Identification, characterization and expression analysis of the VQ motif-containing gene family in tea plant (*Camellia sinensis*)

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**Abstract**

**Background:** VQ motif-containing (VQ) proteins are plant-specific proteins that interact with WRKY transcription factors and play important roles in plant growth, development and stress response. To date, VQ gene families have been identified and characterized in many plant species, including Arabidopsis, rice and grapevine. However, the VQ gene family in tea plant has not been reported, and the biological functions of this family remain unknown.

**Results:** In total, 25 *CsVQ* genes were identified based on the genome and transcriptome of tea plant, and a comprehensive bioinformatics analysis was performed. The *CsVQ* proteins all contained the typical conserved motif FxxhVQxhTG, and most proteins were localized in the nucleus. The phylogenetic analysis showed that the VQ proteins were classified into 5 groups (I, III-VI); the evolution of the *CsVQ* proteins is consistent with the evolutionary process of plants, and close proteins shared similar structures and functions. In addition, the expression analysis revealed that the *CsVQ* genes play important roles in the process of tea plant growth, development and response to salt and drought stress. Furthermore, a potential regulatory network including the interactions of *CsVQ* proteins with *CsWRKY* transcription factors and the regulation of upstream microRNA that is closely related to the above-mentioned processes is proposed.

**Conclusions:** The results of this study increase our understanding and characterization of *CsVQ* genes and their encoded proteins in tea plant. This systematic analysis provided comprehensive information for further studies investigating the biological functions of *CsVQ* proteins in various developmental processes of tea plants.

**Keywords:** Tea plant, VQ gene, Expression profiling, Salt stress, Drought stress

**Background**

The tea plant [*Camellia sinensis* (L.) O. Kuntze] is an important economic crop worldwide that experiences various abiotic stresses during its lifecycle, such as drought, salinity, heavy-metals and soil nutrient deficiency [1]. Generally, plants have developed a series of elaborate mechanisms for adaptation to stress conditions, and many transcription factors, such as WRKY, MYB, NAC, AP2/ERF, bHLH and bZIP, have been implicated in responses to various types of stresses [2]. Among these transcription factors, WRKY has been identified in many plants species, including Arabidopsis, rice, grapevine and tea plants [3–5], and many WRKY transcription factors have been confirmed to be involved in the response to abiotic stresses as regulators of gene expression [6]. In addition, studies conducted over the prior several years have revealed that WRKY transcription factors physically interact with a wide range of proteins that play important roles in plant responses to various stresses [7], which has become a hotspot in the study of WRKY biological functions. More recently, several studies have reported that VQ motif-containing (VQ) proteins can interact with WRKY transcription factors that play crucial roles in plant growth, development,
and stress responses [8]. The VQ proteins constitute a class of plant-specific proteins with a highly conserved and single short FxxhVQxhTG (h: hydrophobic amino acid, x: any amino acid) amino acid sequence motif that includes five conserved amino acids and interacts with WRKY transcription factors via the conserved V and Q residues [9]. Initially, a VQ protein named MKS1 was reported to interact with WRKY25 and WRKY33 to regulate growth and defence responses in Arabidopsis [10]. Subsequently, many interactions among different VQ proteins and WRKY transcription factors were confirmed to be highly important for the growth and development of plants, such as the interaction between AtVQ14 and AtWRKY10, which determines endosperm growth and seed size [11, 12], and AtVQ20 acts as a vital partner of AtWRKY2 and AtWRKY34 in the regulation of male gametogenesis in Arabidopsis [13]. In addition, evidence suggests that mitogen-activated protein kinase (MAPK) is involved in the interaction between VQ proteins and WRKY proteins, which form a trimeric complex that plays important roles in the growth process in plants [14]. For example, MAPK4 can bridge AtVQ21 and WRKY33 to influence plant growth and disease resistance [15, 16], and MAPK3 and MAPK6 participate in the interaction between AtVQ4 and multiple WRKY transcription factors to regulate immune responses in Arabidopsis [11]. Recently, many investigations have revealed that VQ proteins play crucial roles in the plant response to pathogen infections; for example, the AtVQ16 and AtVQ23 genes were induced by Botrytis cinerea, and disease resistance was significantly enhanced in over-expressed plants [17]. In contrast, AtVQ12 and AtVQ29 are negative regulators of the Arabidopsis response to B. cinerea [18, 19]. Furthermore, VQ proteins have been shown to participate in plant abiotic stress responses; for example, the Arabidopsis VQ9 protein acts as a repressor of the WRKY8 transcription factor to maintain an appropriate balance of WRKY8-mediated signalling pathways to establish salt stress tolerance [20], and AtVQ15 negatively regulates the tolerance of Arabidopsis to salt and osmotic stress [21]. However, reports investigating the functions of VQ proteins outside of model plants are lacking.

Recently, VQ gene families have been increasingly identified in diverse plants, including rice, grapevine, maize and soybean [9, 22, 23], and the expression patterns of the VQ genes have been analyzed in response to salt stress, drought stress, pathogen infection, low nitrogen and tissue-specific stimuli [8, 24]. However, the VQ gene family in tea plant has not been reported to date, and the biological functions of this family remain unknown, particularly in the tea plant’s response to abiotic stresses. In this study, 25 CsVQ genes were identified based on genome and transcriptome databases of tea plant, and a comprehensive analysis was performed, including a multiple sequence alignment, phylogenetic analysis, conserved structural motif analysis, functional interaction network analysis and microRNA prediction. In addition, the expression profiles of the CsVQ genes were examined in different tissues during tea leaf development and in response to salt and drought stresses. The results of this study reveal the molecular foundation of the CsVQ gene family and may serve as a theoretical basis for future studies aiming to elucidate the biological functions of CsVQ proteins under abiotic stresses.

**Results**

**Identification and characterization of VQ genes in tea plant**

To identify the complete VQ gene family in tea plant, 108 VQ motif sequences conserved in plants were used as queries to search against the genome and transcriptome database of tea plant using the BLASTP program. In total, 25 CsVQ genes were identified and named CsVQ1 to CsVQ25 (Additional file 1: Table S1). The results of the protein sequence analysis showed that all CsVQ proteins contained the typical conserved motif FxxhVQxhTG (Fig. 1a), and 2 h amino acid residue sites were highly conserved, particularly the second h site, which is almost entirely Leucine (Fig. 1b). In addition, the physiological and biochemical properties of the 25 CsVQ proteins were determined by computing different parameters, including a length range from 89 to 433 amino acids (aa), a molecular weight range from 9.118 to 45.748 kDa, and a theoretical isoelectric point (pI) range from 4.87 to 11.12. Furthermore, bioinformatics predictions of the subcellular localization of the CsVQ proteins were performed, and the results showed that most CsVQ proteins are localized in the nucleus, while a few proteins are localized in the chloroplast or nucleus. And more detailed information, including the grand average of hydropathicity (GRAVY), instability index and aliphatic index, is listed in Table 1.

**Phylogenetic analysis of CsVQ proteins**

To obtain the subfamily classification and evolutionary relationships of the 25 CsVQ proteins, a phylogenetic tree was constructed using the neighbour-joining method (Fig. 2a). The VQ proteins were classified into six groups, and the 25 CsVQs appeared in only five groups. Most CsVQs belonged to group VI, followed by groups IV, I, III and V, and the protein sequences in the same group have a higher homology, meaning they have similar origins and evolutionary relationships. Similarly, the VQ-containing motif of CsVQ proteins in the same group also has extremely high conservatism, which is consistent with the evolutionary relationship of CsVQ proteins (Fig. 2b). In addition, the evolutionary relationships show that the CsVQ proteins have a close affinity...
with the Arabidopsis VQ proteins and a distant affinity
with the rice VQ proteins in the same group.

Conserved motif analysis of CsVQ proteins
To investigate the sequence features of the CsVQ proteins, their conserved motifs were predicted and analysed using the MEME tool. According to the results, 20 motifs describing the details of the CsVQ proteins were predicted (Fig. 3 and Additional file 2: Figure S1), and the regular expression levels of the conserved motifs are listed in Additional file 3: Table S2. Motif 1 corresponds to the VQ-containing motif distributed in all CsVQ proteins, and Motif 3, Motif 15 and Motif 17 are distributed across 16, 15 and 16 CsVQ proteins, respectively. In addition, CsVQ proteins belonging to the same group contain similar motifs, which is consistent with the results of the phylogenetic analysis. Conversely, the type and number of motifs in the different group CsVQ proteins have significant differences that represent the structural basis for the diversity in protein function.

Interaction network of CsVQ proteins
In this study, the functional and physical interactions of the CsVQ proteins were validated by constructing an Arabidopsis association model using STRING software. The results showed that 25 CsVQ proteins associated with 10 known Arabidopsis proteins participated in the interaction network (Additional file 7: Table S6). Several key nodes, including AT1G78310 (corresponding to CsVQ15, CsVQ16 and CsVQ17), AT1G78410 (corresponding to CsVQ3 and CsVQ18), AT1G17147 (corresponding to CsVQ1 and CsVQ2), MKS1 (corresponding to CsVQ14 and CsVQ17) and IKU1 (corresponding to CsVQ15), closely interact with different WRKY transcription factors (Fig. 4). In addition, a complicated interaction was observed between CsVQ14 and CsVQ23 (MKS1) and WRKY25, WRKY33 and MPK4. Further, the tea plant WRKY (CsWRKY) and CsVQ proteins were respectively mapped to the WRKY and VQ proteins of Arabidopsis thaliana (Additional file 8: Table S7 and Additional file 9: Table S8), and the interaction between CsVQs and CsWRKYS were forecasted based on the PAIR tool. As shown in Fig. 5a, 8 CsWRKYS are presumed to interact with different CsVQ proteins, including CsWRKY3, CsWRKY6, CsWRKY12, CsWRKY14, CsWRKY16, CsWRKY20, CsWRKY25 and CsWRKY33. The interaction between CsWRKYS and CsWRKYs belongs to group IIc WRKY transcription factor and the others belong to the group I WRKY transcription factor (Additional file 10: Figure S2). Sequence analysis results showed that the core binding domain of the above CsWRKY proteins is highly consistent with the corresponding Arabidopsis WRKY proteins that has been proven to be the key to the interaction between VQ and WRKY proteins (Fig. 5b).

Expression profiles of CsVQ genes in response to salt and drought stresses
To understand the response of the CsVQ genes to salt and drought stresses, the expression levels of all CsVQ
genes were detected by performing qRT-PCR. Under salt stress, the expression of eight CsVQ genes (i.e., CsVQ4, CsVQ14, CsVQ15, CsVQ16, CsVQ18, CsVQ20, CsVQ21 and CsVQ25) was significantly up-regulated, although the expression levels of certain genes briefly decreased at 2 and 24 h; the expression of six CsVQ genes (i.e., CsVQ1, CsVQ2 CsVQ3, CsVQ6, CsVQ17 and CsVQ19) was up-regulated at 4, 12 and 48 h and down-regulated at the other time points; and the expression of the remaining 11 CsVQ genes, including CsVQ23, CsVQ24 and most group VI members, was significantly down-regulated (Fig. 6a, Additional file 11: Table S9). Interestingly, most CsVQ genes were up-regulated after the exposure to drought stress, although the expression levels of certain genes began to decrease at 24 and 48 h (Fig. 6b, Additional file 12: Table S10). In addition, the expression trends in genes belonging the same group, particularly genes belonging to groups I, IV and VI, were basically consistent under the same stress. Moreover, the expression trends of the genes in groups IV and V were identical under different stresses, but the opposite was observed in genes belonging to groups III and VI, including CsVQ14 and CsVQ23 (Fig. 6). These results reveal that almost all CsVQ genes are involved in the response of tea plant to salt and drought stress and that the response mechanism is complex and diverse.

**Expression profiles of CsVQ genes in different tissues of tea plant**

To elucidate the tissue-specific expression profiles of the CsVQ genes, their expression levels in four tissues were examined using qRT-PCR. The CsVQ genes exhibited differential expression in the root, stem, leaf and flower of tea plant (Fig. 7, Additional file 13: Table S11). Among these genes, 10 CsVQ genes, including CsVQ6, CsVQ7, CsVQ8, CsVQ10, CsVQ11, CsVQ12, CsVQ13, CsVQ15, CsVQ17 and CsVQ21, were highly expressed in the root, leaf and stem but exhibited a lower expression in the flower. The expression levels of CsVQ1, CsVQ2, CsVQ5, CsVQ9, CsVQ16, CsVQ20 and CsVQ22 were higher in the stem and leaf than those in the root and flower. In contrast, six CsVQ genes, particularly CsVQ3,
CsVQ4 and CsVQ18, were more highly expressed in the flower than in other organs. Finally, 2 CsVQ genes (CsVQ14 and CsVQ23) were mainly expressed in the root. In general, almost all CsVQ genes were more highly expressed in the root, stem and leaf, while only a few genes were more highly expressed in the flower, implying that these genes play different roles in the growth and development of the tea plant.

Discussion
The VQ proteins are plant-specific proteins that play important roles in the regulation of plant growth, development and responses to various external environment stresses [25, 26]. To date, VQ gene families have been identified and characterized in Arabidopsis, rice, banana, grapevine, etc. [8, 9, 22, 25, 26, 32] but have not been elucidated in tea plant. Recently, the genome of tea plant has been sequenced and published, rendering the identification and characterization of the CsVQ gene family easy and reliable [30]. In this study, 25 CsVQ genes were identified in tea plant, and the structural and functional framework of the CsVQs was established, which can provide a powerful theoretical foundation for future functional studies.

The VQ protein is named for its highly conserved VQ motif FxxhVQxhTG, where x represents any amino acid, and h denotes a hydrophobic residue [31]. Expectedly, the 25 CsVQ proteins all contained the typical conserved FxxhVQxhTG motif, and the second h amino acid residue site was highly conserved and almost entirely Leucine, resembling VQ proteins in other plants, such as Arabidopsis, rice, soybean, grapevine, Chinese cabbage and maize [8, 9, 22, 25, 26, 32]. Existing studies have confirmed that most VQ proteins in plants are located in the nucleus, and our data show that most CsVQ proteins are localized in the nucleus. Interestingly, 6 CsVQ proteins were predicted to be located in the chloroplast or nucleus, suggesting that certain members of the CsVQ protein family may be dual targeted to both the nucleus and chloroplasts, which is similar to the VQ proteins in Arabidopsis and poplar [27, 31]. Recently, a study about the origin and evolution of VQ proteins showed that the same group VQ proteins from different species underwent different evolutionary histories and evolved into different subgroups, among dicots and monocots were clustered in different class, which is highly consistent with the evolutionary process of plants [33]. Similarly, our phylogenetic analysis results showed that the CsVQ proteins were classified into five groups.
and each group proteins had a higher affinity with the VQ proteins in Arabidopsis with a dicot, but a more distant affinity with the VQ proteins in the monocot rice, implying the evolution of the CsVQ proteins depended on the evolutionary process of plants, which is highly consistent with the results of previous studies mentioned above [33]. Furthermore, 20 conserved motifs were identified in the CsVQ proteins, and close proteins shared similar structures, revealing that these CsVQ proteins perform similar functions. Notably, Motif 1 corresponds to the VQ-containing motif, which is the common motif in all CsVQ proteins, and may impart specific functions to the CsVQ proteins [24]. Additionally, the similarities in the motif composition of the CsVQ proteins were consistent with the results of the phylogenetic analysis, while the distinctions among the different groups suggest that the function of the CsVQ members varies [22].

Accumulating evidence has demonstrated that the transcription of VQ genes is regulated by various endogenous and environmental signals, which is consistent with their diverse roles in plant growth and development [32]. For example, the AtVQ14/IKU1 gene, which is involved in seed development, is expressed preferentially in the early endosperm, and its deficiency reduces endosperm growth and results in small seeds [12, 34]; AtVQ8 plays an important role in chloroplast development and photosystem assembly [27]; and the over-expression of AtVQ29 causes hyposensitivity in hypocotyl growth in far-red and low-light conditions and substantially delays the blossom of Arabidopsis [18, 27]. Moreover, AtVQ17, AtVQ18 and AtVQ22 have been shown to highly stunt plant growth [27]. Correspondingly, many studies have shown significant histological specificity in the transcription of plant VQ genes, such as 7, 9 and 10 GmVQ genes were specifically expressed in soybean pod shell, root
11 bamboo VQ genes were highly expressed in leaf, early panicle, advanced panicle, root, and rhizome tissue, but expressed at low levels in shoots [29]. Recently, the VvVQ family genes were confirmed to be involved in regulating the growth and development of grapevine by a specific expression in various tissues and at different developmental stages [9]. Similarly, we also found that most CsVQ genes were differentially expressed in the different tissues of tea plant, including root, stem, leaf and flower, suggesting that the members of the CsVQ gene family are extensively involved in the growth and development of tea plant and play different roles in different organs or tissues, which was confirmed in multiple species such as rice [26, 33], Chinese cabbage [32] and maize [22]. On the other hand, the transcription of the VQ genes was either induced or inhibited by salt, drought, low nitrogen and temperature stress, indicating that these genes are also important for the plant response to various abiotic stresses [8, 9, 22, 26, 32]. In this study, we assessed the expression levels of the CsVQ genes in tea plant under salt and drought stress; the expression levels of approximately half of the CsVQ genes were up-regulated, while the expression levels of the other half were down-regulated under salt stress, and most CsVQ genes were up-regulated after the exposure to drought stress, which is similar to previous reports of VQ genes in rice, Chinese cabbage and grapevine [9, 26, 32]. In addition, we found that close CsVQ genes show a similar expression trend under the same stress but may exhibit the opposite expression trend under different stress, implying that the CsVQ genes are actively involved in the tea plant response to salt and drought stress and that their response mechanism is complex and diverse.

The VQ proteins are generally considered the most important interaction proteins of the WRKY transcription factor, and together, they regulate various physiological and biochemical processes in plants [9, 25, 34, 35]. For instance, AtVQ15 interacts with AtWRKY25 and AtWRKY51 to regulate the tolerance of plants to osmotic stress [11, 27], AtVQ14/Iku1 act as vital partners of AtWRKY10 to determine endosperm growth and seed
size [11, 12], and the interaction between AtVQ9 and AtWRKY8 decreases the DNA-binding activity of AtWRKY8 to mediate plant responses to salt stress [20].

Here, multiple CsVQ proteins, such as CsVQ14, CsVQ15, CsVQ16, CsVQ17, CsVQ18, and CsVQ23, were presumed to closely interact with different WRKY transcription factors by constructing an Arabidopsis association model, implying there may be similar biological functions of CsVQ proteins that are dependent on interaction with WRKY transcription factor in tea plant. In addition, the results of CsWKRY proteins analysis showed that there may be an interaction between 8 CsWRKYs and different CsVQ proteins, this is mutually reinforcing with our speculation above. At present, existing evidence indicated that VQ proteins appeared to interact only with group I and Ile WRKY transcription factors, but failed to interact with group IIa, IIb, IId, Ile, and III WRKY transcription factors [23, 27]. In expectation, there is a similar result in our study that predicted interacting CsWRKY proteins all belong to either group I or group Ile WRKY transcription factor. Simultaneously, the C-terminal WRKY domains of group I CsWRKY proteins and the sole WRKY domains of group Ile CsWRKY proteins are highly consistent with the proven core binding domain in other species that further confirms our conclusion, although we cannot rule out the possibility of other groups I, Ile CsWRKY proteins interacted with CsVQ proteins. Moreover, CsVQ14 and CsVQ23 participated in complicated interactions that not only included several WRKY transcription factors but also included the MAPK protein, which is consistent with previous studies. For example, MAPK4 can bridge AtVQ21 and WRKY33 to influence plant growth and disease

**Fig. 5** Interaction of CsVQ proteins with CsWRKY proteins in tea plant. a The prediction of interaction between CsVQ proteins and CsWRKY proteins by the PAIR website, and the interaction network was drawn in Cytoscape 3.6.0. b Sequence analysis of the C-terminal WRKY domains of group I CsWRKY proteins and the sole WRKY domains of group Ile CsWRKY proteins

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resistance [15, 16], and MAPK3 and MAPK6 participate in the interaction between AtVQ4 and multiple WRKYs to regulate immune responses in Arabidopsis [11]. Furthermore, an appreciable number of specific microRNA target sites in the CsVQ genes, including miRNA396, miRNA172, miRNA157, miRNA857, miRNA319, etc., which are microRNAs known to play important roles in various life processes of plants [36, 37], was identified in this study, implying that the CsVQ genes may be controlled by a diversified microRNA regulatory network. Altogether, these results suggest that the CsVQ gene family is extensively involved in tea plant growth, development and stress response, and these processes are closely related to the interactions between CsVQ proteins and CsWRKY transcription factors, and the regulation of upstream microRNA, although this hypothesis requires a more in-depth investigation.

Conclusions
In conclusion, this study provides the first comprehensive and systematic analysis of the VQ gene family in tea plants. In total, 25 CsVQ genes were identified and classified into 5 groups (I, III-VI), and bioinformatics and expression profile analyses of the CsVQ genes were performed to determine their potential functions in the growth, development and stress response in tea plant. CsVQ genes are actively involved in regulating tea plant growth development and responding to salt and drought stress, and these processes are closely related to the interactions of CsVQ proteins with CsWRKY transcription factors and the regulation of upstream microRNA. These results provide an important foundation for further functional studies investigating the CsVQ proteins in tea plant.

Methods
Plant materials and stress treatments
Two-year-old seedling cuttings of tea plants [C. sinensis cv. ‘Longjingchangye’] were pre-incubated under normal conditions (25 ± 2 °C temperature, 60 ± 10% relative humidity and 12-h light/12-h dark cycle) for 2 weeks in an artificial climate chamber at the Northwest A&F
University (Yangling, China). Then, the salt and drought stress treatments were performed using 200 mM NaCl and 20% (w/v) polyethylene glycol (PEG) 6000, while all other environmental conditions remained constant. The first and second tender leaves of the treated tea plants were randomly collected at 0, 1, 2, 4, 8, 12, 24 and 48 h, and the samples were quickly frozen in liquid nitrogen and stored at −80 °C until analysis.

Identification of the CsVQ gene family in tea plant
To identify the VQ genes in tea plant, 108 VQ motif sequences conserved in plants were downloaded from the Pfam database and used as queries to perform a BLASTP search against the genome and transcriptome (NCBI SRA: SRP128078) database of tea plant [30]. Then, the obtained CsVQ genes were rechecked and confirmed. In brief, the candidate CsVQ gene sequences were assessed by BLAST to ensure that they belonged to the VQ gene family, and a comparative analysis of the 25 CsVQ gene sequences was performed to avoid repetition.

Sequence alignment, phylogenetic analysis and conserved motif analysis
Thirty-four Arabidopsis and 39 rice VQ protein sequences were downloaded from the TAIR database (http://www.arabidopsis.org/) and Rice Data sites (http://www.ricedata.cn/gene/). Multiple alignments of the amino acid sequences of the VQ proteins in tea plant, Arabidopsis and rice were performed using DNA-MAN V6.0 with the default options. A phylogenetic tree was constructed using the neighbour-joining method in MEGA 6.0, and a diagram of the phylogenetic tree was drawn using EVOLVIEW (www.evolgenius.info/evolview). In addition, the conserved motifs of the CsVQ protein sequences were obtained and analysed using the MEME website (version 4.12.0, http://meme-suite.org/) using the parameters of 20 previously identified motifs [38].

Analysis of interaction networks of the CsVQ proteins
Functional interacting network models of CsVQ proteins were integrated using the web STRING (http://string-db.org), and the confidence parameters were set at a 0.40 threshold. Seventy-six protein sequences of CsWRKY transcription factor were obtained from the genome database of tea plant [30], and mapped to the WRKY proteins of A. thaliana by BLASTP tool in the TAIR database. Subsequently, the interaction between CsVQs and CsWRKYs were forecasted based on the PAIR website (http://www.cls.zju.edu.cn/pair/), and the interaction network was drawn in Cytoscape 3.6.0.

Prediction of microRNAs targeting the CsVQ genes
MicroRNA libraries of tea plant (unpublished data) were used as the target reference, and the psRNA Target and psRobot online program (http://plantgrn.noble.org/psRNAATarget/ and http://omicslab.genetics.ac.cn/psRobot/) was used to predict the putative miRNAs targeting the CsVQ genes.

RNA extraction and cDNA reverse transcription
The total RNA from the tea plant leaves was extracted using RNAiso Plus (TakaRa, Japan), and the concentration and quality of the RNA were measured using a NanoDrop ND-1000 spectrophotometer (NanoDrop, Wilmington, DE). Equal amounts of RNA were reverse transcribed to cDNA using a 5 x All-In-One RT MasterMix Kit (ABM, Canada) according to the manufacturer’s protocol.

Quantitative RT-PCR (qRT-PCR) analysis of CsVQ genes
qRT-PCR was performed using SYBR® Premix Ex Taq™ II (TakaRa, Japan) on a Bio-Rad IQ5 Real-Time PCR System (Bio-Rad, USA) according to the manufacturer’s instructions. Briefly, each reaction was performed in a total volume of 20 μL containing 10 μL SYBR Premix Ex Taq II (Tli RNaseH Plus) (2x), 7.4 μL ddH2O, 1 μL diluted cDNA template, and 0.8 μL of each gene-specific primer (Additional file 14: Table S12) using the following PCR program: 95 °C for 3 min, followed by 40 cycles of 95 °C for 30 s and 63 °C for 1 min 10 s. Melting curves were obtained to verify the amplification specificity through a stepwise heating of the amplicon from 63 to 95 °C. CsPTB [39] was used as an internal control gene. The experiments were repeated in triplicate, and the relative gene expression levels were calculated based on the threshold cycle using the 2-ΔΔCT method [40]. The clustering analysis of the CsVQ genes was performed using Cluster 3.0 software, and the heat maps of the gene expression were visualized using Tree View software.

Additional files

| Additional file 1: Table S1. The CDS sequences and deduced amino acid sequences of CsVQ genes. (XLSX 18 kb) |
| Additional file 2: Figure S1. LOGO of twenty conserved motifs among CsVQ proteins. (TIF 6891 kb) |
| Additional file 3: Table S2. Regular expression of conserved motifs in CsVQ proteins. (DOCX 20 kb) |
| Additional file 4: Table S3. List of predicted known microRNA target sites of CsVQ transcripts. (XLSX 14 kb) |
| Additional file 5: Table S4. List of predicted novel microRNA target sites of CsVQ transcripts. (XLSX 17 kb) |
| Additional file 6: Table S5. List of predicted microRNAs targeting CsVQ genes. (XLSX 10 kb) |
| Additional file 7: Table S6. Information regarding the STRING search results of the interaction networks of the CsVQ proteins. (XLSX 11 kb) |
| Additional file 8: Table S7. The CsVQ proteins maps to the VQ proteins in Arabidopsis thaliana. (XLSX 11 kb) |
Additional file 9: Table S8. The CsWRKY proteins maps to the WRKY proteins in Arabidopsis thaliana. (XLSX 14 kb)

Additional file 10: Figure S2. Phylogenetic tree of WRKY transcription factors in tea plant and Arabidopsis constructed using the neighbour-joining method in MEGA 6.0. (TIF 4469 kb)

Additional file 11: Table S9. Expression levels of CsVQ genes following the NaCl treatment in tea plants. (XLSX 16 kb)

Additional file 12: Table S10. Expression levels of CsVQ genes following the PEG 6000 treatment in tea plants. (XLSX 16 kb)

Additional file 13: Table S11. Expression levels of CsVQ genes in four different tissues from tea plants. (XLSX 11 kb)

Additional file 14: Table S12. Primers used for the qRT-PCR of the CsVQ genes. (XLSX 10 kb)

Abbreviations
GRAVY: Grand average of hydropathicity; MAPK: Mitogen-activated protein kinase; MEME: Multiple Em for motif elicitation; PEG 6000: Polyethylene glycol 6000; qRT-PCR: Quantitative real-time polymerase chain reaction; VQ: VQ motif-containing

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Availability of data and materials
All data generated or analysed during this study were included in this published article and the additional files.

Authors’ contributions
WW conceived and designed the study. JG and WW wrote the paper. JG, JC and JY performed the experiments and analyzed the data. YBY and YJY performed the experiments and analyzed the data. YBY and YJY participated in the data analysis and the paper revision. All authors read and approved the final manuscript.

Ethics approval and consent to participate
The tea plants used in this study were from the Tea Sci-tech Demonstration Base of Northwest A&F University, in Xixiang, Shaanxi, China, and no permissions are needed to obtain the material. Our study fully complies with institutional regulations.

Consent for publication
Not applicable.

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

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