TcMYC2a, a Basic Helix–Loop–Helix Transcription Factor, Transduces JA-Signals and Regulates Taxol Biosynthesis in Taxus chinensis

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INTRODUCTION

Taxol is widely used as an essential anticancer medicine for the effective clinical treatment of ovarian cancer, breast cancer, lung cancer, Kaposi sarcoma, cervical cancer, and pancreatic cancer, among others (Schiff et al., 1979). More than 20 enzymes are involved in catalyzing geranylgeranyl pyrophosphate to taxol, and the biosynthesis pathway is highly complex (Figure 1; Croteau et al., 2006). Nearly all known taxol-biosynthesis genes are up-regulated by JA and its derivatives, such as

Abbreviations: BAPT, phenylpropanoyltransferase; bHLH, basic helix loop helix; DBAT, 10-deacetylbaccatin III-10-O-acetyltransferase; DBTNBT, 3′-N-debenzoyltaxol N-benzylotransferase; IAA, jasmonate acid; PAM, phenylalanine aminomutase; T10H, 5-alpha-taxadienol-10-beta-hydroxylase; T13H, 13-alpha-hydroxylase gene; T5H, taxadiene 5-alpha hydroxylase; TASY, taxadiene synthase; TAT, taxadienol acetyltransferase; TBT, taxane 2-alpha-O-benzoyltransferase; zip, leucine zipper.
JA-responsive TcMYC2a regulates taxol biosynthesis

TASY, DBAT, DBTNBT, T7H, T2H, etc., but its mechanism is poorly understood (Figure 1) (Zhang et al., 2015b; Du et al., 2017).

Jasmonate acid (JA) is a vital endogenous hormone that regulates taxol biosynthesis and other biological processes, especially (a)biotic stress tolerance in various plants (Roberts, 2001). The physical interactions of JASMONATE ZIM-domain (JAZ) with several downstream transcription factors reportedly control the JA signal-transduction system, also called the JA signaling pathway (Deshaies, 1999; Thines et al., 2007; De Geyter et al., 2012). These downstream transcription factors are functionally disabled by physical interaction with JAZ until JA signaling molecules release them (Chini et al., 2016; Zhang et al., 2017). Among all JA downstream transcription factors, MYC2 is considered as the fatal point of the entire JA signaling pathway (Kazan and Manners, 2013).

MYC2, which belongs to clade IIIe of the basic helix–loop–helix (bHLH) superfamily that is one of the largest group of transcription factors in plants, is the master JA regulator involved in secondary metabolite biosynthesis (Heim et al., 2003; Chini et al., 2007, 2016; Schweizer et al., 2013; Berens et al., 2017). MYC2 is essential for JA response to secondary-metabolite accumulation in several plants. In Catharanthus roseus, CrMYC2 activates the expression of ORCA3 gene, which responds to JA signal and functions as a crucial activator of STR gene, resulting in vinblastine accumulation (van der Fits and Memelink, 2000; Zhang et al., 2011). In Nicotiana tabacum, NtMYC2a/b, which can bind with NtJAZ1/2/3, is able to activate the expression of the following key genes of nicotine biosynthesis: putrescine N-methyltransferase (PMT), quinolinate phosphoribosyltransferase (QPT2), and NtA622. NtMYC2a/b can also bind ERF189 and ERF221, which are the major JA-responsive activators of PMT2 (De Boer et al., 2011; Shoji and Hashimoto, 2011; Zhang et al., 2012). In Artemisia annua, AaMYC2 binds with AaJAZ1-4 and activates CYP71A1V and DDBR2 genes, which encode enzymes of artemisinin biosynthesis (Shen et al., 2016). In Salvia miltiorrhiza, SmMYC2a/b and SmJAZ1/2 can form a dimer complex, and the contents and expression of genes of tanshinones and phenolic acids are repressed in SmMYC2a/b knockdown plant lines. Thus, SmMYC2a and SmMYC2b are positive regulators (Zhou et al., 2016). In rice, JA-inducible OsMYC2 drastically enhances the activity of naringenin 7-O-methyltransferase (OsNOMT) promoter and is essential for JA-inducible sakuranetin production (Ogawa et al., 2017b). All these studies indicate that MYC2 is a vital regulator of secondary-metabolite biosynthesis and that the JAZ–MYC2 complex is critical to the JA signaling pathway.

Taxadiene synthase (TASY) promoter has been obtained and characterized, and the fragment containing the GCC-box at −142 bp is found to be the minimum region responding to JA signals. Thus, TcERF12 and TcERF15 are JA-responsive ERF factors that negatively and positively regulate TASY gene expression by binding with the GCC-box (Zhang et al., 2015b). However, how TcERF12 and TcERF15 respond to JA signals is unclear. Our previous study has shown that TASY promoter has a T/G- (AACGTG; −256 bp) and a G-box (CACGTG, −158 bp), both of which are MYC2 binding cis-elements. The two cis-elements contribute to JA responses, indicating that MYC2 is a candidate regulator of taxol biosynthesis responding to JA (Zhang et al., 2015b).

In the present work, a MYC2 transcription factor was screened in Taxus chinesis and named as TcMYC2a. Homolog search and phylogenetic analysis identified it is a typical MYC2 transcription factor, and conserved domains and motifs showed its potential functions in JA signaling pathway. Yeast two-hybrid experiments verified the interaction of TcMYC2a with Taxus JAZ protein. Then, a series of experiments would be performed to determine the function of TcMYC2a in taxol biosynthesis and explain the regulation system.

MATERIALS AND METHODS

RNA Isolation and cDNA Synthesis

Taxus chinesis cell line #48 was established using the callus cultures derived from a budding Taxus stem in May 2003 and was maintained in modified Gamborg’s B5 medium (Zhang et al., 2000). RNA isolation were conducted by RNAprep Pure Plant Kit (TIANGEN, Beijing, Cat: DP441), and the first chain of cDNA were transcribed by ReverTra Ace® qPCR RT Master Mix with gDNA Remover kit (Toyobo, Japan, Code No. FSQ-301).

Screening, Sequence Alignment and Phylogenetic Analyzing of TcMYC2a

The HMM files of HLH (Accession NO.: PF00010), bHLH-MYC_N (Accession NO.: PF14215) and ACT (Accession NO.: PF01842) domains were retrieved from the Pfam database1. Then, the HMMsearch program of HMMER package2 was used to identify the MYC2 proteins from our previous transcriptome datasets in T. chinesis (Zhang et al., 2015a,b; Liao et al., 2017). Specific primers (Supplementary Table S1) were designed to amplify TcMYC2a from the cDNA of T. chinesis under the following PCR parameters: initial denaturation at 95°C for 6 min; 30 cycles of denaturation at 94°C for 30 s; annealing at 58°C for 30 s; extension at 72°C for 2 min; and a final extension at 72°C for 10 min. The PCR products were subcloned into pMD18-T (TaKaRa, Japan) for sequencing. BLAST3 was used for the homology search from Swiss-prot database. ClustalW online4 at the default setting was used to align and evaluate the percentage identity of TcMYC2a with known MYC2s. The alignment results were sent to ESPript 3.05 for coloring (Robert and Gouet, 2014). Phylogenetic analysis was performed with a neighbor-joining (NJ) method by using 1000 bootstrap resamplings in MEGA 5.05.

Subcellular Localization

TcMYC2a was inserted into pCAMBIA1300-sGFP with Kpn I and Sal I restriction sites, then recombiant vector was transformed into Taxus chinensis suspension cultures derived from a budding Taxus stem in May 2003 and was maintained in modified Gamborg’s B5 medium (Zhang et al., 2000). RNA isolation were conducted by RNAprep Pure Plant Kit (TIANGEN, Beijing, Cat: DP441), and the first chain of cDNA were transcribed by ReverTra Ace® qPCR RT Master Mix with gDNA Remover kit (Toyobo, Japan, Code No. FSQ-301).

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1http://pfam.sanger.ac.uk
2http://hmmer.org/
3http://www.ncbi.nlm.nih.gov/BLAST/
4https://npsa-prabi.ibcp.fr/cgi-bin/npsa_automat.pl?page=/NPSA/npsa_clustalw.html
5http://espript.ibcp.fr/ES Pript/cgi-bin/ES Pript.cgi
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**FIGURE 1** | Transcription regulation of taxol biosynthesis. Current progress of transcription regulation of taxol biosynthesis. TASY (taxadiene synthase), TSH (taxadiene 5-alpha hydroxylase), TAT (taxadiene 5-alpha hydroxylase), T10H (6-alpha-taxadienol-10-beta-hydroxylase), T13H (13-alpha-hydroxylase gene), TBT (taxane 2-alpha-O-benzoyltransferase), DBTNBT (3’-N-debenzoyl taxol N-benzoyltransferase). BAPT (phenylpropionaldehyde), DBAT (10-deacetylbaccatin III-10-O-acetyltransferase) and PAM (phenylalanine aminomutase).

into LBA4404. 1 cm × 1 cm size onion epidermal cells were infected by positive clones in 1/2 MS liquid medium at 25°C, dark for 2 days. The infected onion epidermal cells were then plated on MS medium for 2 days at 25°C in darkness. The GFP fluorescence was observed by Laser scanning confocal microscope FV1000 (OLYMPUS, Japan).

**Yeast One-Hybrid and Yeast Two-Hybrid**

TcMYC2a was amplified using primers as TcMYC2ayF and TcMYC2ayR (**Supplementary Table S1**), then purified PCR product and pGADT7-Rec2 plasmid were digested by EcoRI and XmaI enzymes at the same time. T4 ligase was used to link the two fragments to construct the prey vector, TcMYC2a-pGADT7R2. For constructing bait vector, sense and antisense sequences of quadrupled T/G-box, G-box and E-box were artificially synthesized, respectively (**Supplementary Table S1**). The cohesive ends of EcoRI and SpeI would expose after annealing, then the double strand were linked with digested pHIS2.1 vector to form pbaitT/G and pbaitG, respectively. Finally, TcMYC2a-pGADT7R2 were co-transformed with two bait vector, respectively, into the yeast strain Y187, and Y187 cells were plated onto SD/Leu-/Trp-/His-deficient medium with 40 mM 3-AT for 3–5 days.

TcMYC2a-pGADT7R2 and pGADT7 were digested by EcoRI and XmaI, the coding region of TcMYC2a were inserted into pGADT7 to construct TcMYC2a-pGADT7. TcJAZ3 (Zhang et al., 2017) was inserted into pGBKT7 with EcoRI and BglII restriction sites for TcJAZ3-pGADT7. Then TcMYC2a-pGADT7 and TcJAZ3-pGBK7 were co-transformed into AH109 using LiAc conversion protocol. The transformants were screened on SD/Leu-/Trp-/His-deficient medium with 40 mM 3-AT for 3–5 days.

**Construction of Overexpression, RNAi Vector and Reporter Vectors**

The coding region of TcMYC2a was inserted into the pCAMBIA1303 vector with NcoI and BstEII restriction sites under control of the CaMV 35S promoter, resulting in the overexpression of TcMYC2a in Taxus cells. To construct TcMYC2a-RNAi vector, a 400 bp 3′ ORF sequence of TcMYC2a was inserted into pCAMBIA1300-35S-X forward (BamHI) and reverse (SpeI) so that SPLRNAi fragments were stuck between them. The primers were showed in **Supplementary Table S1**. The construct was transformed into the *T. chinensis* cells while the pCAMBIA1303 was used as the control.

**Analysis of TcERF12 and TcERF15 Promoters**

Firstly, the *T. baccata* genome sequences (Nystedt et al., 2013) were made a local database by the software BLAST+. Then, the ORF sequences of TcERF12/15 were used as the

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4ftp://ftp.ncbi.nlm.nih.gov/blast/executables/blast+/LATEST/
reference gene to blast with the local database at the default parameters. Therefore, the homolog sequences containing 5' upstream sequences and the ORF sequences were obtained. According to the homolog sequences, the specific forward primers were designed, and the specific reverse primer were designed according to ORF (Supplementary Table S1). Genome DNA was extracted by SDS-CTAB method and then used as a template. The specific product was T/A cloned, sequenced and aligned with designed partial sequence of TcERFs' ORF to decide the 5'-flanking region. The 5'-flanking sequence was analyzed online by PlantCare and PLACE at default settings.

**Transient Transformation and GUS Activity Analysis**

For overexpression of TcMYC2a, the TcMYC2a-p1303 vector was transformed into T. chinensis suspension cells mediated by Agrobacterium LBA4404, the detailed method referred to Zhang et al. (2015b). For co-overexpression, pTG, pG, pmTG, and pmG were co-transformed with TcMYC2a-p1303 into Taxus cells, respectively, quantitative analysis of GUS activity was conducted as our previous reports (Li et al., 2013). Each construction had three independent experiments. The transient transformations were conducted three times independently to make sure there were three independent biological repeats.

**Quantitative Real-Time PCR**

1 μl synthesized cDNA (diluted 1:5) was used as template for real-time PCR. Beta-actin expression was used to normalize all values in the RT-PCR assays in T. chinensis. Primers for RT-PCR were designed using Primer3 software (Supplementary Table S1). SYBR Green Realtime PCR Master Mix (Toyobo, Japan) was used to conduct qRT-PCR, StepOnePlus (Applied Biosystems, United States) was used to detect and analyze the reactions. To enable statistical analysis, three fully independent biological replicates were obtained, each experiment was conducted three times, and we used Student’s T-test.

**HPLC (High Performance Liquid Chromatography)-MS (Mass Spectrum) Analysis of Taxanes**

The transformation cells were 125 rpm shake at 25°C darkness for 6 days. Preparation of samples and HPLC analysis procedures were conducted as described previously (Song et al., 2014; Dong et al., 2015).

**RESULTS**

**TcMYC2a Shares Common Domains With MYC2 in Other Plants**

After homolog searching in the Swiss-Prot database, TcMYC2a was found to share the highest similarity with OsMYC2, which is a transcriptional activator involved in JA signaling pathway during spikelet development (Cai et al., 2014; Ogawa et al., 2017a). Sequence alignment showed that TcMYC2a contained all of the essential domains and motifs of MYC2 (Figure 2). Three conserved domains/motif were highly conserved in both TcMYC2a and other MYC2 factors: bHLH-zip-, bHLH-MYC_N-, and BIF/ACT-like-domain (Pauwels and Goossens, 2011).

Compared with GL3, EGL3, and TT8 (clade IIIf), TcMYC2a shared a more similar bHLH-zip domain and 14 aa co-adjacent N-terminal peptides with other MYC2s (clade IIIe) and clade IIId bHLHs. Clade IIIId and IIIe are always considered as one subgroup because they function complementarily and redundantly (Feller et al., 2011; Tominaga-Wada et al., 2011). Additionally, nearly all bHLH-zip domains of MYC2 factors had a SELS peptide (414 aa–417 aa) similar to the SDHS peptide of AtMYC2, which was assumed to be the phosphorylation site. This 4aa-length peptide was only highly similar among MYC2s but highly variable in clades IIId and IIIf (Figure 2D) (Kazan and Manners, 2013).

Except for TcJAMYC2 and TcJAMYC4, the other clade IIIId/eff bHLHs including TcMYC2a contained the bHLH-MYC_N-domain (Figure 2B). Our results showed that TcMYC2a had a Gly-Asp-Gly motif, as well as other JAZ-interactive bHLHs (IIIId-β), indicating that TcMYC2a can bind with JAZ proteins (Goossens et al., 2015). Additionally, non-MYC2 factors (IIIId/β) contained a RSLAKSA5 motif, in which K and A residues determined the transactivation strength of proteins, whereas all MYC2s had the RA(K/R)QAQQ(F/Y)LG-motif at the corresponding positions in the bHLH-MYC_N-domain. Meanwhile, TcMYC2a contained a RAQQAQLG-motif that was highly homologous with MYC2 factors, indicating that TcMYC2a was a MYC2 homolog.

The aspartokinase, chorismate, and TyrA (ACT) domain was first found in a wide range of metabolic enzymes and formed the ββαβ secondary structures (Chiapan and Shaanan, 2001). The ACT domain of bHLH was predicted to have a ββαββ secondary structure that differed from the ββαββ structure (Chang et al., 2017). According to our results, the BIF/ACT-like domain was present in all subgroup IIIId-IIIIf bHLHs, and the similarity of BIF/ACT-like domain was low among the three clades (Figure 2D). TcMYC2a contained the BIF/ACT-like domain that was more highly similar to MYC2 proteins, indicating that TcMYC2a was more closely related to MYC2s.

All these results showed that TcMYC2a was preferred as a MYC2 transcription factor and that highly conserved domains enabled TcMYC2a to share similar functions to other MYC2s.

**Phylogenic Analysis Grouped TcMYC2a With Other MYC2 Proteins**

Thirteen MYC2s, three clade IIIf (TT8, EGL3, and GL3), and three clade IIIId bHLH were selected for phylogenic analysis. Twenty proteins were grouped into three clades, the MYC2 clade (IIIe), clade IIId, and clade IIIf (Figure 3). TT8, EGL3 and GL3, and AtbHLH3/13/17 were grouped together.
in accordance with previous reports (Tominaga-Wada et al., 2011). In the present work, TcMYC2a was grouped with MYC2 proteins and was most closely related to TcJAMYC4, which was identified from *T. cuspidate* although TcJAMYC4 lacked most of the entire bHLH_MYC-N domain (Lenka et al., 2015).
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FIGURE 3 | Phylogenetic tree of TcMYC2a. Phylogenetic analysis of full-length TcMYC2a and other clade IIId/IIf bHLHs; TcMYC2a was framed by a solid box. Accession NO. in GenBank: TcMYC2a (MG494378), TcJAMYC1/2/4 (ACM48567.1/AGO03813.1/AGO03814.1), TmMYC2 (AHL44338.1), AtMYC2/3/4 (OAO98324.1/OAO98324.1/OAO98324.1), CrMYC2 (OAO98324.1), OsMYC2 (OAO98324.1), AaMYC2 (AKO62850.1), NtMYC2a/b (NP_001313001.1/NP_001312960.1), AtbHLH3/13/17 (AAL55710.1/AHL44338.1/Q9ZPY8.2), GL3 (NP_193864.1), EGL3 (AHL44338.1) TT8 (OAO98324.1) and DYT1 (OAO98340.1).

FIGURE 4 | Subcellular localization of TcMYC2a. The fused protein of TcMYC2a and GFP were transformed into onion epidermal cells. GFP fluorescence were observed 2 days after infecting by Laser scanning confocal microscope FV1000 (OLYMPUS, Japan) with 10×20 magnification.

TcMYC2a Localized in Nucleus

Transcription factors always function in nucleus, and MYC2 factors confirm that a nucleus localized signal peptide exists in the JID domain (Uji et al., 2016). Thus, we investigated TcMYC2a localization by expressing TcMYC2a-GFP in onion epidermal cells. Results showed that TcMYC2a actually localized in nucleus (Figure 4).

TcMYC2a Was a Target of Taxus JAZ Protein

To verify the roles of TcMYC2a in the JA-response system, TcJAZ3, which was a homolog of AtJAZ3, was used to detect interaction in a yeast two-hybrid system (Zhang et al., 2017). TcMYC2a-AD and TcJAZ3-BD vectors were constructed and then co-transformed into yeast AH109. The vacant AD and TcJAZ3-BD were also co-transformed as a control.

Apparently, the positive clones containing both TcMYC2a-AD and TcJAZ3-BD vectors grew on Trp-, Leu-, and His-deficiency media, whereas control clones did not, indicating that TcMYC2a was a target of TcJAZ3 (Figure 5).

TcMYC2a Can Bind With the T/G-, G-Box, and E-Box of TASY and TcERF12/15 Gene Promoters

Previously, a T/G-box (5′-AACGTG-3′; −256 bp to −251 bp) and a G-box (5′-CACGTG-3′; −158 bp to −153 bp) were found in the promoter of TASY gene, and deletion analysis indicates that T/G-box is related to JA response (Zhang et al., 2015b). In the present study, 5′-flanking sequences of TcERF12 and TcERF15 were obtained and analyzed with PlantCare. Results showed a T/G-box (−454 bp to −459 bp) in the 5′-flanking sequence of TcERF12 and G-boxes (CATATG, −736 bp to −741 bp) in the 5′-flanking sequence of TcERF15 (Figure 6).

Using quadruple T/G-box, G-box, and E-box as baits, Y187 cells containing bait vector and vacant pGADT7-Rec2 were inhibited when the concentration of 3-AT reached 40 mM in deficiency medium. The yeast Y187 with TcMYC2a-pGADT7-Rec2 vector and bait vector grew well on an SD/-Trp/-Leu/-His+40 mM 3-AT plate (Figure 6). These results showed that TcMYC2a can bind with T/G-, G-box, and E-box, indicating that TASY, TcERF12, and TcERF15 were the regulating targets of TcMYC2a.

TcMYC2a Activated the GUS Reporter Gene That Was Under Control of TASY Promoter by Functioning With Either T/G-Box or G-Box

Yeast one-hybrid experiment showed that TcMYC2a can bind with T/G- and G-box of TASY promoter in vitro, indicating that TASY gene was a direct target of TcMYC2a. Thus, to confirm the ability of TcMYC2a to bind with TASY promoter in vivo and analyze its ability in regulating transcription, T/G- and G-box of TASY promoter (−259 bp ~ 0 bp) were mutated and inserted into pBI121 by substituting 35S promoter, respectively, named
FIGURE 6 | TcMYC2a could bind with T/G-, G- and E-box in yeast. (A) The sketch map of promoters of TASY, TcERF12, and TcERF15 genes, the red boxes indicated T/G-box (5′-AACGTG-3′), blue hexagon was G-box (5′-CACGTG-3′) and green pentagon was E-box (5′-CATATG-3′). (B) Sketch map of the prey vector and bait vectors. The lighter gray boxes indicated T/G-, G-, or E-box, they were synthesized for four times and ligated into pHis2.1; while TcMYC2a was fused with GAL4 AD in pGADT7-Rec2. (C) The binding capability tests of TcMYC2a and cis-elements in yeast. Related bait vector with each cis-element was co-transformed with vacant pGADT7-Rec2 into Y187 as the control.

as mTtsp and mGtsp (Figure 7). Moreover, the vector with the fragment containing both mutated T/G- and G-box were used as control and named as mTgtsp. Each of the three vectors were co-transformed with TcMYC2a-p1303 into Taxus cells, and GUS activity was detected. Results showed that GUS activity increased 1.78- and 2.18-fold when TcMYC2a was co-expressed with mTtsp and mGtsp, respectively, indicating that TcMYC2a can activate downstream genes by functioning with either T/G-box or G-box (Figure 7).

TcMYC2a Induced Taxol Biosynthesis

To identify TcMYC2a function in taxol-biosynthesis genes, TcMYC2a was overexpressed in Taxus cells. The expression levels of taxol-biosynthesis genes were examined by qRT-PCR. Results showed that TASY, TAT, BAPT, DBTNBT, and T13H significantly increased in all three TcMYC2a-overexpression cell lines. Meanwhile, the other genes T5H, T10H, DBAT, and PAM were likely to be invariable in TcMYC2a-overexpression cell lines (Figure 8A). Thus, TcMYC2a function was limited compared with JA treatment, which increased nearly all taxol-biosynthesis genes, indicating that TcMYC2a was not the only way for JA signal transduction.

Additionally, we used HPLC-MS to evaluate the content of taxol in TcMYC2a-overexpression cells (Figure 8C). The content of taxol was found to be higher in TcMYC2a-overexpression cells than in control samples (Dong et al., 2015).

TcMYC2a Was an Activator of TcERF15 in Cells

TcERF12 and TcERF15 are reportedly regulators of TASY gene and are downstream modules in JA signal response (Zhang et al., 2015b). To clarify the relationship between TcMYC2a and TcERFs in the JA signaling pathway, the expression levels of TcERFs genes were verified in TcMYC2aOE cell lines. qRT-PCR results showed that both TcERF12 and TcERF15 were induced in overexpression cell lines (Figure 8B). When TcMYC2a was interfered with by RNAi (RNA interference), only TcERF15 showed a reduction, whereas TcERF12 expressed high stability in transgenic Taxus cells (Figure 9). All these results indicated that TcERF15 was the dominant target when TcMYC2a transduced JA signals to regulate taxol biosynthesis.

DISCUSSION

Jasmonate acid signaling pathway is proved to be one of the most important regulating systems in land plants; it is involved in growth, development, (a)biotic defense, and
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FIGURE 7 | TcMYC2a can transactivate either T/G- or G-box-mediated gene expression in Taxus cells. (A) Schematic diagram of mutant fragments of TASY promoter. TASY promoter revealed the original T/G- and G-box, and mTtsp indicated that the T/G-box was mutated by the sequence in the dotted boxes as well as the mGtsp and mTGtsp. (B) Schematic diagram of the reporter and effector plasmids used in transient assays. Each of mTtsp, mGtsp, and mTGtsp was fused to the GUS gene, respectively. Effector plasmids were under the control of the CaMV 35S promoter. (C) Relative GUS activities in TcMYC2a and related reporters co-transformed Taxus cells.

secondary metabolism (Roberts, 2001). The response mechanism to JA signal has been clarified in Arabidopsis, Oryza, Nicotiana, Solanum, and other angiosperms, but reports focusing on gymnosperm are few (van der Fits and Memelink, 2000; De Boer et al., 2011; Shoji and Hashimoto, 2011; Zhang et al., 2011, 2012; Zhou et al., 2016). Taxus spp. is an ancient gymnosperm conifer that provides us a precious anticancer drug, taxol. Moreover, the JA signaling pathway is one of the most efficient methods regulating taxol biosynthesis (Patil et al., 2012; Onrubia et al., 2013; Zhang et al., 2015b). Therefore, the mechanism by which taxol biosynthesis regulate JA signals can promote our understanding on the significance of the JA signaling pathway in both gymnosperm and angiosperm.

MYC2, which is a bHLH transcription factor, is the key element for transferring JA signals to downstream genes. MYC2 plays important roles in the crosstalk of the JA signaling pathway with other hormone signals (Hong et al., 2012). Recently, MYC2 factors have been discovered as crucial regulators of the JA-mediated activation of secondary-metabolite biosynthesis in medicinal plants, such as NtMYC2a/b, CrMYC2, and AaMYC2. The present study showed that TcMYC2a, first identified in T. chinensis, shared high similarity with these known MYC2 factors. Three conserved domains/motif were highly conserved in both TcMYC2a and other MYC2 factors: bHLH-zip-, bHLH-MYC_N-, and BIF/ACT-like-domain. Thus, TcMYC2a was a typical MYC2 factor with similar functions (Pauwels and Goossens, 2011).

The bHLH-zip domain, the common domain of almost all bHLH factors, consists of three conserved motifs, i.e., the basic-region motif responsible for DNA binding (Carretero-Paulet et al., 2010), the Helix-loop-helix motif, and the zip motif involved in homo-/hetero-dimerization
FIGURE 8 | Taxol-biosynthesis related genes were induced in TcMYC2a overexpression cells. The expression of (A) taxol biosynthesis genes, (B) regulator genes and (C) taxol contents (µg/ml) were quantified in TcMYC2a overexpression cell lines by qRT-PCR. Actin was used as reference gene, each experiment was conducted three repeats, and totally three biological repeats were used; the statistics method of p-value was Student’s T-test. Stars indicated the significance, * means that 0.01 < p-value < 0.05, ** means that p-value < 0.01. TASY (taxadiene synthase), T5H (taxadiene 5-alpha hydroxylase), TAT (taxadienol acetyltransferase), T10H (5-alpha-taxadienol-10-beta-hydroxylase), T13H (13-alpha-hydroxylase gene), TBT (taxane 2-alpha-O-benzoyltransferase), DBTNBT (β-N-debenzytaxol N-benzoyltransferase), BAPT (phenylpropanoyltransferase), DBAT (10-deacetylbaccatin III-10-O-acetyltransferase) and PAM (phenylalanine aminomutase).

(Fernandez-Calvo et al., 2011; Zhou et al., 2016; Lian et al., 2017). Apart from these conserved motifs, TcMYC2a contained the 14-aa co-adjacent peptide, which was conserved only in MYC2 (IIIe) and IIIb bHLHs, and a putative phosphorylation site (SELS in TcMYC2a that was homologous with SDHS residues in AtMYC2) was found only in MYC2 factors.

The bHLH-MYC_N-domain was also called MYB interactive region (Pattanaik et al., 2008) and JAZ-interactive domain, which was essential for binding with JAZ and MYB proteins (Fernandez-Calvo et al., 2011; Li et al., 2017). A transcriptional activation domain reportedly required for binding with MED25 in Arabidopsis is present in all MYC2 factors (Chen et al., 2012; Zhai et al., 2013). Conserved bHLH-MYC_N-domain factors show similar protein–protein interactions between TcMYC2a and other MYC2, indicating that TcMYC2a has similar functions. The BIF/ACT-like domain is necessary for homo-/heterodimerization and function as a switch that permits distinct configurations of a regulatory complex to be tethered to different promoters (Zhang et al., 2003; Kong et al., 2012).

Overall, results of molecular structural analysis of both gymnosperm and angiosperm plants showed that MYC2 transcription factors evolved extremely conservatively, indicating that MYC2 played crucial functions in the plant kingdom. As a homolog of OsMYC2, CrMYC2, AaMYC2a, and NtMYC2a/b, the potential of TcMYC2a in secondary-metabolite biosynthesis in T. chinensis was high.

In the JA signaling pathway, JAZ and MYC2 formed the most important heterodimer to control a series of downstream
bioactivities by protein–protein interaction, such as AtMYC2 and AtJAZs, NtMYC2b and NtJAZ1/2/3, and AaMYC2 and AaJAZs (Pauwels and Goossens, 2011; Shoji and Hashimoto, 2011; Shen et al., 2016). Previously, we have identified JAZ proteins from several transcriptome datasets of T. chinensis. TcJAZs and PsJAZs are found to have several evolutionary differences from AtJAZs. Among nine TcJAZ proteins, TcJAZ3 was most closely related to AtJAZs (Zhang et al., 2015b). The present results showed that TcMYC2a can bind with TcJAZ3 in yeast, indicating that TcMYC2a was a target of JAZ proteins and that the JA signaling pathway in Taxus spp. was similar to that in angiosperm plants.

Our previous work has identified the JA-responsive element of tasy gene promoter and confirmed that the GCC-box (TGCCGCT; from −143 to −136 bp), T/G-box (AACGTG; from −256 to −251 bp), and G-box (CACGTG; from −153 to −158 bp) were the effective response regions (Zhang et al., 2015b). AACGTG (T/G-box) and CACGTG (G-box) are classic MYC2-binding sequences (Qi et al., 2015; Du et al., 2017). In the present work, we found that TcMYC2a can bind with both T/G- and G-box in yeast, indicating TcMYC2a can regulate tasy gene expression. Results of GUS activity analysis showed that TcMYC2a can positively regulate downstream gene expression by binding with both T/G- and G-box. Subsequently, TcMYC2a overexpression in T. chinensis cells increased the expression of most taxol biosynthesis genes, especially acyltransferases. HPLC-MS analysis also showed that taxol contents increased in overexpression cells. All these results indicated that TcMYC2a was a positive regulator of tasy gene and even most taxol-biosynthesis genes.

However, TcMYC2a was not able to induce all taxol-biosynthesis genes, in contrast to JA stimuli. According to previous studies on taxol induction, JA application can generally promote the expression of taxol-biosynthesis genes in 12 h; for example, TASY, PAM, T2H, T13H, and T7H can increase several dozen times; DBAT, DBTNBT, and BAPT also increase (Patil et al., 2012; Onrubia et al., 2013). However, the present results showed that only several genes encoding acyltransferases and a few hydroxylation genes were up-regulated in TcMYC2a-overexpression cells, indicating that the JA signaling pathway had more than one way of TcMYC2a dependence (Hu et al., 2013).

TcMYC2a overexpression can also increase the expression of TcERF12 and TcERF15, which encode a negative and a positive regulator of tasy gene, respectively (Zhang et al., 2015b). RNAi analysis revealed that TcERF12 was not influenced by the down-regulation of TcMYC2a, indicating the possibility of other ways for TcERF12 to respond to JA stimuli. After 5’-flanking sequence analysis, TcERF12 gene was found to contain many JA-responsive cis-elements, such as a W-box (TGGACC; −505 to −500 bp), a T/G-box (CACGTT/AACGTG; −454 to −459 bp), and MYB-binding sites (CAGTTA; −63 to −58 bp). Considering the presence of these elements, TcMYC2a was capable of activating TcERF12 directly or indirectly in accordance with the results of TcMYC2a-overexpression analysis. In tobacco, MYC2 factors simultaneously activate negative and positive regulators, and down-regulation is observed for ERF189 as a negative regulator and other positive clade 2 ERFs in NtMYC2-RNAi plants (Shoji and Hashimoto, 2011). To prevent hypersensitive JA response, many researchers have reported that negative and positive factors are up-regulated simultaneously because they are essential to the balance of various bioactivities in plants (Nakata et al., 2013). Meanwhile, distinct preferences in DNA binding with the GCC-box are possessed by ERF repressors and activators, indicating that they regulate different targets in natural plants (Fujimoto et al., 2000; Zhang et al., 2015b).

In wild-type Taxus cells, TcERF15 seemed to be the key regulating target of TcMYC2a when transducing JA signals to taxol-biosynthesis genes. All these results indicated that TcMYC2a can transduce JA signals and regulate taxol biosynthesis by activating TcERF15 expression. In other plants, MYC2 factors regulate the biosynthesis of secondary metabolites.
in several ways. NtMYC2a/b can up-regulate biosynthesis genes and regulators, whereas CrMYC2 up-regulates biosynthesis genes by up-regulating regulators (De Boer et al., 2011; Zhang et al., 2011). Interestingly, physical interactions and cooperation between OsMYC2 and other factors enhance their transactivation activity (Ogawa et al., 2017a, b). Our results indicated that TcMYC2a had a similar functional mode with NtMYC2a/b. Additionally, interactors and cooperative regulators of TcMYC2a and known MYC2s should be further identified, so the dual functions of MYC2 factors should be clarified.

All these results indicated two regulating pathways of TcMYC2a (Figure 10): (i) TcMYC2 can directly bind with T/G-, G-, and E-box present in promoters and then activate the expression of taxol-biosynthesis genes; and (ii) TcMYC2a can activate the expression of transcriptional factors to regulate downstream genes. These results were highly accordance to those observed in tobacco, C. roseus, and tomato (Shoji and Hashimoto, 2011; Zhang et al., 2011; Du et al., 2017).

Furthermore, TcMYC2a was proved to be the crucial element for transducing JA signals to downstream responsive genes and then positively regulate taxol biosynthesis either directly or by regulating ERF regulators. Our results indicated that Taxus had a similar JA signaling pathway to angiosperm plants, supposing that the JA signaling pathway was highly conservatively evolved land plants.

DATA AVAILABILITY

All sequences obtained in this paper has submitted into GenBank, and their accession numbers are MG494378 (TcMYC2a), MG494379 (TcERF12p), and MG494380 (TcERF15p).

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AUTHOR CONTRIBUTIONS

MZ, CF, and LY conceived and designed the research. MZ, XJ, YC, SZ, MW, and WL conducted the experiments. MZ analyzed the data and wrote the manuscript. All authors read and approved the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fpls.2018.00863/full#supplementary-material

TABLE S1 | Primers and synthesized fragments used in this paper. All primers used in this paper. The primers names started with d were used for qRT-PCR. The other primers were used for vector construction and gene amplification, and the restriction sites indicated the restriction enzymes used in vector constructing.

mgGtsp was synthesized by Sangon Biotech (Shanghai) Co., Ltd. The restriction sites in TqGF, TqGR, GbF, GbR, EbF, and EbR were designed since the cohesive end would expose after annealing.

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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