A 2-Cys peroxiredoxin gene from *Tamarix hispida* improved salt stress tolerance in plants

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**Abstract**

**Background:** Peroxiredoxins (Prxs) are a large family of antioxidant enzymes that respond to biotic and abiotic stress by decomposing reactive oxygen species (ROS). In this study, the stress tolerance function of the *Th2CysPrx* gene was further analysed. It lays a foundation for further studies on the salt tolerance molecular mechanism of *T. hispida* and improved salt tolerance via transgenic plants.

**Results:** In this study, the stress tolerance function of the *Th2CysPrx* gene was further analysed. The results of transgenic tobacco showed higher seed germination rates, root lengths, and fresh weight under salt stress than wild-type tobacco. Simultaneously, physiological indicators of transgenic tobacco and *T. hispida* showed that *Th2CysPrx* improved the activities of antioxidant enzymes and enhanced ROS removal ability to decrease cellular damage under salt stress. Moreover, *Th2CysPrx* improved the expression levels of four antioxidant genes (*ThGSTZ1*, *ThGPX*, *ThSOD* and *ThPOD*).

**Conclusions:** Overall, these results suggested that *Th2CysPrx* enhanced the salt tolerance of the transgenic plants. These findings lay a foundation for further studies on the salt tolerance molecular mechanism of *T. hispida* and improved salt tolerance via transgenic plants.

**Keywords:** Salt stress, *Tamarix hispida*, Th2CysPrx

**Background**

Abiotic stresses, such as drought, salinity, and extreme temperature, among others, negatively affect the growth and yield of plants, resulting in very large economic losses. Among the harmful environmental stresses, salt stress in particular leads to slower plant growth and declines in cultivated plant production [1]. In the long-term evolutionary process, some plants, such as *Atriplex canescens* [2], *Halostachys caspica* [3], *Suaeda salsa* [4], and *Salicornia brachiata* [5], among others, have gradually adapted to a salt stress environment and can grow in dry and saline land.

*Tamarix hispida* is a typical woody halophyte that can form natural forests in saline alkali soil with 1% salt content. In addition, it can also endure drought stress, which makes it an ideal material to clone genes related to drought and salt tolerance and to study the salt tolerance mechanism of woody halophytes [6]. Many previous studies have been conducted to examine the mechanism of salt and drought tolerance and the function of stress resistance genes of *T. hispida*. For example, *ThelF1A, ThDREB, ThZFP1, ThGSTZ1* and *ThPOD3* in *T. hispida* increase salt and drought tolerance by regulating superoxide dismutase (SOD) and peroxidase (POD) activities to reduce reactive oxygen species (ROS) accumulation [7–11]. These results all suggest that scavenging of ROS plays an important role in the response of *T. hispida* to salt stress.
POD, SOD, CAT (Catalase), GPX (Glutathione peroxidase) and GST (Glutathione S-transferases) are important ROS-scavenging genes. Gao et al. [12] found that ThGSTZI improved tolerance to abscisic acid ABA and methyl viologen (MV) stress by augmenting the activities and expression levels of ThGPX, ThSOD, and ThPOD and the ROS-scavenging capacity. The NtSOD, NtAPX, NtCAT, NtPOX, and NtGST genes play roles in eliminating ROS and increasing stress tolerance in plant under stress. Fortunately, overexpressed wheat TaWRKY44 was found to activate the above five genes [13]. In Betula platyphylla, BplMYB46 could increase the ROS-scavenging capacity and proline content to improve salt and osmotic tolerance by affecting the expression of genes, including POD, SOD and P5CS (Δ1-pyrroline-5-carboxylate synthetase) [14]. Zhang et al. [15] found that under NaCl stress, ALA in tomatoes increased the activity of ROS-scavenging antioxidant enzymes and the expression of the SOD, APX and POD genes encoding these enzymes.

Peroxiredoxins (Prxs) are a family of non-haem peroxidases that are widely present in animals, plants and microorganisms. The biological function of these peroxidases is to regulate the balance of ROS and intracellular signal transduction through hydrogen peroxide (H₂O₂), alkyl hydroperoxides, and peroxynitrite [16–19]. In plants, based on the number and position of conserved Cys residues, Prxs are grouped into four classes: 1CysPrx, 2CysPrx, type-II Prx and PrxQ. The 1CysPrxs have only one conserved Cys residue, and PrxQ contains two cysteine residues that are catalytically active and linked by intramolecular disulphide bonds [20]. Both 2-Cys Prx and type-II Prx have two conserved Cys residues, but the difference between them is that 2-Cys Prx is a stromal protein [21], and Prx-II has various isoforms [22].

Prxs operate in a particular way during plant growth, development and stress tolerance. For example, Kim et al. [23] found that 2CysPrx could eliminate H₂O₂ by participating in an alternative water-water cycle and protect photosynthetic structures against oxidative damage under environmental restriction. Pea chloroplast 2CysPrx and mitochondrial Prx IIF affect the structure and peroxidase activity of photosynthetic structures [24]. Kim et al. [25] found that 2CysPrx from Oryza sativa could increase tolerance to ROS-induced oxidative stress by improving cellular redox homeostasis. Overexpression of 2CysPrx in tall fescue plants increases resistance to oxidative stress and antioxidant activity [26]. Prx also detoxifies ROS and modulates signalling responses [27]. Interestingly, there is no direct evidence that the Prxs gene of T. hispida is involved in ROS scavenging and the abiotic stress response. In a previous study, Gao et al. [28] cloned four Prxs genes from T. hispida. Real-time quantitative PCR (qRT-PCR) analysis indicated that these genes could respond to several abiotic stresses and ABA application. However, Th2CysPrx displayed a unique expression pattern under the studied stress conditions. In the present study, the role of Th2CysPrx in salt stress was further demonstrated, and elucidated the molecular mechanism of this gene under salt stress. It also provided potential application prospects for molecular breeding to improve salt tolerance.

### Results

#### Overexpression of Th2CysPrx improves salt stress tolerance in transgenic tobacco

To study the biological role of Th2CysPrx in the salt stress response, the 14 resistant lines were obtained. The qRT-PCR results showed that Th2CysPrx gene were overexpressed in all 14 lines (Additional file 1: Figure S1). Two representative T₃ homzygous transgenic lines (Line 7 and Line 11) were randomly selected for further salt tolerance analysis. The germination and growth of transgenic and wild-type (WT) tobacco were compared during exposure to normal and 125 mM NaCl stress conditions. The results showed that, under normal conditions, there was no obvious difference in the germination rates and seedling growth between transgenic and WT plants (Fig. 1a-c). Under salt stress, however, the germination rate and seedling growth of transgenic plants was significantly better than those of WT plants. Under NaCl stress, the germination rates of transgenic lines were 89.7% (Line 7) and 84.5% (Line 11), while that of WT plants was only 53.5% (Fig. 1d). The chlorophyll content of the transgenic lines under NaCl treatment was 1.54- and 1.68-fold greater, respectively, than that of the WT plants (Fig. 1e). The average fresh weight of the transgenic lines under NaCl treatment was 2.15- and 2.10-fold greater, respectively, than that of the WT plants (Fig. 1f). The average root length of transgenic lines under NaCl treatment was 1.99- and 1.87-fold greater, respectively, than that of WT plants (Fig. 1g). It was apparent that salt stress significantly inhibited the growth of transgenic and WT plants. However, the growth of transgenic plants was significantly superior to that of WT plants. These results indicated that the expression of Th2CysPrx in tobacco could significantly increase the salt tolerance of the plants.

#### The Th2CysPrx gene significantly improves ROS scavenging and reduces cell damage in transgenic tobacco

The H₂O₂ and O₂⁻ concentration in the transgenic lines and WT plants were examined by 3,3′-diaminobenzidine (DAB) and nitro blue tetrazolium (NBT) staining, respectively. The results showed that under normal condition, there was no significant variation in the ROS generated by the three lines. In contrast, under salt
stress, the WT plants exhibited deeper staining, suggesting much greater ROS accumulation in the WT than the transgenic lines under stress conditions (Fig. 2a). 2,7-dichlorofluorescin diacetate (H$_2$DCF-DA) staining of the leaves of the transgenic and WT lines was also performed after 1 and 2 h of salt stress, respectively. There were no obvious differences in ROS levels in intact guard cells under normal conditions. However, after salt stress, the WT plants showed increased ROS accumulation in guard cells compared with the transgenic lines (Fig. 2b). The transgenic lines exhibited lower ROS content than the WT plants. These results indicated that Th2CysPrx led to a significant reduction of ROS accumulation in plant cells under salt stress.

Furthermore, SOD and POD activities in the transgenic and WT plant lines were measured under salt stress. The results revealed no significant differences between two independent transgenic lines and WT plants under normal conditions. In contrast, SOD and POD activities were higher in the two transgenic lines than in
the WT plants under NaCl stress. Specifically, the SOD activities of the transgenic lines were 1.17- and 1.10-fold higher than that of WT (Fig. 3a). And the POD activity values in the transgenic lines were 1.40- and 1.46-fold higher, respectively, than those in the WT plants (Fig. 3b).

Similarly, there were no obvious differences between the transgenic and WT plant lines based on Evans blue staining, malondialdehyde (MDA) contents and electrolyte leakage (EL) rates under normal conditions. Under salt stress conditions, deeper Evans blue staining was observed, and the MDA content and electrolyte leakage were significantly increased. However, Evans blue staining was weaker and the MDA content and EL rate were lower in the transgenic than in the WT plants (Fig. 3b).

Taken together, these results indicated that Th2CysPrx in tobacco improved salt stress tolerance by increasing ROS scavenging and preventing cell damage to maintain better plant growth. The transgenic lines exhibited lower ROS content and less damage than the WT plants, suggesting that Th2CysPrx directly affected ROS scavenging and cell protection during NaCl treatment.

**Use of transient overexpression of Th2CysPrx in T. hispida to further evaluate the results in transgenic tobacco**

To confirm the results of heterologous expression in tobacco, Th2CysPrx was transiently transferred into T. hispida. qRT-PCR analysis showed that the expression of the Th2CysPrx gene in 35S::Th2cys overexpression T. hispida was 12.38-fold that of the control, indicating that the transient overexpression line of T. hispida was successfully obtained (Fig. 4a). Then, biochemical staining and related physiological indexes were analysed and compared between 35S::Th2cys and CK plants. The DAB and NBT staining results showed no significant difference between 35S::Th2cys and CK plants prior to salt stress. However, CK plants showed darker staining than 35S::Th2cys plants after salt stress, with darker staining after 2 h than 1 h (Fig. 4b).
Similarly, the SOD and POD activity levels also showed no difference between 35S::Th2cys and CK plants under normal conditions. Under salt stress, SOD activity in CK plants was 80.69% of that in 35S::Th2cys plants (Fig. 4c), and POD activity in CK plants was 70.51% of that in 35S::Th2cys plants (Fig. 4d). Under normal conditions, there was no significant difference in terms of MDA or electrolyte leakage content between 35S::Th2cys and CK plants. However, 35S::Th2cys plants showed significantly lower MDA level and electrolyte leakage than CK plants, and the MDA content and electrolyte leakage in 35S::Th2cys plants was 74.85 and 67.70% of that in CK plants (Fig. 4e, f). Altogether, these results showed that Th2CysPrx conferred salt tolerance to the transgenic tobacco and T. hispida plants.

Additionally, the expression levels of four antioxidant genes (ThGSTZ1, ThGPX, ThSOD and ThPOD) in transiently transfected Th2CysPrx T. hispida were analysed by qRT-PCR. The results showed that these genes shared similar expression profiles with Th2CysPrx in transiently transfected T. hispida seedlings, all of which were upregulated after NaCl stress. The expression levels of these genes were 1.22, 1.55, 1.50, and 1.86-fold that of CK plants, respectively (Fig. 5). These results indicated that Th2CysPrx overexpression affected ThGSTZ1, ThGPX, ThSOD and ThPOD gene expression. Thus, these results indicated that the expression of Th2CysPrx altered the expression of other stress-related genes, suggesting that salt stress tolerance regulation involves a complex network.

**Discussion**

Under adversity, plants usually accumulate ROS, which leads to damage to protein synthesis and stability to produce cellular macromolecules and membrane lipids and the generation of oxidative stress [29]. The 2CysPrx plays an important role in scavenging ROS and regulating signal transduction, acting as a molecular chaperone and DNA damage response. In previous studies, 2CysPrx served as a barrier for H$_2$O$_2$. Rapeseed 2CysPrx activates chloroplast fructose-1,6-bisphosphatase (FBPase) and participates in the Calvin cycle [30]. Kim et al. [31] showed that *Arabidopsis* 2CysPrx can protect citrate synthase (CS) from heat-induced aggregation and
function as a molecular chaperone. Banerjee et al. [32] found that overexpression of cyanobacterium *Anabaena* 2CysPrx could decrease intracellular ROS levels under abiotic stress to reduce cell damage. Overexpression of 2CysPrx from mungbean can efficiently eliminate cellular ROS to improve tolerance to MV stress in *Arabidopsis* [33]. However, in the process of preventing DNA damage, some organisms use the ROS pathway, and some plants protect the cells from oxidative damage through the non-ROS pathway. For example, 2CysPrx-induced tobacco gene silencing largely affects the regeneration of ascorbic acid and thus the biosynthesis of abscisic acid (ABA) [17]. Above researches showed the 2CysPrx has different functions, most of which are research on glycogen plants, but little exploration in woody halophytes. Therefore, we need to explore whether *Th2CysPrx* prevents cell damage through the ROS pathway in woody halophyte *T.hispida*.

Vidigal et al. [17] demonstrated that 2CysPrx was the key to H2O2 clearance and regulated ABA signalling downstream of ROS genes. Many studies have shown that ABA has important biological functions and is an important signalling molecule under abiotic stresses [34–36]. The expression of genes that participate in the ABA signalling pathway can increase plant abiotic tolerance [37, 38]. In this study, compared with WT plants, transgenic *Th2CysPrx* tobacco exhibited increased germination rates, root lengths and chlorophyll contents under salt stress (Fig. 1). Overexpression of *Th2CysPrx* resulted in better SOD and POD activities, and reduced ROS accumulation. Additionally, the expression levels of the *ThGSTZ1*, *ThGPX*, *ThSOD* and *ThPOD* genes were markedly upregulated in transgenic *T. hispida* under salt stress, which indicated that the *Th2CysPrx* gene could improve salt tolerance by increasing antioxidant enzyme and strengthening ROS scavenging activities, leading to reduced ROS accumulation.
We also found that the activities and expression levels of SOD and POD in CK and transgenic *T. hispida* were not consistent under normal condition. However, the overall trend between the activities and expression levels is consistent under NaCl stress. The results showed the studied ThSOD and ThPOD gene may be key effective gene among SOD and POD family genes responding to NaCl stress (Figs. 2, 3, 4, 5).

It remains unclear whether the Th2CysPrx gene improves plant ROS scavenging by activating the expression of ABA signalling pathway genes. So, in future studies, we will further analyze the salt tolerance mechanism of the Th2CysPrx gene in *T. hispida*, while comparing the differences in abiotic stress in plants between 2CysPrx in tobacco and *T. hispida*.

**Conclusion**

Prxs have an important place in plant growth and development, as well as in responses to stress. However, there is no direct evidence to demonstrate that the Prxs gene of *T. hispida* is involved in ROS scavenging and abiotic stress responses. In the present study, a Th2CysPrx was isolated from *T. hispida*, and transgenic tobacco and *T. hispida* showed advantages with respect to morphological, physiological, and biochemical traits under salt stress. Additionally, the expression levels of four antioxidant genes (*ThGSTZ1*, *ThGPX*, *ThSOD* and *ThPOD*) were significantly higher in overexpressed Th2CysPrx of *T. hispida* than in the control. Altogether, the results indicated that Th2CysPrx increased salt tolerance via increasing the expression and activity of antioxidant enzymes and improved ROS scavenging ability. Future studies should assess whether Th2CysPrx participates in the ABA signalling pathway.

**Methods**

**Plant materials and growth conditions**

Tobacco seeds (Longjiang 911) were obtained from the Tobacco Science Research Institute of Heilongjiang Province [39]. And kept in our laboratory. Tobacco seeds stored at 4 °C for 3–5 days were sterilized with 70% (w:v) ethanol for 1 min followed by 3% sodium hypochlorite for 5 min, rinsed eight times with sterile water, and sown onto plates containing half-strength.

![Fig. 5](image-url) The relative expression levels of *ThGSTZ1*, *ThGPX*, *ThSOD*, *ThPOD* in WT and 35::Th2CysPrx exposed to NaCl stress. a The relative expression levels of *ThGSTZ1*. b The relative expression levels of *ThGPX*. c The relative expression levels of *ThSOD*. d The relative expression levels of *ThPOD*. CK: *T. hispida* plants transformed with empty pROKII; 35::Th2cys: *T. hispida* overexpressing Th2CysPrx; The error bars were calculated from three independent replicates of the qRT-PCR.
Murashige and Skoog (1/2 MS) medium. They were then placed in a cabinet with a controlled environment (16 h light: 8 h dark) at 22 °C. One week later, the seedlings were transplanted to pots containing a mixture of perlite and soil (1:3 v/v) and grown in a greenhouse (14 h light: 10 h dark) at 24 °C with 70% relative humidity.

_T. hispida_ seeds (the Turpan Desert Botanical Garden, Xinjiang, 293 China) was planted in a greenhouse in Harbin (China). Seeds for propagation of plant material were harvested from these _T. hispida_ plants, and planted on 1/2 MS medium and grown in a greenhouse (14 h light: 10 h dark) at 24 °C with 70% relative humidity and a photon flux density of 250 μE m⁻² s⁻¹.

### Generation of transgenic plants

The ORF of _Th2CysPrx_ was amplified and cloned into pROKII vector (referred to as 35S::Th2cys). The primers are shown in Table 1. The 35S::Th2cys was introduced into _Agrobacterium tumefaciens_ EHA105 by electroporation, and the transgenic tobacco was further obtained by the agrobacterium-mediated leaf disc transformation method [40]. Fifteen lines were generated in the T₀ generation. The two T₃ homozygous transgenic lines (Line 7 and Line 11) were randomly selected for further analysis.

Concurrently, _Th2CysPrx_ was also transiently overexpressed in _T. hispida_. Specifically, it was transiently transformed into one-month-old _T. hispida_ seedlings with 35S::Th2cys to overexpress _Th2CysPrx_ and with empty pROKII plasmid (used as a control, CK) according to Zheng et al. [41] with some modifications. In particular, single colonies of _A. tumefaciens_ strain EHA105 harbouring 35S::Th2cys or empty pROKII were cultured to an OD₆₀₀ = 0.8, and the cells were harvested by centrifugation. The transformants used [1/2 MS + 3% (w/v) sucrose + 150 μM acetosyringone + 0.01% (w/v) Tween 20] were adjusted to an OD₆₀₀ = 0.8. _T. hispida_ was immersed in transformation solution and incubated at 25 °C for 4 h. Subsequently, 2% sucrose was used to quickly wash the seedlings once for 1 min, after which they were planted vertically on 1/2 MS solid medium for 12, 24, 36 or 48 h. Total RNA was isolated from every samples using the CTAB method [42], and 1 mg of RNA was reverse transcribed with the PrimeScript™ RT reagent Kit (TaKaRa, China). The resulting cDNA product was diluted to 10x and used as a template for the qRT-PCR analyses. Real-time RT-PCR was conducted using a Bio-Rad (MJ) Opticon™ System (Bio-Rad, Hercules, USA) Actin (FJ618517), α-tubulin (FJ618518), and β-tubulin (FJ618519) were used as internal controls to normalize the amount of total RNA present in each reaction. The primers used are listed in Table 1. The PCR conditions were 94 °C for 30 s, 45 cycles of 94 °C for 12 s, 58 °C for 30 s, 72 °C for 40 s, and 80 °C for 1 s for plate reading. A melting curve was generated for each sample at the end of each run to assess the purity of the amplified products. To determine the expression of _Th2CysPrx_ and several antioxidant genes. The primers are listed in Table 1. The reaction system and procedure for qRT-PCR were performed according to Gao et al. [28]. The 2⁻ΔΔCt method was used to calculate the relative gene expression [43].

### Analysis of stress tolerance

The T₃ generation seeds of _Th2CysPrx_ transgenic tobacco were surface sterilized and sown on 1/2 MS agar medium or 1/2 MS with 125 mM NaCl. Germination was recorded after 10 d. In addition, 3-day-old seedlings sown on 1/2 MS were transferred to 1/2 MS medium or 1/2 MS with 125 mM NaCl for 2 weeks to compare the fresh weight and root length between transgenic and wild type (WT) lines.

### Physiological analysis

Seven-day-old tobacco seedlings sown on 1/2 MS were transferred to a mixture of perlite and soil (1:3 v/v). After 6 weeks, the tobacco seedling roots were watered with the solution of 200 mM NaCl. Simultaneously, the seedlings were watered with water as a control. After 7 d, the leaves of stressed and control tobacco were harvested.

One-month-old seedlings of the transient transgenic _T. hispida_ seedlings (35S::Th2cys and CK) exposed to 150 mM NaCl for 12 h were harvested, and the physiological index was measured, respectively. The SOD, POD activities and MDA contents were measured following the method of Wang et al. [44]. The EL was measured according to Ben-Amor et al. [45]. The chlorophyll contents were measured following the method of Lichtenthaler [46].

### Detection of ROS and cell death

The transgenic and WT/control young leaves were collected after 0, 1 or 2 h of NaCl treatment, and histochemical staining analysis was carried out immediately.

### Table 1 List of primers and their applications

| Gene symbol | Forward Primers (5’→3’) |
|-------------|-------------------------|
| pROKII-Th2CysPrx-F | ATCG TCTAGAATGGCGTGCCGACGCCAACT |
| pROKII-Th2CysPrx-R | AGCTGAAGCTCTAAATTGCAAGGAGTACTC |
| Th2CysPrx-F | TGAGATCAGTTCCTAGT |
| Th2CysPrx-R | TGATAACCAATCCTTGAG |
| Actin-F | AAACAAATGCGTGATGCTG |
| Actin-R | ACAATACCGTGCTCATAGG |
| a-tubulin-F | CACCACCGTGTCCTCAG |
| a-tubulin-R | ACCGTGTCATCTTCCACC |
| β-tubulin-F | GGAAGCCCATAGAAAGACC |
| β-tubulin-R | CAACAAATGGGGATGCT |
To detect superoxide accumulation, hydrogen peroxide accumulation, and cell death, leaves were infiltrated with DAB, NBT and Evans blue staining according to the method described in detail by Zhang et al. [47] and Kim et al. [48]. Evaluation of ROS production in intact guard cells was performed by staining with H$_2$DCF-DA (Sigma-Aldrich) [49].

Statistical analysis
Each experiment was repeated at least three times independently. Error bars represent standard deviations. Differences were compared using Student's t-test. $P < 0.05$ was considered significant, which is indicated by *.

Supplementary information
Supplementary information accompanies this paper at https://doi.org/10.1186/s12870-020-02562-6.

Additional file 1 Figure S1. The relative expression levels of Th2CysPrx gene in the WT and transgenic tobacco as measured by qRT-PCR. The data was processed using the $2^{-\Delta\Delta CT}$ method. WT: the wild type tobacco. Line1–14: different transgenic tobacco lines. All experiments were repeated three times. The error bars represent the standard deviation.

Abbreviations
SOD: Superoxide dismutase; POD: Peroxidase; ROS: Reactive oxygen species; CAT: Catalase; GPX: Glutathione peroxidase; GST: Glutathione S-transferases; MV: Methyl violox; PSCS: Δ1-pyrroline-5-carboxylate synthetase; Pnxs: Peroxidoxins; H$_2$O$_2$: Hydrogen peroxide; qRT-PCR: Real-time quantitative PCR; WT: Wild-type; DAB: 3,3'-Diaminobenzidine; NBT: Nitro blue tetrazolium; H$_2$DCF-DA: 2,7-Dichlorofluorescin diacetate; MDA: Malondialdehyde; EL: Electrolyte leakage; FBPase: Fructose-1,6-bisphosphatase; CS: Citrate synthase; ABA: Abscisic acid; 1/2 MS: Half-strength Murashige and Skoog medium; MV: Methyl viologen; qRT-PCR: Real-time quantitative PCR; WT: Wild-type; DAB: 3,3'-Diaminobenzidine; NBT: Nitro blue tetrazolium; H$_2$DCF-DA: 2,7-Dichlorofluorescin diacetate; MDA: Malondialdehyde; EL: Electrolyte leakage; FBPase: Fructose-1,6-bisphosphatase; CS: Citrate synthase; ABA: Abscisic acid; 1/2 MS: Half-strength Murashige and Skoog medium.

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Authors’ contributions
CG designed the research and revised the manuscript; YW, XL, JW and WD mainly conducted data analysis and wrote the paper and all authors read and approved the manuscript.

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Availability of data and materials
All data and materials generated or analyzed during this study are included in this article or are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
Not applicable.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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