other developmental and physiological processes [49]. AtMYB52 is much more abundantly expressed in roots and siliques compared to flowers and leaves [50]. In rice, OsMYB48-1 is expressed most abundantly in roots at both seedling stage and reproductive stage but expressed at a low level in the sheath at seedling stage [51]. The highest expression level of OsMYB2 is measured in leaves, which was followed by roots and shoots [52]. In soybean, GsMYB15 is mainly detected in the roots, leaves and stems of transgenic plants and was a notably strong expression in the pods and flowers [53]. In the present research, tissue expression pattern analysis exhibited that IbMYB48 had higher expression levels in leaf and root (Figure 2A). Compared to other MYB genes in the above species, differential tissue expression patterns of IbMYB48 implied that it might have different functions. Furthermore, the relevant physiological parameters of IbMYB48-overexpression transgenic Arabidopsis under salt and drought stresses are consistent with those of ZmMYB3R, ZmMYB48 and FtMYB9, which were described to be engaged in salt or drought tolerance in plants. The ZmMYB3R-overexpression Arabidopsis plants show increased activities of the antioxidant enzymes CAT, POD and SOD after salt and drought treatments compared to WT plants [54]. The contents of proline and MDA are elevated in both ZmMYB48-overexpression Arabidopsis plants and WT under drought stress; however, the transgenic plants have markedly higher proline content compared to WT as well as significantly lower MDA content than WT [55]. When plants are subjected to salt and drought treatments, the MDA content of FtMYB9-overexpression Arabidopsis plants are lower than WT [56]. In addition, the proline contents of FtMYB9 transgenic plants are shown to be higher than those of WT during normal and abiotic stress conditions [56]. In the present research, we found that the contents of proline and SOD activities were significantly increased, and the contents of MDA and H2O2 were significantly decreased in transgenic plants under salt and drought stresses, suggesting that transgenic plants maintain intracellular osmotic balance and cell membrane integrity (Figure 7D–G).

In plants, ABA biosynthetic pathway related to salt and drought stress have been elucidated [57]. ABA is biosynthesized de novo from a C40 carotenoid precursor β-carotene by the oxidative cleavage of neoxanthin as well as a two-step conversion of xanthoxin to ABA via ABA-aldehyde [57,58]. There are some key enzymes that are essential for this series of synthetic reactions, like ZEP, 9-cis-epoxycarotenoid dioxygenase (NCED), AAO, molybdenum cofactor sulfurase (MCSU) [58]. ABA1 encodes a ZEP that has an important role in ABA biosynthesis [59]. When plants are exposed to salt stress, the ABA concentration is distinctly increased [60]. Meantime, ABA can increase the proline accumulation in plants [61]. Some identified MYB proteins could response to abiotic stress through ABA biosynthetic pathways, such as OsMYB48-1, OsMYB91, ZmMYB3R and TaMYB33. The reaction catalyzed by NCED to produce xanthoxin from neoxanthin in plastid is commonly
ThMYB9 was strongly induced by JA treatment, and its expression reached the maximum at 10 mM MeJA (Figure 2). The JA content was increased under salt and drought stresses with WT after salt and drought treatments (Figure 8H,I), which could improve the intra-GUS expression. Additionally, MeJA could induce ABA biosynthetic genes ABA1, ABA2 and NCED3 are higher in ZmMYB3R-overexpression transgenic lines than in WT plants after salt and drought treatments [54]. Similarly, in the present research, it was demonstrated that the contents of ABA in transgenic plants are higher than the WT under salt and drought stresses (Figure 7B). Overexpression of IbMYB48 up-regulated the genes associated with the ABA biosynthetic pathway (AAO, ZEP, ABA1) in overexpression lines compared with WT under salt and drought treatments (Figure 8A–C). These results indicate that IbMYB48 may participate in ABA biosynthetic pathway along with additional factors to enhance the abiotic stress resistance. It is generally considered that plant salt and drought-responsive genes are mainly involved in both ABA-independent and ABA-dependent pathways [1]. RD29A and ERD1 play an important role in ABA-independent pathway [66,67]. In wheat, the expression level of RD29A and ERD1 is significantly up-regulated in the TaMYB30-B-overexpression transgenic plants [19]. In GmMYB118-overexpression plants, the expression levels of DREB2A, RD29A are induced under drought stress, indicating that GmMYB118 could improve plant tolerance to drought stress by activating both the ABA-independent and ABA-dependent pathway genes [68].

JA is a compound derived from lipids that function as a signal molecule during plant stress response and development [69]. In Arabidopsis, there are numerous enzymes involved in the JAs biosynthesis have been extensively characterized, such as phospholipases (PLA), LOX, allene oxide synthase (AOS), AOC, OPC-8:0 CoA ligase 1 (OPCL1) and thioesterase (TE) [69–71]. In the present research, we demonstrated that the expression of IbMYB48 was the highest at 24 h when the sweet potato mutant JS6-5 were treated with 10 mM MeJA (Figure 2). The JA content was increased under salt and drought stresses in IbMYB48-overexpression transgenic plants compared with WT (Figure 7C). Similarly, ThMYB9 was strongly induced by JA treatment, and its expression reached the maximum at 72 h [72]. JcMYB2 expression level is gradually increased as well as also reached a peak level at 6 h under MeJA treatment, then dramatically decreased and back to the primal level [73]. Additionally, MeJA could induce GUS expression in JcMYB2pro::GUS transgenic plants via GUS histochemical staining [73]. In the present research, under salt and drought stresses, the JA biosynthesis and signaling genes LOX2, OPR3, AOC1 and MYC2 were up-regulated in the IbMYB48-overexpression transgenic plants compared with WT (Figure 8D–G). These results imply that the genes of JA biosynthesis pathway and JA signaling transduction are activated by IbMYB48 and improve abiotic stress tolerance of IbMYB48-overexpression transgenic plants.

Proline is one of the osmotic stress regulation substances that respond to cell dehydration [74]. Under drought stress, plants can accumulate more proline in response to stress [75]. In the present research, the contents of proline were considerably increased in transgenic lines under salt and drought treatments (Figure 7D). The expression of proline synthase genes P5CS and P5CR was significantly increased in transgenic plants compared with WT after salt and drought treatments (Figure 8H,I), which could improve the intracellular osmotic balance and cell membrane integrity of the transgenic plants, thereby enhancing the stress resistance of transgenic plants.

ROS plays a double role in response to abiotic stresses as toxic by-products of stress metabolism, as well as important signal transduction molecules in plants [76]. Singlet oxygen (¹O₂), superoxide (O²⁻), H₂O₂ and hydroxyl radicals (·OH) are one of the main forms of ROS in plants [77]. ROS accumulation mainly depends on the dynamic balance between ROS production and ROS scavenging under salt and drought stresses [78]. It is imperative
for plants to control ROS toxicity effects through ROS scavenge system. SOD and CAT were essential for ROS-scavenging mechanism [76,77]. In the present research, SOD activity and H$_2$O$_2$ content were significantly increased and decreased in ibMYB48 overexpression transgenic plants than those in WT under salt and drought stresses, respectively (Figure 7F,G). The expression of genes encoding DHAR, SOD, and CAT, members of the ROS scavenging system, was all increased in transgenic plants compared with WT (Figure 8J–L). These results indicated that overexpression of ibMYB48 activates the genes of ROS scavenging system under salt and drought stresses and improves the salt and drought tolerance.

Taken together, these results reveal that the ABA biosynthetic pathway, JA biosynthesis and signaling pathway and ROS-scavenging system are activated by overexpression of ibMYB48, thus improving abiotic stress tolerance of transgenic plants (Figure 9). However, ibMYB48 is only heterologously expressed in Arabidopsis for correlated functional validation. In the future, we will further investigate the potential function of ibMYB48 in salt and drought tolerance by using gene-editing technology.

![Figure 9. The proposed regulatory network model of IbMYB48 conferring salt and drought tolerance.](image)

Under salt or drought stress, the expression level of IbMYB48 gene is induced. Elevated expression of IbMYB48 up-regulates the genes involved in the ABA, JA and proline biosynthetic or signaling pathways. As a result, the contents of ABA, JA, and proline are elevated, meanwhile, the ROS scavenging system is activated, thus improving salt and drought resistance in plants through maintaining osmotic balance and membrane integrity.

5. Conclusions

In conclusion, we cloned and characterized IbMYB48 from sweet potato mutant line JS6-5. IbMYB48 was found to be a nuclear-protein, consistent with functioning as a TF. Self-transcriptional activation domain of IbMYB48 located in the C-terminal region. Ectopic expression of IbMYB48 in Arabidopsis results in increased ABA, JA, proline contents and SOD activity, indicating that IbMYB48 positively regulates tolerance to salt and drought stresses. Our results suggest IbMYB48 may be a candidate gene for improving plant tolerance to abiotic stress.
**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/genes13101883/s1, Table S1: The primers used in this study. Figure S1: The pCAMBIA1300-IbMYB48-GFP vector used in this study.

**Author Contributions:** Conceptualization, H.Z. (Hong Zhai); validation, Q.L.; formal analysis, H.Z. (Hong Zhai); formal analysis, H.Z. (Hongyuan Zhao), H.Z. (Haqiang Zhao) and S.G.; investigation, H.Z. (Hongyuan Zhao), Y.H., H.Z. (Huan Zhang), S.Z. and N.Z.; writing—original draft preparation, H.Z. (Hongyuan Zhao) and H.Z. (Haqiang Zhao); writing—review and editing, H.Z. (Hong Zhai) and S.G.; funding acquisition, S.H., H.Z. (Hongyuan Zhao), N.Z. and Q.L. All authors have read and agreed to the published version of the manuscript.

**Funding:** This work was supported by the National Key R&D Program of China (grant numbers 2018YFD1000704 and 2018YFD1000700), the National Natural Science Foundation of China (31872878 and 31901599), the National Key R&D Program of China (grant numbers 2019YFD1001301 and 2019YFD1001300), and the earmarked fund for CARS-10-Sweetpotato.

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Conflicts of Interest:** The authors declare no conflict of interest.

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