CsPrx25, a class III peroxidase in Citrus sinensis, confers resistance to citrus bacterial canker through the maintenance of ROS homeostasis and cell wall lignification

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Abstract
Citrus bacterial canker (CBC) results from Xanthomonas citri subsp. citri (Xcc) infection and poses a grave threat to citrus production. Class III peroxidases (CIII Prxs) are key proteins to the environmental adaptation of citrus plants to a range of exogenous pathogens, but the role of CIII Prxs during plant resistance to CBC is poorly defined. Herein, we explored the role of CsPrx25 and its contribution to plant defenses in molecular detail. Based on the expression analysis, CsPrx25 was identified as an apoplast-localized protein that is differentially regulated by Xcc infection, salicylic acid, and methyl jasmonate in the CBC-susceptible variety Wanjincheng (C. sinensis) and the CBC-resistant variety Calamondin (C. madurensis). Transgenic Wanjincheng plants overexpressing CsPrx25 were generated, and these transgenic plants exhibited significantly increased CBC resistance compared with the WT plants. In addition, the CsPrx25-overexpressing plants displayed altered reactive oxygen species (ROS) homeostasis accompanied by enhanced H₂O₂ levels, which led to stronger hypersensitivity responses during Xcc infection. Moreover, the overexpression of CsPrx25 enhanced lignification as an apoplastic barrier for Xcc infection. Taken together, the results highlight how CsPrx25-mediated ROS homeostasis reconstruction and cell wall lignification can enhance the resistance of sweet orange to CBC.

Introduction
Plants possess an intricate repertoire of cell-based defense systems to maintain their resistance to potentially harmful pathogens. As an immediate pathogen recognition response, oxidative bursts produced in apoplasts induce reactive oxygen species (ROS), including superoxide (O₂⁻) and H₂O₂, as a first line of defense. The current models of plant responses include ROS and other radicals as catalysts of covalent cell-wall modifications, as signals for cell-death reactions, and as regulators of resistance-associated genes. In plants, high concentrations of ROS act to strengthen the cell wall and inhibit pathogen growth, which results in the enhancement of host resistance to pathogens via hypersensitive responses (HRs) and the modulation of gene expression via signaling molecules. However, high accumulation of ROS can be toxic to plant cells by inhibiting plant growth and development. Thus, ROS homeostasis needs to be maintained by antioxidant compounds and enzymes. In plant cells, ROS are produced by NADPH oxidase resident at the cell surface, class III peroxidases (CIII Prxs, or POD) and their associated pathways, including photosynthesis, photorespiration, and respiration. In addition, the ROS scavengers superoxide dismutase (SOD), catalase (CAT), and glutathione s-transferase (GST) cooperate with ROS producers to maintain ROS homeostasis. Moreover, antioxidant enzyme activities and ROS homeostasis are regulated by important plant hormones, including jasmonic acid (JA) and salicylic acid (SA).
CIII Prxs are heme-binding proteins that are ubiquitously expressed in all plants and comprise large multigene families\(^{18-21}\). For example, a total of 73 CIII Prxs are present in Arabidopsis thaliana\(^{22,23}\), and 138, 374, 93, 94 and 72 have been found in Oryza sativa\(^{24}\), Triticum aestivum\(^{25}\), Populus trichocarpa\(^{26}\), Pyrus bretschneideri\(^{27}\) and Citrus sinensis\(^{28}\), respectively. CIII Prxs regulate the loosening of cell walls, lignification and suberization\(^{29-32}\) and participate in ROS and RNS metabolism during abiotic and biotic stress responses\(^{33-35}\). CIII Prxs are key to the innate resistance of many plants to both fungal and bacterial pathogens and mediate both passive and active defense mechanisms\(^{6,36,37}\), and the efficiency of this mediation determines their susceptibility to pathogenic infections\(^{38}\). Rapid ROS production is one such exemplar defense strategy that leads to \(O_2^-\) generation and \(H_2O_2\) production in apoplasts. \(H_2O_2\) is tightly regulated by CIII Prxs as both producers and scavengers depending on whether the enzyme participates in peroxidative cycles and hydroxylc cycles, respectively\(^{12,13}\). In French bean and tobacco plants, apoplastic CIII Prxs produce ROS and act as catalysts for covalent cell-wall modifications\(^8\) and cell death regulators\(^6\). Based on these functions of CIII Prxs, an increasing number of studies have identified links between this enzyme and pathogen attack and have improved host resistance due to CIII Prxs. Radwan and colleagues reported that bean yellow mosaic virus infection leads to increased levels of monodihydroascorbate (MDA) and \(H_2O_2\) in Vicia faba leaves\(^{39}\). Enhanced CIII Prx and SOD activities have also been observed in leaves infected by yellow mosaic virus, which suggests that enzymatic antioxidants regulate ROS generation in response to pathogen infection\(^{39}\). Increasing the expression of a peroxidase in plants can effectively increase the resistance of the plants to disease. For example, the overexpression of HvPrx40 and TaPrx10\(^{39,41}\) leads to higher levels of resistance to Blumeria graminis (wheat powdery mildew) in wheat (\(T. aestivum\)).

Xanthomonas citri subsp. citri (Xcc) pathogen is the causative agent of citrus bacterial cancer (CBC), a known cause of citrus yield losses in an array of citrus-producing regions\(^{42,43}\). In our previous studies of the citrus transcriptomes induced by Xcc, we found that CIII Prxs were differentially expressed and explored the relationship between CBC and CIII Prxs, and our results revealed CsPrx25 as a potential gene for improving CBC resistance\(^{28}\). Here, we performed both a structural and functional characterization of CsPrx25. We also developed transgenic sweet orange overexpressing CsPrx25 that displayed enhanced tolerance to CBC due to ROS homeostasis accompanied by high levels of \(H_2O_2\) and high lignification of the apoplastic barrier. We herein describe the utility of transgenic plants overexpressing CsPrx25 for enhancing CBC resistance.

**Results**

**CsPrx25 encodes a CIII Prx in citrus**

We amplified and sequenced the complete transcript of CsPrx25 using cDNA from Wanjincheng leaves as the PCR template. The primary sequences were searched in PeroxiScan, which is built in RedoxiBase\(^{44,45}\). The findings revealed that CsPrx25 belonged to the CIII Prx family (PeroxiScan accession: PS52045), a subgroup of non-animal peroxidases (PeroxiScan accession: PS50873). The CsPrx25 sequence was further analysed by the Blast tool built in RedoxiBase and CAP\(^{46}\), and the results revealed that CsPrx25 was clustered with the CIII Prx sequence ID 8898 in RedoxiBase and Cs3g21730 in CAP due to 100 and 98% sequence similarities, respectively. CsPrx25 is a 344-residue CIII Prx (molecular weight: 38.06 kD; isoelectric point: 8.55) present on chromosome 3 of C. sinensis (Fig. 1a) that possesses two introns (1515 bp and 659 bp, respectively) (Fig. 1b). The N-terminus of CsPrx25 contains a signal peptide of 27 residues that is required for correct trafficking to the apoplast. Throughout the sequence, eight cysteine residues were detected (C1–C8) (Fig. 1c), and these form a total of four disulfide bonds (DB) that maintain thermal stability. These 4-DB structures are common to almost all plant CIII Prxs and impart distinction from ascorbate and other plant peroxidases\(^{37}\). The three-dimensional (3D) structures also showed that the cysteines that form disulfide bonds are close to each other (Fig. 1d). To study the evolutive scenario of CIII Prxs between organisms, the phylogeny of CIII Prxs orthologs was assessed, and close relationships between CsPrx25 and AtPrx12 were found (Fig. 1e).

**CsPrx25 is an apoplast-localized protein that is induced by Xcc and phytohormones**

To elucidate the localization of CsPrx25, software predictions and transient expression systems were investigated. CELLO V2.5 displayed extracellular loci values of 2.46, which were larger than other loci (Supplementary Table S1). The signal peptide detected by SignalP V4.0 suggests that CsPrx25, as most of the CIII Prxs, is extracellular. To validate these predictions, the transient expression of CsPrx25 was assessed with 35S::CsPrx25-GFP (Fig. 2a). Relative to the controls, both cytoplasmic and nuclear fluorescence were observed before and after plasmolysis (Fig. 2b). In epidermal onion cells, CsPrx25-GFP showed robust cell surface expression (Fig. 2c), confirming that CsPrx25 localizes to apoplasts.

Pathogens and phytohormones can mediate gene expression changes that occur in response to plant disease\(^{48,49}\). In Calamondin, CsPrx25 was upregulated, and maximal expression (~5-fold) was observed at 36 hpi. In contrast, Wanjincheng CsPrx25 showed little-to-no expression changes in response to Xcc infection (Fig. 2d). To detect the effect of drought during in vitro...
inoculation, we tested the inducibility of CsPrx25 under drought stress. The results indicated that CsPrx25 was hardly induced by drought in both varieties, which indicated that it was specifically induced by Xcc (Supplementary Fig. S1). CsPrx25 is therefore likely to represent an Xcc resistance gene. To reveal the molecular mechanisms through which CsPrx25 mediates disease resistance, CsPrx25 transcripts were assessed in SA- and MeJA-treated leaves. The expression of CsPrx25 rapidly increased in Calamondin in response to SA. In contrast, CsPrx25 expression was downregulated in Wanjincheng (Fig. 2e). The expression of CsPrx25 induced by MeJA increased and then decreased over time in both Wanjincheng and Calamondin, and the times to maximal expression in these varieties was different (Wanjincheng: 24 hpt vs Calamondin: 6 hpt) (Fig. 2f). The different expression patterns of CsPrx25 induced by phytohormones indicate the different roles of CsPrx25 in disease resistance signaling in Calamondin and Wanjincheng.

**CsPrx25 overexpression in sweet orange induces resistance to CBC**

Transgenic citrus constructs overexpressing CsPrx25 were used to fully dissect the role of CsPrx25 during Xcc resistance. CsPrx25 was overexpressed using exogenous expression plasmids driven by the 35S promoter (Fig. 3a). The generation of four CsPrx25-overexpressing plants 1–4 (OE1–OE4) that successfully integrated CsPrx25 was confirmed by qRT-PCR, GUS assay and Southern blot. Through PCR, we detected an 1874-bp fragment that was...
not present in the wild-type (WT) lines (Fig. 3b), and the GUS assay revealed blue color on the periphery of the leaf discs (Fig. 3c). As determined by Southern blot, OE1 and OE2 contain two copies of \( CsPrx25 \), and OE3 and OE4 harbor only one copy (Fig. 3d). We confirmed that all lines expressed high levels of \( CsPrx25 \) (550-fold, 589-fold, 401-fold and 395-fold of the WT levels, respectively) by qRT-PCR analysis (Fig. 3e). According to the Southern blot assay, a certain positive correlation exists between copy number and expression (Fig. 3d). With respect to phenotypes, the four transgenic lines showed normal growth rates compared with the WT lines (Fig. 3f).

Acupuncture is an effective method for quantitatively assessing the resistance to CBC and can be used to accurately quantify CBC resistance, which would allow the assessment and comparison of resistance between varieties. To assess the CBC resistance of \( CsPrx25 \)-OE plants, in vitro assays were performed with acupuncture inoculation at 10 dpi. Smaller lesion sizes, which are indicative of less-severe symptoms, were observed in the OE leaves compared with the WT leaves (Fig. 3g). This finding suggested that \( Xcc \) pustules are reduced by \( CsPrx25 \) overexpression, and OE2 showed the highest levels of resistance. Compared with the WT plants, OE2 showed smaller lesions (45.8% of the WT levels), OE1 exhibited comparable lesions (47.0% of the WT levels), and OE3 and OE4 displayed larger lesions (65.8% and 68.8% of the WT levels) (Fig. 3h). The disease severity...
decreased by 29.2% (OE3) to 50.7% (OE2) in the OE plants compared with WT plants (Fig. 3i). Using infiltration assays after 10 dpi, symptoms of canker (including pustules) were observed in then WT lines, but these symptoms were markedly reduced in the OE plants (Fig. 3j). We therefore conclude that CsPrx25 overexpression enhances Xcc resistance in the OE transgenic citrus lines.

**CsPrx25 overexpression modulates the enzymatic antioxidant system**

Plants possess a well-developed ROS homeostasis enzymatic system that efficiently regulates the ROS levels, and this system includes CIII Prx, SOD, CAT and GST. To assess the changes in the antioxidant system following the induction of CsPrx25-mediated resistance to Xcc, the antioxidant activity in transgenic lines in these lines was compared with that in the WT plants at 12 hpi. OE plants with higher resistance (OE1 and OE2) to CBC were selected for analysis. The activities of both CIII Prx and SOD were upregulated by CsPrx25 overexpression (Fig. 4a, b). The overexpression of CsPrx25 conferred antioxidant defenses and led to the induction by Xcc infection. In contrast to CIII Prx and SOD, the activities of CAT in OE plants were downregulated, and the Xcc-induced profiles were altered compared with those observed in the WT plants (Fig. 4c). In contrast to SOD, CIII Prx and CAT, the overexpression of CsPrx25 did not affect the activity of GST compared with that found in the WT plants (Fig. 4d).

**CsPrx25 overexpression establishes ROS homeostasis to confer a more sensitive HR to Xcc infection**

In response to pathogen infection, ROS production intricately controls many responses, including apoptotic cell death and oxidative damage. To confirm the involvement of ROS homeostasis in CsPrx25-mediated Xcc resistance, the levels of H$_2$O$_2$ and O$_2$-$^-$ in WT vs. CsPrx25-OE lines were assessed. We observed higher levels of H$_2$O$_2$ in the OE lines. Of interest, Xcc infection did not significant change the levels of H$_2$O$_2$ but also reversed the inducible patterns of H$_2$O$_2$ during Xcc infection. The levels of O$_2$-$^-$ also increased in response to CsPrx25 overexpression (Fig. 5b). The cell membrane is first affected by lipid peroxidation, and MDA is the final product. A spectroscopic analysis of the transgenic and WT plants...
revealed elevated levels of MDA, and these levels were modestly reduced in response to \textit{Xcc} infection (Fig. 5c), which indicate lower levels of damage following \textit{Xcc} infection in both the transgenic and WT plants. These data indicate that Wanjincheng has the ability to suppress the oxidative damage caused by \textit{Xcc} infection, and this suppression is strengthened by \textit{CsPrx25} overexpression. 

\H2O2 is a key mediator of an early HR. Because \textit{CsPrx25} overexpression regulates \H2O2 modulation, the immediate question was whether the HR is also altered in the transgenic plants. To investigate the relationship between the increased CBC resistance induced by \textit{CsPrx25} and HR, we assessed the HR of the transgenic plants before and after \textit{Xcc} infection. The expression of the HR marker gene, \textit{CsHSR203}–56–58 was significantly upregulated in the transgenic plants infected with \textit{Xcc} but only modestly increased in the \textit{Xcc}-infected WT plants. No obvious changes in the expression of \textit{CsHSR203} were observed between the transgenic and WT plants in the absence of \textit{Xcc} infection (Fig. 5d). We therefore conclude that the transgenic plants are more sensitive to a HR following \textit{Xcc} infection, which increases the early resistance of transgenic plants to CBC.

\begin{figure}[h]
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\includegraphics[width=\textwidth]{fig4}
\caption{CsPrx25 overexpression modulates the enzymatic antioxidant system. CIII Prx (a), SOD (b), CAT (c) and GST (d) in both OE and WT plants inoculated with mock (ddH2O) (filled bars) and \textit{Xcc} (open bars) for 12 h. FW: fresh weight. In a–d, the values were compared to those found for the WT lines. The differences between the mock- and \textit{Xcc}-infected samples were analyzed using Fisher’s LSD test, *\textit{P}< 0.05; **\textit{P}< 0.01. Tukey’s HSD test was used to compare the WT and OE plants (\textit{P}= 0.05; \textit{n}= 3).}
\end{figure}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig5}
\caption{CsPrx25 reconstructs ROS homeostasis and imparts HR sensitivity during \textit{Xcc} infection. The levels of \H2O2 (a), \textit{O2}– (b) and MDA (c) in OE and WT plants were assayed at 12 h after mock inoculation (ddH2O) (filled bars) and \textit{Xcc} infection (open bars). In a–c, FW: fresh weight. d \textit{CsHSR203} transcript levels in WT vs. OE plants at 12 h after mock (ddH2O) (filled bars) or \textit{Xcc} inoculation (open bars). The data were normalized to the \textit{CsActin} levels. In a–d, the differences between the mock- and \textit{Xcc}-infected samples were analyzed by Fisher’s LSD test, *\textit{P}< 0.05; **\textit{P}< 0.01. The differences between the OE and WT plants were analyzed using Tukey’s HSD test (\textit{P}= 0.05; \textit{n}= 3).}
\end{figure}
CsPrx25 overexpression enhances lignification as an apoplastic barrier for Xcc infection

CIII Prxs regulate cell wall lignification, which suggests a direct role of CIII Prxs on the cell walls of plants. To investigate the effects of lignification on Xcc resistance in transgenic plants, the role of CsPrx25 in lignification was assessed. The transcript levels of lignin biosynthetic genes, namely, hydroxycinnamoyl transferase (CsHCT), cap ID: Cs1g14450), cinnamyl alcohol dehydrogenase (CsCAD, cap ID: Cs1g20590) and caffeoyl-CoA O-methyltransferase (CsCCoAOMT, cap ID: Cs4g13430), were elevated in the leaves of the transgenic lines after mock inoculation and Xcc infection (Fig. 6a–c). These findings highlight the role of CsPrx25 in lignin biosynthesis. All the data were confirmed through lignin assays, which showed higher values in the transgenic compared with the WT plants (Fig. 6d). These data therefore reflect the role of CsPrx25 in the polymerization of lignin during its biosynthesis and highlight its importance in CBC resistance through enhanced lignification.

CsPrx25 enhances CBC resistance, and this effect is associated with ROS homeostasis reconstruction and lignification

CsPrx25 overexpression confers ROS homeostasis to the transgenic lines through modulation of the enzymatic antioxidant system (Figs. 4–5). The levels of lignin were also higher in the transgenic lines than in the WT plants, and some lignin biosynthetic genes were more highly expressed in the transgenic lines (Fig. 6). Based on these results, we proposed a model to explain how Calamondin and CsPrx25-OE transgenic Wanjincheng acquired CBC resistance (Fig. 7). In Calamondin, Xcc infection improves
the levels of CsPrx25, and this effect enhances the H2O2 levels and HR sensitivity and induces lignification, resulting in CBC resistance. The overexpression of CsPrx25 in CBC-susceptible Wanjincheng establishes ROS homeostasis, and higher H2O2 levels confer HR sensitivity in response to Xcc infection. In the transgenic plants, CsPrx25 overexpression also enhanced lignin biosynthesis, reinforcing the apoplastic barrier for Xcc infection. Through these two mechanisms, CsPrx25 promotes CBC resistance.

**Materials and methods**

**Plants, bacteria and growth conditions**

All the plants were obtained from the National Citrus Germplasm Repository. Wanjincheng (*C. sinensis*) was used for gene transformations. All the plants were grown at 28 °C in a greenhouse. The Xcc variants were derived from citrus leaves that are susceptible to natural infections. The Xcc cultures were grown at 28 °C in peptone-yeast extract-malt extract containing 1.5% (w/v) D-glucose.

**In silico characterization of CsPrx25**

The complete transcript sequence of CsPrx25 was amplified from Wanjincheng leaves using the primers Fclone (ATGGCAACTGCTTCAGCTTCT) and Rclone (TTAGATAATCCCAGACCAAGCC). PeroxiScan was used for the family classification of CsPrx25. Blast tools built in RedoxiBase were used to confirm the sequence of CsPrx25 retrieved by PCR. The chromosomal loci and the locations of exons and introns were defined using GSDS V2.0 based on the genome assembly of *C. sinensis* in CAP. SignalP V4.0 was used for signal peptide predictions, and CELLO V2.5 was used for cellular localization prediction. Phyre V2.0 was used for the 3D assessments of CsPrx25. The gene, protein, and coding sequences (CDSs) of CsPrx25 are shown in Table S2.

**Transient expression of GFP-tagged CsPrx25**

The coding sequence (CDS) of CsPrx25 lacking a stop codon was amplified with flanking restriction sites using the primers Fs (CGGGATCCTAGCTTTCAATCCTGTA) and Rs (ATGGCAACTGCTTCAGCTTCT) (KpnI) and Rs (TCCCCGGGTCTAC TGGTTGAAATTAAAGGATCT) (SmaI), digested, recovered and cloned into pHNe-GFP driven by the 3SS promoter to construct the recombinant plasmid pHNe-CsPrx25-GFP. The pHNe-CsPrx25-GFP plasmid encodes a fusion protein composed of CsPrx25 and GFP. The plasmids were heat-shocked into Agrobacterium EHA105. The transformed EHA105 was infiltrated into onion epidermal cells, and the GFP fluorescence signals were observed at 48 hpi by laser-scanning confocal microscopy (LSM 510 Meta, Zeiss).

**Treatments with Xcc and phytohormones**

The expression of CsPrx25 in excised leaves maintained in culture plates for 16 h of light and 8 h of darkness was assessed. Diluted Xcc (OD600: 0.8) was inoculated onto the leaves at 28 °C, and after defined durations, the expression of CsPrx25 was assessed by qRT-PCR. For phytohormone assessments, leaf discs were soaked in 10 μmol L−1 SA or 100 μmol L−1 MeJA and collected for qRT-PCR assays of exogenous phytohormones. The primers used for CsPrx25 detection were FRT (CCCCACTTCGGATCCAGCAAC) and RRT (CAACCCCTGTCGGTTACATCA).

**Overexpression vector construction and plant transformation**

For the generation of overexpression lines, full-length CsPrx25 was PCR amplified using F0EC (GGGGTACCA TGGCAACTGCTTCAGCTTCT) and R0EC (CGGATCTT CTTAGATAATCCCAGACCAAGCC) and cloned into pLGNe to yield the recombinant plasmid pLGNe-CsPrx25. Wanjincheng shoot transformations were performed using *Agrobacterium tumefaciens* as previously described by Li and He.

**Validation of the transgenic lines by PCR and GUS assays**

PCR assays were used to confirm the presence of the transgenic gene with the primers F0ED (CGACACGCTTACGTTCAAGCTT) and R0ED (CGGGATCC TCTAGATAATCCCAGACCAAGCC) and cloned into pLGNe-CsPrx25. The DS was assessed by qRT-PCR. For phytohormone assessments, leaf discs were soaked in 10 μmol L−1 SA or 100 μmol L−1 MeJA and collected for qRT-PCR assays of exogenous phytohormones. The primers used for CsPrx25 detection were FRT (CCCCACTTCGGATCCAGCAAC) and RRT (CAACCCCTGTCGGTTACATCA).

**Southern blot assay**

Total genomic DNA (gDNA) was extracted from the leaves of the transgenic plants and WT plants using a CTAB kit (Zoonbio, China). The gDNA was fragmented using the restriction enzyme EcoRI, and the DNA fragments were separated on a 0.7% agarose gel and transferred to a Hybond-N+ membrane (Amersham, UK). The NPTII coding gene labeled by digoxin (DIG) was used to hybridize the membrane-bound DNA (Roche, Switzerland). The nylon membrane was then exposed using nonradioactive probe detection. In the Southern blot assay, the pLGNe-CsPrx25 plasmid was used as the positive control.

**Assessment of CBC resistance**

CBC resistance analyses were performed as previously described. Briefly, six punctures were made in six healthy mature leaves of each transgenic line via 0.5-mm pins, and 1 μL of each Xcc suspension (1 × 10⁵ cfu mL−1) was subsequently inoculated. CBC development was assessed at 10 dpi, and both the disease severity (DS) and lesion size (LS) of the diseased spots were used for the assessment of CBC resistance. The DS was calculated as
previously described\textsuperscript{14,70}. CBC resistance was further evaluated through Xcc infiltration assays (1 × 10\textsuperscript{5} cfu mL\textsuperscript{-1}), and canker symptoms were imaged at 10 dpi.

**Biochemical analysis**

The activities of CIII Prx, SOD, CAT and GST and the concentrations of H\textsubscript{2}O\textsubscript{2}, O\textsubscript{2}\textsuperscript{−}, MDA, and lignin were measured via SinoBestBio assays (Shanghai, China). The experiments were repeated three times, and the results are shown as the means ± SEs.

RNA isolation, cDNA synthesis and qRT-PCR assay

Miniprep kits (AidLab) were used for RNA isolation, and cDNA was synthesized using TaKaRa kits. qRT-PCR was performed using QuantStudio 7. The values were normalized to the CsActin levels (GenBank accession: GU911361.1, CAP ID: Cs1g05000) obtained using F\textsubscript{Actin} (CATCCCTCAGCACCTTCC) and R\textsubscript{Actin} (CCAACCT TAGCATTCTCC). The qRT-PCR parameters were as follows: 95 °C for 5 min followed by 40 cycles of 95 °C for 10 s and 56 °C for 30 s. The reaction mixtures (total volume of 12 μL) contained 50 ng of cDNA, 0.5 μM primers and 6 μL of the PCR mix. The relative gene expression levels were assessed using the 2\textsuperscript{−ΔΔCT} method\textsuperscript{71}. NCBI was used for qRT-PCR primer design (Supplementary Table S3). The data are presented as the means from three independent biological repeats.

**Statistics**

The data were analyzed using SPSS V22. Gene expression was compared by analysis of variance (ANOVA). The statistical significance was analyzed by Fisher’s LSD test. *P < 0.05 and **P < 0.01 indicate significant and extremely significant differences, respectively. The plant lines were compared using Tukey’s HSD test (P = 0.05).

**Discussion**

CIII Prxs belong to a plant-specific multigene family that promotes disease resistance\textsuperscript{18,33,34}, lignification, the flexibility of cell walls and suberization\textsuperscript{29,30}. In sweet orange, 72 CIII Prxs have been identified\textsuperscript{28}. The expression of each isoform varies across tissues and can be influenced by environmental factors, which suggests that different peroxidase isoenzymes regulate distinct processes\textsuperscript{72}. The distribution of enzymes to either the cell walls or vacuoles and their destinations reflect their specific functions\textsuperscript{31}. In CBC-resistant and CBC-susceptible varieties, CsPrx25 exhibits altered expression patterns (Fig. 2d–f), which suggests its role during CBC development. The importance of CIII Prxs for the resistance of plants to pathogenic diseases was identified through reverse genetics. CIII Prxs mediate innate resistance both passively and actively\textsuperscript{6}. HvPrx40\textsuperscript{40} and TaPrx10\textsuperscript{39,41} enhance the resistance of wheat against wheat powdery mildew. Here, CsPrx25 was found to mediate protection against Xcc pathogenesis, which confirmed its role as a CIII Prx and further highlighted the importance of this family in pathogen immunity in sweet orange. We explored its functional role using overexpression strategies and found that CsPrx25 strongly conferred CBC resistance to the transgenic plants (Fig. 3g–j).

Oxidative bursts, particularly the production of H\textsubscript{2}O\textsubscript{2} and O\textsubscript{2}\textsuperscript{−}, are common innate responses in plant cells in response to pathogen infection\textsuperscript{38}. As key enzymes for ROS homeostasis in plants, CIII Prxs have multiple functions and are proposed to serve as key regulators of the extracellular H\textsubscript{2}O\textsubscript{2} and O\textsubscript{2}\textsuperscript{−} levels depending on peroxidative cycles (ROS scavenging) or hydroxyl cycles (ROS production)\textsuperscript{72}. Plant defense responses are governed by the ROS levels and peroxidase-generated radicals, which mediate cell wall reinforcement, damage repair\textsuperscript{1,6} and apoptotic responses to induce plant resistance\textsuperscript{5,6}. In this study, the molecular mechanisms of CsPrx25 were explored. Based on our analysis of ROS homeostasis and enzymatic antioxidant activities in the transgenic plants, we concluded that CsPrx25 overexpression enhances CIII Prx activities and leads to a simultaneous improvement in the H\textsubscript{2}O\textsubscript{2} and O\textsubscript{2}\textsuperscript{−} content (Fig. 5a, b). In plants, the HR is directly related to plant disease resistance and represents the classic response to pathogen infection\textsuperscript{6}. These reactions lead to both rapid and localized necrosis of the infected tissues and thus prevent the spread of infection\textsuperscript{56,57}. H\textsubscript{2}O\textsubscript{2} is key to the HR and is related to programmed cell death (PCD) in infected plants\textsuperscript{58}. To investigate the relationship between the CBC resistance induced by CsPrx25 and the HR, we assessed the HR of the transgenic plants before and after Xcc infection (Fig. 5d). HSR203 is upregulated by plant HRs and is used as a marker for the HR levels\textsuperscript{56,58}. In this study, the links among CsPrx25 activity, ROS content and HR level were established. Cell wall lignification was further shown to mediate CBC resistance, which was also demonstrated in rice due to the enhancement in Xanthomonas oryzae resistance conferred by CIII Prx-mediated lignification\textsuperscript{73}.

Due to the evolutionary diversity and functional diversity of CIII Prxs, different studies have drawn different links between CIII Prx and disease resistance. Increased LePrx06 makes tomato more susceptible to Pseudomonas syringae infection. In contrast to CsPrx25, the suppression of LePrx06 can enhance resistance to this pathogen\textsuperscript{74}. Long-term studies of the relationship between the ROS levels and the development of CBC have revealed increased peroxidase activity and thus a reduced ROS content. Furthermore, the reduction in the ROS levels was associated with CBC resistance. These effects parallel the overexpression of MdATG18a and can enhance resistance to Diplolaron mali infection via H\textsubscript{2}O\textsubscript{2} scavenging\textsuperscript{75}. Cybrids of grapefruit with a kumquat plastid genome
exhibit increased CBC resistance through an early upregulation of ROS-controlling genes upon Xcc infection.\textsuperscript{76} These findings illustrate potential links between ROS homeostasis mediated by plastid ROS-controlling genes and CBC resistance. However, this study revealed that CsPrx25 is an apoplast-localized enzyme rather than a plastid enzyme (Fig. 2b, c), and this knowledge expands the list of ROS-controlling enzymes that can upregulate CBC resistance.

In this study of CsPrx25, regulation of the ROS levels by CsPrx25 and improvements in HR sensitivity were the major mechanisms through which transgenic citrus developed resistance to CBC. Although CsPrx25 overexpression greatly improved the resistance of Wanjincheng to CBC, CsPrx25-overexpressing Wanjincheng cells were still not as resistant as Calamondin cells, which might be due to the fact that Calamondin also has other mechanisms to maintain an even higher level of CBC resistance. Anyway, this study explores new insights into the mechanisms of CIII Prxs in CBC resistance and provides potential clues for breeding CBC-resistant citrus.

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Author contributions
Q.L. and S.C. conceived the experiments; Q.L., J.Q., X.Q., and W.D. conducted the experiments; Q.L. and C.D. conducted the bioinformatics analysis; Q.L. and W.D. were responsible for the data analyses; and Q.L. and Y.H. wrote the paper.

Conflict of interest
The authors declare that they have no conflict of interest.

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