NAC (NAM, ATAF1/2, and CUC2) proteins play critical roles in many plant biological processes and environmental stress. However, NAC proteins from *Tamarix hispida* have not been functionally characterized. Here, we studied a NAC gene from *T. hispida*, ThNAC13, in response to salt and osmotic stresses. ThNAC13 is a nuclear protein with a C-terminal transactivation domain. ThNAC13 can bind to NAC recognized sites and calmodulin-binding NAC (CBNAC) binding element. Overexpression of ThNAC13 in *Arabidopsis* improved seed germination rate and increased root growth and fresh weight gain under salt or osmotic stress. Transgenic *T. hispida* plants transiently overexpressing ThNAC13 and with RNAi-silenced ThNAC13 were generated for gain- and loss-of-function experiments. Following exposure to salt or osmotic stress, overexpression of ThNAC13 induced superoxide dismutase (SOD) and peroxidase (POD) activities, chlorophyll and proline contents; decreased the reactive oxygen species (ROS) and malondialdehyde levels; and reduced electrolyte leakage rates in both transgenic *Tamarix* and *Arabidopsis* plants. In contrast, RNAi-silenced ThNAC13 showed the opposite results in transgenic *Tamarix*. Furthermore, ThNAC13 induced the expression of SODs and PODs in transgenic *Arabidopsis*. These results suggest that ThNAC13 improves salt and osmotic tolerance by enhancing the ROS-scavenging capability and adjusting osmotic potential.

**Keywords:** abiotic stress, NAC transcription factor, ROS-scavenging, stress resistance, *Tamarix hispida*

**INTRODUCTION**

Transcriptional regulation plays crucial roles in controlling plant biological processes and signaling pathways, and transcription factors specifically interact with promoter elements of target genes to activate or repress their transcription (Pérez-Rodriguez et al., 2009; Shang et al., 2013). In plants, several families of stress-responsive transcription factors have been functionally characterized in stress regulation networks, such as NAC, bZIP, WRKY, MYB/MYC and AP2/ERF (Le Hénaff et al., 2013; Shiriga et al., 2014).

The NAC (NAM, ATAF1/2, and CUC2) transcription factors are plant-specific proteins, which comprise one of the largest transcription factors families, and are involved in various plant biological processes, such as lateral root formation, cell division, wood formation and xylary
Most NAC proteins contain a highly conserved N-terminus domain and a diversified C-terminal domain (Puranik et al., 2012). To date, some NAC genes have been shown to be involved in abiotic stress. In Arabidopsis and rice, the stress-responsive NAC gene, ANAC096, was reported to have a synergistic relationship with ABRE binding factor and increased the plant survival rate under osmotic and drought stresses (Xu et al., 2013). AtNAC019, AtNAC055, and AtNAC072 could specifically bind to the NAC recognized site (NACRS) in the promoter of the EARLY RESPONSIVE TO DEHYDRATION STRESS 1 (ERD1) gene to enhance drought tolerance (Tran et al., 2004; Hickman et al., 2013). The Arabidopsis NAC016 can repress the transcripts of ABCISIC ACID-RESPONSIVE ELEMENT BINDING PROTEIN 1 (AREB1), which is a key transcription factor in the ABA-dependent stress-signaling pathway, and then promote responses to drought stress (Sakuraba et al., 2015). OsNAC5 expression is induced by drought, high salinity and ABA, enhancing tolerance to drought, salinity and low temperature in transgenic rice (Jeong et al., 2013). Hong et al. (2016) demonstrated that ONAC022 expression was a response to abiotic stress, and transgenic rice overexpressing ONAC022 showed enhanced tolerance to drought and salt stresses by modulating the ABA signaling pathway. In addition, many NAC proteins have been shown to play important roles in modulating ROS levels. For example, Fang et al. (2015) reported that SNAC3 was ubiquitously expressed under drought, salinity and ABA treatments. Overexpression of SNAC3 in transgenic rice conferred tolerance to high temperature, drought and oxidative stresses through modulation of ROS scavenging pathways. Oilseed rape NAC36 and NAC55 could induce the expression of genes involved in ROS scavenging to decrease ROS accumulation, leading to improved abiotic stress tolerance (Niu et al., 2016; Chen et al., 2017). In contrast, sweet potato ThNAC1 could increase ROS in response to jasmonic acid and plays a role in reprogramming the transcriptional response to wounding via the JA-mediated pathway (Chen et al., 2016). In soybean, GmNAC20 and GmNAC11 are involved in different abiotic stresses, and their expression in Arabidopsis conferred both salt and freezing tolerance (Hao et al., 2011). Overexpression of horseradish MuNAC4 in transgenic groundnut plants reduces the damage to membrane structures and enhances osmotic adjustment and antioxidative enzyme regulation under drought stress (Pandurangaiyah et al., 2014). TaNAC29 and TaNAC67 are up-regulated under different abiotic stresses, and transgenic Arabidopsis plants expressing these genes showed increased salt and drought tolerance (Mao et al., 2014; Huang et al., 2015). In addition, a NAC-type transcription factor from chickpea, CarNAC4, induces the expression of genes involved in stress tolerance, including DREB2A, COR15A, RD29A, and KIN1, which increased tolerance to salt and drought stresses (Yu et al., 2016).

Although different NAC transcription factors have been successively identified in several plant species and reported to be involved in abiotic stress, the specific biological functions of NAC transcription factors from halophyte woody plants have not been fully elucidated. Tamarix (Tamaricaceae) is a family of woody halophyte species, which includes small trees or shrubs, and is extensively distributed in saline soils of drought-stricken areas of Central Asia and China. Tamarix hispida, a species of the genus Tamarix, is highly tolerant to salinity, drought and extreme temperatures. These characteristics make this species a suitable plant for investigation of the mechanism of stress tolerance. We previously identified 21 full-length coding region NAC genes from T. hispida and found that ThNAC13 was predominantly up-regulated by salt and drought treatments (Wang et al., 2014b). In this study, we performed a detailed functional analysis of ThNAC13 in response to salt and osmotic stresses. ThNAC13 protein is localized in the cell nucleus and has transactivation activity at the C-terminus. T. hispida plants transiently overexpressing and with RNAi-silencing of ThNAC13 were generated to investigate the tolerance to salt and osmotic stresses. Furthermore, Arabidopsis plants overexpressing ThNAC13 were also generated and used to further confirm the results obtained from T. hispida plants. Our results suggest that ThNAC13 is a potential candidate for enhancing tolerance to salt and osmotic stresses via genetic breeding.

**MATERIALS AND METHODS**

**Plant Materials and Growth Conditions**

*Tamarix hispida* plants were used for gene cloning, transgenic analysis and expression analysis. The seedlings of *T. hispida* were cultured in tissue culture bottles containing half-strength Murashige-Skoog (1/2 MS) solid medium [2% (w/v) agar] in a culture room at average temperature of 24°C with 14 h light/10 h darkness photocyte conditions. *Arabidopsis thaliana* (ecotype Columbia) was used for genetic transformation and functional analysis. Seven-day-old seedlings were grown in pots filled with soil and perlite in a controlled chamber (16 h light/8 h darkness photocycle; 70–75% relative humidity; 22°C).

**Isolation and Bioinformatic Analysis of ThNAC13**

The cDNA sequence of ThNAC13 (GenBank number: JQ974967) was identified from the transcriptomes of *T. hispida* (Wang et al., 2014b). Multiple sequence alignments of the deduced protein sequence of ThNAC13 together with other reported stress-responsive NACs from different plant species were deduced using ClustalW software, and the phylogenetic tree was constructed with MEGA (version 5.1) program (Tamura et al., 2011). The nuclear localization signal of ThNAC13 was predicted using the WoLFPSORT algorithm (Horton et al., 2007). The conserved motifs among the full length NAC protein sequences were predicted using MEME online program1 (Bailey et al., 2009).

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1http://meme-suite.org
Subcellular Localization Analysis of ThNAC13 Protein

The full-length coding region of ThNAC13 (without stop codon) was cloned into the pBI121 vector containing the GFP reporter gene driven by the CaMV 35S promoter to generate the 35S:ThNAC13-GFP construct. The 35S:ThNAC13-GFP and the 35S:GFP (control) constructs were separately introduced into live onion epidermal cells using the particle bombardment method and visualized by a confocal laser scanning microscope (Wang et al., 2014a). The nuclei were stained with 4,6'-diamidino-2-phenylindole (DAPI, 10 μg.mL⁻¹) in phosphate-buffered saline for 3 min and analyzed.

Transcriptional Activation Assay

To study the transactivation activity of ThNAC13, the full coding region and a series of truncated ThNAC13 genes were cloned into the pGBK7 vector (Clontech, USA) to fuse with a GAL4 DNA binding domain. Fusion plasmids and the empty pGBK7 vector (negative control) were individually transformed into the yeast strain Y2HGold and grown on SD medium lacking tryptophan (negative control) were individually transformed into the yeast strain Y187. The effector construct was co-transformed together with each reporter construct into the yeast strain Y187. The DNA-protein interactions were evaluated according to the growth status of the transformants. The transcriptional activation activities were evaluated according to the growth status of the transformants.

Yeast One-Hybrid (Y1H) Assay

Two tandem copies of the NACRS with the core sequence “CGT[A/G]” and calmodulin-binding NAC (CBNAC) binding site with the core sequence “GCTT” were separately inserted into the pHIS2 vector (Clontech, USA) as the reporter construct. The coding region of ThNAC13 gene was cloned into the pGADT7-Rec2 vector as the effector construct. The effector construct was co-transformed together with each reporter construct into the yeast strain Y187. The DNA-protein interactions were evaluated according to the growth ability of the cotransformants on SD medium lacking leucine (Leu) and tryptophan (Trp; SD/-Leu/-Trp, DDO) and SD medium without leucine (Leu), tryptophan (Trp) and histidine (His; SD/-Leu/-Trp/-His, TDO) containing 3-AT (3-amino-1,2,4-triazole).

Generation of Transgenic Plants

The full coding sequence (CDS) of ThNAC13 was cloned into pROKII under the control of the CaMV 35S promoter (35S:ThNAC13) and transformed into A. thaliana according to the protocol described by Clough and Bent (1998). The positive transgenic lines were selected on kanamycin (50 mg/L) plates, and further identified by genomic DNA PCR, and the ThNAC13 expression level in leaves of each transgenic line was examined by quantitative reverse transcription polymerase chain reaction (qRT-PCR). The homozygous lines of T3 generation plants were used for study.

The sense and antisense sequences of the specific coding region of ThNAC13 with 259 bp in length were cloned into the RNAi vector (pFGC5941) to generate pFGC:ThNAC13 and silence the expression of ThNAC13. Transient transformation of 6-week-old T. hispida plants was performed according to Ji et al. (2014) with some modifications. Briefly, the whole seedlings were soaked in the transformation solution [1/2 MS + 150 μM acetylsyringone + 3% (w/v) sucrose + 0.01% (w/v) Tween-20, pH 5.8] with Agrobacterium tumefaciens EHA105 strain at 0.6 OD₆₀₀ and incubated with shaking at 100 r/min for 4 h at 25°C. Then, the seedlings were washed with distilled water twice and whipped with sterile facial tissue to remove excessive water. The whole seedlings were grown vertically on 1/2 MS agar medium [150 μM acetylsyringone + 2.5% (w/v) sucrose, pH 5.8] in tissue culture bottles. Three types of transient transgenic T. hispida plants were generated: T. hispida overexpressing ThNAC13 (transformed with 35S:ThNAC13, OE), T. hispida RNAi-silencing ThNAC13 (transformed with pFGC:ThNAC13, IE) and plants transformed with empty pROKII vector as a control (VC). The expression of ThNAC13 in the leaves of these transformed T. hispida plants was studied using qRT-PCR.

Analysis of Stress Tolerance

Two representative transgenic lines overexpressing ThNAC13 and wild type (WT) Arabidopsis plants were selected for abiotic stress tolerance assays. For germination rate assays, the seeds were surface-sterilized and sown on 1/2 MS solid medium supplemented with 100 mM NaCl, 200 mM mannitol or 2.5 μM ABA and incubated at 22°C. The rates of seeds were calculated to determine when the seedling opened green cotyledons. Next, 5-day-old seedlings were cultured vertically on 1/2 MS solid medium supplemented with 100 mM NaCl, 200 mM mannitol or 10 μM ABA. After 7 days of these treatments, the fresh weight and the length of primary roots of each plant were measured.

NBT and DAB Staining

NBT and DAB in situ staining for detection of hydrogen peroxide (H₂O₂) and superoxide (O₂⁻) were performed. The transformed seedlings of T. hispida were exposed to 150 mM NaCl, 200 mM mannitol or 30 μM ABA for 24 h. Four-week-old soil-grown Arabidopsis plants were treated with the same concentration of NaCl, mannitol or ABA for 2 h, and approximately 10 young branches harvested from T. hispida and 10 rosette leaves harvested from Arabidopsis were incubated with nitroblue tetrazolium (NBT) or 3,3’-diaminobenzidine (DAB) solutions for staining (Zhang et al., 2011).

Measurement of Physiological Changes

The transformed T. hispida plants grown on 1/2 MS solid medium supplemented with 150 mM NaCl, 200 mM mannitol or 30 μM ABA for 24 h were used to assess physiological changes. Four-week-old soil-grown Arabidopsis seedlings were treated with same concentration of NaCl, mannitol or ABA for 2 h (for H₂O₂ content measurement) and 48 h (for other physiological analyses). H₂O₂ levels were determined according to Thordal-Christensen et al. (1997). Malondialdehyde (MDA) contents, SOD and POD activities were measured according to Wang et al. (2010). Electrolyte leakage was determined following...
FIGURE 1 | Bioinformatics analysis of the ThNAC13 protein. (A) Phylogenetic tree was constructed from ThNAC13 and other different NAC proteins using MEGA 5.1 with 1000 bootstrap replicates. Bootstrap support values are indicated on each node. All protein sequences were retrieved from NCBI. Their corresponding accession numbers are as the follows: Tamarix hispida ThNAC13 (JQ974967); Oryza sativa SNAC2 (CBX55846), OsNAC5 (BAA89799), OsNAC6 (BAA89800), OsNAC052 (NP_001061889), and OsNAC063 (NP_001061900); Glycine max GmNAC3 (AAP44250), GmNAC4 (AAP44251), GmNAC11 (AAP44249), and GmNAC20 (AAP44252); Triticum aestivum TaNAC2 (AAU08786) and TaNAC69 (AAU08785); Arabidopsis thaliana ANAC019 (NP_175697.1), ANAC021 (NP_175697.1), and ANAC055 (NP_175697.1). (B) Schematic of putative conserved motifs among these NAC protein sequences was obtained using the MEME program. Different predicted motifs are represented by numbered boxes (annotations of individual motifs are listed in Supplementary Table S2).

The full CDS of ThNAC13 is 1122 bp in length and encodes a 373 amino acid protein with a molecular weight of 40.74 kDa. Multiple sequence alignments showed that ThNAC13 contains a highly conserved DNA-binding domain at N-terminal, which is a typical NAC domain N-terminal with five subdomains (A, B, C, D, and E), and a highly variable C-terminal transcriptional regulation domain (Supplementary Figure S1). The putative nuclear localization signal of ThNAC13 was identified in the D subdomain. Phylogenetic analysis revealed that ThNAC13 shares high sequence similarities with the plant stress-responsive NAC members (Figure 1), indicating that ThNAC13 may have similar functions to these NACs. Ten conserved motifs (termed motif 1–10) were predicted based on the sequence features of NAC proteins using the MEME program, which are listed in Supplementary Table S2. Motif 1 corresponds to the A and B subdomains, and motifs to be significant if $P < 0.05$. Asterisks indicates significant difference (*$P < 0.05$).

RESULTS

Bioinformatics Analysis of ThNAC13 from T. hispida
The full CDS of ThNAC13 is 1122 bp in length and encodes a 373 amino acid protein with a molecular weight of 40.74 kDa. Multiple sequence alignments showed that ThNAC13 contains a highly conserved DNA-binding domain at N-terminal, which is a typical NAC domain N-terminal with five subdomains (A, B, C, D, and E), and a highly variable C-terminal transcriptional regulation domain (Supplementary Figure S1). The putative nuclear localization signal of ThNAC13 was identified in the D subdomain. Phylogenetic analysis revealed that ThNAC13 shares high sequence similarities with the plant stress-responsive NAC members (Figure 1), indicating that ThNAC13 may have similar functions to these NACs. Ten conserved motifs (termed motif 1–10) were predicted based on the sequence features of NAC proteins using the MEME program, which are listed in Supplementary Table S2. Motif 1 corresponds to the A and B subdomains, and motifs...
FIGURE 2 | Nuclear localization and transcriptional activation of the ThNAC13 protein. (A) Nuclear localization of the ThNAC13 protein. Schematic of the GFP and ThNAC13-GFP fusion constructs. The GFP (I–III, control) and ThNAC13-GFP (IV–VI) fusion gene were expressed transiently in onion epidermal cells and observed with confocal microscopy. DAPI, DAPI for nuclear staining image; GFP, GFP green fluorescence image; Merge, the merged images of bright-field, GFP and DAPI staining. (B) Transcriptional activation activity analysis of ThNAC13 in yeast. Diagram showing a full-length (construct 2) and a series of deletion mutations of the ThNAC13 gene (constructs 3–14) fused to the downstream of the GAL4 DNA binding domain in the yeast vector pGBKT7. The pGBKT7 vector (control, constructs 1) and the fusion proteins (constructs 2–14) were expressed in yeast strain Y2HGold. The positive transformants were grown on SD/-Trp medium supplemented with or without X-α-Gal and applied to β-gal assay.

Subcellular Location and Transactivation Activity Assay

To determine the localization of the ThNAC13 protein, the 35S::GFP (control) and 35S::ThNAC13-GFP constructs were transformed into live onion epidermal cells. The GFP fluorescence of ThNAC13-GFP was exclusively observed in the nucleus, whereas the fluorescence of GFP was distributed in the whole cells (Figure 2A). These results clearly indicate that ThNAC13 is a nuclear protein.

To determine the transcriptional activation ability of ThNAC13, the full CDS and serial deletions of the ThNAC13 CDS were cloned into the pGBK77 vector (Figure 2B, construct 2–14). The different constructs were then transformed into the Y2HGold yeast cells and were assayed for their transcriptional activation. All transformants grew well on the SD/-Trp medium (left panel), indicating that these constructs had been successfully transformed into Y2HGold cells. On the SD/-Trp medium containing X-α-Gal (SD/-Trp/X), the cells harboring full-length CDS of ThNAC13 (construct 2, amino acid 1–373) or empty pGBK77 (construct 1, control) were not blue (right panel), suggesting that the full CDS of ThNAC13 does not have transcriptional activation ability. However, the yeast cells containing the C-terminal putative activation domain (construct 4, amino acid 166–373) of ThNAC13 grew well and were blue, whereas the N-terminal domain (construct 3, amino acid 1–165) was not blue, suggesting that only the C-terminal
FIGURE 3 | Identification of the NACRS and CBNACRS elements recognized by ThNAC13. (A) Oligonucleotides used in construction of the recombinant report vectors. (B) Scheme of the reporter and effector vector. (C) Analysis of the interactions between the ThNAC13 and NACRS/CBNACRS elements using the yeast one-hybrid. The reporter and effector constructs were co-transformed into the yeast strain Y187. The transformants were further identified by spotting serial dilutions (1, 10⁻¹ and 10⁻²) of yeast onto SD/-Leu/-Trp (DDO) and SD/-His/-Leu/-Trp (TDO) plates containing 3-AT (3-amino-1,2,4-triazole). P, positive control (pGAD-Rec2-53 + p53HIS2); N, negative control (pGAD-ThNAC13 + p53HIS2); R1–R6 and RM1–RM2, The positive transformants containing pGAD-ThNAC13 and pHIS2-NACRS/CBNACRS.

FIGURE 4 | Comparison of the germination rate between ThNAC13-transformed and WT Arabidopsis plants. (A) Seed germination assay between ThNAC13-transformed and WT Arabidopsis plants. The surface-sterilized seeds were sowed on 1/2 MS solid medium supplemented with 100 mM NaCl, 200 mM mannitol or 2 µM ABA and incubated at 22°C. (B) Germination assays of ThNAC13-transformed Arabidopsis plants. Data are represented as mean ± SD of at least three independent replications. Asterisk indicates significant difference (*P < 0.05) between transgenic lines and WT plants.

ThNAC13 Binds to NACRS and CBNACBS

To investigate whether ThNAC13 protein can bind to the NACRS and CBNACBS, two tandem copies of these motifs or their mutants were tested for interaction with ThNAC13 using a Y1H system. The sequences for construction of the recombinant report vectors are shown in Figure 3A, and the scheme of the reporter and effector vectors used in Y1H are shown in Figure 3B. On the SD/-Leu/-Trp medium, all transformants grew well (left panel). On the SD/-Leu/-Trp/-His medium containing 50 mM 3-AT, yeast cells carrying pGAD-ThNAC13/pHIS2-NACRS (R1–R6) grew well, while the mutants RM1 and RM2 and the negative control could not grow (Figure 3C). These results indicate...
that ThNAC13 can bind to both NACRS and CBNACBS but failed to bind to the mutants, which further suggest that ThNAC13 can specifically bind to the NACRS and CBNACBS motifs.

**ThNAC13 Increases the Tolerance to Salt and Osmotic Stresses**

Twelve lines of T3 homozygous ThNAC13-transformed Arabidopsis were generated by selection on medium containing kanamycin and further verified by genomic DNA PCR and qRT-PCR (Supplementary Figure S2). Two independent transgenic Arabidopsis lines (lines 7 and 12) with high transcript levels of ThNAC13 were selected for further study. We first evaluated the stress tolerance of transgenic and wild-type Arabidopsis seeds with seed germination tests. There was no difference in germination rates between WT and ThNAC13-transformed plants on 1/2 MS medium. However, the seed germination rates of ThNAC13-transformed lines were significant higher than those of WT plants under NaCl, mannitol or ABA stress (Figure 4), suggesting that overexpression of ThNAC13 improves salt and osmotic tolerances during the germination stage. Next, 5-day-old Arabidopsis seedlings were grown vertically on 1/2 MS solid medium supplemented with or without NaCl, mannitol or ABA. After growth for 7 days, the phenotype was similar between ThNAC13-transformed and WT plants under normal conditions (Figure 5), suggesting that overexpression of ThNAC13 did not influence the growth and phenotype of transgenic plants. Under NaCl, mannitol or ABA stress conditions, the root length and fresh weight gain were significantly increased in ThNAC13-transformed plants compared with WT plants, indicating that ThNAC13 improves the tolerances to NaCl, mannitol or ABA stress.

**FIGURE 5 | Comparison of root length and fresh weight between ThNAC13-transformed and WT Arabidopsis plants.** (A) The growth and phenotype of ThNAC13-transformed and WT Arabidopsis plants. (B) Effects of salt, drought stresses and ABA treatment on root length and fresh weight. Five-day-old seedlings were cultured vertically on 1/2 MS solid medium supplemented with 100 mM NaCl, 200 mM mannitol or 10 µM ABA for 7 days. The photographs were taken, and the root length and fresh weight of each plant were measured. Data are represented as the mean ± SD of at least three independent replications. Asterisk indicates significant difference (∗P < 0.05) between transgenic lines and WT plants.
FIGURE 6 | Histochemical staining and physiological analyses of ThNAC13-transformed and WT Arabidopsis plants. Rosette leaves from ThNAC13-transformed and WT Arabidopsis plants untreated or treated with 150 mM NaCl, 200 mM mannitol or 30 µM ABA for 2 h were used for (A) NBT and (B) DAB staining. (C–I) Physiological analyses among the Arabidopsis plants. Four-week-old seedlings of ThNAC13-transformed and WT Arabidopsis plants were irrigated with 100 mM NaCl, 200 mM mannitol or 30 µM ABA for 2 h (for H$_2$O$_2$ measurement) and for 48 h (to measure MDA content, SOD and POD activities, electrolyte leakage rate, chlorophyll and proline contents). Data are represented as the mean ± SD of at least three independent replications. Asterisk indicates significant difference (*P < 0.05) between transgenic lines and WT plants.

FIGURE 7 | Histochemical staining and physiological analyses in transgenic T. hispida plants. (A) NBT staining for detecting O$_2^-$; (B) DAB staining for detecting H$_2$O$_2$. (C–I) Physiological analyses among the different transgenic T. hispida plants. (C) H$_2$O$_2$ contents; (D) MDA contents; (E) SOD activity; (F) POD activity; (G) electrolyte leakage; (H) chlorophyll contents; (I) proline contents. The T. hispida plants were grown on 1/2 MS medium or 1/2 MS medium containing 150 mM NaCl, 200 mM mannitol or 30 µM ABA for 24 h and used for analysis. VC, the empty pROKII-transformed T. hispida plants; OE, transiently overexpressing ThNAC13 in T. hispida plants; IE, ThNAC13 RNAi-silenced T. hispida plants transiently. Data are represented as the mean ± SD of at least three independent replications. Asterisk indicates significant difference (*P < 0.05) between transgenic (OE and IE) and control (VC) plants.
**Generation of the Transgenic *T. hispida* Plants with Overexpression and RNAi-Silencing of ThNAC13**

For further investigation of the function of ThNAC13 using gain- and loss-of-function methods, *T. hispida* plants transiently overexpressing ThNAC13 (OE), RNAi-silenced for ThNAC13 (IE) and controls (transformed with empty pROKII, VC) were generated. The expression of ThNAC13 in these 3 transformed *T. hispida* plants under different stresses was studied using qRT-PCR. The transcript levels of ThNAC13 were significantly increased in OE plants and substantially decreased in IE plants compared with control (VC) plants under all the conditions, especially with salt, mannitol or ABA stress for 24 h (Supplementary Figure S3). These results indicate that the genetically transformed *T. hispida* plants are suitable for gain- and loss-of-function studies.

**ThNAC13 Mediates ROS Scavenging Capability**

Abiotic stress can induce oxidative stress through generation of reactive oxygen species (ROS), and the accumulation of ROS damages the cell membrane by oxidizing proteins, lipids and DNA (Mittler et al., 2004). NBT and DAB staining were first used for detecting the cellular levels of O$_2^.$ and H$_2$O$_2$ in three types of transiently transformed *T. hispida* plants. The results showed that the transgenic *Tamarix* plants overexpressing ThNAC13 (OE) had the lowest O$_2^.$ and H$_2$O$_2$ accumulation, and the transgenic *Tamarix* plants with knockdown of ThNAC13 (IE) had the highest O$_2^.$ and H$_2$O$_2$ accumulation, although they had similar O$_2^.$ and H$_2$O$_2$ levels under normal conditions (Figures 6A,B). To confirm the results from *T. hispida*, we further determined the ROS accumulation in Arabidopsis plants using *in situ* histochemical staining with NBT and DAB. There was no obvious difference in NBT and DAB staining between ThNAC13-transformed and WT plants under normal conditions. However, compared with WT plants, the transgenic plants showed substantially reduced O$_2^.$ and H$_2$O$_2$ accumulation under salt, mannitol and ABA conditions (Figures 7A,B).

Additionally, we further measured H$_2$O$_2$ and MDA contents in transgenic *T. hispida* plants. There was no difference in H$_2$O$_2$ and MDA contents among OE, VC, and IE plants under normal conditions. However, under stress conditions, IE plants had the highest level, followed by IE plants, and OE had the lowest H$_2$O$_2$ levels (Figures 6C,D). Consistent with these results, transgenic Arabidopsis plants overexpressing ThNAC13 displayed low H$_2$O$_2$ and MDA contents compared with WT plants under stress conditions (Figures 7C,D). As ROS levels were decreased by expression of ThNAC13, we investigated whether SOD and POD activities were regulated by ThNAC13. Among all the studied *Tamarix* plants, there was no significant difference in SOD and POD activities under normal conditions. The SOD and POD activities were also markedly elevated in OE plants and reduced in IE plants compared with VC plants (Figures 6E,F). We further studied SOD and POD activities in Arabidopsis. There was no significant difference in the activities of SOD and POD between transgenic and WT plants under normal conditions. Compared with WT Arabidopsis plants, the activities of POD and SOD in ThNAC13-transformed Arabidopsis plants were significantly increased under the different stress conditions (Figures 7E,F), which was consistent with the results obtained from *T. hispida* plants. Therefore, overexpression of ThNAC13 significantly activates SOD and POD activities under abiotic stress conditions.

**Analysis of Electrolyte Leakage Rate and the Contents of Chlorophyll and Proline**

To estimate the level of abiotic stress damage to the studied plants, several stress-related physiological changes were measured. Under normal conditions, all transgenic *T. hispida* plants displayed similar electrolyte leakage rates, chlorophyll and proline contents (Figures 6G–I). However, the electrolytic leakage rates in IE plants were the highest, followed by VE, and were the lowest in OE plants (Figure 6G). Meanwhile, OE plants had the highest contents of chlorophyll and proline, followed by VC plants, and IE plants had the lowest contents of chlorophyll and proline (Figures 6H,I). The electrolyte leakage rates in transgenic Arabidopsis plants overexpressing ThNAC13 were decreased compared with WT plants under salt, osmotic or ABA stress (Figure 7G). Furthermore, the contents of chlorophyll and proline in ThNAC13-transformed Arabidopsis were both significantly higher than those in WT plants (Figures 7H,I). These results are consistent with the results from *T. hispida*, indicating the reliability of these findings.

**ThNAC13 Induces the Expression of SODs and PODs**

As the activities of POD and SOD were both markedly elevated in the Arabidopsis plants overexpressing ThNAC13, we further analyzed the transcription levels of POD and SOD genes, which were shown to have enzymatic activities in previous studies. The expression of many studied genes was highly increased in ThNAC13-transformed Arabidopsis under NaCl and mannitol treatments (Figure 8). These results suggested that ThNAC13 can induce the transcription levels of SOD and POD genes, which will enhance the activities of SOD and POD.

**DISCUSSION**

**ThNAC13 Is a Member of the NAC Transcription Factors**

Most NAC proteins share a well-conserved N-terminal DNA binding domain and a diversified C-terminal domain (Puranik et al., 2012; Wang et al., 2014b). In this study, we found that the ThNAC13 protein has the conserved NAC domain at N-terminus. Multiple sequence alignments showed that ThNAC13 has high amino acid sequence identity with the reported stress-responsive NAC proteins of Arabidopsis (AtNAC019, AtNAC055, and AtNAC072) (Supplementary Figure S1), suggesting that ThNAC13 may also be a stress-responsive NAC. Previous studies have shown that NAC proteins could bind to the NACRS containing the core
FIGURE 8 | Analysis of the expression of the POD and SOD genes in ThNAC13-transformed Arabidopsis plants. Four-week-old seedlings of ThNAC13-transformed and WT Arabidopsis plants were treated with 100 mM NaCl or 200 mM mannitol for 24 h, and were harvested for qRT-PCR analysis. The transcription levels of SODs or PODs in WT plants under the same condition were set as a calibrator to 1 to calculate their expression in transgenic plants. The error bars were calculated from three replicates. The Tair locus numbers of Arabidopsis PODs and SODs are shown in Supplementary Table S1.

sequence “CGT[A/G]” (Duval et al., 2002; Tran et al., 2004; Olsen et al., 2005; Xue, 2005; Takasaki et al., 2010; Hao et al., 2011). However, Kim et al. (2007) found that the Arabidopsis transcriptional repressor CBNAC can bind to CBNACBS with a “GCTT” sequence but not the “CGT[A/G]” sequence, indicating that different NAC transcription factors may bind to different element sequences. Our study suggested that ThNAC13 can bind to the NACRS element; at the same time, it can also bind to a motif sequence containing the “GCTT,” similar to CBNAC (Figure 3C). These results suggested that ThNAC13 could bind to NACRS and CBNACBS to regulate gene expression.

Transactivation assays showed that the C-terminal of ThNAC13 has a high transactivation ability, although the full-length protein does not have this capacity (Figure 2B). However, when the N-terminal of ThNAC13 is present, the transactivation is abolished. These results indicated that the N-terminal of ThNAC13 may have a transcriptional repression domain, which leads to abolished transactivation. Previous studies also support our hypothesis. For example, the conserved hydrophobic LVFY motif was a transcription repression domain that is present in the N-terminal of GmNAC20 protein (Hao et al., 2010). We also found the LVFY motif in N-terminal of ThNAC13 (Supplementary Table S2). However, ThNAC13 could regulate the expression of POD and SOD genes (Figure 8). This is probably due to its C-terminal domain, which has high transcriptional activation activity, or the interplay among the C-terminal activation domain, the DNA-binding domain and NARD (NAC Repression Domain). Therefore, ThNAC13 may act as a transcriptional repressor or activator, depending on interactions with other transcription factors or conformational changes.

**ThNAC13 Improves Tolerance to Salt, Osmotic, and ABA Stresses**

Recently, numerous reports have shown that NAC transcription factors play important roles in the response to different abiotic stresses and function as positive stress response TFs (Jeong et al., 2013; Mao et al., 2014; Fang et al., 2015). In the present study, our results showed that ThNAC13 can induce a series of physiological changes to improve tolerance to salt, osmotic and ABA stresses (Figures 4, 6, 7). These results suggested that ThNAC13 functions as a stress-responsive protein and enhances the salt and osmotic tolerance. In addition, ThNAC13 promotes tolerance to ABA, and NADPH oxidases are involved in abiotic stress tolerance and regulated by ABA, which plays an important role in ABA-mediated stomatal closure (Acharya et al., 2013). Therefore, there may be a relationship between ThNAC13 and NADPH oxidase activity, which deserves further study.

**ThNAC13 Improves ROS-Scavenging Capability**

Plants are commonly exposed to various adverse environments, which result in accumulation of ROS, damage to protein synthesis and stability, cellular macromolecules and membrane lipids in plant cells and generation of oxidative stress (Wang et al., 2005). Therefore, ROS-scavenging is important for plant tolerance. Two prominent ROS species, $O_2^{•−}$ and $H_2O_2$, are considered to be important signaling molecules in plant cells. In this study, biochemical staining (NBT and DAB) showed that the ROS accumulation ($O_2^{•−}$ and $H_2O_2$) in transgenic Tamarix and Arabidopsis plants overexpressing ThNAC13 was lower than that in control (VC or WT) plants under abiotic stress and ABA treatment (Figures 6A,B, 7A,B). MDA, the product of
lipid peroxidation caused by ROS, is also used to evaluate ROS-mediated injuries in plants (Moore and Roberts, 1998). We found that the H$_2$O$_2$ and MDA contents were both decreased by expression of ThNAC13 under abiotic stress and ABA treatment (Figures 6C,D, 7C,D). These results suggested that ThNAC13 can reduce ROS accumulation and decrease lipid peroxidation to improve tolerance to salt and osmotic stresses.

The antioxidant enzymes, SOD and POD, are known to play important roles in ROS scavenging, which influences the cellular ROS levels. We found that the activities of SOD and POD were significantly induced by ThNAC13 under different stress conditions. Furthermore, most of the studied SOD and POD genes were activated in ThNAC13 transgenic Arabidopsis plants, which may lead to increased SOD and POD activities in plants overexpressing ThNAC13 (Figures 6E,F, 7E,F, 8). These results suggest that ThNAC13 improves stress tolerance at least partly through enhancing ROS-scavenging capability, which will decrease the degree of membrane lipid peroxidation. In addition, we found that the core sequences of NACRS and CBNACBS elements were both enriched in the promoters (<1.5 kb) of the studied SODs and PODs (Supplementary Table S3), suggesting that ThNAC13 may bind to these elements to directly regulate the expression of these genes. These results suggest that ThNAC13 improves tolerance to salt and osmotic stresses by regulating the transcript levels of SODs and PODs to increase the ROS-scavenging capability.

**ThNAC13 Induces Proline Biosynthesis**

Plant proline could protect cells by serving as an osmotic agent and also as a radical scavenger under biotic or abiotic stress. In addition, when stress is relieved, the accumulated proline could be degraded as a supply of energy for plant growth (Kavi Kishor and Sreenivasulu, 2014). In this study, proline content could be degraded as a supply of energy for plant growth (Kavi Kishor and Sreenivasulu, 2014). In this study, proline content was positively correlated with the transcripts of ThNAC13 (Figure 6I), and Arabidopsis plants overexpressing ThNAC13 displayed increased proline contents under different stresses (Figure 7I), indicating that ThNAC13 induces the biosynthesis of proline, and the increased proline enhances both osmotic potential and ROS-scavenging capability to improve abiotic stress tolerance.

**CONCLUSION**

A stress-responsive NAC gene from *T. hispida*, ThNAC13, was identified and functionally characterized. ThNAC13 functions as a transcription factor that improves salt and osmotic tolerance by increasing ROS-scavenging capability and osmotic potential. This study improves our understanding of the biological functions of NAC proteins in response to abiotic stress in woody plants.

**AUTHOR CONTRIBUTIONS**

LW and ZL carried out all the experiments and data analysis. LW, ZL, and YW conceived the project, designed the experiments and drafted the manuscript. ML and YW supervised the analysis and critically revised the manuscript. All authors read and approved the final manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: http://journal.frontiersin.org/article/10.3389/fpls.2017.00635/full#supplementary-material

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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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