PeCHYR1, a ubiquitin E3 ligase from *Populus euphratica*, enhances drought tolerance via ABA-induced stomatal closure by ROS production in *Populus*

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**Summary**

Drought, a primary abiotic stress, seriously affects plant growth and productivity. Stomata play a vital role in regulating gas exchange and drought adaptation. However, limited knowledge exists of the molecular mechanisms underlying stomatal movement in trees. Here, PeCHYR1, a ubiquitin E3 ligase, was isolated from *Populus euphratica*, a model of stress adaptation in forest trees. PeCHYR1 was preferentially expressed in young leaves and was significantly induced by ABA (abscisic acid) and dehydration treatments. To study the potential biological functions of PeCHYR1, transgenic poplar 84K (*Populus alba* × *Populus gandulosa*) plants overexpressing PeCHYR1 were generated. *PeCHYR1* overexpression significantly enhanced *H₂O₂* production and reduced stomatal aperture. Transgenic lines exhibited increased sensitivity to exogenous ABA and greater drought tolerance than that of WT (wild-type) controls. Moreover, up-regulation of *PeCHYR1* promoted stomatal closure and decreased transpiration, resulting in strongly elevated WUE (water use efficiency). When exposed to drought stress, transgenic poplar maintained higher photosynthetic activity and biomass accumulation. Taken together, these results suggest that *PeCHYR1* plays a crucial role in enhancing drought tolerance via ABA-induced stomatal closure caused by hydrogen peroxide (*H₂O₂*) production in transgenic poplar plants.

**Introduction**

Drought stress is a common abiotic stress, affecting plant growth, seed production, gas exchange, water relations and cellular homeostasis in plants (Shinozaki and Yamaguchi-Shinozaki, 2007; Zhu, 2016). To adapt to hostile environments, plants have developed intrinsic mechanisms to mitigate drought stress, such as closing stomata, reducing transpiration, generating abscisic acid (ABA) and accumulating hydrogen peroxide (*H₂O₂*) (Dietz et al., 2016; Zhu, 2016). ABA-induced stomatal closure is a defensive action that is usually accompanied by the production of *H₂O₂* (Li et al., 2017; Zhang et al., 2001). NADPH oxidase (RbohF) can generate *H₂O₂* in guard cells, which crucially modulates ABA-induced stomatal closure and plant drought tolerance (Gudesblat et al., 2007; Kolla et al., 2007; Ma et al., 2016). Previous studies have indicated that AtrRbohF can be phosphorylated by OST1 (an ABA-induced SnRK2.6 protein kinase) (Joshi-Saha et al., 2011; Sirichandra et al., 2009; Wege et al., 2014). *H₂O₂* is a reactive oxygen species (ROS) that acts as a major signalling molecule in the ABA response pathway in guard cells (Bright et al., 2006; Li et al., 2017). High concentrations of ROS lead to cell injury or even hypersensitive cell death, whereas low concentrations of ROS function as developmental signals, controlling all aspects of plant biology (Ahmad et al., 2008; Dietz et al., 2016; Huang et al., 2016; Karuppanpanandian et al., 2011; Pitzschke et al., 2006). OsASR5 and DCA1, which enhance drought tolerance in rice, control stomatal closure by adjusting the concentration of *H₂O₂* (Cui et al., 2015; Li et al., 2017). GHR1 modulates stomatal movement associated with *H₂O₂* signalling in Arabidopsis (Hua et al., 2012).

In stomata, ABA-mediated ROS generation elevates cytosolic calcium ion levels and gives rise to stomatal closure (McAinsh et al., 1996; Pei et al., 2000; Xing et al., 2008). Recently, it was reported that ROS can induce stomatal closure through an ABA-independent pathway in rice (*Oryza sativa*). The SNAC1-targeted gene OsSRO1c positively controls *H₂O₂*-induced stomatal closure by regulating hydrogen peroxide level in rice, and the *E3* ligase OsHTAS can promote heat tolerance by modulating *H₂O₂*-induced stomatal closure in rice (Liu et al., 2016; You et al., 2013). However, 35S:OsASR5 plants were more sensitive to exogenous ABA treatment, activating the production of *H₂O₂* (Li et al., 2017).

RING (really interesting new gene) finger proteins consist mainly of conserved ‘CxHY’ domains (McDowell, 2007). These conserved ‘CxHY’ motifs are part of the CHY zinc-finger domain, which plays a role in physical interaction and ubiquitination (Lee and Kim, 2011; Woff et al., 2004; Yu et al., 2016), contains 12 histidines and cysteines, and binds with three zinc ions, composing a unique zinc-finger motif (Cayrol et al., 2007; Ding et al., 2015; Esposito et al., 2006). Proteins such as ubiquitin (ubi) *E3* ligases, a core component of ubiquitination pathway containing a CHY zinc-finger domain, possess a universal function including ubiquitylation of substrate proteins (Lim et al., 2013a; Ning et al., 2016; Stone et al., 2005). The ubiquitination process plays a crucial role in plant development and response to environmental stresses (Lim et al., 2013a; Stone et al., 2005). RING ubi *E3*
ligases play decisive roles in almost all plant growth processes, including photomorphogenesis, flower development, phytohormone signalling and regulation of senescence (Bae et al., 2011; Ding et al., 2015; Lim et al., 2013b, 2017). RING E3 ligases have been found in diverse plants, such as hot pepper (Capsicum annuum), maize (Zea mays), Arabidopsis and rice, positively or negatively regulating abiotic stress (Lyzena and Stone, 2012; Park et al., 2015; Zhao et al., 2014). Recently, a ubiquitin E3 ligase, Arabidopsis CHYR1, was found to positively facilitate ABA and drought-mediated stomatal closure, ROS production and plant drought tolerance via SnRK2.6-mediated phosphorylation (Ding et al., 2015). However, the expression of a homolog of CHYR1, rice OsRZFP24, enhances stomatal opening, leaf cooling and ABA insensitivity (Hsu et al., 2014).

Although the roles of CHYR1 family proteins have been widely reported in herbaceous plants, their possible functions in ligneous plants, especially in stomatal movement, remain unknown. Several mechanisms of plant response to drought exist, including avoidance, escape and tolerance strategies. Tolerance mechanisms attempt to maintain plant functions at the same level as in unstressed conditions, while avoidance mechanisms are used to adjust the balance between water loss and water uptake (Moradi, 2016). Populus, a pioneer genus in forest ecosystems, possesses great economic and ecological value (Sterky et al., 2004). Most poplar plants are fast-growing and have poor drought tolerance (Larchevque et al., 2011; Tuskan et al., 2006; Windt et al., 2006). However, one exception, P. euphratica, is the only large tree species that forms forests in desert areas, playing a very important role in the maintenance of the local ecological balance. Due to its strong adaptability to extreme temperature, drought and salinity, P. euphratica has served as a model woody plant to explore the mechanisms of stress resistance (Chen et al., 2013; Li et al., 2011; Ma et al., 2013; Tang et al., 2013; Yan et al., 2012). From an analysis of a transcriptome from P. euphratica under drought stress, we selected PeCHYR1, which is up-regulated in drought-stressed P. euphratica leaves, and performed further functional analysis of this gene in poplar (Tang et al., 2013).

In the current study, PeCHYR1 was cloned from P. euphratica and transferred into poplar B4K (P. alba × P. glandulosa) (Feng et al., 2015; Ke et al., 2016). Next, we verified the function of PeCHYR1 by means of molecular biology and plant physiological indexes. We found that PeCHYR1 was a functional homolog of CHYR1. Overexpression of PeCHYR1 enhanced drought tolerance by promoting H$_2$O$_2$-mediated stomatal closure in poplar.

**Results**

**Isolation and sequence analysis of PeCHYR1**

To elucidate the potential function of PeCHYR1 in the abiotic stress response of ligneous plants, we took advantage of P. euphratica, which naturally grows in deserts, to clone PeCHYR1. The coding sequence length of PeCHYR1 was 876 bp, encoding 291 amino acids. Homology analyses of the amino acid sequence showed 52% identity to the amino acid sequence of PeCHYR1, P. alba, P. glandulosa and A. thaliana. The evolutionary tree of related genes was divided into four parts (I, II, III and IV). The nucleic acid sequence of PeCHYR1 is mostly identical to that of PtrCHYR1 (Potri.009G005700), sharing 96.7% sequence identity (Figure 1a). A multiple sequence alignment revealed that the amino acid sequences of PeCHYR1, P. alba, P. glandulosa and A. thaliana (AT5G22920) contained the same conserved domains (Figure 1b). Furthermore, the PeCHYR1 promoter contains CAAT and TATA motifs that are involved in drought stress response (Figure S1).

**PeCHYR1 involvement in ABA and water stress response**

To study the potential biological functions of PeCHYR1, we applied quantitative real-time polymerase chain reaction (RT-qPCR) analysis to determine the relative transcript abundance in various tissues of P. euphratica. We observed that PeCHYR1 was mainly expressed in the leaf rather than the other tissues (Figure 2a). P. euphratica plants were treated with exogenous ABA and drought stress to investigate the expression pattern of PeCHYR1. The results showed that the transcript abundance of PeCHYR1 is transiently increased in the leaf under water stress, reaching a maximum of 3.5 times higher than control at 3 h, dropping to approximately 3 times at 6 and 9 h, and decreasing to 1.2 times at 12 h (Figure 2b). The abundance of PeCHYR1 was gradually augmented by ABA treatment, significantly increasing at 0.5 h, maintaining a stable level until 6 h and reaching a maximum of 9 times higher than control at 12 h (Figure 2c).

Drought stress-induced ABA generation leads to the accumulation of ROS. Young leaves of 1.5-month-old poplar were treated with 100 μM ABA for 0, 1, 2 and 3 h, and then, the leaves were immersed in diaminobenzidine (DAB) overnight. We observed that the red-brown colour of the leaves darkened gradually as the time of ABA treatment progressed (Figure 2d).

**Subcellular localization of PeCHYR1**

Previous studies indicated that Arabidopsis CHYR1, a ubiquitin E3 ligase, was generally localized in the nucleus, cytoplasm and endoplasmic reticulum (ER) (Ding et al., 2015). Thus, P. euphratica PeCHYR1, a functional homolog of CHYR1, might localize to the same cellular structures. To determine the subcellular localization of PeCHYR1, a 35S:PeCHYR1-GFP (green fluorescent protein) fusion protein, together with a 35S: HDEL-RFP (red fluorescent protein) fusion protein, was transiently transfected into tobacco leaves and Arabidopsis leaf protoplasts. Colocalization of PeCHYR1-GFP and HDEL-RFP was distinctly detected in the ER (Figure 3a,b). Simultaneously, colocalization with 4', 6-diamidino-2-phenylindole (DAPI) dye proved that the 35S: PeCHYR1-GFP fusion protein was localized to the nucleus (Figure 3a,b). Overall, the results indicated that PeCHYR1 was localized to the nucleus and ER.

To study the potential biological functions of PeCHYR1, transgenic poplar B4K (P. alba × P. glandulosa) plants overexpressing PeCHYR1 were generated. Every transgenic line was verified by PCR, RT-qPCR and histochemical staining with β-glucuronidase (GUS) (Figure S2). In the literature, these three methods are usually used to verify transgenic plants (Jin et al., 2017).

**PeCHYR1 promotes ABA-induced stomatal closure via ROS production**

Abscisic acid and ROS can induce stomatal closure (Bright et al., 2006), so we considered whether PeCHYR1 could regulate ABA-induced ROS accumulation. We treated 35S:PeCHYR1 plants with ABA, and WT plants with 100 μM ABA for 0, 1, 2 and 3 h, and then performed DAB staining on their leaves. When the leaves of the different lines were treated with ABA, we found that the brown colour of 35S: PeCHYR1 leaves was deeper than that of WT leaves (Figure 4a), even after multiple hours of treatment. These results indicated that 35S: PeCHYR1 plants had higher ROS levels than WT plants in the instantaneous ABA treatment.
To confirm whether PeCHYR1 participated in ABA-induced H$_2$O$_2$ signalling, we measured the endogenous H$_2$O$_2$ levels in the stomata of WT and 35S:PeCHYR1 poplars. We used 2,7-dichlorodihydrofluorescein diacetate (H$_2$DCF-DA), a unique fluorescent probe, to detect the H$_2$O$_2$ levels within guard cells. In the absence of ABA, we found that 35S:PeCHYR1 plants showed slightly elevated H$_2$O$_2$ content compared with WT plants (Figure 4b,c). In the presence of ABA, the levels of H$_2$O$_2$ were increased in both WT and 35S:PeCHYR1 poplar, while 35S:PeCHYR1 poplar exhibited significantly higher H$_2$O$_2$ levels than WT plants (Figure 4b,c).

To explore whether PeCHYR1 promotes ABA-induced stomatal closure, stomatal movement in response to ABA treatment was observed in WT and 35S:PeCHYR1. Then, leaf stomata were observed by scanning electron microscopy. Meanwhile, we observed a significant difference when the leaves were treated with 5 μM ABA for 1 and 2 h; the stomatal apertures closed more quickly in the leaves of the transgenic plants than in the leaves of WT poplar (Figure 4b,c).

Figure 1 Amino acid sequence alignment and phylogenetic tree of different CHYR1 protein family members. (a) Phylogenetic analysis of the PeCHYR1 homologs from Populus, Amborella, Eucalyptus, Malus, Prunus, Theobroma, Picea, Pinus, Salix, Arabidopsis, rice and maize. (b) Multiple alignment of the amino acid sequences of CHYR1 proteins from Populus, Salix and Arabidopsis.

Figure 2 PeCHYR1 expression patterns in different tissues and under different treatments. (a) Relative expression levels of the PeCHYR1 gene in different tissues of P. euphratica. Young leaf; Adult leaf; Old leaf; Phloem; Xylem; Root. (b) Transcript levels of PeCHYR1 were measured by RT-qPCR in response to dehydration. (c) Transcript levels of PeCHYR1 were measured by RT-qPCR in response to ABA. (d) DAB staining in WT seedling leaves during ABA treatment. Error bars are means ± SE (n = 20). Asterisks denote significant differences: *P < 0.05; **P < 0.01.
the WT plants (Figure 4d,e). Conclusively, these data indicated that PeCHYR1 actively regulated ABA-induced ROS production.

PeCHYR1 overexpression enhances WUE by reducing stomatal conductance

As H$_2$O$_2$ can also induce stomatal closure (Bright et al., 2006), we hypothesized that overexpression of PeCHYR1 in transgenic poplar plants may decrease stomatal opening. To determine whether PeCHYR1 could affect photosynthesis in plants, we measured photosynthesis–light curves in WT and 35S:PeCHYR1 poplars. The data indicated that the patterns of photosynthesis were roughly similar in WT and 35S:PeCHYR1 poplars (Figure 5a). The stomatal conductance (Gs) data indicated that WT and 35S:PeCHYR1 poplars had altered Gs responses to light, while the Gs of 35S:PeCHYR1 poplars was lower than that of WT (Figure 5b).

At the same time, leaf transpiration in 35S:PeCHYR1 poplars was significantly lower than that in WT poplars (Figure 5c). In general, instantaneous WUE values in 35S:PeCHYR1 plants were higher than those in WT plants (Figure 5d). Vapour pressure deficit (VPD) was measured, and no significant differences were found between WT and 35S:PeCHYR1 poplars (Figure 5e). This lack of a significant difference in VPD indicates that the differences in transpiration rate were not caused by VPD.

35S:PeCHYR1 poplars possess drought tolerance

Because ABA and drought stress can induce the expression of PeCHYR1 (Figure 2b,c), we hypothesized that PeCHYR1 played a vital role in drought response. Afterwards, 35S:PeCHYR1 poplars and WT poplars were cultured under the same conditions and then subjected to a drought treatment in which the soil relative water content (RWC) was reduced from 70% and rewatered again for 2 days. On day 3, most leaves of the WT poplars were slightly wilted, while the transgenic poplars still appeared normal. On day 7, the leaves of the WT poplars were seriously wilted, while those of the transgenic poplars remained turgid. Furthermore, when the plants were watered again after drought stress, the WT poplars were unable to recover completely, whereas the transgenic poplars returned to normal and continued to grow (Figure 6a). Control plants were kept under the same conditions, except that the soil RWC was maintained at 70%, and no significant difference in physiological indicators and phenotype

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**Figure 3** PeCHYR1 is targeted to the endoplasmic reticulum (ER) and nucleus. (a) Subcellular localization of 35S: GFP and 35S: PeCHYR1-GFP in transiently expressed tobacco leaves. HDEL-RFP was used as an ER localization marker fused with red fluorescent protein (RFP). The nuclear dye DAPI (blue) was applied to mark the nucleus. Bar = 10 μm. (b) Subcellular localization of 35S: GFP and 35S:PeCHYR1-GFP in transiently expressed Arabidopsis leaf protoplasts. 35S: HDEL-RFP was cotransformed with 35S: PeCHYR1-GFP to verify the ER localization of PeCHYR1. The nuclear dye DAPI (blue) was applied to mark the nucleus. Bars = 5 μm.
Figure 4 PeCHYR1 promotes ABA-induced stomatal closure via ROS production. (a) DAB staining shows different levels of ABA-induced ROS production in the leaves of WT and 35S:PeCHYR1. Scale bars = 1 cm (b) Representative confocal images of 200 μM ABA-induced H$_2$O$_2$ production (10 min) in the guard cells of WT poplar and transgenic lines coloured with H2DCFDA. (c) Quantification of H$_2$O$_2$ production in the guard cells of WT poplar and transgenic lines. (d) Detection of ABA-induced stomatal closure in the leaves of the WT and transgenic lines; leaves were separated and immersed under light in stomata-opening solution (OS) for 2 h and then treated with 5 μM ABA for 2 h (OS + ABA). (e) Stomatal closure was observed at 0, 1 and 2 h of ABA treatment with scanning electron microscopy of stomatal aperture. Scale bars = 50 μm. Error bars are means ± SE (n = 50). Asterisks denote significant differences: **P < 0.01.

Figure 5 Light response curves were measured in WT, OXPeCHYR1-1 and OXPeCHYR1-8. Light response curves were measured in the same greenhouse conditions. (a) A- light curve. (b) Gs - light curve. (c) Transpiration- light curve. (d) Instantaneous WUE- light curve. (e) VPD- light curve. Data are means ± SE (n = 25). Asterisks denote significant differences: **P < 0.01.

was found between 35S:PeCHYR1 poplars and WT poplars in control conditions (Figure 6).

Leaf RWC reflects the degree of water deficit imposed. When exposed to short-term drought stress, 35S:PeCHYR1 poplar retained observably higher leaf RWC values than WT poplar (Figure 6b). As malondialdehyde (MDA) content is a valid indicator of cytomembrane oxidative damage, we measured the variations in MDA content caused by drought stress in the leaves of WT and transgenic poplars. The results indicated that the 35S:PeCHYR1 poplars had lower levels of MDA than the WT poplars did when exposed to short-term drought stress (Figure 6c). The REC (relative electrical conductance) of both 35S:PeCHYR1 and WT poplars increased after drought treatment. After drought stress, the transgenic poplars showed less membrane damage.
H2O2 content in WT poplars was significantly higher than that in 57.6% ion leakage) (Figure 6d).

Corresponded with the H2O2 levels, the results indicated that the PeCHYR1 expression in plants influences the response to long-term water deficit, 35S:PeCHYR1 transgenic and WT poplars were subjected to a 32-day water-deficit assay by holding the soil RWC at stable values (45% and 70%). After 32 days, in both shoot and root, the conditions of the WT and 35S:PeCHYR1 poplar were approximately identical at 70% soil RWC. After 32 days of drought treatment, 35S:PeCHYR1 poplar reached greater heights and appeared in better condition than WT at 45% soil RWC (Figure 8a). Furthermore, at 70% soil RWC, the chlorophyll level of 35S:PeCHYR1 poplar was similar to that of WT plants. However, after the 32-day water-deficit assay at 45% soil RWC, 35S:PeCHYR1 poplar had higher chlorophyll content than WT plants (Figure 8b). In addition, 35S:PeCHYR1 poplar showed higher Maximal PSII quantum yield (Fv/Fm) and higher leaf RWC than WT poplar did (Figure 8c,d).

To explore whether PeCHYR1 would affect growth and development in the absence of water stress (70% soil RWC) and under mild water stress (45% soil RWC), plant height and stem height growth rate were monitored. The 35S:PeCHYR1 poplar had similar plant height and stem height growth rate to those of wild-type plants in the absence of water stress (Figure 9a,b). At 45% soil RWC, the 35S:PeCHYR1 poplar had a significantly greater plant height and stem height growth rate than those of WT plants (Figure 9a,b). These results suggested that 35S:PeCHYR1 poplar maintained higher growth under long-term water-deficit conditions.
of 35S:PeCHYR1: water stress, and the shoot weight, root weight and total biomass accumulation in the shoot and root under mild water stress. However, compared with that of WT plants, transgenic plants had more biomass accumulations in all parts of the plant (Figure 9c–f).

Under no water stress, transgenic and WT plants had similar total biomass, and root weight, and mild water stress (45% soil RWC), shoot weight, root weight, and 35S:PeCHYR1 poplar had a larger root–shoot ratio than that of WT after 16 days of drought treatment (Figure 9f). After 32 days of drought treatment, the biomass of 35S:PeCHYR1 poplar was 36.9% and 43.4% higher, respectively, than that of WT (Figure 9e), and the root–shoot ratio of 35S:PeCHYR1 poplar was 15.3% and 8.9% higher, respectively, than that of WT (Figure 9f).

To observe the difference in biomass accumulation between WT and 35S:PeCHYR1 poplar under no water stress (70% soil RWC) and mild water stress (45% soil RWC), shoot weight, root weight, total biomass, and root–shoot ratio were monitored every 8 days. Under no water stress, transgenic and WT plants had similar biomass accumulations in all parts of the plant (Figure 9c–f). However, compared with that of WT plants, transgenic plants had more biomass accumulation in the shoot and root under mild water stress, and the shoot weight, root weight and total biomass of 35S:PeCHYR1 poplar were obviously higher than those of WT plants after 8 days of drought treatment (Figure 9c–e). The root–shoot ratio reflects the relationship between the aboveground and underground parts of the plant. Both 35S:PeCHYR1 and WT plants had greater root–shoot ratios, and 35S:PeCHYR1 poplar had a larger root–shoot ratio than that of WT after 16 days of drought treatment (Figure 9f). After 32 days of drought treatment, the biomass of 35S:PeCHYR1 poplar was 36.9% and 43.4% higher, respectively, than that of WT (Figure 9e), and the root–shoot ratio of 35S:PeCHYR1 poplar was 15.3% and 8.9% higher, respectively, than that of WT (Figure 9f). In summary,
under water-deficit conditions, 35S:PeCHYR1 poplar maintained higher growth than the WT.

**PeCHYR1 mediated the expression of upstream and downstream genes involved in adversity**

The genes *LEA14, RbohD, RbohF* and *SnRK2.6* are known to be drought responsive and have positive functions in drought tolerance (Ding et al., 2015). Thus, WT and 35S:PeCHYR1 poplars were treated with 2 h of dehydration stress to investigate changes in *PeLEA14, PeRbohD, PeRbohF* and *PeSnRK2.6* expression. The results showed that the expression levels of *PeLEA14, PeRbohD, PeRbohF* and *PeSnRK2.6* in transgenic poplars were greater than those in WT poplars after 2 h of dehydration stress (Figure 10a–d). These results indicated that the improved drought tolerance of 35S:PeCHYR1 transgenic poplars may depend on the regulation of downstream genes.

**Discussion**

*PeCHYR1* was cloned from *P. euphratica*. A multiple alignment of amino acid sequences showed that *PeCHYR1, PtrCHYR1, SpCHYR1* and *AtCHYR1* all contain a RING domain and a CHY zinc-finger domain (Figure 1b). The conserved “CxHY” motif lies in the CHY zinc-finger domain, which plays a role in physiological
interaction and ubiquitination (Lee and Kim, 2011; Leng et al., 2003) and consists of 12 histidines and cysteines that bind with three zinc ions, composing a unique zinc-finger motif (Cayrol et al., 2007). Phylogenetic analysis indicated that the sequences of PeCHYR1, PtCHYR1 and AChYR1 had higher homology (Figure 1a), potentially sharing a conserved and crucial function. Based on the evolutionary tree of related genes, PeCHYR1, PtCHYR1 (Potri.009G005700) and AChYR1 (ATSG22920) belonged to the first part (I) in the same branch (Figure 1a), which may characterize relative stomatal aperture and enhance drought tolerance under stress control conditions (Ding et al., 2015). However, OsRZFP34, AtCHYR2 (ATSG25560) and PtCHYR2 (Potri.006G245400) belonged to the third part (III) in the same branch (Figure 1a), which may control stomatal opening even with ABA treatment (Hsu et al., 2014). PeCHYR1 also possesses a particular function; this protein is probably phosphorylated by PeSnRK2.6 and augments drought and ABA-induced responses in plants, which is similar to the function of CHYR1 in Arabidopsis (Ding et al., 2015). Furthermore, transgenic poplar plants overexpressing PeCHYR1 showed increased drought tolerance via ABA-induced stomatal closure and ROS production.

Previous studies have indicated that plants regulate ABA-induced stomatal closure by ROS accumulation (Bright et al., 2006; Hua et al., 2012; Li et al., 2017). H$_2$O$_2$ is a ROS that acts as a second messenger in ABA signalling in guard cells (Bright et al., 2006; Dietz et al., 2016; Kohler et al., 2003). The content of H$_2$O$_2$ in plants increases considerably under treatment with exogenous ABA (Figure 2d). This increased H$_2$O$_2$ level is an early signalling event in ABA-induced signal transduction (Bright et al., 2006; Dietz et al., 2016; Kreslavski et al., 2012). In addition, the level of H$_2$O$_2$ increased more under treatment with exogenous ABA in 35S:PeCHYR1 transgenic plants than in WT plants (Figure 4a-c). The altered relative expression levels of PerbohD and PerbohF (Figure 10c,d), which are involved in producing H$_2$O$_2$, were in agreement with the above results. Overexpression of PeCHYR1 augmented ABA responsiveness, thereby activating stress-inducible gene expression. These results indicated that PeCHYR1 may promote ABA-induced H$_2$O$_2$ production.

Abscisic acid is involved in abiotic stress response and regulates the expression of many stress-responsive genes (Danquah et al., 2014; Kobayashi et al., 2005; Zhu, 2016). Whether these genes are induced in response to exogenous ABA treatment depends on their involvement in the ABA-independent or ABA-dependent stress response pathways (Yang et al., 2011). The protein DST generates an increase in hydrogen peroxide and promotes stomatal closure by an ABA-independent pathway (Huang et al., 2009). However, 35S:OsASR5 plants were more sensitive to exogenous ABA treatment than WT plants, suggesting that OsASR5 was involved in an ABA-dependent pathway (Li et al., 2017). In our study, treatment with exogenous ABA did observably affect the expression of PeCHYR1 (Figure 2c), and significant differences were present between the stomatal apertures of the WT and 35S:PeCHYR1 lines under ABA treatment (Figure 4d,e), demonstrating that PeCHYR1 increased sensitivity to exogenous ABA and reduced stomatal aperture. At the same time, PeCHYR1 also participates in biological stress responses (Figure S3a,b).

Stomatal control gas exchange, and they can regulate transpiration by limiting water loss and affect photosynthesis by facilitating CO$_2$ uptake (Hetherington and Woodward, 2003; Vahisalu et al., 2008). Stomatal conductance is a direct result of stomatal density and aperture (Bussis et al., 2006; Chaerle et al., 2005; Qin et al., 2016). In our study, changes in transpiration were consistent with changes in stomatal conductance. As same as stomatal conductance, transpiration in 35S:PeCHYR1 plants was significantly lower than that in WT plants (Figure Sb,c). However, the change of photosynthesis in WT plants was roughly similar to that in 35S:PeCHYR1 plants. Accordingly, the instantaneous WUE in the transgenic plants was elevated (Figure 5d), consistent with the normal-temperature water loss rate of 35S:PeCHYR1 plants, which showed less water loss than WT plants did (Figure 7d). Previous studies have indicated that the decreased stomatal conductance in 35S:PeCHYR1 transgenic plants would not necessarily lead to decreased photosynthesis under well-watered conditions (Wang et al., 2016). Rubisco and ribulose-1,5-bisphosphate are nonstomatal factors that play roles in modulating photosynthetic rate (Peary and Seemann, 1990; Qin et al., 2016), and thus, stomatal conductance does not solely control changes in foliar photosynthetic rate (Ninemets, 2002). These results indicate that a modest decrease in stomatal conductance in 35S:PeCHYR1 transgenic plants can increase WUE but not affect photosynthesis under unstressed conditions.

When the plants were in a strictly water-controlled environment, the reduced stomatal conductance of the 35S:PeCHYR1 transgenic plants possibly reduced water evaporation and played an important part in maintaining a higher leaf RWC (Figure 7b). Under ABA and dehydration treatment, the expression level of PeCHYR1 gradually rose (Figure 2b,c), consistent with the characteristics of drought tolerance in 35S:PeCHYR1 transgenic plants. The evidence indicated that 35S:PeCHYR1 plants had a stronger drought tolerance phenotype than WT plants did. Under drought conditions, 35S:PeCHYR1 transgenic plants could assimilate more carbon dioxide (Figure 7c). Generally, the level of H$_2$O$_2$ in plants can significantly increase under stress conditions (Bhattacharjee, 2012; Bright et al., 2006; de la Garma et al., 2015). Abrupt increases in plant H$_2$O$_2$ content are regarded as a marker of various stresses that can induce the antioxidant defence system (Ding et al., 2010; Jiang and Zhang, 2002). Nevertheless, exceedingly high levels of H$_2$O$_2$ can give rise to oxidative injury (Kuge et al., 2010; Wang et al., 2008). In our study, both in the short-term ABA treatment and early in the drought treatment, the endogenous hydrogen peroxide levels in 35S:PeCHYR1 transgenic plants suddenly increased and became higher than those in the WT plants (Figure 4a-c), thereby activating the native antioxidant system to cope with the stress. However, when the plants were in extreme drought conditions, the accumulation of H$_2$O$_2$ in the WT was significantly higher than that in the overexpression lines (Figure 6e), the stomatal conductance in the WT lines almost reached zero (Figure 7b), and photosynthesis was markedly lower than that in the overexpression lines (Figure 7c). The major ROS-scavenging mechanisms include APX (ascorbate peroxidase), POD, SOD and CAT (catalase) (Ahmad et al., 2008; Dietz et al., 2016; Willekens et al., 1997), and 35S:PeCHYR1 plants showed higher activity levels of POD and SOD under drought stress conditions. POD and SOD, which are beneficial to the maintenance of ROS levels during long-term drought, were activated in transgenic plants (Figure 6f,g). Our observations demonstrated that 35S:PeCHYR1 transgenic plants underwent a sudden, dramatic increase in H$_2$O$_2$, which activated the antioxidant system to reduce oxidative damage after drought treatment.

In addition, the level of MDA indirectly indicates the extent of membrane lipid peroxidation (Shi et al., 2013, 2014). In our research, drought stress led to mass accumulation of ROS in WT
plants, elevating the content of MDA compared with that in 35S: PeCHYR1 transgenic plants, seriously damaging membrane permeability, causing intracellular ion leakage, and thereby exacerbating plant senescence and death (Figure 6c,d). Thus, WT plants continued to receive more damage than transgenic plants, and we conclude that overexpression of PeCHYR1 in plants is beneficial for drought tolerance.

Maximal PSI quantum yield (Fv/Fm), which reflects the potential maximum light energy conversion efficiency of plants, is 0.8–0.85 under normal conditions in the vast majority of higher plants (Kitajima and Butler, 1975). Photosystem II efficiency declines in plants under environmental stress (Ke et al., 2016; Winter and Lesch, 1992). Although drought stress affects plant photosystem II electron transport and reduces chlorophyll content, 35S: PeCHYR1 poplar exhibited higher chlorophyll content and photosystem II efficiency than WT poplar did (Figure 8b,c), leading to less injury under water-deficit conditions (Figure 8a). Furthermore, we found that leaf RWC in 35S:PeCHYR1 poplar was obviously higher than that in WT poplar (Figure 8d), which also ensured that plant photosystem II could transport electrons properly and that plants could assimilate carbon dioxide normally under severe drought stress. Our experiment clearly demonstrated that poplars with decreased stomatal conductance could increase in plant height and biomass under water-deficit conditions (Figure 9). 35S:PeCHYR1 poplar showed decreased transpiration, increased photosynthetic rate, enhanced WUE, elevated plant height and boosted biomass under water-deficit conditions. Drought is the primary abiotic stress responsible for negatively influencing poplar growth (Dash et al., 2017; Ke et al., 2016; Yin et al., 2004). We suggest that 35S:PeCHYR1 poplar, a novel and drought-tolerant forestry species, can improve WUE and increase biomass under drought stress.

In summary, a ubiquitin E3 ligase from P. euphratica, PeCHYR1, increased sensitivity to exogenous ABA, enhanced ROS production and reduced stomatal aperture, thereby reinforcing plant drought tolerance. Moreover, the up-regulation of PeCHYR1 reduced transpiration by decreasing the stomatal conductance without influencing photosynthesis under normal conditions. 35S:PeCHYR1 poplar achieved higher photosynthesis and biomass accumulation along with reduced levels of MDA and REC under drought stress. Therefore, the PeCHYR1 gene family will be conducive to further exploration of the mechanisms of drought tolerance, and we obtained a novel line of poplars with increased drought tolerance.

Experimental procedures

Plant materials

One-year-old seedlings of P. euphratica with heights of 15 cm were obtained from Yuli County, Korla, Xinjiang province, China (41°19'37.06" N, 86°15'44.78" E) in March, which lies in a warm temperate zone and has a continental desert climate. The average annual rainfall is 43 mm, and the average annual evaporation is 2700 mm. One-year-old seedlings of P. euphratica were transplanted in individual pots (10 L) containing sandy soil (approximately 70% sand) and placed in a seed plot [light cycle: 16.0 h of light (06:00 am–10:00 pm); temperature (20–24 °C)] at Haidian, Beijing, China (40°00′00″N, 116°20′00″E; 49 m above sea level). Potted P. euphratica were watered on the basis of evaporation demand and irrigated with 1 L of Hoagland nutrient solution every 2 weeks for 2 months before treatment (Li et al., 2011; Wang et al., 2016). To analyse gene expression, similarly grown seedlings of P. euphratica (40–50 cm high, with 30–40 leaves) were subjected to ABA treatment and dehydration. The seedlings were treated with a solution containing 200.0 μM ABA (Sigma, A1049) and 1 g TWEEN® 40 (Sigma, P1504-500 ML), which was sprayed on the leaves of P. euphratica. Meanwhile, the control was treated with 1 g TWEEN® 40 in ultrapure water. Dehydration treatment was applied to seedlings that had been removed from the soil and exposed to air with 70% RH at 23 °C under dim light for 12 h (Ma et al., 2010; Wang et al., 2016). To obtain complete plants, we gently pulled out each seedling and carefully washed the roots to remove the soil. For every test, leaves were separated from plants at the given time periods and promptly immersed in liquid nitrogen. We collected different organs and tissues of 2-month-old P. euphratica, including YL, young leaf; AL, adult leaf; OL, old leaf; Pe, petiole; X, xylem; and R, root at the same time and immediately immersed them in liquid nitrogen.

cDNA cloning of PeCHYR1 from P. euphratica

Total RNA was extracted from the collected materials using the cetyltrimethylammonium bromide (CTAB) method (Chang et al., 1993). In the last step, potentially contaminating DNA was removed by treatment with DNase I. We used a NanoDrop 2000 Spectrophotometer (Thermo, West Palm Beach, FL) to measure the quality and quantity of RNA. Two micrograms of total RNA was used for the reverse transcription reaction with a Thiengan FastQuant RT Kit (with gDNase) (Qagen, Düsseldorf, Germany) according to the protocol. The cDNA of PeCHYR1 was cloned by PCR, and the primers are displayed in Table S1.

Quantitative real-time polymerase chain reaction (RT-qPCR) analysis

Twenty microlitres of cDNA was diluted 1 : 10 with nucleic-free water. Each reaction contained 10 μL SuperReal PreMix Plus (Tiangen Bio Inc., Beijing, China), 2 μL ROX Reference Dye (Qiagen), 1 μL cDNA (single-stranded circular DNA (sscDNA), corresponding to 10 ng of total RNA), 5.8 μL nucleic-free water and 0.6 μL of each primer. The cycling parameters were 95 °C for 15 min and then 45 cycles of 20 s at 95 °C and 60 s at 60 °C (Wang et al., 2014). RT-qPCR was performed with the ABI StepOnePlus Real-Time PCR System (ABI, Foster City, CA) according to the manufacturer’s specifications. Each experiment was performed in 20 replicates (five biological replicates × four technical replicates), and all primers used are displayed in Table S1. We used the software tool Primer Premier 6 to design primers. We calculated the relative expression level of PeCHYR1 by the ratio = (Et)ACTR/(Et)ACTR method (Wang et al., 2014, 2016).

Phylogenetic and domain analysis of PeCHYR1

Homologous amino acid sequences were obtained from the Phytozome database (http://www.phytozome.net/), and a phylogenetic tree of PeCHYR1 was estimated with MEGA 7. DNAMAN was used to examine the amino acid sequences and confirm the conserved structures. The sequence data from this study can be found in the Phytozome database (http://www.phytozome.net/). The accession numbers of the genes used are displayed in Table S2.

Plasmid construction and genetic transformation of poplar

The PeCHYR1 sequence was inserted into the Smal and SacI sites in the pCAMBIA-1301 vector. The construct was transformed into
Agrobacterium tumefaciens (EHA105) and then transformed into poplar 84K using the leaf disc method (Hsu et al., 2011; Yao et al., 2016). The leaves of poplar 84K were incubated on substrates (pH 5.80) containing 0.020 mg/L thidiazuron (TDZ), 0.50 mg/L 6-benzylaminopurine (6-BA), 0.050 mg/L 1-naphthaleneacetic acid (NAA), 250 mg/L cefotaxime, 4 mg/L hygromycin and 0.80% (w/v) agar for shoot induction and selection. The regenerated shoots were individually detached from the callus and inserted into a rooting culture [1/2 MS substrate including 0.050 mg/L NAA, 250 mg/L cefotaxime, 150 mg/L hygromycin phosphotransferase and 0.9% (w/v) agar]. One-month-old regenerated seedlings were transplanted to small pots (12 cm width × 12 cm height) with the same soil potting soil, turfy soil, and vermiculite 1:1:1) and then placed in a greenhouse [light cycle: 16.0 h of light, 8.0 h of dark; temperature: 24 °C (light)/20 °C (dark); relative humidity: 40%–45%].

Subcellular localization

To determine the subcellular localization of PeCHYR1, 35S: PeCHYR1-GFP fusion protein and 35S: HDEL-RFP fusion protein were transiently transfected together into tobacco leaves by the previously published protocol (Cui et al., 2012). Cotransformation of 35S: HDEL-RFP in combination with 35S: PeCHYR1-GFP was also performed in Arabidopsis leaf protoplasts by means of polyethylene glycol (PEG) treatment (Yoo et al., 2007). HDEL-RFP was used as an ER localization marker (Cui et al., 2012; Tian et al., 2013). The nuclear dye DAPI (10 mg/mL; Sigma-Aldrich) was applied to mark the nucleus. The transformed protoplasts and tobacco leaves were observed by laser confocal fluorescence microscopy (Leica TCS SP8). The LAS-AF software was used to record the images. The following lines of the argon ion laser were used: 488 nm for GFP, 584 nm for RFP, 380 nm for DAPI and 488 nm for chlorophyll. Fluorescence was detected at 495–515 nm for GFP, 575–590 nm for RFP, 430–450 nm for DAPI and 650 nm for chlorophyll.

Molecular verification of transgenic plants

Total DNA was extracted from every line using the CTAB method (Tiangen FastQuant RT Kit (with gDNase) (Qiagen, Dusseldorp, Germany), according to the manufacturer’s instructions. qRT-PCR was performed with the ABI StepOnePlus Real-Time PCR System (ABI, Foster City, CA) according to the manufacturer’s specifica-
tions. The internal control was a combination of two reference genes, PeActin and PeUBQ, which were screened previously (Bustin et al., 2009; Czechowski et al., 2005; Rao et al., 2013; Wang et al., 2014). The primers used are shown in Table S1.

Histochemical staining of GUS activity

Histochemical staining of GUS was handled as described previ-
ously (Lee et al., 2010). Briefly, isolated leaves were immersed at 37.0 °C in a mix including 2 mM 5-bromo-4-chloro-3-indolyl-β-D-glucoronide, 0.1 mM sodium phosphate buffer (pH 7.0), 0.50 mM each of potassium ferri- and ferrocyanide, 10.0 mM EDTA (pH 7.0) and 0.10% Triton X-100 for 12 h. Then, the isolated leaves were incubated in 70% ethanol for 12 h and photographed.

DAB staining

DAB staining was applied to detect the production of H2O2, as described previously (Ding et al., 2015). The leaves of 2-month-old transgenic and WT plants were sprayed with 100 μM ABA (Sigma, A1049) and 1 g TWEEN® 40 (Sigma, P1504-500 ML) for 0, 1, 2 and 3 h and infiltrated in prepared 0.1 mg/mL DAB solution (Beyotime, ST033) in the dark for 8 h at 28 °C. Meanwhile, the controls were treated with 1 g TWEEN® 40 in ultrapure water. To remove chlorophyll after DAB staining, the stained leaves were incubated in a solution of 70% ethanol, and then, they were photographed.

H2O2 detection in guard cells

A unique fluorescent dye, H2DCF-DA (Sigma, D6883 HZB1212), was used for H2O2 detection in the guard cells of the WT and transgenic plants (Huang et al., 2009). To observe transient ABA responses, the leaves of two-month-old seedlings (WT, OXPe-
CHYR1) were immerged in 50 mM H2DCF-DA for 12 min at 25 °C in the dark. Then, the leaves were steeped in liquid 1/2 MS and treated with 200 μM ABA. Confocal measurements were taken every 30 s for 5 min of ABA stress in xyt mode. The amount of H2DCF-DA fluorescence in guard cells was measured with Image-Pro Plus6 (Shi et al., 2013).

Stomatal movement analysis

Stomatal movement tests were performed as described previously (Cui et al., 2015; Hsu et al., 2014; Li et al., 2017), with slight amendments. Leaves from two-month-old seedlings (WT and OXPeCHYR1) were detached into wire netting and immediately fixed in liquid nitrogen. Then, the fixed samples were immediately put into an –80 °C Ultra-low Freeze Dryer (Biosafer-1BA, Jiangsu, China (Mainland)) and fully dried for 24 h. The fully dried sections were coated with gold using an Ion Sputter Coater (Eiko Engineering, IB-2, Japan). All observations were collected using a scanning electron microscope (Hitachi S-4700, Japan). For the ABA-induced stomatal closure experiments, the same parts of the leaves were immersed in stomata-opening solution (OS) contain-
ing 0.01 mM KCl, 0.1 mM CaCl2 and 0.01 mM MES-KOH for 0.5 h in the dark and 2.0 h in the light, and then, 5 μM ABA was added. Then, stomata were fixed after 0, 1 and 2 h of ABA treatment. The stomatal images were collected using a Hitachi S-4700 scanning electron microscope. More than 100 guard cells from each sample were measured to derive the stomatal aperture.

Drought tolerance experiment

Fifteen 35S:PeCHYR1 and five WT poplar plants grown in a greenhouse [light cycle: 16.0 h of light (06:00 am–10:00 pm); temperature (20 °C–24 °C)] were subjected to a short-term drought experiment. All of the plants (height 18–20 cm) were grown in suitably sized pots (12 cm × 12 cm × 12 cm), and each pot had a tray. All of the plants encountered drought treatment, in which the soil RWC was reduced from 70%. The control plants were kept in the same conditions, except that the soil RWC was maintained at 70%. A soil RWC of 70% induces no stress and is the most suitable environment for plant growth (Wang et al., 2016).

Physiological and biochemical analysis

A Li-Cor portable photosynthesis meter (LI-COR 6400) was applied to detect net photosynthetic rate, transpiration and stomatal conductance in the ninth to eleventh leaves of 35S:PeCHYR1 and...
WT poplar plants. We used the LI-COR 6400 portable photosynthesis analysis system to measure light curves under normal conditions of poplars grown in a greenhouse for 1.5 months. Twenty plants were measured. The experiment covered net CO₂ assimilation (A), Gs, transpiration, VPD and instantaneous WUE (A/Transpiration). Photosynthetic light response curves were determined at photosynthetically active radiation (PAR) levels of 1500, 1200, 1000, 800, 600, 400, 200, 150, 100, 80, 50, 20 and 0 μmol/m²/s with 450 μmol/m²/s external CO₂, which was controlled by the LI-COR 6400 (Wang et al., 2016).

Measurement of H₂O₂ content and antioxidant enzyme activities

Based on the continuous changes of photosynthesis under drought stress, H₂O₂ and antioxidant enzyme activities were measured at 0, 3 and 7 days. Samples of 0.5 g fresh leaves were detached from the drought treatment and control plants at 0, 3 and 7 days and ground into a fine powder in liquid nitrogen. The plant extracts were isolated in 50 mM sodium phosphate buffer (pH 7.8) mixed thoroughly with 1 mL of 0.1% (w/v) titanium sulphate in 20%, v/v H₂SO₄ for 10 min (Shi et al., 2012, 2014). H₂O₂ content and the activities of antioxidant enzymes (SOD and POD) were measured using previously published protocols (Shi et al., 2014). The levels of H₂O₂, SOD and POD were expressed as μmol (g * fresh weight (FW))⁻¹. These experiments consisted of 20 replicates (4 biological replicates × 5 technical replicates) under identical conditions.

MDA level and relative electrical conductance (REC) assay

Materials detached from drought-stressed plants at 0 and 7 days were measured to determine MDA level and REC using previously published protocols (Shi et al., 2013, 2014). MDA (mol/g FW) = (6.45*(A532–A600) – 0.56 * A450) * 9 μVs⁻¹FW⁻¹ [Vt: total volume of extract (mL); Vs: extract volume was determined with (mL)]. The electrical conductivity of the supernatant (L1) was detected using a DDS-307 Conductivity Meter (Leici-DDS-307A, Shanghai, China). These experiments were independently replicated 6 times under identical conditions.

Analysis of chlorophyll fluorescence and content

Photosynthetic activity in the 10th–13th leaves of transgenic and WT plants was monitored by Maximal PSII quantum yield (Fv/Fm) values, which reflect the potential maximum light energy conversion efficiency of plants, using a PAM chlorophyll fluorometer (PAM1000) after 20 min of dark adaptation. Chlorophyll content was detected in the 10th–13th leaves of transgenic and WT plants using a portable chlorophyll meter (SPAD-502Plus, Konica Minolta, Japan). Total chlorophyll contents were measured at 70% soil RWC and 45% soil RWC.

Long-term drought experiment

Drought stress was implemented in transgenic and WT poplar plants by maintaining the soil RWC at 45% for 32 days. Control plants were maintained at a soil RWC of 70%. A soil RWC of 70% causes no stress and is a suitable environment for plant growth (Wang et al., 2016). The transgenic and WT poplar plants (five poplars per line under various treatment) were grown in a greenhouse [light cycle: 16.0 h of light (06:00 am–10:00 pm); temperature (20 °C–24 °C)], all plants (height 40–43 cm) were kept in suitably sized pots (volume, 15 L), and every pot had a tray. The containers were weighed daily, and lost water was supplemented. We measured the height, shoot biomass and root biomass of each poplar every 7 days. After 32 days, chlorophyll content, leaf RWCA and Maximal PSII quantum yield (Fv/Fm) were measured.

Relative water content (RWC)

The 10th–13th leaves were detached and used for RWC measurements. In short, we determined the leaf FW (removed fresh leaves were weighed), leaf turgid weight (TW, leaves were measured after submerging in water for 8 h) and leaf dry weight (DW, leaves were measured after drying at 80 °C for 72 h). The RWC was calculated as (FW – DW)/(TW – DW) × 100%.

Statistical analysis

The experimental data were subjected to analysis of variance by Statistical Product and Service Solutions 17.0 (SPSS). For statistical analyses, Student’s t-test was used to generate every P value (one-way analysis of variance: *P < 0.05; **P < 0.01). Both one-way and two-way analyses of variance were used to determine significance. The data were normalized, and all samples were normally distributed with homogeneity of variance.

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Supporting information

Additional Supporting Information may be found online in the supporting information tab for this article.
Figure S1 Analysis of Cis elements of the PeCHYR1 gene.
Figure S2 Analysis of the transgenic poplar plants overexpressing PeCHYR1.
Figure S3 35S: PeCHYR1 plants exhibited increased disease resistance under anthracnose fungus treatment.
Table S1 Primer sequences used for cloning of PeCHYR1 cDNA and RT-PCR.
Table S2 Accession numbers of gene.