Isolation and Characterization of a Class III Peroxidase cDNA from Cucumber under Salt Stress

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ABSTRACT. Salt stress reduces the fresh weight, dry weight, and relative growth rate of cucumber (Cucumis sativus) seedlings and results in serious quality loss in cucumber production. Our previous study indicated that the netting-associated peroxidase (CsaNAPOD) protein in cucumber seedling roots was induced by salt stress. Here, we amplified the coding sequence of CsaNAPOD from a cDNA isolated from the roots of cucumber seedlings. Sequence analysis indicated that the coding sequence of CsaNAPOD is 1035 bp, encoding a deduced protein of 344 amino acids, with a predicted molecular weight of 37.2 kD and theoretical isoelectric point of 5.64. The deduced amino acid sequence of CsaNAPOD showed high sequence similarity to peroxidases (PODs) from other plant species. Moreover, CsaNAPOD possesses the typical sequence structures of class III PODs and indicated that CsaNAPOD belongs to this subfamily. CsaNAPOD was highly expressed in the roots and was weakly expressed in the stems and leaves of cucumber seedlings. Salt stress significantly increased the expression of CsaNAPOD in the leaves during the entire experimental period compared with the control, and the expression of CsaNAPOD in roots was reduced at 6 hours and induced at 48 and 72 hours by salt treatment. In stems, the expression of CsaNAPOD declined at 48 and 72 hours as a result of the salt treatment compared with the control. These results indicate that the expression of CsaNAPOD responded to salt stress in cucumber seedlings, and the expression patterns under salt stress in different tissues were not identical. Our research suggests that CsaNAPOD may have potential function during the plant response to salt stress.

Peroxidases, which comprise an important class of oxidoreductases, are widely distributed in all living organisms. PODs use H₂O₂ as their electron acceptor and iron porphyrin as their prosthetic group to catalyze substrate oxidation (Valério et al., 2004; van Huystee, 1987).

Plant PODs belong to a superfamily that consists of three different classes of PODs (Smulevich et al., 2006). Class I, the intracellular POD called catalase-peroxidase, is found in plants, bacteria, fungi, and Archaea. The primary function of class I PODs in the cell involves the detoxification of excess H₂O₂ (Shigeoka et al., 2002). Class II contains extracellular PODs secreted by fungi such as lignin PODs and manganese PODs that are involved in the degradation of soil debris (Martínez et al., 2005). Class III includes the large family of secreted PODs in plants that are secreted into the cell wall or the surrounding medium and the vacuole. In the standard peroxidative cycle, these PODs catalyze the reduction of H₂O₂ using electrons from various donor molecules. Class III PODs have been implicated in various functions, including auxin catabolism and the biosynthesis of secondary metabolites; the defense against pathogen penetration, wounding, and other abiotic stresses; cell wall hardening through the immobilization and extension of matrix polysaccharide crosslinking; and lignin and suberin deposition (Cosio and Dunand, 2009; De Gara, 2004). Recently, class III PODs have also been considered as potentially important components of plant signal transduction pathways (McInnis et al., 2005).

Genes encoding class III PODs belong to a large multigenic family with 73 members in Arabidopsis thaliana (Tognolli et al., 2002), 138 members in rice (Oryza sativa (Passardi et al., 2004)), and 173 members in the typical grass model Brachypodium distachyon (International Brachypodium Initiative, 2010). However, the presence of multiple isoforms and the complexity of the physiological processes regulated by these enzymes create difficulty in understanding the unique function of a specific class III POD member in plant development and in adapting to the environment. Information regarding the expression pattern during different developmental stages in tissues and under stress conditions aids in the understanding of the unique role of a specific POD.

The expression of class III POD genes is organ-dependent and influenced by abiotic stresses. Class III PODs in A. thaliana, Sesbania rostrata, and Senecio squaricus show obvious organ specificity, suggesting a highly specific role in the corresponding organ (Cosio and Dunand, 2009). The expression of class III POD genes is regulated by various environmental factors, including drought (Csiszár et al., 2012; Mohammadi et al., 2008), aluminum stress (Kumari et al., 2008), air pollutants and ultraviolet radiation (Kim et al., 2007), wounding (Gerchikov et al., 2008), and damage from plant disease and pests (Ansari et al., 2007; Severino et al., 2012). However, to date, information regarding class III POD gene expression mediated by salt is very limited.

Cucumber is one of the most important cucurbit (Cucurbitaceae) vegetables worldwide and is sensitive to a saline environment. The physiological responses of cucumber seedlings to salt stress have been well characterized. Specifically, salt stress destroyed the steady-state balance of reactive oxygen species...
(ROS) in cucumber seedlings, resulting in the loss of relative water content and net photosynthetic rates as well as a decrease in plant growth rates (Du et al., 2010a; Duan et al., 2008; Fan et al., 2012; Shi et al., 2007). Therefore, studies on salt stress are important to further understand the developmental mechanisms affected by salt for cucumber production.

In our previous study, to identify components of salt stress signaling, we analyzed the protein expression profiles of the roots of cucumber seedlings treated with 50 mmol NaCl for 3 d using two-dimensional gel electrophoresis, and we identified a few differentially expressed proteins using matrix-assisted laser desorption/ionization time-of-flight mass spectrometry and liquid chromatography electrospray ionization tandem mass spectrometry (Du et al., 2010b). We found that one of the identified proteins, named netting-associated peroxidase [NAPOD (high peptide coverage with netting-associated peroxidase in *Cucumis melo*)], was induced in cucumber seedling roots under salt stress and may be involved in the defense against stress in cucumber seedlings (Du et al., 2010b). In the present study, we further analyzed the protein sequence characteristics and designated the protein as CsaNAPOD. The expression pattern of CsaNAPOD in different cucumber seedling tissues over time under salt treatment was also investigated. The objective of the present research was to investigate whether CsaNAPOD is regulated by salt stress at the transcriptional level.

**Materials and Methods**

**Plant materials and treatment.** The experiments were conducted in an environment-controlled greenhouse. Cucumber cultivar Jinchun 2 seeds were allowed to germinate in the dark in a thermostatically controlled chamber at 28 to 30 °C for ≈30 h. The germinated seeds were sown in 50-cell plug trays containing peatmoss and perlite (1:1, v/v) at 1200 μmol·m⁻²·s⁻¹ photosynthetic photon flux and a 14/10 h (25 to 30/16 to 20 °C) day/night regime. Relative aerial humidity fluctuated between 60% and 75%. When the cotyledons expanded, the seedlings were supplied with half-strength Hoagland’s nutrient solution [pH 6.4 to 6.6, electrical conductivity (EC) 2.0 to 2.2 dS·m⁻¹]. After the full development of the second leaf of each plant, 12 seedlings of uniform size were transplanted into a plastic container (51 × 33 × 20 cm) filled with full-strength Hoagland’s nutrient solution. The nutrient solutions were kept at 20 to 25 °C and continuously aerated using an air pump at an interval of 20 min to maintain the dissolved oxygen at 7.8 to 8.2 mg·L⁻¹. After 3 d of pre-culture, the treatments were started. The cucumber seedlings were treated as follows: 1) control plants were grown in Hoagland’s solution; and 2) salt-treated plants were grown in Hoagland’s solution plus 50 mM NaCl. NaCl was added directly into the nutrient solution at the beginning of the treatment, and the plastic containers were arranged in a completely randomized block design with three replicates for each treatment (for a total of six plastic containers, each with 12 seedlings).

**Growth determination.** To understand the effects of salt treatment on cucumber development, we treated the pre-cultured cucumber seedlings with 50 mM NaCl for 96 h. At 96 h after NaCl treatment, six seedlings were prepared and were washed with tap water two to three times, rinsed twice with distilled water, gently blotted dry with a paper towel, and weighed to determine the weight of the seedlings when fresh. After measuring the fresh weight, the samples were killed in an oven at 105 °C for 15 min, dried at 75 °C for 72 h, and the dry weight was measured. The relative growth rate was defined as
The alignment of the predicted amino acid sequences of Cucumis sativus netting-associated peroxidase (CsaNAPOD) and three homologous peroxidases (PODs) with the highest similarity from different plant species. The accession numbers of the used amino acid sequences are as follows: C. sativus [Csa (XP_004151583.1)], Cucumis melo [Cm (AAPR19041.1)], Cucurbita pepo [Cp (ABF68751.1)], and Theobroma cacao [Tc (EOY09519.1)]. The three horizontal red lines correspond to the conserved distal heme-binding domain (B), the proximal heme-binding domain (F), and central conserved domains of unknown function (D). The predicted N-terminal signal peptide is italicized, the heme-binding sites are marked by a red asterisk, the calcium binding sites are marked by the inverted blue “V” symbol, and S-S-bridge-forming cysteines are noted by the inverted black triangle. The two green boxes indicate the putative glycosylation sites.

**PCR amplification of the CsaNAPOD coding sequence.**  
Using the Protein-BLAST tool in combination with the cucumber genomic database (Chinese Academy of Agricultural Sciences, 2009), we found only one protein, Csa023919, denoted also as POD 2 in C. sativus, with 100% identity to the CsaNAPOD amino acid sequence, indicating that CsaNAPOD and Csa023919 are the same protein. cDNA isolated from the roots of cucumber seedlings was used as the template for the PCR. The CsaNAPOD coding sequence was amplified corresponding to the coding sequence of Csa023919 in the cucumber genomic database. The primers used were as follows: forward, 5′-ATGGGCTCTCTCTCCGCCC-3′, reverse, 5′-TTACATAAACCAGCTGCGGCT-3′. The PCR was conducted according to the following program: 94 °C for 5 min followed by 30 cycles of 30 s at 94 °C, 30 s at 60 °C, 60 s at 72 °C, with a final extension step at 72 °C for 10 min.

The PCR product was separated through a 1.2% agarose gel and purified with the Agarose Gel DNA Extraction Kit (TaKaRa Bio). The fragment was inserted into a pMD18-T vector (TaKaRa Bio) and sequenced by Biological Engineering Co., Ltd., Shanghai, China.

The CsaNAPOD coding sequence is 1035 bp and encodes a deduced 344 amino acid sequence, which is identical to the Csa023919 sequence.

**Bioinformatics analysis.**  
The Protein-BLAST program at the National Center of Biotechnology Information (NCBI, Bethesda, MD) was used to search for sequence homology. The accession numbers of the amino acid sequences are as follows: C. sativus (XP_004151583.1), C. melo (AAPR19041.1), Cucurbita pepo (ABF68751.1), Theobroma cacao (EOY09519.1). Multiple sequence alignment was determined using the BioEdit programs. To predict the isoelectric point and molecular weight, Expasy (Swiss Institute of Bioinformatics, 2005) software was used. Signal peptides of the deduced proteins were determined by Signal-3L (Shen and Chou, 2007). The cell localization of the protein was predicted based on its amino acid sequence by PSORT (Nakai, 2007).

**Phylogenetic analysis.**  
All of the POD sequences (Supplementary material) mentioned in the text are available in the peroxidase databases (Laboratoire de Recherche en Sciences Végétales, 2004) and were used to build the phylogenetic tree. Multiple sequence alignment based on the full-length POD proteins was determined using the ClustalW method in MEGA Version 3.1 (Kumar et al., 2004). The phylogenetic trees were
constructed using the neighbor-joining method. The bootstrap analysis was set at 1000 replicates to evaluate the reliability of the different phylogenetic groups. The tree files were viewed and edited by MEGA 3.1 software.

**Quantitative real-time PCR (qRT-PCR) analysis.** To understand the effects of salt stress on CsaNAPOD expression, we treated cucumber seedlings with 50 mM NaCl stress for 96 h. The expression of CsaNAPOD in the leaf, stem, and root was determined by quantitative real-time PCR (qRT-PCR) at 0, 6, 24, 48, 72, and 96 h of treatment. qRT-PCR was performed in a 96-well plate format ABI 7300 Real-Time PCR System (Applied Biosystems, Foster City, CA). The reactions (20 μL) were conducted using the SuperReal PreMix Plus (SYBR Green) kit (Tiangen Biotech, Beijing, China) according to the manufacturer’s protocol. For qRT-PCR, we designed primers specific for CsaNAPOD. The primer sequences were as follows: for CsaNAPOD, forward, 5-TGAGCTGCCTGTTCGATGCTATAT-3, reverse, 5-CAAAACAAAGATGAGCCTACC-3; for CsaActin, forward, 5-CCGGTTCTGCTCTTCACTACCTGGTT-3, reverse, 5-GGAACTGCTCTTTGCACTGCCAG-3. As an internal standard, the expression level of CsaActin in control and salt-treated cucumber roots was consistent and relatively stable (Du et al., 2010b; Kabała and Kłobus, 2008). The PCR conditions were optimized for each primer set. The PCR amplification was conducted after denaturing the cDNA at 95 °C for 15 min followed by 40 cycles of 95 °C for 10 s, annealing at 58 °C for 20 s, and extension at 72 °C for 31 s. The experimental samples, internal standards, and no template controls were run in triplicate. The relative gene expression was quantified using the comparative cycle threshold (Ct) method (referred to as the 2−ΔΔCt method) (Livak and Schmittgen, 2001). All data were statistically analyzed with SAS software (Version 8.2). The t test was used for the statistical calculation of comparison data between the control and salt treatment. Duncan’s multiple range test was used for the statistical calculation of the data obtained by comparisons among the three tissues or six time points.

**Results**

As shown in Figure 1, compared with the control plants, the plant fresh weight (Fig. 1B), dry weight (Fig. 1C), and RGR (Fig. 1D) of salt-treated plants at 96 h of treatment were decreased by 36.2%, 16.4%, and 64.6%, respectively (P < 0.05). This result indicates that salt stress significantly inhibited the growth of the cucumber seedlings.

Sequence analysis showed that the deduced amino acid sequence of CsaNAPOD harbors three conserved domains: the conserved distal heme-binding (B) and the central conserved (D) and proximal heme-binding (F) domains of unknown function (Fig. 2). The CsaNAPOD amino acid sequence also contains eight conserved cysteine residues leading to four disulfide bridges. These characteristics are common in most class III PODs. The analysis of the translated polypeptide also revealed the presence of two putative glycosylation sites and indispensable amino acids required for the heme-binding coordination of Ca²⁺ ions (Fig. 2).

According to the predicted localization by PSORT, CsaNAPOD may be secreted extracellularly. A predicted 23 amino acid signal peptide was identified by Signal-3L at the 5' end of the amino acid sequence, which is consistent with an apoplastic protein.
Fig. 3. Phylogenetic relationships of *Cucumis sativus* netting-associated peroxidase (*CsaNAPOD*) with other class III peroxidase (POD) amino acid sequences. The phylogenetic tree file was produced by MEGA 3.1 (Kumar et al., 2004). The bootstrap values were obtained by the neighbor-joining method and indicate the divergence of each branch. The scale indicates the branch length. All the protein sequences (Supplementary material) are available in the peroxidase database (Laboratoire de Recherche en Sciences Végétales, 2004); *C. sativus* (Csa), *Cucumis melo* (Cm), *Cucurbita pepo* (Cp), *Gossypium hirsutum* (Gh), *Ricinus communis* (Rc), *Manihot esculenta* (Me), *Glucine max* (Gm), *Arabidopsis thaliana* (At), *Ipomoea batatas* (Ib), *Citrus sinensis* (Cs), *Vitis vinifera* (Vv).

Using Expasy software, the theoretical isoelectric point and molecular weight of the deduced *CsaNAPOD* protein were predicted to be 5.64 and 37.2 kDa, respectively. The alignment of the deduced amino acid sequences with homologs from other species showed that *CsaNAPOD* has the highest similarity to the netting-associated anionic POD of *C. melo* [AAR19041.1 (93%)] followed by the PODs of *C. pepo* [ABF68751.1 (74%)] and *T. cacao* [EEOY9519.1 (57%)].

The analysis of the phylogenetic relationships of *CsaNAPOD* with other class III POD proteins indicated that the POD homologs from the Cucurbitaceae were positioned on a well-supported branch clearly separated from the PODs present in other families (Fig. 3). The percentage of identity decreased greatly with sequences from *Citrus sinensis* and other non-Cucurbitaceae species. In *A. thaliana*, the closest ortholog was AtPrx53 with 56% identity to the *CsaNAPOD* protein sequence.

The results showed that among the three tested organs, *CsaNAPOD* was highly expressed in the root with lower expression in the stem and leaf (Table 1). The expression of *CsaNAPOD* increased within 24 h and decreased after 48 h in the leaves of the cucumber seedlings in the control treatment (Fig. 4); however, the overall trend of *CsaNAPOD* expression declined within the 96 h of treatment in the roots and stems of the cucumber seedlings. *CsaNAPOD* in the leaves of the cucumber seedlings was significantly induced after 6 h of NaCl treatment. The *CsaNAPOD* transcript levels were significantly enhanced by 7.9-, 3.9-, 7.2-, 6.2-, and 5.5-fold of control at 6, 24, 48, 72, and 96 h of treatment, respectively (Fig. 4A). As shown in Figure 4B, compared with the control, salt stress caused a decrease in *CsaNAPOD* transcripts in the roots of cucumber seedlings after 6 h of treatment and caused an increase at 48 and 72 h, and no obvious change was observed at 24 and 96 h. The *CsaNAPOD* transcript level in the stems of cucumber seedlings under NaCl treatment decreased compared with the control at 48 and 72 h of treatment, respectively, and no significant change in the transcript level was observed at 6, 24, and 96 h after salt treatment (Fig. 4C).

**Discussion and Conclusions**

Salt stress causes adverse effects on plant growth and productivity (Parida and Das, 2005), and our results showed that cucumber seedling growth was reduced by salt stress (Fig. 1). This result is in agreement with the previous reports regarding salt-treated cucumbers (Du et al., 2010a; Duan et al., 2008; Fan et al., 2012; Shi et al., 2007). Duan et al. (2008) suggested that cucumber seedlings most likely increase the protein level of antioxidant enzymes and the content of osmotics to enhance resistance to salinity. The molecular mechanism of this process remains unclear; however, it has been well established that any stress condition or significant change in the environment is associated with the up- or downregulation of several hundreds of genes, including PODs (Cheeseman, 2007).

We identified a protein termed NAPOD, which was induced by salt stress in cucumber seedling roots (Du et al., 2010b). In this study, our result showed that this peroxidase belongs to class III PODs. The predicted amino acid sequence of the *CsaNAPOD* gene was compared with other known class III POD sequences (Fig. 2). The alignment analysis showed that the *CsaNAPOD* protein harbored conserved structural features and activity sites shared by the majority of the class III PODs, including the predicted N-terminal signal peptide; the highly conserved B, D, and F domains essential to enzyme catalysis and protein folding; and eight cysteine residues involved in the formation of four disulfide bridges. Two putative glycosylation sites also found in the *C. melo* CmNAPOD (Keren-Keiserman et al., 2004) suggest that modifications of the mature *CsaNAPOD* occur.

As a result of the recent plant genome project activities, many POD gene sequences or homologous sequences have been identified. Phylogenetic analyses have been conducted for PODs from several plants, including *A. thaliana* (Tognolli et al., 2002), rice (Zhang et al., 2001), cotton (*Gossypium hirsutum* (Delannoy et al., 2006)), buffalograss (*Buchloe dactyloides* (Gulsen et al., 2007)), and apple (*Malus sieversii* (Gulsen et al., 2010)). Here, the analysis of the phylogenetic relationships of *CsaNAPOD* with other class III POD proteins confirmed that *CsaNAPOD* is a member of the class III POD subfamily. The *CsaNAPOD* grouped with Cucurbitaceae PODs to form a branch clearly separated from the other species (Fig. 3).

A few POD genes exhibit organ-dependent expression; however, other POD genes are active in the entire plant (Tognolli et al., 2002; Welinder et al., 2002). In the *A. thaliana* genome, 73 class III POD genes have been annotated, 65 of which were expressed in various tissues, and only three were identified as specific to roots (Valério et al., 2004). Delannoy et al. (2006) reported the expression profiles of 10 cotton class III POD genes (pod1 to pod10) from several different cotton tissues examined. None of the 10 pod genes was found to be
organ-specific. Class III PODs generally do not appear to have a high level of tissue specificity (Tognolli et al., 2002). However, in recent years, class III PODs with organ/tissue specificity were identified (Cosio and Dunand, 2009; Mei et al., 2009). In *S. squalidus*, a stigma-specific class III POD was identified that may be involved in the defense against pathogens or in pollen–stigma signaling (McInnis et al., 2005). Three PODs are specifically expressed in stigmas but are not found in other floral tissues/organs in *A. thaliana* (Swanson et al., 2005). In the present study, *CsaNAPOD* showed no organ specificity and was expressed in each organ evaluated (root, leaf, and stem); however, the transcript abundance of *CsaNAPOD* was different in these organs. For example, *CsaNAPOD* demonstrated high expression in roots and lower expression in the leaves and stems (Table 1), suggesting that, at least to a certain extent, the role of *CsaNAPOD* may be distinct in different organs. The relatively high expression of *CsaNAPOD* in roots may be used for eliminating excessive ROS generated by salt stress, which suggests that *CsaNAPOD* may be associated with the most powerful defense against salinity in roots.

PODs have been associated with high salinity stress in a few plants. Ritter et al. (1993) demonstrated a peroxidase mRNA was induced 3-fold in the young leaves of salt-stressed cotton plants. Kumar et al. (2011) indicated that the expression levels of *CrPrx3* and *CrPrx4*, two new class III POD genes, were down-regulated in *Catharanthus roseus* under salt stress. In addition, the overexpression of the sweetpotato (*Ipomoea batatas*) swpa4 peroxidase significantly enhanced the salt stress tolerance of tobacco plants [Nicotiana tabacum (Kim et al., 2008)]. In our study, *CsaNAPOD* protein in cucumber seedling roots was induced at 3 d under NaCl stress (Du et al., 2010b), which is in agreement with the transcript expression in roots at 3 d (72 h) after starting NaCl treatment.

In our study, salt stress greatly up- and downregulated *CsaNAPOD* expression in the leaves and roots of cucumber seedlings, respectively, at 6 h after the start of treatment, indicating the effects of salt stress in these two organs at the early stage of treatment in the experiment. *CsaNAPOD* expression in the stem was affected by salt at 48 h of treatment. Salt stress caused different effects on *CsaNAPOD* transcript accumulation in the leaves, roots, and stems of cucumber seedlings, suggesting that the response to salt stress was organ-specific. The differential expression of *CsaNAPOD* among different tissues may be considered to dominate the steady-state level of ROS; therefore, neutralizing the toxic effects of salt stress is crucial. Our results indicate that *CsaNAPOD* expression in leaves and roots is very rapidly induced and suppressed in cucumber seedlings under salt stress, respectively, suggesting that *CsaNAPOD* in leaves and roots may play an important role in the cucumber plant against salinity. Moreover, the upregulated expression of *CsaNAPOD* in leaves during the entire experimental period and in the roots at 48 and 72 h after starting treatment under salt stress may be directly involved in a variety of protective or adaptive biological mechanisms controlling distinct cellular functions under different environmental conditions.

Plant class III PODs have been proposed as key regulators of the level of extracellular H$_2$O$_2$ and as producers of the extremely reactive hydroxyl ions (OH•) and hydroperoxyl (OOH•) radicals (Bolwell et al., 2002), depending on whether the peroxidative or the hydroxylic cycles of the enzyme are operating (Kawano, 2003; Passardi et al., 2005). In this study, the study of salinity-induced *CsaNAPOD* regulation is an essential issue, directly implicated in the understanding of the biological functions involved in plant defense response. A

![Fig. 4](image-url) The effect of salt stress on the expression of *Cucumis sativus* netting-associated peroxidase gene (*CsaNAPOD*) in the leaf (A), stem (B), and root (C) of cucumber seedlings. The transcript levels of *CsaNAPOD* in cucumber seedlings under NaCl stress after 0, 6, 24, 48, 72, and 96 h of treatment were detected using the quantitative real-time polymerase chain reaction. Control = control plants; NaCl = 50 mM NaCl-treated plants. The relative expression level value was normalized to an internal standard *CsaActin*. The expression level of *CsaNAPOD* in control plants at 0 h of treatment was defined as 1. Each histogram represents the mean value of three independent experiments, and the vertical bars indicate SE (n = 3). Asterisks indicate the significant differences between control and NaCl stress at each time point calculated using the *t* test (*P < 0.05).
broad spectrum of defense responses includes early and late responsive genes geared toward immediate defense and long-lasting defense strategies. Different peroxidase isoenzymes likely play an essential role in the normal development and lasting defense strategies. Different peroxidase isoenzymes responsive genes geared toward immediate defense and long-lasting defense strategies. Regardless of whether or when the expression of the gene was inhibited or increased by salt stress, our observations suggest that CsaNAPOD has a potential defense response in cucumber plants against salt stress. PODs are activated in response to stress-related signals of a biotic or abiotic nature and are believed to play key roles in these processes. To better understand the function of peroxidases, further experiments using the overexpression and suppression of each gene in transgenic plants are necessary.

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