Identification and Expression Analysis of Multiple Ferric Chelate Reductases in *Citrus junos*

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ABSTRACT. Ferric chelate reductase (FRO) is a critical enzyme for iron absorption in strategy I plants, reducing Fe^{3+} to Fe^{2+} . To identify *FRO* family genes in the local *Citrus junos* cultivar Ziyang Xiangcheng and to reveal their expression model, the citrus (*Citrus* sp.) genome was searched for homologies of the published sequence *CjFRO1*. Five *FROs* were found, including *CjFRO1*; these were named *CjFRO2*, *CjFRO3*, *CjFRO4*, and *CjFRO5*, respectively, and cloned via reverse transcription polymerase chain reaction (RT-PCR) and rapid amplification of cDNA ends (RACE) PCR. The deduced amino acid sequences of five *CjFROs* contained flavin adenine dinucleotide (FAD)-binding motifs, nicotinamide adenine dinucleotide (NAD)-binding motifs, and 6–10 transmembrane domains, with isoelectric points between 6.73 and 9.46, and molecular weights between 67.2 and 79.9 kD. *CjFRO1* and *CjFRO2* were predominantly found in the aboveground parts of *C. junos*, with *CjFRO1* highly expressed in leaves, and *CjFRO2* largely expressed in stems and leaves. *CjFRO3* was less expressed in roots, stems, and leaves. *CjFRO4* and *CjFRO5* were predominately found in roots. Under iron-deficient conditions, *CjFRO4* was significantly and specifically increased in the roots of *C. junos*, whereas *CjFRO1* was upregulated in the roots and leaves.

Iron is an essential element for plant growth, which is involved in numerous vital biological processes, including chlorophyll biosynthesis, photosynthetic electron transfer, DNA synthesis, and nitrogen reduction. Although abundant in soil, iron is one of the most common nutrients to limit plant growth and development because of its particularly low solubility in aerobic soils at neutral and alkaline pH (Mimmo et al., 2014). Based on their mechanism for Fe acquisition, plants can be categorized into strategy I or strategy II plants (Eide et al., 1996; Robinson et al., 1999; Romheld, 1987; Romheld and Marschner, 1986). Strategy II plants include grasses with roots that secrete compounds known as phytosiderophores (PS), which chelate Fe^{3+} from the rhizosphere. Subsequently, the yellow stripe 1 carrier protein in the plasmalemma introduced the Fe^{3+} -PS complex into the root cell. By contrast, strategy I plants are all dicots and non-graminaceous monocots, and the acquisition of Fe^{3+} occurs in reducing reaction processes: 1) proton excretion via a self-phosphalated type of adenosine triphosphatase, thus acidifying the surrounding soil to increase iron solubility; 2) reduction of Fe^{3+} to Fe^{2+} via FRO; and 3) transport of Fe^{2+} via iron-regulated transporter through the plasmalemma membrane.

The FRO protein is a key enzyme for Fe^{3+} acquisition from soil, which has been confirmed through its functionality for chlorosis tolerance and RNA-interference in plants (Gama et al., 2017; Lee et al., 2016; Martínez-Cuenca et al., 2016).

The first *FRO* gene was identified in the model plant *Arabidopsis thaliana* (Robinson et al., 1999). Further *FRO* genes have been reported in a variety of plants such as *Arachis hypogaea* (Ding et al., 2009), *Cucumis sativus* (Waters et al., 2007), *Lotus japonicus* (Klein et al., 2012), *Malus xiaojinensis* (Wu et al., 2012), *Medicago truncatula* (Andaluz et al., 2009; Orozco-Mosqueda et al., 2012), *Oryza sativa* (Gross et al., 2003), *Pisum sativum* (Waters et al., 2002), and *Solanum lycopersicum* (Kong et al., 2013; Li et al., 2004). Eight *FROs* have been described in the genome of *A. thaliana* and shown to exhibit tissue-specific expression (Mukherjee et al., 2006; Wu et al., 2005). *FRO* family members regulate metal ion homeostasis in different locations in the plant. *AtFRO2* is mainly expressed in epidermal cells of roots where it participates in the reduction of Fe^{3+} to Fe^{2+} in the rhizosphere. *AtFRO3* is predominantly expressed in the vascular cylinder of roots where it might be involved in the transport of Fe. *AtFRO7* is located in the chloroplast membrane, retaining iron homeostasis in the chloroplast (Connolly et al., 2003; Jeong et al., 2008; Jeong and Connolly, 2009; Mukherjee et al., 2006). Furthermore, *FROs* have been reported to have different regulatory mechanisms, although their overexpression in plants can rescue lime-induced chlorosis (Connolly et al., 2003; Li et al., 2011). *AtFRO2* has been reported to be positively regulated by transcriptional factor FIT (FER-like iron-deficiency-induced transcription factor), and POPEYE (bHLH047) has been shown to upregulate the expression of *AtFRO3* (Hindt and Guerinot, 2012; Long et al., 2010).

The local *Citrus junos* cultivar Ziyang Xiangcheng was reported to be tolerant to lime-induced iron chlorosis in both field and laboratory experiments, and it could efficiently uptake iron from calcareous soils. Until now, only *CjFRO1* (GenBank

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no. DQ985810) had been cloned from *C. junos*. To explore whether additional *FRO* genes were present and functional in the *C. junos* genome, an in silico search was conducted for genes that encode *FRO* functions. With the exception of *CjFRO1*, the results showed that at least four additional novel genes were present, which indicate features of characterized *FRO* proteins. These features include the existence of highly conserved FAD-binding motif, NAD-binding motif, transmembrane regions, and oxidoreductase signature motifs (Andaluz et al., 2009; Mukherjee et al., 2006). In addition, *CjFRO* gene expression was confirmed via real-time PCR in different tissues and in conditions of Fe sufficiency and deficiency. The identification of the *CjFRO* gene family enables an improved understanding of Fe uptake, translocation, distribution, and usage mechanisms in citrus plants.

Materials and Methods

IN SILICO SEARCH AND IDENTIFICATION OF *FRO* GENE COPIES. To identify unknown *FRO* members, a homology Basic Local Alignment Search Tool (BLAST) search was performed, employing the previously identified *CjFRO1* gene (GenBank no. DQ985810). The BLASTs were conducted in the database of the *Citrus sinensis* Genome Sequencing Resources (Xu et al., 2012). Based on the predicted cDNA sequences of *FROs* in the database, sense and antisense primers for each *CjFRO* were designed, and the coding sequences were amplified from the total RNA of 1-year-old ‘Ziyang Xiangcheng’ seedlings via RT-PCR or RACE (634858; Clontech, Beijing, China). They were purified with the Biospin Gel Extraction Kit (BSC02M1; Bioer Technology Co., Hangzhou, China), subsequently cloned into a pMD19-T vector (6013; TaKaRa, Beijing, China), and sequenced. The sequences were predicted with the software packages Protparam (Gasteiger et al., 2005) and Pfam (Finn et al., 2016). Transmembrane domains of putative *CjFRO* proteins were predicted with the TMHMM program (Technical University of Denmark, 2017). Multiple alignments of sequences were performed using the ClustalX2.1 software package (Thompson

et al., 2002), and the phylogenetic tree was obtained with the software MEGA 5.0 (Tamura et al., 2011).

PLANT MATERIALS AND GROWTH CONDITIONS. ‘Ziyang Xiangcheng’ seeds were germinated on wet filter paper in the dark for 1 week and were planted in jars with 1/2 Hoagland’s solution. The nutrient solution was replaced every 2 weeks. After 2 months, the seedlings were transferred into either iron-sufficient (50 μ M FeEDTA) or iron-deficient (0 μ M FeEDTA) Hoagland’s solutions for 2 d because of the *FRO* enzyme activity and gene expression could be induced after 2 d iron starvation in other plants (Mukherjee et al., 2006; Wu et al., 2012). Then, roots and leaves were harvested for iron starvation analysis.

One-year-old ‘Ziyang Xiangcheng’ seedlings were grown in the germplasm resources garden of the Citrus Research Institute (Chongqing, China). For analyzing the *CjFROs* expression in different tissues, the root tips (1.5 cm), stems (between fifth and sixth leaves), and the first fully developed leaves were collected from five plants.

RNA EXTRACTION. Total RNA was extracted with the RNA Extraction Kit (RN0902; Aidlab Biotechnologies Co., Beijing, China) and the RNA was treated with RNase-free DNase I to remove residual genomic DNA. RNA samples were run in a 1.2% agarose gel and stained with ethidium bromide to verify RNA quality. Each RNA sample was quantified with a spectrophotometer (NanoDrop 2000c; Thermo Fisher Scientific, Shanghai, China) at 260 nm.

FULL-SEQUENCE CLONING OF *CjFROs*. Full sequences were obtained via PCR using primer pairs listed in Supplemental Table 1. The PCR conditions were 95 °C for 1 min; 94 °C for 30 s, 58 °C for 30 s, 72 °C for 2 min 30 s for 40 cycles; followed by 72 °C 10 min. The PCR products were cloned into the pMD19-T vector and transformed into competent *Escherichia coli* DH5 α cells. PCR confirmed positive clones were sequenced in the Beijing Genomics Institute (Shenzhen, China).

REAL-TIME PCR. cDNA was prepared from 1 μ g total RNA with the iScriptTM cDNA Synthesis Kit following the manufacturer’s instructions (170–8891; Bio-Rad Laboratories, Hercules, CA). Amplification with the primers in Supplemental

Fig. 1. Polymerase chain reaction (PCR) results of ferric chelate reductase genes of *Citrus junos* (*CjFROs*). *CjFRO2* (A), 5' terminal PCR products of *CjFRO3* (B), and *CjFRO4* (C), were cloned with primers FROafS/FROafR, FROcfS2/FROcfR2, FROdfS/FROdfR, respectively. *CjFRO5* (D) were cloned with two primer pairs FROedS/FROeqR3 (lane 1) and FROeqS3/FROefR. The sequences of primer pairs could be found in Supplemental Table 1. Lane M was DNA Marker DL 2000 (2000, 1000, 750, 500, 250, and 100 bp).

Table 1. Summarized features of the ferric chelate reductase genes of *Citrus junos* (*CjFROs*) from ‘Ziyang Xiangcheng’ according to their nucleic acid sequences and amino acid sequences.

Name	Nucleic acids (no.)	Amino acids (no.)	MW (kD) ^z	pI ^y	Transmembrane regions (no.)	Introns (no.)	Exons (no.)
<i>CjFRO1</i>	2103	700	79.7	9.23	10	13	14
<i>CjFRO2</i>	2097	698	77.3	6.73	6	7	8
<i>CjFRO3</i>	2112	703	79.9	9.46	10	7	8
<i>CjFRO4</i>	1809	602	67.2	9.10	6	5	6
<i>CjFRO5</i>	2112	703	79.2	9.06	9	7	8

^zMolecular weight.

^yIsoelectric point.

Table 2 was performed using the SYBR Green PCR Master Mix (1725271; Bio-Rad Laboratories) and the iCycler iQ Multicolor Real-Time PCR Detection System (Bio-Rad Laboratories). The following protocol was used for amplification: 95 °C for 1 min; 94 °C for 10 s, 60 °C for 15 s, and 72 °C for 15 s, for 40 cycles. After completion of the amplification reaction, melt curves were compared to verify the amplification of a single peak. Each experiment was performed in at least three technical replications and three biological replicates were run per sample (using three independent sets of RNA). The data analysis used the 2^{-ΔCt} method, and *actin* was used as a reference gene.

Results

IDENTIFICATION AND BIOINFORMATIC ANALYSIS OF *CjFROs*.

The genome sequence of citrus was searched through BLASTx against the amino acid sequence of *CjFRO1*. Five predicted sequences of *C. sinensis* (Cs3g01120, Cs9g19350, Cs5g06040, orange1.1t00399, and Cs9g19360) were found. Cs9g19350 corresponds to *CjFRO1*, and the other genes of ‘Ziyang Xiangcheng’ were designated as *CjFRO2* (Cs3g01120), *CjFRO3* (Cs5g06040), *CjFRO4* (orange1.1t00399), and *CjFRO5* (Cs9g19360). These revealed high-sequence identity among each other at the amino acid level (Supplemental Table 3).

Based on the predicted open reading frame sequences of the citrus genome database, sense and antisense primers were designed for all four novel *CjFRO* genes, and the cDNAs of these genes were amplified with the primers listed in the Supplemental Table 1, using total RNA that was isolated from 1-year-old ‘Ziyang Xiangcheng’ seedlings. Via RT-PCR, *CjFRO2* was obtained with the primer pairs FROafS/FROasR of leaves, and *CjFRO4* was obtained with FROdfS/FROdfR of roots. For the remaining two *FRO* genes, it was difficult to obtain the whole open-reading-frame sequence. The upstream nucleic acid sequence of *CjFRO5* was obtained via PCR with the primer pair FROefS/FROeqR3, and the downstream sequence was obtained with the primer pair FROeqS3/FROefR. For *CjFRO3*, the upstream nucleic acid sequence was obtained via PCR, using the primer pair FROcfS2/FROcfR2, and the downstream sequence was obtained via 3′ RACE with the primer pairs FROcfS3/3′ RACE outer primer and the FROcfS4/3′ RACE inner primer, which were subsequently merged (Fig. 1). These all contained the conserved features of Fe(III) chelate reductases described by Waters et al. (2002) and were classified into the *FRO* gene family of citrus. Analysis with the TMHMM program indicated that the deduced amino acid sequences of *CjFROs* contained six (*CjFRO2* and *CjFRO4*), nine (*CjFRO5*), or ten (*CjFRO1* and *CjFRO3*) integral transmembrane domains. Most *CjFROs* belong to the

basic protein excluding *CjFRO2*, which is almost neutral with an isoelectric point of 6.73 (Table 1).

A neighbor-joining tree was constructed based on the deduced amino acid sequences of five *CjFROs* from citrus, eight *AtFROs* from *A. thaliana* (Wu et al., 2005), two *MtFROs* from *M. truncatula* (Andaluz et al., 2009), *LjFRO1* from *L. japonicus* (Klein et al., 2012), *LeFRO1* from *S. lycopersicum* (Kong et al., 2013), *MxFRO* from *M. xiaojinensis* (Wu et al., 2012), and *PsFRO1* from *P. sativum* (Waters et al., 2002). The phylogenetic relationship of these five *CjFROs* could be clustered into four groups (Fig. 2). *CjFRO4* and *CjFRO5* were closer related to the well-characterized Fe(III) chelate reductase genes *LeFRO1* from a herbaceous plant and *MxFRO* from a woody plant, respectively, all of which could be grouped with *AtFRO2* and *AtFRO3* of *A. thaliana*. *CjFRO1* clustered with

Fig. 2. Phylogenetic relationship of plant ferric chelate reductases (FRO). The complete predicted protein sequences of the five *CjFROs* in *Citrus junos* ‘Ziyang Xiangcheng’ were aligned with the other plant *FROs* through MEGA 5.0 (Tamura et al., 2011). *AtFROs* from *Arabidopsis thaliana* (*AtFRO1*, NM_100041; *AtFRO2*, NM_100040; *AtFRO3*, NM_102150; *AtFRO4*, NM_122303; *AtFRO5*, NM_122304; *AtFRO6*, NM_124351; *AtFRO7*, NM_124352; *AtFRO8*, NM_124395), *MtFROs* from *Medicago truncatula* (*MtFRO1*, AY439088; *MtFRO2*, XM_003594382), *LjFRO1* from *Lotus japonicus* (JF290367), *LeFRO1* from *Solanum lycopersicum* (AY224079), *MxFRO* from *Malus xiaojinensis* (EF577061), and *PsFRO1* from *Pisum sativum* (AF405422). The bootstrap *P* values were obtained by the neighbor-joining method and indicate the divergence of each branch. *FROs* from *C. junos* were highlighted with diamonds on the tree.

Fig. 3. The expression of ferric chelate reductase genes of *Citrus junos* (*CjFROs*) in different tissues of 1-year-old 'Ziyang Xiangcheng' seedlings. The transcript levels of *CjFROs* in citrus seedlings were detected using the real-time polymerase chain reaction. The relative expression level value was normalized to an internal standard *actin*. Each histogram represents the mean value of three independent experiments, and the vertical bars indicate SE. Dashed frame enlarges the results of the lower-expressed *CjFROs* in the dotted rectangle.

AtFRO4 and *AtFRO5*; *CjFRO3* grouped with *AtFRO8*; *CjFRO2* shared a group with *AtFRO6* and *AtFRO7*.

EXPRESSION PROFILES OF CjFROs IN CITRUS TISSUES. Real-time PCR was performed to evaluate the expression of all five *CjFROs* in roots, stems, and leaves of 1-year-old 'Ziyang Xiangcheng' seedlings. The sense terminal primers and anti-sense terminal primers for real-time PCR (Supplemental Table 2) were designed in different exons, to avoid genomic DNA contamination. According to tissue expression (Fig. 3), *CjFRO2* and *CjFRO1* were expressed at higher levels in stems and leaves. Compared with *CjFRO2* expression in roots, its accumulation increased 396-fold in stems and 189-fold in

leaves. *CjFRO1* expression in stems and leaves was 21-fold and 296-fold higher than in roots. However, *CjFRO4* expression in roots was 31-fold higher than in stems and 7-fold higher than in leaves. *CjFRO5* was a gene with a root-specific expression and was barely detectable by real-time PCR in stems and leaves. *CjFRO3* showed lower expression in roots, stems, and leaves.

CjFROs RESPONSE TO THE Fe DEFICIENT CONDITION. To analyze *CjFROs* expression in both Fe sufficiency and Fe deficiency conditions, citrus roots and leaves of 2-month-old seedlings, cultured in Hoagland's nutrient solution were tested via real-time PCR (Fig. 4). After 2 d of iron deficiency stress, *CjFRO1* was upregulated 3.0-fold in roots and 1.9-fold in leaves. The expression *CjFRO2* and *CjFRO3* remained unchanged in roots and leaves (change fold ≤ 1.5). *CjFRO4* showed a specifically and significantly higher accumulation in roots (4.5-fold) under the Fe starvation and not changed in the leaves. *CjFRO5* were not detectable in leaves and roots of 2-month-old seedlings under Fe-sufficient and Fe-starvation conditions.

Discussion

Online plant genomics offer greatly valuable databases for the identification and further characterization of genes that are involved in diverse cellular processes, such as Fe-starvation responses. Eight *FROs* had been found in the *A. thaliana* genome and six *FROs* in the *M. truncatula* according to their genomic databases (Orozco-Mosqueda et al., 2012; Wu et al., 2005). Just like *A. thaliana* and *M. truncatula*, *C. junos* belongs to the plants using strategy I for Fe acquisition, which can reduce rhizospheric iron, thus increasing the bioavailability of iron. In the present in silico study, four putative sequences with high identity to *FROs* of other plants were identified and cloned (apart from *CjFRO1*, which was previously described.). All sequences described herein showed highly conserved characteristics of *FRO* proteins, such as FAD and NAD binding domains, transmembrane regions, and oxidoreductase signature motifs. *FROs* are present in various plant species and are

Fig. 4. Real-time polymerase chain reaction (PCR) analysis of *Citrus junos* ferric chelate reductase (*FRO*) gene-expression in root (left) and leaf (right) of iron-sufficient and iron-deficient plants. The seedlings were initially cultured for 2 months in Hoagland's solution and then transferred to either iron-sufficient (50 μ M FeEDTA) or iron-deficient (0 μ M FeEDTA) Hoagland's solution for 2 d of treatment. The *FRO* relative expression level value was normalized to an internal standard *actin*. Each histogram represents the mean value of three independent experiments, and the vertical bars indicate SE.

responsible for the reduction of Fe^{3+} to Fe^{2+} , a more soluble form that can be absorbed by the roots. Gene expression and posttranslation of *FRO2* is positively regulated by carbon monoxide, nitric oxide, ethylene, and bHLH transcription factors on Fe-deficient conditions (Du et al., 2015; Hindt and Gueriot, 2012; Kabir et al., 2012; Yang et al., 2016; Yuan et al., 2008; Zhai et al., 2016).

In plants, *AtFRO2* was the first FRO gene that was characterized (Robinson et al., 1999). Subsequently, seven other functional members of the same family have been identified in the model plant *Arabidopsis* (Mukherjee et al., 2006; Wu et al., 2005). Similarly, six *MtFROs* were reported for *M. truncatula* (Orozco-Mosqueda et al., 2012) and five *CjFROs* for *C. junos* in our experiment. The reason why plants contain multiple *FRO* genes in their genomes has already been summarized in excellent reviews: *FROs* are well known for their role in iron uptake by roots and furthermore, may play critical roles in the iron uptake by leaves as well as for iron homeostasis in chloroplasts and mitochondria (Jain et al., 2014; Jeong and Connolly, 2009). Moreover, *FROs* are not only found in strategy I plants, but also in strategy II plants, such as *O. sativa* that contains two *FRO* genes (Ishimaru et al., 2007). The *O. sativa FRO1* mutant revealed that *FRO* might have a new function and might be involved in iron trafficking under Fe-toxic conditions (Siriphat et al., 2015).

In this study, multiple members of *FROs* in *C. junos* were found to be present in roots and shoots and are induced by Fe deficiency. These results agree with *FRO* studies in *A. thaliana*, which demonstrated that genes such as *AtFRO2* and *AtFRO3* are mainly expressed in roots under low-Fe conditions, whereas *AtFRO5*, *AtFRO6*, *AtFRO7*, and *AtFRO8* are exclusively found in shoots and not regulated by Fe supply (Wu et al., 2005). Furthermore, the expression of each gene is tissue specific; for example, *AtFRO2* and *AtFRO3* are mainly expressed in roots, whereas *AtFRO5* and *AtFRO6* were detected in shoots and flowers, and *AtFRO7* in cotyledons and trichomes. *AtFRO6* and *AtFRO7* were located on plasma membranes and chloroplast membranes, respectively, whereas *AtFRO8* was specifically expressed in the veins of leaves (Jeong et al., 2008; Wu et al., 2005). In *C. junos*, *CjFRO4* and *CjFRO5* were clustered with the well-characterized plant *FROs*, which were highly or specifically expressed in the roots. *CjFRO1–3* was clustered with the shoot-specific-expressed *AtFROs* (Fig. 2). The real-time PCR analysis for the five *CjFROs* confirmed the result of cluster grouping (Fig. 3). *CjFRO4* accumulated significantly higher levels in roots under iron starvation (Fig. 4). The lack of *CjFRO5* observation in leaves and roots was unexpected, although it was highly expressed in the roots of 1-year-old ‘Ziyang Xiangcheng’ seedlings. This phenomenon has been described in *Arabidopsis* for *AtFRO3*, which was expressed at high levels in seedlings, but at lower levels in mature, soil-grown plants (Mukherjee et al., 2006).

In conclusion, at least five *FRO* genes have been found within the citrus genome. Here, we detected the four novel genes *CjFRO2*, *CjFRO3*, *CjFRO4*, and *CjFRO5*, which had the transmembrane domain, FAD-binding domain, and NAD binding domain of ferric reductase. The genes *CjFRO5* and *CjFRO4* were highly expressed in roots, whereas *CjFRO2* and *CjFRO1* were highly expressed in leaves. *CjFRO3* had lower expression in roots, stems, and leaves. Fe stress significantly increased *CjFRO4* expression in roots.

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Supplemental Table 1. Primers used for full cDNA sequence cloning of ferric chelate reductase genes of *Citrus junos* (*CjFROs*). FROafS and FROafR were used for *CjFRO2* PCR. Primer pairs FROcfS2/FROcfR2 were used to obtain the upstream sequence of *CjFRO3*, FROcfS3/3' RACE outer primer and FROcfS4/3' RACE inner primer for the downstream sequence of *CjFRO3*. FROdfS and FROdfR were used for *CjFRO4* PCR. Primer pairs FROefS/FROeqR3 and FROeqS3/FROefR were used to obtain the upstream and downstream sequences of *CjFRO5*, respectively.

Genes	Sense primers	Antisense primers
<i>CjFRO2</i>	FROafS: 5'-ATGTTGGAAGTACGCGGACTT-3'	FROafR: 5'-CTACAGATCAAACTATGGCTGTGG-3'
<i>CjFRO3</i>	FROcfS2: 5'-ATGTTGATGATTTTGATATGCGCTG-3'	FROcfR2: 5'-AATCTGTATTTACGGTTGCTCTGTGC-3'
	FROcfS3: 5'-AGCATGCAGGCTTAAAGTTCACT-3'	3' RACE outer primer: 5'-TACCGTCGTTCCACTAGT ATTT-3'
	FROcfS4: 5'-CGAGTATTTCCAAATTTCACTGGCA-3'	3' RACE inner primer: 5'-CGCGGATCCTCCACTAGT ATTTCACTATAGG-3'
<i>CjFRO4</i>	FROdfS: 5'-ATGCTGGTGAAAGGCCCACT-3'	FROdfR: 5'-TCACCAGCTAAAGCTGATAGATTC-3'
<i>CjFRO5</i>	FROefS: 5'-ATGGAGACCATTAAATCAGCTATAAAG-3'	FROeqR3: 5'-ATTATTGGCCAAAGAGAGCAGC-3'
	FROeqS3: 5'-CCATTTTACAGGGCACGACAT-3'	FROefR: 5'-TCACCAGCTAAAGGCTATGGATT-3'

PCR = polymerase chain reaction; RACE = rapid amplification of cDNA ends.

Supplemental Table 2. Primers used for analyzing the expression of *Citrus junos* ferric chelate reductase genes (*CjFROs*) in different tissues and responding to the iron starvation.

Genes	Sense primers	Antisense primers	cDNA length (bp)
<i>CjFRO2</i>	5'-AGGCACCCTTTACTACCTCCTAC-3'	5'-GACAAGACTTCCCTTCGTTGATA-3'	181
<i>CjFRO1</i>	5'-CATGCTGGGCATTCTCTTCTT-3'	5'-GTTTCGCTCCATTCTAGCACCT-3'	202
<i>CjFRO3</i>	5'-CCAGCTTTGAGATTCTGTTGGT-3'	5'-GGTACTTGAGTTGCCATGTGTT-3'	135
<i>CjFRO4</i>	5'-GACCTGAGCTAAAGAGGATGCT-3'	5'-GCCAGACCAGATGAACAGATTG-3'	115
<i>CjFRO5</i>	5'-CCATTTTACAGGGCACGACAT-3'	5'-ATTATTGGCCAAAGAGAGCAGC-3'	100
<i>actin</i>	5'-CATCCCTCAGCACCTTCC-3'	5'-CCAACCTTAGCACTTCTCC-3'	198

Supplemental Table 3. Distributions of percentage protein sequence identity for pairs of *Citrus junos* ferric chelate reductases (*CjFROs*).

	CjFRO1	CjFRO2	CjFRO3	CjFRO4	CjFRO5
CjFRO1		35	30	58	54
CjFRO2			32	35	33
CjFRO3				32	31
CjFRO4					69
CjFRO5					