**PtiCYP85A3, a BR C-6 Oxidase Gene, Plays a Critical Role in Brassinosteroid-Mediated Tension Wood Formation in Poplar**

Yanli Jin\(^1,2,3\), Chunyan Yu\(^1,3\), Chunmei Jiang\(^1,4\), Xiaotong Guo\(^1,3\), Bei Li\(^1,3\), Cuiting Wang\(^2\), Fanjing Kong\(^5\), Hongxia Zhang\(^1,2,3\)* and Haihai Wang\(^2\)*

\(^1\) College of Agriculture, Ludong University, Yantai, China, \(^2\) National Key Laboratory of Plant Molecular Genetics, Institute of Plant Physiology and Ecology, Shanghai Institutes for Biological Sciences, Chinese Academy of Sciences, Shanghai, China, \(^3\) The Key Laboratory of Molecular Module-Based Breeding of High Yield and Abiotic Resistant Plants in the Universities of Shandong, Institute for Advanced Study of Coastal Ecology, Ludong University, Yantai, China, \(^4\) Hubei Collaborative Innovation Center for Green Transformation of Bio-Resources, College of Life Sciences, Hubei University, Wuhan, China, \(^5\) Ministry of Natural Resources Key Laboratory of Saline Lake Resources and Environments, Institute of Mineral Resources, Chinese Academy of Geological Sciences, Beijing, China

In angiosperm trees, the gelatinous layer (G-layer) takes a great part of the fiber cell wall in the tension wood (TW). However, the mechanism underlying G-layer formation in poplar is largely unknown. In this work, we demonstrate that G-layer formation in poplar TW cells is regulated by brassinosteroid (BR) and its signaling. **PtiCYP85A3**, a key BR biosynthesis gene, was predominantly expressed in the xylem of TW, accompanied with a relatively higher castasterone (CS) accumulation, than in the xylem of opposite wood (OW). A wider expression zone of BZR1, a key transcriptional factor in BR singling pathway, was also observed in G-fiber cells on TW side than in wood fiber cells on the OW side, as indicated by immunohistochemistry assays. Transgenic poplar plants overexpressing **PtiCYP85A3** produced thicker G-layer with higher cellulose proportion, and accumulated more BZR1 protein in the xylem of TW than did the wild type (WT) plants. Expression of most TW-associated CesAs, which were induced by 2, 4-epibrassinolide, an active BR, and inhibited by brassinazole, a BR biosynthesis inhibitor, were also up-regulated in the xylem of TW in transgenic plants compared to that in WT plants. Further studies with dual-luciferase assays demonstrated that the promoters of **PtiCesAs** were activated by **PtiMYB128**, a TW specific transcription factor, which was then regulated by BZR1. All these results indicate that BR plays a crucial role in the G-layer formation of TW fiber cells by regulating the expression of **BZR1**, **PtiMYB128**, and **PtiCesAs** in poplar.

**Keywords:** brassinosteroids, G-layer, poplar, **PtiCYP85A3**, tension wood, xylem

**INTRODUCTION**

Tension wood (TW) is a kind of reaction wood formed in angiosperm plants as a response to gravity. When plants are subjected to environmental forces, such as landslide, typhoon, flood, and snow storm, TW is formed on the upper side of leaning stems to generate a strong tensile force which helps to pull the inclined stems back to the vertical position (Okuyama et al., 1994;
Yoshida et al., 1999, 2000; Yamamoto et al., 2002; Clair et al., 2006; Ruelle et al., 2006; Coutand et al., 2007; Clair et al., 2010. Tension wood is generally characterized by the presence of gelatinous xylem fiber cells with a thick inner gelatinous cell wall layer (G-layer), which contains a high proportion of cellulose, and a low proportion of lignin, various non-cellulosic polysaccharides and glycosylated proteins (Timell, 1969; Goswami et al., 2008; low proportion of lignin, various non-cellulosic polysaccharides (G-layer), which contains a high proportion of cellulose, and a wood is generally characterized by the presence of gelatinous Yoshida et al., 1999, 2000; Yamamoto et al., 2002; Clair et al., 2006; Clair et al., 2010). In Arabidopsis, CESA proteins are categorized into six classes: CESA1, CESA3, and CESA6 (including CESA2, CESA5, CESA6, and CESA9) responsible for primary cell wall cellulose biosynthesis, and CESA4, CESA7, and CESA8 responsible for secondary cell wall cellulose biosynthesis (Taylor et al., 2003; Desprez et al., 2007; Persson et al., 2007). In the poplar genome, a total number of 17 CESA genes have been indentified and the nomenclature for CesAs, PtiCesA1-A/B, PtiCesA3-A/B/C/D, PtiCesA4, PtiCesA6-A/B/C/D/E/F, PtiCesA7-A/B, and PtiCesA8-A/B, was updated based on the alignments of Populus CesA gene family with the Arabidopsis CesAs (Kumar et al., 2009). Consistent with the CesAs in Arabidopsis, three classes of CesAs, CESA4, CESA7a/b, and CESA8a/b, are involved in cellulose biosynthesis in secondary wall and G-layer formation.

Cellulose, composed of linear chains of β-1,4-linked Glc units, is synthesized by the cellulose synthase complexes (CSCs) consisting of cellulose synthases (CESAs) (Zhang et al., 2018). In Arabidopsis, CESA proteins are categorized into six classes: CESA1, CESA3, and CESA6 (including CESA2, CESA5, CESA6, and CESA9) responsible for primary cell wall cellulose biosynthesis, and CESA4, CESA7, and CESA8 responsible for secondary cell wall cellulose biosynthesis (Taylor et al., 2003; Desprez et al., 2007; Persson et al., 2007). In the poplar genome, a total number of 17 CESA genes have been indentified and the nomenclature for CesAs, PtiCesA1-A/B, PtiCesA3-A/B/C/D, PtiCesA4, PtiCesA6-A/B/C/D/E/F, PtiCesA7-A/B, and PtiCesA8-A/B, was updated based on the alignments of Populus CesA gene family with the Arabidopsis CesAs (Kumar et al., 2009). Consistent with the CesAs in Arabidopsis, three classes of CesAs, CESA4, CESA7a/b, and CESA8a/b, are involved in cellulose biosynthesis in secondary wall and G-layer formation.

Previously studies have shown that plant hormones such as auxin, ethylene, and gibberellin (GA) were involved in TW formation (Little and Pharis, 1995; Mellerowicz et al., 2001; Pilate et al., 2004; Kwon, 2008). In poplar, although the balance of endogenous auxin level was not significantly altered, the expression of several Aux/IAA genes changed during TW formation, indicating that auxin may not directly regulate the production of TW (Moyle et al., 2002; Hellgren et al., 2004). Ethylene has been confirmed as a key regulator in TW formation (Andersson-Gunnerärs et al., 2003; Love et al., 2009). A poplar ACC oxidase gene, which was induced by gravitational irradiation, displayed an asymmetric expression between TW and OW (Andersson-Gunnerärs et al., 2003). Upon treatment with 1-methyleclopprobe (1-MCP), an ethylene perception inhibitor, TW formation was inhibited (Love et al., 2009). On the other hand, application of exogenous ethylene or its precursor 1-aminoacyclopropa1-carboxylic acid (ACC) induced G-layer formation and altered cellulose microfibril angle in absence of gravitational stimulus in aspen (Felten et al., 2018). In addition, many genes related to cell expansion and cell wall modification for gelatinous layer induction were regulated by ethylene signaling (Felten et al., 2018). In a weeping type of Prunus spachiana, GA promoted the formation of TW on the upper side of branches, and prevented the bending of its branches (Nakamura et al., 1994; Baba et al., 1995; Yoshida et al., 1999). In several species of angiosperm trees, exogenous application of GA to their vertical stems also induced TW formation in the absence of gravitational stimulus (Funada et al., 2008). In transgenic poplar, GA mediated TW formation via the regulation of fasciclin-like arabinogalactan protein gene expression (Wang et al., 2017).

As one of the six classes of phytohormones, BRs also play a key role in plant growth, reproduction, and response to biotic and abiotic stresses (Kim et al., 2005). In Arabidopsis, BRs are synthesized by a list of enzymes step by step, including deetiolated2 (DET2) (Li et al., 1996), constitutive photomorphogenesis and dwarfism (CPD) (Szekeres et al., 1996), dwarf4 (DWF4) (Choe et al., 1998), BR-6-oxidase1 (BR6ox1) (Bishop et al., 1999), and rotundfolia3 (ROT3) (Kim et al., 2005). To date, more than 70 BRs have been identified from the entire plant kingdom (Saleh et al., 2006). Among them, castasterone (CS) and brassinolide (BL) were shown to be the most important BRs (Altmann, 1999). A cytochrome P450, AtCYP85A2, known as a BR C-6 oxidase, catalyzes the conversion of CS to BL. When the function of AtCYP85A2 was knocked out, the Arabidopsis cyp85a2-2 mutant produced dark green curled leaves with shortened petioles compared with those of the wild type plants. On the other hand, when AtCYP85A2 was overexpressed, both vegetative and reproductive growth was enhanced in transgenic Arabidopsis plants (Kim et al., 2005). The functions of BRs in other herbaceous and woody plants such as rice, tomato, grape, and pea were also examined in some details (Mori et al., 2002; Nomura et al., 2005; Symons et al., 2006; Jager et al., 2007). However, the biological functions of CYP85A2 in TW formation in woody plants are still not fully clarified.

Previously, we reported that overexpression of PtiCYP85A3, the homology of AtCYP85A2, promoted the endogenous CS content, and enhanced the growth and biomass production in transgenic poplar (Jin et al., 2017). In this work, we examined the possible function of BR in TW formation by manipulating the expression of PtiCYP85A3 in transgenic poplar plants. We found that, by activating BR signaling, PtiCYP85A3 positively regulated G-layer formation of TW fiber cells in poplar.

**MATERIALS AND METHODS**

**Plant Materials and Growth Conditions**

*Populus trichocarpa* genotype Nisqually-1, a commercial clone Shaxin yang (*P. davidiana Dode × P. bolleana Lauche*) and transgenic Shaxin yang overexpressing *PtiCYP85A3* (lines L3, L5, and L8) were used in this study (Jin et al., 2017). Generally, *in vitro*-grown plants were sub-cultured monthly by aseptically transferring shoot apices to fresh MS medium supplemented with 0.1 mg l⁻¹ of 1-naphthaleneacetic acid (Murashige and Skoog, 1962). Plantlets were then transferred into individual pots and grown in a greenhouse under a 12 h light/12 h dark photoperiod. The temperature was kept at 21–25°C in daytime and 15–18°C at night. All plants were well watered according to the evaporation demands during different growth stages, and fertilized biweekly with water-soluble fertilizers (Plant-Soul, China). Nisqually-1 was used for gene cloning only. Wild type and transgenic Shaxin yang were used for all the other experiments.

**TW Induction and Histochemical Staining**

Before TW induction, 2-month-old poplar plants grown in greenhouse were fixed to sticks to make sure all plants grow upright. To induce TW formation, poplar plants were tilted
to a 45° angle for 14 days. Over 10 plants from wild type and each transgenic line were used for each treatment, and three replicates were carried out for each treatment. In this way, TW and OW were produced on the upper and under sides of leaning stems, respectively. The middle part of stems under the bended point was used to isolate the xylem tissues (without any pith) of tension wood side (TW-X) and opposite wood side (OW-X). Phloem tissues of tension wood side (TW-P) and opposite wood side (OW-P) in the barks were also isolated.

Tension wood staining was performed as described previously (Wang et al., 2017). Briefly, stem segments were cut from the tilted plant, ~50 stem sections were fixed with 2% formaldehyde and subsequently passed over a gradient ethanol series. Then the sections were embedded in paraffin, and about 10 µm thick sections were cut out with a rotary microtome. After dewaxing, they were stained with 1% aqueous safranin-O (MP, United States) and subsequently with 1% aqueous astra-blue (Santa Cruz, United States) as described previously (Srebotnik and Messner, 1994). High power images were captured under bright field using an ECLIPSE 80i microscope (Nikon, Japan) and low power images were taken under the SMZ800 microscope (Nikon, Japan).

**BR Content Assays**

The xylem tissues of TW or OW (TW-X or OW-X), and the bark of TW or OW (TW-P or OW-P) isolated from WT plants were grounded into fine powder in liquid nitrogen for BR content determination as described previously (Jin et al., 2017). Three replicates were carried out with over 5 g dry material for each sample and each experiment. The variability was indicated with the standard deviation (SD).

**Quantitative Real-Time RT-PCR**

For the expression analysis of *PtiCYP85A3*, cellulose synthase genes (*CesA*), and MYB transcription factor genes (*MYBs*) during TW formation, total RNA was extracted from TW-X, TW-P, OW-X, and OW-P of WT plants using RNAiso Reagent (Takara, Japan). After treated with DNase I (Promega, United States), a total amount of 2 µg total RNA was subjected to reverse transcription using the HiScript® II Q RT SuperMix for qPCR (+gDNA wiper) (Vazyme, China). Quantitative real-time RT PCR (qRT-PCR) was performed using an AceQ qPCR SYBR Green Master Mix (Vazyme Biotech, China) and a CFX Connect Real-Time System (Bio-Rad, United States). Three independent replicates of measurements were performed for each sample.

For the expression analyses of *CesA* genes, TW-X of the middle parts of inclined stems from WT and transgenic plants were used for RNA extraction. The relative expression of each target gene was normalized using the housekeeping gene *Pd β-Tubulin*. Gene specific primers used in this study were designed according to the sequences of *P. trichocarpa* and listed in Supplementary Table S1. For genes cloned from *P. trichocarpa*, prefix *Pti* was used in gene names. For gene investigation in *P. davidiana x bolleana*, prefix *Pd* × *I* was used in gene names.

**Exogenous EBL and Brassinazole Treatments**

To analyze the responses of *CesAs* and *MYB128* to BR, stem segments of the 3th to 4th internodes of 2-month-old WT (Shangxin yang) plants were cut into 1 mm slices, and at least 60 stem sections from 20 stem segments were soaked in 100 nM EBL for each treatment (10, 30, 60, or 220 min). Then, the stem sections were used for RNA extraction and qRT-PCR analyses.

For analyses of the effects of brassinazole (Brz, one of BR biosynthesis inhibitors) on the expressions of *CesAs* and *MYB128*, wild type Shangxin yang plants were tilted to 45° for 10 days and painted with lanolin containing 5 mM brassinazole (Sigma, United States) once every 2 days. Then, TW-X of the middle part of inclined stems of 20 plants was used for RNA extraction and further qRT-PCR analyses.

**Cellulose Quantification**

TW-X separated from the middle parts of 20 inclined stems of WT and each transgenic lines (L3, L5, and L8) were cut into small pieces, ground into fine powder in liquid nitrogen. Then, cell wall material (CWM) was isolated by sequentially washing the samples with 70% (v/v) ethanol, chloroform:methanol (1:1), and acetone as described previously (Foster et al., 2010). Starch was removed from the pellet by incubating the sample in 1 ml of 0.1 M sodium acetate buffer (pH 5.0) with amylase (50 µg ml⁻¹, Sigma, United States) and pullulanase (Sigma, United States) at 37°C for 12 h. After washed three times with water, the resultant CWM was suspended with acetone and dried at 35°C for 12 h. Cell wall material was used to determine the contents of cellulose.

To determine the cellulose content, CWM was incubated in 1 ml of Updegraff reagent for 30 min at 100°C, and then washed three times with 1 ml of acetone. The pellet (crystalline cellulose) was completely hydrolyzed into glucose in 175 µl of 72% sulfuric acid at room temperature for 45 min. After the addition of 825 µl water, 10 µl of each sample of the supernatant and 90 µl of water was pipetted into separate cells of 96-well polystyrene microtiter plates before the addition of 200 µl of freshly prepared Anthrone reagent. The plate was heated for 30 min at 80°C, and the absorption at 625 nm was measured after being cooled to room temperature (Foster et al., 2010). Three independent replicates of measurements were performed for each sample.

**Antibody Preparation**

For Western blot analyses, OsBZR1 polyclonal antibody was purchased from Beijing Genomics Institute (BGI)¹. Arabidopsis *CesA7* (*At5g17420*) and *CesA8* (*At4g18780*) polyclonal antibody were bought from Agrisera². Plant β-actin antibody and secondary antibodies were purchased from ABclonal³.

**Western Blotting Analyses**

For Western blot analyses, total proteins were extracted from the TW-X tissues of inclined WT and transgenic plants

¹http://www.genomics.org.cn
²http://www.agrisera.com
³http://www.immunogen.com.cn
using RIPA buffer consisting of 1 mM PMSF, and separated on 10% SDS-PAGE gel. After electrotransfering of the proteins onto polyvinylidene difluoride membranes, the membranes were blocked with TBST buffer (10 mM Tris/HCl, pH 7.5, 0.1% NaCl, 0.05% Tween 20) supplemented with 5% non-fat dried milk for 1 h. The membranes were incubated with primary antibody (diluted at 1:1000) overnight at 4°C. Afterward, the membranes were rinsed three times with TBST buffer and incubated with the secondary antibodies (peroxidase-labeled anti-rabbit antibodies) at a dilution of 1:5000 for 1 h. Finally, the samples were rinsed three times with PBS buffer for 30 min each, the membranes were incubated in LumiGLO for chemiluminescence detection (KPL, United States) and then imaged with a Tanon 5500 electrophoresis system (Tanon, China).

**Immunolocalization of BZR1**

Immunohistochemical analyses of BZR1 in TW of WT plants were performed as described previously (Wang et al., 2015). Stem segments of inclined WT plants were fixed overnight in 0.1 M PBS (pH 7.5) containing 4% paraformaldehyde, and embedded in paraffin. The slides were spread with polylysine before the sections were fixed. After deparaffinization and dehydration, the sections were washed twice with PBS buffer. The samples were blocked with 5% bovine serum albumin (BSA) in PBS for 1 h at room temperature. Subsequently, they were incubated with anti-OsBZR1 antibodies (diluted at 1:30 with 0.1 M PBS containing 0.1% BSA) at room temperature for 1 h. In the negative control, BZR1 antibody was omitted. After rinsed three times in PBS, the samples were incubated with FITC secondary antibody (diluted at 1:50 in the same buffer) at room temperature for 1 h. Finally, the samples were rinsed with PBS buffer for three times and mounted with a cover glass for photographing. Images were captured under a confocal microscope Zeiss LSM 510 (META, Germany).

**Transient Transcription Dual-Luciferase Assays**

For dual-luciferase assays, the LUC reporter constructs were generated by cloning the promoters of PtiCesA4, PtiCesA7-A, and PtiMYB128 into the pGreenII0800-LUC, respectively (Hellens et al., 2005). The CaMV 35S promoter-driven transcriptional factor effector constructs were generated by inserting PtiMYB128 or AtBZR1 into the pGreenII62-SK, respectively (Hellens et al., 2005). To detect the induction of PtiMYB128 to the promoters of PtiCesAs, the effector of PtiMYB128 and each LUC reporter construct of PtiCesAs were respectively co-expressed in poplar leaf protoplasts. To test the transcription activity of AtBZR1 to the PtiMYB128 promoter, the effector of AtBZR1 was co-transfected with PtiMYB128 LUC reporter construct in poplar leaf protoplasts. Poplar leaf protoplast extraction and transformation were performed as described previously (Wang et al., 2013). After 16 h, transfected cells were collected and homogenized in 300 µl of passive lysis buffer. The crude extract (20 µl) was mixed with 40 µl of luciferase assay buffer and firefly luciferase (LUC) activity was measured using GLOMAX 20/20 luminometer (Promega, Wisconsin, United States). Stop and Glow Buffer (40 µl) was then added to the solution and renilla luciferase (REN) activity was measured. LUC/REN ratio was used to represent the relative activity of the transcriptional factors on the driving promoters. Three replicates were carried out for each assay, and the variability was indicated with the standard deviation (SD).

**Statistics**

All data were obtained from at least three independent experiments with three biological replicates each. For statistical analyses, Student’s t-test was used to generate every P-value. All the tests were two-tailed.

**RESULTS**

**Castasterone Is Accumulated in the Xylem of TW**

Since TW can be induced by mechanical stress, as done by gravitational stimulation (Jourez et al., 2001; Jourez and Avella-Shaw, 2003; Déjardin et al., 2010), we inclined wild type poplar plants to a 45 degree when grown in greenhouse to induce TW formation (Figure 1A). After 2 weeks, TW was obviously produced on the upper side of leaning stems, and was stained into blue color with safranin/astra blue double staining (Figure 1B). To explore the possible function of BR in TW formation, we first examined the expression of genes associated with BR biosynthesis and metabolism. Poplar CPD, DWF4, CYP85A1, CYP85A3, and BAS1, the homologs of Arabidopsis CPD, DWF4, CYP85A2, and BAS1, were selected. Although no significant increase was observed in the expression of Pd × ICPD, increased expression of Pd × IDWF4 in TW, including both xylem and phloem (TW-X and TW-P), was detected, as compared with that in OW (Figures 1C,D). Both Pd × CYP85A1 and Pd × ICYP85A3 were predominately expressed in the xylem of TW (Figures 1E,F), whereas Pd × lBAS1, the homolog of Arabidopsis BAS1, encoding a cytochrome P450 monooxygenase (CYP734A1, formerly CYP72B1) which inactivates active brassinosteroids (BRs) such as brassinolide (BL) and castasterone (CS), was highly expressed in the xylem of OW (Figure 1G; Turk et al., 2005). The up-regulated expression of BR synthesis genes and the down-regulated expression of BR metabolic genes in TW may have promoted BR accumulation in the xylem of TW. Therefore, we examined the content of BRs in the xylem and phloem of both TW and OW. We found that although no significant difference was detected in the content of BL, asymmetric distribution of CS was observed in the tissues of TW and OW. A higher CS content was detected in the phloem of OW and the xylem of TW (Figures 1H,I), suggesting that CS may be the major BR involved in TW formation.

**BR Signaling Is Enhanced in the Xylem of TW**

BZR1/BES1 protein is the key positive transcription factor in the signaling pathway of BRs. The increased CS content was detected in the xylem of TW (Figure 1H).
in the xylem of TW could be connected with an enhanced BR signaling. To confirm this hypothesis, we investigated the distribution of BZR1 protein during TW formation in WT plants by fluorescence immunohistochemistry test using an OsBZR1 antibody. A very strong BZR1/BES1 protein signal was detected in the G-fiber cells on the TW side (a wide band), but only a weak BZR1/BES1 protein signal was detected in the wood fiber cell on the opposite side (a narrow band) (Figures 2A–C). No fluorescence signal was detected in the control section (Figures 2D–F). These results indicate that BR signaling is activated during TW formation.
**PtiCYP85A3 Promotes G-Layer Formation in the Xylem Fiber Cells of TW**

Previously, we reported that overexpression of *PtiCYP85A3* in poplar promoted the accumulation of CS, and significantly improved the growth and biomass of transgenic plants (Jin et al., 2017). To understand the possible effects of BRs on TW formation, we compared the fiber cells in the TW of wild type and transgenic plants by histochemical staining. Although the stem diameter of transgenic plants was greater than that of WT plants, the ratio of TW to OW in the xylem area was not significantly changed in transgenic plants (Supplementary Figures S1A,B). We further examined fiber cells of TW, and found that although the cell number per area was not significantly changed, the G-layer in the xylem fiber cell walls of transgenic plants was obviously thicker than that of WT (Figures 3A–J).

As the most typical feature of TW, G-layer is in rich of cellulose (Scurfield and Wardrop, 1963; Norberg and Meier, 1966; Cronshaw and Morey, 1968). Therefore, we determined the content of cellulose in the TW xylem of both WT and transgenic plants. Consistent with the thickened G-layer, the content of cellulose in the TW xylem of transgenic plants was significantly higher than that in the wild type (Figure 3K). We also confirmed the overexpression of *PtiCYP85A3* and accumulation of BZR1 protein in the TW xylem of transgenic plants. Again, consistent with the overexpression of *PtiCYP85A3*, BZR1 protein was more abundantly accumulated in the TW xylem of all transgenic plants than in that of WT plants (Figures 3L,M).

**TW Associated *PtiCesAs* Are Regulated by BR**

In the genome of *Populus trichocarpa*, a total number of 17 cellulose biosynthesis genes were identified (Kumar et al., 2009). We examined the expression patterns of these *CesAs* during TW formation in Shanxin yang, and found that seven of them, *Pd × lCesA3-A*, *Pd × lCesA3-B*, *Pd × lCesA4*, *Pd × lCesA7-A*, *Pd × lCesA7-B*, *Pd × lCesA8-A*, and *Pd × lCesA8-B*, were highly expressed in the TW xylem tissues (Supplementary Figure S2). These results imply that these *CesA* genes may play an important role in the cellulose biosynthesis during G-layer formation. To confirm this hypothesis, the stems of WT plants were treated with 100 nM 2, 4-epibrassinolide (EBL) for different time periods, and the expression levels of these *CesA* genes were examined. As expected, expression of all these *CesA* genes were up-regulated by EBL (Figure 4A). We further tested the effect of brassinazole (Brz), a BR biosynthesis inhibitor, on the expression of these TW-induced *CesAs*. WT plants grown in...
greenhouse were inclined at 45° for 10 days and painted with 5 mM Brz on the upper side of stems. Then, the expression levels of these CesAs in TW xylem tissues were analyzed by qRT-PCR. Again, as expected, the transcription of these CesA genes were all down-regulated at different levels (Figure 4B). These results indicate that expressions of these TW associated CesA genes are regulated by BRs.

Expression of TW Associated CesAs Are Up-Regulated in TW Xylem of PtiCYP85A3 Transgenic Poplar

Based on the observations that TW associated CesAs were regulated by BRs, we speculated that the thickened G-layer in the TW fiber cell walls of transgenic plants maybe be due to the up-regulated expression of CesA genes. Therefore, we analyzed the expression levels of these CesAs in the TW xylem tissues of transgenic poplar plants overexpressing PtiCYP85A3. Indeed, all the tested CesAs were up-regulated in TW xylem tissues of transgenic plants (Figure 5A). We further checked the protein levels of CesA proteins recognized by AtCesA7 and AtCesA8 antibodies by Western blotting. Consistently, the levels of proteins recognized by both antibodies were significantly higher in the TW xylem of transgenic plants compared with that of WT plants (Figure 5B). These results imply that BRs could promote G-layer formation by regulating the expression of TW-associated CesAs.

MYB128 Is Highly Expressed During TW Formation and Regulated by BR Signaling

Previous studies have shown that poplar CesAs, including some TW-associated CesAs, participating in xylem development were regulated by a list of MYB transcription factors (Ohtani et al., 2011; Zhong et al., 2011; Lin et al., 2013; Ye and Zhong, 2015). We examined the expression patterns of xylem-related MYBs during TW formation in Shanxin yang plants and found that
FIGURE 4 | qRT-PCR analyses of cellulose biosynthesis genes. Stems of wild type Shangxin yang plants were treated with 100 nM 2, 4-epibrassinolide (EBL) for 0, 30, 60, 220 min, or with 5 mM brassinazole (Brz) for 10 days, and the relative expressions of $Pd\times CesA$s were determined. (A) Expressions of $Pd\times CesA$s in the stems treated with EBL. (B) Expressions of $Pd\times CesA$s in the tension wood xylem tissues of the stems treated with Brz. The relative expression of $Pd\times CesA$s was normalized using the housekeeping gene $Pd\times EF1\beta$. Gene expression value in the control was set to 1. Data were means ± SD ($n = 6$) of three independent biological replicates with three technical replicates each. Gene Potri ID numbers were provided in Supplementary Table S1.

except $Pd \times IMYB121$, all xylem-related $Pd \times IMYB$s were highly expressed in the xylem tissues of TW or OW side. However, the expression levels of $Pd \times IMYB10$ and $Pd \times IMYB128$ were significantly higher in the TW xylem than in the OW xylem (Supplementary Figure S3). Further treatments with EBL or Brz demonstrated that both $Pd \times IMYB10$ and $Pd \times IMYB128$ were repressed by Brz, but only $Pd \times IMYB128$ was significantly induced by EBL (Figures 6A,B and Supplementary Figure S4). Therefore, TW-associated $Pd \times IMYB128$ could be involved in BR-mediated G-layer formation in poplar.

We further investigated whether $Pd \times IMYB128$ was regulated by BR or BR signaling by transient transcription dual-luciferase assays. We found that AtBZR1, the key transcription factor in BR signaling, significantly activated the promoter of $PtiMYB128$ (Figure 6C), suggesting that $Pd \times IMYB128$ could be regulated by BZR1 protein.

PtiCesAs Are Regulated by PtiMYB128

Since the expression of $Pd \times IMYB128$ was responsive to EBL and Brz treatments, we further performed dual-luciferase assays to see if it can activate the promoters of TW-associated PtiCesAs. As we have expected, PtiMYB128 successfully activated the promoters of PtiCesA4 and PtiCesA7-A, showing a higher LUC/REN ratio than the control (Figure 6D). Therefore, cellulose synthesis facilitated by CesAs in G-layer formation is regulated by PtMYB128.

DISCUSSION

As a class of plant specific polyhydroxylated steroid hormones, BRs have important functions in the growth and development, as well as in response to biotic and abiotic stresses of plants. In zinnias (Zinnia elegans), low concentration of BL promoted the differentiation of mesophyll cells into tracheary elements (Iwasaki and Shibaoka, 1991). In Cress plants (Lepidium sativum), Brz treatment inhibited the growth of secondary xylem (Nagata et al., 2001). In Arabidopsis, BR-deficient mutants displayed dwarfed phenotypes and abnormal development of vascular tissues, with excessively proliferated phloem cells and dramatically reduced xylem cells (Caño-Delgado et al., 2004, 2010; Xie et al., 2011; Hossain et al., 2012). However, the roles of BRs in TW formation are not well illustrated.

To date, among the more than 70 BRs identified in higher plants (Li et al., 1996), only BL and its immediate precursor castasterone (CS) have detectable biological activity (Yokota et al., 1982). The P450 protein CYP85A1 acts as a C-6 oxidase, which catalyzes the multiple C-6 oxidation reactions including 6-deoxo CS to CS (Bishop et al., 1999;
Shimada et al., 2001). CYP85A1 and CYP85A3 in *Solanum lycopersicum*, AtCYP85A1 and AtCYP85A2 in *Arabidopsis thaliana*, CYP85A1 and CYP85A6 in *Pisum sativum*, BR6ox1 in *Vitis vinifera*, DWARF in *Oryza sativa* and *Hordeum vulgare*, and BRD1 (BRASSINOSTEROID DEFICIENT DWARF 1) in *Zea mays* and *Brachypodium distachyon*, were all involved in the multiple C-6 oxidation reactions, but only AtCYP85A2 in *Arabidopsis* and CYP85A3 in tomato could convert CS to BL (Bishop et al., 1996, 1999; Schultz et al., 2001; Shimada et al., 2001; Hong et al., 2002; Mori et al., 2002; Kim et al., 2005; Nomura et al., 2005; Symons et al., 2006; Jager et al., 2007; Gruszka et al., 2011; Makarevitch et al., 2012; Xü et al., 2015). In poplar, three BR-C6-oxidase encoding genes, named as *PtiCYP85A1*, *PtiCYP85A3*, and *PtiCYP85A4*, have been identified (Kim et al., 2008). *PtiCYP85A1* and *PtiCYP85A3* shared as high as 91.9 and 96.98% homology in DNA and amino acid sequence, respectively. The homologies of *PtiCYP85A4* and *PtiCYP85A3* were lower, 89.48% (DNA sequence) and 67.38% (protein sequence), respectively (Jin et al., 2017).

In order to understand the possible function of *PtiCYP85A3* in TW formation in poplar, we first examined the expression of genes involved in BR biosynthesis and metabolism, and the contents of CS and BL during TW formation in Shanxi yang (Figures 1A–H). *Pd × ICYP85A1* and *Pd × ICYP85A3* showed the same expression pattern, which was different from the expression pattern of *Pd × ICYP85A4* (Figures 1E,F). Therefore, *PtiCYP85A1* and *PtiCYP85A3* may have the same functions. The predominate expression of *Pd × ICYP85A3*
in the TW-X tissue was accompanied by an increased CS content. Consistent with these observations, BZR1 protein was also abundantly transcribed in the xylem fiber cells undergoing G-layer development (Figures 2A, C). All these results imply that PtiCYP85A3 is involved in TW formation, possibly by affecting the production and distribution of BR in poplar. This is further confirmed with transgenic poplar plants overexpressing *PtiCYP85A3*, which produced thicker G-layer with increased cellulose and BZR1 protein contents in the xylem fiber cell walls than did the WT plants (Figures 3A–M).

In Arabidopsis, BRs regulate cellulose biosynthesis by controlling the expression of CESA genes (Xie et al., 2011). Since the major part of TW fiber cell walls were composed of cellulose, we investigated the expressions of all the 17 cellulose biosynthesis genes identified in the poplar genome and found that seven of them were highly transcribed in the TW xylem (Supplementary Figure S2). Further studies indicated that their expressions were up-regulated by 2, 4-epibrassinolide (EBL) and down-regulated by brassinazole (Brz) (Figures 4A,B). Similar results were also observed in transgenic poplar plants overexpressing *PtiCYP85A3* (Figures 5A, B). These results demonstrate that BRs could improve G-layer formation by regulating CesA gene expression.

Previously, it was reported that *AtMYB103* was involved in secondary cell wall synthesis, and was predominantly expressed in TW than in opposite wood (Zhong et al., 2008; Chen et al., 2015). We found that expression of *Pd × IMYB128*, one of the homologs of *AtMYB103* in poplar, was up-regulated in the TW fiber cells and induced by EBL (Supplementary Figure S3 and Figure 6A). Based on the observations that *Pd × IMYB128* was induced by BR treatment and suppressed by BR biosynthesis inhibitor Brz, we deduced that it may be involved in BR signaling (Figures 6A, B). The poplar genome contains seven homologs of *BZR1/BES1*, a key positive transcriptional factor in Arabidopsis (Supplementary Figure S5). However, their possible functions are still unknown. Therefore, we performed dual-luciferase assays with the Arabidopsis BZR1, and found that BZR1 activated the promoter of *PtiMYB128*, which further activated the promoters of *PtrCesA4* and *PtrCesA7-A* (Figures 6C, D). This is consistent with previous report that MYB128 could regulate the promoter activity of promoters of *PtrCesA4*, *PtrCesA8*, and *PtrCesA17*, the orthologs of *PtiCesA4*, *PtiCesA8-A*, and *PtiCesA7-B*, respectively (Zhong et al., 2011).

The released strain of growth stress was tightly correlated with the production of reaction wood (Yamamoto et al., 1991; Sugiyama et al., 1993; Okuyama et al., 1994). In TW, the longitudinally contractive released strain increased with cellulose content and crystallinity. It is negatively correlated with the Klassen lignin content and microfibrillar angle (Sugiyama et al., 1993; Okuyama et al., 1994). The greater the contractive released strain was, the greater the tensile growth stress became, leading to the generation of thicker G-layers (Yoshida et al., 2002). Auxin, ethylene and gibberellin have been identified as important factors that regulate the formation of TW (Cronshaw and Morey, 1965; Morey and Cronshaw, 1968; Nakamura et al., 1994; Yoshida et al., 1999; Andersson-Gunnerås et al., 2003; Du and Yamamoto, 2003, 2007; Du et al., 2004; Clair et al., 2006; Funada et al., 2008; Love et al., 2009; Wang et al., 2017; Felten et al., 2018). Interestingly, most of the BR-related genes, except for the LRR receptor kinase precursors, were down-regulated in TW as compared to opposite wood after 6 h of mechanical bending treatment (Jin et al., 2011). Since the size of fiber cells in TW was much smaller than that in opposite wood (Jin and Kwon, 2009), the down-regulation of BR-related genes may be correlated with the reduced size of fiber cells in TW, thereby implying the possible involvement of BR in TW development (Jin et al., 2011). However, the molecular mechanism how BRs regulate TW formation still lacks accuracy. Our study demonstrated, for the first time, that BRs increased the thickness of G-layer by regulating the expression of *PtiMYB128*, one of the homologous genes of *AtMYB103*, to activate the expression of cellulose synthase genes, and consequently affected the formation of TW in poplar. A model was proposed to
illustrate the biological roles of BRs in TW formation in *Populus* (Supplementary Figure S6). Under the action of external force or gravity, some secondary signal activated the biosynthetic and signal transduction pathways of BRs, thus more activated BZR1/BES1 moved into the nucleus to promote the transcription of *PtiMYB128*, which up-regulated the expressions of TW-associated cellulose synthesis genes, and as a result, facilitated the thickness of G-layers during TW formation. Taken together, our findings provide a new insight into the biological functions of *PtiCYP85A3* in the TW formation in poplar.

**CONCLUSION**

In this study, we demonstrated that *PtiCYP85A3* plays a pivotal role in TW formation in poplar. Overexpression of *PtiCYP85A3* promoted G-layer formation, accompanied with augmented BZR1 accumulation and up-regulated CesA gene expression, in the xylem of the TW in transgenic plants. In addition, CesA gene promoters were promoted by *PtiMYB128*, which was activated by BZR1.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/Supplementary Material.

**AUTHOR CONTRIBUTIONS**

YJ, CY, CJ, XG, BL, CW, and FK performed the experiments and analyzed the data. YJ, CY, CJ, XG, BL, CW, and FK performed the experiments and analyzed the data. YJ, CY, HZ, and HW conceived the study. YJ, CY, CJ, XG, BL, CW, and FK performed the experiments and analyzed the data. YJ, CY, HZ, and HW wrote the manuscript. All authors read and agreed at the last version of 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.2020.00468/full#supplementary-material
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Conflict of Interest: 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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