Abstract: The CHYR (CHY ZINC-FINGER AND RING FINGER PROTEIN) proteins have been functionally characterized in iron regulation and stress response in Arabidopsis, rice and Populus. However, their roles in soybean have not yet been systematically investigated. Here, in this study, 16 GmCHYR genes with conserved Zinc_ribbon, CHY zinc finger and Ring finger domains were obtained and divided into three groups. Moreover, additional 2–3 hemerythrin domains could be found in the N terminus of Group III. Phylogenetic and homology analysis of CHYRs in green plants indicated that three groups might originate from different ancestors. Expectedly, GmCHYR genes shared similar conserved domains/motifs distribution within the same group. Gene expression analysis uncovered their special expression patterns in different soybean tissues/organs and under various abiotic stresses. Group I and II members were mainly involved in salt and alkaline stresses. The expression of Group III members was induced/repressed by dehydration, salt and alkaline stresses, indicating their diverse roles in response to abiotic stress. In conclusion, our work will benefit for further revealing the biological roles of GmCHYRs.

Keywords: CHYR; soybean; genome-wide identification; expression analysis; abiotic stress

1. Introduction

As one of the most widely grown crops in the world, soybean (Glycine max) provides an important source of plant-based protein and edible oil [1]. However, its yield and quality are enormously hindered by germplasm resources and diverse environmental factors, especially water deficiency, high salt, and alkaline [2]. Drought is one of the major natural disasters for world’s agricultural production. Globally, this extreme weather phenomenon has led to cereal loss of 1820 million Mg during the past four decades [3,4]. Soil salinity-alkalinization is another worldwide abiotic stress restraining land utilization, grain yield and local economic development. According to official statistics, more than 6% of the world’s soil resources are affected by saline and alkaline. Furthermore, continuous drought has a great influence on soil salinization and salt accumulation in root zone [5]. Utilization and management of the saline-alkaline soil is requisite to alleviate the ever-growing population’s demand for food. Consequently, it is meaningful to focus on uncovering the molecular mechanism of plant response to abiotic stress and cultivating crops with stress resistance.
The RING E3 (Really Interesting New Gene) proteins were found to play critical roles in abiotic stress response via protein ubiquitination degradation [6,7]. Previously, a C3H2C3 RING (Really Interesting New Gene) zinc finger domain containing protein from Arabidopsis was characterized and named as MIEL1 (MYB30-Interacting E3 Ligase1) [8]. According to the conserved RING zinc finger domain, they were also called CHYR (CHY Zinc-FINGER AND RING FINGER PROTEIN) and RZFP (RING ZINC-FINGER PROTEIN) [9,10]. Protein sequence alignment has proved that MIEL1, RZFP and CHYR were in the same family, with conserved CHY zinc-finger, C3H2C3-type ring finger and rubredoxin-type fold domain [9]. In addition, when hemerythrin domains appeared in the N-terminus of CHY zinc finger domain, they were designated BTS/BSTL (BRUTUS/BRUTUS-like) in Arabidopsis, but HRZ (Hemerythrin motif-containing RING-and Zinc-finger protein) in rice [11–14]. Above all, we could uniformly define these proteins containing CHY zinc-finger, C3H2C3 -type ring finger and rubredoxin-type fold domain as the CHYR family.

Increasing evidence has shown the diverse roles of CHYR genes in plant growth, development and stress responses. MIEL1 was first found to control protein stability of MYB96 and MYB30 in balancing cuticular wax biosynthesis and defense [8,15,16]. AtCHYR1 was reported to enhance ABA and drought responses by elevating ROS production and stomatal closure [9]. Homologous gene of Populus euphratica (PeCHYR1) showed similar phenotypes, enhancing drought tolerance, stomatal closure, and H2O2 production [17]. However, overexpression of OsRZF34 (AtCHYR1 homologous gene in rice) enhanced stomatal opening, leaf cooling and ABA insensitivity [10]. CHYR proteins with 2–3 additional hemerythrin domains (also known as BTS/BTSL/HRZ) were found to regulate iron response in Arabidopsis and rice [11,12,18].

Though several CHYR genes have been identified with diverse names, they have not yet been systematically analyzed at the gene family level. In particular, their roles in soybean development and stress response have not been uncovered. Here, in this study, 16 CHYR genes were identified through an extensive search of soybean genome (Wm82.a2.v1). Furthermore, their chromosome localization, phylogeny, conserved domains and expression patterns, especially in response to abiotic stress were comprehensively analyzed. These results will provide valuable clues for further functional studies on GmCHYR genes and their potential roles in abiotic stress.

2. Results

2.1. Identification and Phylogenetic Analysis of CHYR Genes from Soybean and Arabidopsis

To identify soybean CHYR genes, protein sequences of published Arabidopsis CHYRs [8,9,11,12,19] were used to construct a Hidden Markov Model (HMM) [20]. Whole soybean and Arabidopsis protein sequences were downloaded from Phytozome to carry out the local search. Finally, 16 soybean and 7 Arabidopsis CHYR genes were identified. The 23 proteins were proven to contain at least three conserved domains, including CHY zinc-finger (PF05495), C3H2C3-type ring finger (PF13639) and zinc ribbon domain (PF14599) according to Pfam and SMART analysis. For convenience’s sake, soybean CHYR genes were renamed GmCHYR1 to GmCHYR16 based on their order on the chromosomes, and genes from Arabidopsis were relabeled as AtCHYR1 to AtCHYR7. Their involved information (including sequence length, hydropathicity, predicted protein location, classification, alternative name and functions) were listed in Table S1. As we could see from Table S1, amino acid numbers of GmCHYRs and AtCHYRs ranged from 234 to 1262. Their grand average of hydropathicity were all negative, indicating that GmCHYRs and AtCHYRs are hydrophilic proteins. Furthermore, these CHYR proteins were predicted to localize in the cytoplasm, or nucleus, or chloroplast. The cytoplasm and nucleus distribution of AtCHYR6/MIEL1 in Arabidopsis cells could support this result [8].

To further investigate the phylogenetic relationship of GmCHYRs, their protein sequences were aligned with 7 AtCHYRs. All 23 CHYR proteins contained conservation CHY zinc-finger (PF05495), C3H2C3-type ring finger (PF13639) and zinc ribbon 6 domain (PF14599) (Figure S1). Then, a phylogenetic tree was generated basing on this multiple
alignment by using MEGA 7.0 with the Maximum-Likelihood (ML) method with 1000 bootstrap replications. As shown in Figure 1A, soybean and *Arabidopsis* CHYRs could be classified into three groups according to their topological analysis and bootstrap values. In details, both Group I and Group II consisted of 5 *GmCHYR*s and 2 *AtCHYR*s. The rest, 6 *GmCHYR*s and 3 *AtCHYR*s, were allocated to Group III.

Figure 1. The phylogenetic tree and conserved domains and motifs analysis of CHYR genes in soybean and *Arabidopsis*. (A) Phylogenetic tree of soybean and *Arabidopsis* CHYR proteins, constructed by using MEGA 7.0 with the maximum-likelihood (ML) method under 1000 replications. (B) Conserved domains in GmCHYR proteins were identified by combining the SMART, PFAM, and NCBI CD database, represented by different colors. Green: Zinc_ribbon domain; Yellow: CHY-zinc finger domain; Pink: Ring finger domain; Dark green: Hemerythrin/Hemerythrin-like domain. The conserved motifs of GmCHYR proteins were analyzed by using the MEME tool. Schematic of the conserved domains and motifs were integrated by employing TBtools. The motif number was displayed below each motif.

Furthermore, their conserved domains and motifs were analyzed. As expected, all 16 *GmCHYR*s and 7 *AtCHYR*s contained CHY zince-finger, C3H2C3-type ring finger and zinc ribbon (Figure 1B). Besides, there were 2-3 hemerythrin domains in the N terminus of Group III members. Group III members were also called BTS/BTSL in *Arabidopsis*, and HRZ in rice [12,18]. This is consistent with former reported results that there were 2 BTSL (*AICHYR2/3*) and 1 BTS (*AICHYR4*) in *Arabidopsis* [12]. All of them have been reported to regulate iron homeostasis [11]. Meanwhile, we employed the MEME program to predict conserved motifs (Figure 1B). In accordance with conserved domains, *GmCHYR*s within each group displayed similar motif distribution. Among the detected 15 motifs, motif 1, 5, 9, 12 in the N terminus made up CHY-zinc finger. Motif 3 and 4 formed the Ring finger domain. Motif 2 served as Zinc_ribbon domain. Additionally, hemerythrin domain of Group III members constitutes of motif 7, 10, 11, 14, 15. Additionally, a conserved motif 6 and 8, which was closely to hemerythrin domain, could be found in Group III members. However, their function still needs further investigation.

2.2. Identification and Classification of CHYR Members in Green Plants

Above results showed that only Group III members contained 2–3 additional hemerythrin domains in the N terminus, which are of great importance in regulating iron homeostasis. We wondered whether Group III CHYR proteins gained these hemerythrin domains during evolution, or Group I and II lost these domains. Therefore, the local proteome se-
quences of 21 representative plant species, including Dicots, Monocots, Basal Angiosperms, Pteridophyta, Bryophyta, Chlorophyta and Gymnosperm were searched to identify potential CHYR genes by using the former Arabidopsis HMM. At last, a total of 107 nonredundant sequences were obtained from 21 detected plant species (Tables 1 and S2). Pfam and SMART were further used to detect the three conserved domains for CHYR proteins, including CHY zinc-finger domain, C3H2C3-type ring finger domain and zinc ribbon domain.

Table 1. Overview of genes encoding CHYR proteins in plants.

| Major Lineage          | Species                | Group I | Group II | Group III |
|------------------------|------------------------|---------|----------|-----------|
| Dicots                 | Vitis vinifera         | 3       | 2        | 3         |
|                        | Arabidopsis thaliana   | 2       | 2        | 3         |
|                        | Glycine max            | 5       | 5        | 6         |
| Monocots               | Zea mays               | 3       | 2        | 1         |
|                        | Oryza sativa           | 3       | 2        | 2         |
|                        | Ananas comosus         | 1       | 2        | 1         |
|                        | Musa acuminate         | 1       | 1        | 3         |
|                        | Spirodea polyrhiza     | 1       | 0        | 0         |
|                        | Zostera marina         | 0       | 1        | 2         |
| Basal angiosperms      | Amborella trichopoda   | 1       | 1        | 1         |
| Gymnosperm             | Pinus parviflora       | 4       | 0        | 1         |
|                        | Pinus radiata          | 4       | 0        | 1         |
|                        | Pinus jefferi          | 4       | 0        | 1         |
|                        | Pinus ponderosa        | 4       | 0        | 1         |
|                        | Picea engelmannii      | 3       | 0        | 0         |
| Pteridophyta           | Selaginella moellendorfii | 1     | 0        | 2         |
| Bryophyta              | Marchantia polymorpha  | 1       | 0        | 1         |
|                        | Physcomitrella patens  | 5       | 0        | 3         |
|                        | Sphagnum fallax        | 5       | 0        | 2         |
| Chlorophyta            | Chlamydomonas reinhardtii | 0     | 1        | 1         |
|                        | Volvox carteri         | 0       | 1        | 1         |

To explore their evolutionary relationship, 107 CHYR members were aligned using ML (Maximum-likelihood), NJ (Neighbor-joining), and ME (Minimum-evolution) methods to construct unrooted phylogenetic trees based on their protein sequences (Figures 2, S2 and S3). As the three phylogenetic trees depicted, three methods presented a similar topology. According to their evolutionary relationship, 107 CHYR members could be further divided into three groups (Group I, II, III) as well. Though Group I and Group II were clustered together, CHYR members from Bryophyta, Pteridophyta and Gymnosperms could be only found in Group I, implying the possibility of gene acquisition during evolution. From this result, we speculated that Group II might appear after Group I. Group III did coexist with the other two groups, but was far away from the others in topology, which indicated that they might come from different ancestors. Interestingly, there were only 4 CHYR members in Chlorophyta, two of them were from Chlamydomonas reinhardtii, the others were from Volvox carteri. While CreCHYR2 and VocarCHYR1 were clustered with Group I and Group II, CreCHYRI and VocarCHYR1 were grouped together Group III, indicating the existence of CHYR members throughout green plants evolution. Previous study has reported the up regulation of CreCHYRI under iron deficiency [21], suggesting the conserved role of Group III members in iron regulating. The above findings implied the early emergence of CHYR members and their persistence in the evolution of green plants.
suggesting the conserved role of Group III members in iron regulating. The above findings implied the early emergence of CHYR members and their persistence in the evolution of green plants.

Figure 2. The Maximum-likelihood phylogenetic tree of CHYR genes in green plants. One hundred and seven CHYR protein sequences from 21 detected plant species were aligned with ClustalW and a phylogenetic tree was generated by using MEGA7 with the maximum-likelihood method under 1000 replications. The tree was divided into three groups with green shadow in Group I, blue shadow in Group II, and red shadow in Group III. Confidence values were listed on each node.

2.3. Homology Analysis of CHYR Genes from Soybean and Arabidopsis

According to their phylogenetic relationship, the number of GmCHYRs is more than twice that of AtCHYRs. Particularly, GmCHYRs appeared in pairs. The big genome size and whole genome duplication might be two critical reasons for gene expansion [22], such as gene duplication in soybean LRR-RLK genes [23]. The homologous relationship of GmCHYRs and AtCHYRs was further analyzed by comparing G. max and A. thaliana.
genomic sequence through OrthoVenn2 [24]. As depicted in Figure 3, 15 orthologous gene pairs were identified from Arabidopsis and soybean (green line in Figure 3). Nineteen paralogous gene pairs were characterized from soybean (red line in Figure 3), but only one paralogous gene pair exist in Arabidopsis (blue line in Figure 3), which might be derived from gene expansion during whole genome duplication occurred in soybean, or gene loss in Arabidopsis [25].

![Figure 3](image-url) Chromosomal distribution and homology analysis of CHYR genes in the genomes of soybean and Arabidopsis. Paralogous and orthologous CHYR genes were mapped onto soybean and Arabidopsis chromosomes. Red lines connected soybean paralogous genes. Green lines indicated orthologous genes between Arabidopsis and soybean. Blue lines connected Arabidopsis paralogous genes.

To trace their duplication time, Ka (non-synonymous rate), Ks (synonymous rate) and Ka/Ks ratios of 19 soybean paralogous genes were analyzed (Table S3). All Ka/Ks ratio of GmCHYRs were less than 1, varied from 0.12 to 0.4, indicating that they have undergone strong purify selection. Furthermore, their duplication time was calculated. The duplication time of Group I members varied from 9.5–43.6 Mya (million years ago) and Group II was around 11.5–46.4 Mya. This period is consistent with the latest twice whole genome duplication of soybean [25]. However, the duplication time of GmCHYR3/GmCHYR8, GmCHYR5/GmCHYR8, GmCHYR7/GmCHYR8, GmCHYR8/GmCHYR9 pairs in Group III were greater than 155.6 Mya, which was just in line with the specific γ duplication of dicotyledon [25]. These results uncovered that GmCHYR expansion derived from whole genome duplication, resulting in conserved domains and motifs.

2.4. Expression Pattern of Soybean CHYR Genes in Different Tissues and Organs

To further look into GmCHYRs roles in soybean development, their expression profiles were analyzed based on published data of nine tissues/organs collected in Phytozome, including flowers, nodules, leaves, roots, root hairs, stems, shoot apical meristem, pods, and seeds [26]. As Figure 4 depicted, except that GmCHYR1 showed almost no expression, the rest 15 GmCHYRs displayed specific expression across nine detected tissues/organs. Compared with Group III, Group I and II members were more likely to be expressed in
all detected tissues/organs and had much higher expression values. This suggested their potential roles in soybean growth and development. Group II genes showed relative higher expression in the flowers, suggestive of their roles in reproduction. In particular, paralogous gene GmCHYR6 and GmCHYR14 were all highly expressed in nine detected tissues/organs. However, Group III members preferred to be expressed in nodules, indicating their roles in nitrogen fixation. In general, paralogous gene GmCHYR4/12/16, GmCHYR6/11/13/14 and GmCHYR3/7 shared similar expression patterns. GmCHYR5/8/9 were also paralogs of GmCHYR3/7, but they displayed opposite expression from GmCHYR3/7. This might result from some special regulatory elements, or modification in their promoters, or just functional segregation during evolution.

To uncover the roles of GmCHYRs in response to abiotic stress, their transcriptome data under different stress treatments were analyzed by using published database (including drought, salt (GSE57252) [27] and alkaline [28]). In accordance with tissue expression data, GmCHYR1 showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs showed little expression in root and its expression values were zero under abiotic stresses.

![Figure 4](image-url)  
**Figure 4.** Tissue expression profiles of GmCHYRs in soybean. The transcriptional levels of GmCHYR genes in nine tissues/organs of soybean were analyzed based on published data collected in Phytozome. A heatmap was generated by TBtools. Five to thirty were artificially set with the color scale limits according to their expression values. The color scale shows increasing expression levels from blue to red.

2.5. **Transcription Patterns of GmCHYRs in Response to Dehydration, Saline, Alkaline Stresses**

To uncover the roles of GmCHYRs in response to abiotic stress, their transcriptome data under different stress treatments were analyzed by using published database (including drought, salt (GSE57252) [27] and alkaline [28]). In accordance with tissue expression data, GmCHYR1 showed little expression in root and its expression values were zero under abiotic stresses (Figure 5). As we could see from Figure 5, GmCHYRs displayed various response strategies to diverse abiotic stresses. Group I members were involved in all detected stresses, and they were dramatically increased under alkaline stress. Among Group I, GmCHYR16 was significantly up regulated by three stresses. However, its paralogous gene, GmCHYR12 were only induced by alkaline stress. Another paralogous gene GmCHYR4 exhibited similar expression patterns with GmCHYR10, which were repressed by dehydration, but induced by saline and alkaline stresses. Group II members exhibited similar expression profiles under abiotic stresses. Besides GmCHYR6 showed increased expression under three detected stresses, most of Group II were increased under alkaline stress, but decreased under dehydration and saline stresses. Compared
with Group I and II, genes from Group III were dramatically up regulated by salt stresses. Paralogous gene GmCHYR3/7/5/9 exhibited similar expression profiles. However, the expression of GmCHYR8 and GmCHYR15 were repressed by three stresses. Above all, Group I members might play vital roles in dehydration, salt and alkaline stresses. Group II and III members were participated in salt and alkaline response.

Figure 5. Expression profiles of GmCHYRs under dehydration, salt, and alkaline stress. The transcriptional levels of GmCHYR genes in response to dehydration (abbreviated as de), salt (100 mM NaCl) and alkaline (50 mM NaHCO$_3$) stresses were investigated based on the published transcriptome data. The expression of GmCHYR were normalized by TBtools. According to the normalized value, −2.5 to 2.5 was artificially set with the color scale limits. The color scale shows increasing expression levels from blue to red. The differentially expressed genes (DEGs) were highlighted by red (up-regulation) and blue (down-regulation).

2.6. qRT-PCR Verification of GmCHYRs under Dehydration, Saline and Alkaline Stresses

To confirm that they were indeed involved in the three stresses, the expression of 7 genes from three groups under dehydration, salt and alkaline stresses were validated by qRT-PCR, including GmCHYR16/10 (Group I), GmCHYR2/6 (Group II), and GmCHYR3/5/15 (Group III) (Figure 6). These 7 genes were chosen according to their homology relationship and specific stress expression patterns. Their expression trends were basically consistent with transcriptome results. GmCHYR16 (Group I) was dramatically induced more than 93-fold under alkaline stress and 49-fold under saline stress, respectively. Under dehydration stress, the expression fold change of GmCHYR16 was only 2.5. This suggested the vital roles of GmCHYR16 in salt and alkaline stress response. GmCHYR10 (Group I) displayed similar expression patterns with GmCHYR16. In accord with transcriptome results, the expression level of GmCHYR10 was lower than that of GmCHYR16. Moreover, GmCHYR2/6 (Group II) were induced by salt and alkaline stresses, but did not respond to dehydration. GmCHYR3/5 (Group III) were dramatically induced by three stresses, while GmCHYR15 (Group III) depicted down-regulated trends under three stresses, suggesting their functional differentiation during evolution.
3. Discussion

For a long time, soybean is one of the staple crops and the most important legume in the world, which provides a major source of vegetable protein and edible oil [29]. Nowadays, with the continuous improvement of living level, the aging, the change of population structure and the fast development of stock farming, the consumption of protein and oil products has increased day by day, driving the consumption of soybean. However, such as other crops, soybean yield is restricted by natural environmental conditions. Previous studies have shown that adverse stresses had negative effects on plant growth and development, including abnormal metabolism, protein misfolding and so on [30]. Molecular design breeding is one of the effective methods to improve crops stress resistance. The key is to mine stress-resistance genes.

Previously, CHYR genes with three conserved domains (CHY-zinc finger, Ring finger domain and Zinc_ribbon domain) have been reported in Arabidopsis [9,11], rice [10] and Populus [17] in response to adverse stresses. However, little was known about them in soybean. In this study, a total of 16 CHYR genes were identified through searching against the released genome database of G. max [26] by using AtCHYRs protein sequences as queries [9,11]. According to their phylogenetic analysis, CHYRs could be classified into three groups (Figures 1A, 2, S2 and S3). This result was further supported by conserved domain and motif distribution. An exploration of conserved domain and motif distribution confirmed that all GmCHYR proteins contained conserved CHY-zinc finger, Ring finger domain and Zinc_ribbon domain (Figures 1 and S1). In addition, there were 2–3 hemerythrin domains in the N-terminus of Group III members (Figures 1 and S1). The similar results had been previously reported in Arabidopsis [9,11], rice [10] and Populus [17]. Meanwhile, conserved motifs 6 and 8 were found in Group III members, between the CHY-zinc finger domain and hemerythrin domain (Figure 1), while its function was still unknown. The above results suggested that genes shared similar conserved domain and motif distribution within the same group.

As known to all, the conserved zinc finger domains consisted of several conserved cysteine and histidine residues (Cys - X2 - Cys - X (9–39) - Cys - X (1–3) - His - X (2–3) -
Cys/His - X2 - Cys - X (4–48) - Cys - X2 - Cys) to bind zinc ions (Figure S1), or interact with partner proteins, or catalyze E3 ubiquitin ligase activity. For example, the RING domain of AtCHYR1 (Group I) was essential for ubiquitin E3 ligase activity [9]. The CHY-zinc finger domain of AtCHYR6/MIEL1 (Group II) was responsible for MYB96 interacting and degrading [16]. At the same time, Group III members containing hemerythrin domains could interact with transcription factor PYEL proteins and exert E3 ligase activity via the C-terminal RING domain [31]. The hemerythrin domain in plants was reported to play essential roles in iron binding and protein stability [13]. The removal of hemerythrin domain could make Group III CHYR (also termed as BTS) stable in the existence of iron and also complement hemerythrin-containing CHYR loss, while deletion of the RING domain could not [13,31]. AtCHYR2/3/4 (Group III) and OsCHYR1/5 (Group III) were reported to act as negative regulators in iron deficiency response [11,12,32]. However, E3 ubiquitin ligase activity and iron regulating function of GmCHYRs still need further experiments to validate.

It was worth noting what drove the difference between Group I/II and Group III in gene structure and physiological roles. Further phylogenetic analysis of CHYRs in green plants confirmed that CHYR genes appeared during green plant evolution (Figure 2). This was consistent with reported result that Group III members exist among photosynthetic organisms as well [13]. However, the divergence of Group I and Group II members might occur after angiosperm differentiation (Figures 2, S2 and S3). The number of CHYR family in G. max is more than two folds that of Arabidopsis (7 CHYR genes) and rice (7 CHYR genes), but is relative stable in detected species within Eudicots, Monocots, Gymnosperm, and Chlorophyta (Tables 1, S1 and S2). A similar phenomenon has been found in other ubiquitin ligase families. For example, U-box [33], HETC (homologous to the E6AP carboxyl terminus) [34] family in soybean are more than twice that of Arabidopsis and rice. Whole genome duplication (WGD) is a key reason for gene expansion, gene loss and new functionalization. As previously reported, G. max has gone through three whole genome duplications (117, 59, and 13 Mya) [25], which might lead to the emergence of 19 paralogous gene pairs (Figure 3). A further Ka/Ks calculation also confirmed that purification selection was the main driving force in the evolution of GmCHYRs (Table S3).

In terms of GmCHYRs' biological function, the expression of Group I and II members were much higher than that of Group III in the nine detected tissues/organs. Group I and II members were expressed in all tissues/organs, while Group III members preferred to be expressed in nodules (Figure 4). According to this result, we inferred that GmCHYRs might be involved in nitrogen-fixing genes via ubiquitination. Tissue-specific and stress expression pattern analysis of GmCHYRs were helpful to uncover their potential roles in physiology and development. Previously, studies about CHYRs were mainly focused on iron regulation. For example, AtCHYR4 (BTS) and AtCHYR2/3 (BTSL1/2) from Group III were induced by iron deficiency [12,19]. The expression of rice Group III CHYR1/5 (also known as HRZ1/2) were up regulated by iron insufficiency [14]. Even Group III member of C. reinhardtii, CreCHYR1 was up regulated by iron deficiency as well [21]. These results suggested the conserved roles of Group III members in iron regulating.

Our lab has long been committed to the study of crop abiotic stress response. The expression of GmCHYRs under salt, alkaline and drought stresses was further examined. According to the published transcriptome data, only 15 GmCHYRs (except for GmCHYR1) were detected and they were all up regulated/down regulated by dehydration, salt and alkaline stresses, suggesting their potential role in stress response (Figure 5). Due to gene homology, the expression of seven genes (GmCHYR10/16 (Group I), GmCHYR2/14 (Group II), and GmCHYR3/5/15 (Group III)) from three groups were further confirmed by using qRT-PCR (Figure 6). We could confirm that the expression of GmCHYR15 was repressed by dehydration, salt and alkaline stresses, while the expression of GmCHYR3/5 was induced by these stresses by integrating transcriptome data and qRT-PCR. Though GmCHYR3/5 and GmCHYR15 belong to Group III, their opposite expression patterns might derive from functional differentiation during evolution or cis-regulatory elements in promoter regions.
GmCHYR10/16 (Group I) and GmCHYR2/14 (Group II) were all significantly upregulated by salt and alkaline stresses, up to 93-fold, but only 2.5-fold under dehydration stress, indicating their special roles in salt and alkaline response. In particular, GmCHYR16 might be a key regulator in salt and alkaline stress response.

Above all, our analysis of 16 GmCHYRs showed that they share highly conserved domains and residues, indicating their potential conserved structure and biochemical function. Most of GmCHYRs were found to respond to various stresses, especially Group I and II members might play positive regulators in abiotic stress response. However, their internal regulatory mechanisms are still misty. Whether GmCHYRs function as an E3 ubiquitin ligase also requires further experimental verification.

4. Materials and Methods

4.1. Identification of CHYR Genes from Green Plants

A Hidden Markov Model (HMM) basing on published CHYR protein sequences from Arabidopsis were constructed to search against 21 green plants genomes [8,9,11,12,18–20]. These genomes were collected from NCBI (https://www.ncbi.nlm.nih.gov/genome/, last accessed on 6 June 2020), Phytozome(https://phytozome.jgi.doe.gov/pz/portal.html, last accessed on 6 June 2020) [26] and Congenie (https://congenie.org/, last accessed on 10 June 2020) [35]. CHY zinc-finger domain (PF05495), C3H2C3-type ring finger domain (PF13639) and zinc ribbon domain (PF14599) of putative CHYR genes were then verified by using CD search (https://www.ncbi.nlm.nih.gov/Structure/cdd/wrpsb.cgi, last accessed 10 June 2020), Pfam (http://pfam.xfam.org/, last accessed on 10 June 2020) [36] and SMART (http://smart.embl-heidelberg.de/, last accessed on 10 June 2020) [37] with default parameters.

4.2. Phylogenetic Relationship, Sequence Alignments and Protein Localization Analysis

The phylogenetic relationship of CHYR genes were analyzed by using MEGA 7 through Neighbor Joining (NJ), Maximum Likelihood (ML) and/or Minimum evolution (ME) methods with 1000 bootstrap. Multiple sequence alignments were carried out by using Clustal Omega (https://www.ebi.ac.uk/Tools/msa/clustalo/, last accessed on 1 July 2020) [38] under default parameters. Protein locations were predicted by integrating SoftBerry (http://linux1.softberry.com/all.htm, last accessed on 10 June 2020), PSORT (https://www.genscript.com/psort.html, last accessed on 10 June 2020), and CELLO (http://cello.life.nctu.edu.tw/), last accessed on 10 June 2020).

4.3. Chromosomal Distribution, Homology and Motif Analysis

The position information of CHYR genes on chromosome were picked up from soybean and Arabidopsis annotation [39] by using TBtools [40]. Their homology was then analyzed via OrthoVenn2 (https://orthovenn2.bioinfotoolkits.net/home, last accessed on 7 June 2021) [24]. MEME (http://meme.nbcr.net/meme/cgi-bin/meme.cgi, last accessed on 28 June 2021) was further carried out to investigate their motif [41]. Finally, TBtools was applied to make these information visualization [40]. The $K_s$ (non-synonymous rate), $K_s$ (synonymous rate), and $K_a/K_s$ ratios were calculated by using TBtools according to their coding sequence. The duplication time was calculated according to published method by using the following formula: Time = $K_s/(2 \times$ substitution rate) and the substitution rates of soybean and Arabidopsis are $6.1 \times 10^{-9}$, and $1.5 \times 10^{-8}$ site per year, respectively [42,43].

4.4. Expression Analysis of GmCHYR during Soybean Development and Response to Abiotic Stresses

The transcription data of GmCHYR from nine tissues/organs and under abiotic stresses (GSE57252 for drought and salt stress) [27,28] in soybean were collected from Phytozome, the 1KP Project (http://www.onekp.com/, last accessed on 6 July 2020) and NCBI GEO DataSets (https://www.ncbi.nlm.nih.gov/gds/, last accessed on 6 July 2020). Then, the correlation heatmap was analyzed by using TBtools [40].
4.5. Quantitative Real-Time PCR Analyses

Consistent with the transcription data, *G. max* seeds were cultured in distilled water for 1 day. Then, the swelled seeds were removed to a new Petri dish and covered with wet gauze. Once the roots grew to 1 cm, they were transferred into 1/4 Hoagland’s solution with 60% relative humidity, 24 °C, and 16 h light/8 h dark. When soybean reached the v1 stage (first trifoliolate stage), seedlings were transferred into 100 mM NaCl solution (salt stress) and 50 mM NaHCO$_3$ (alkaline stress). For dehydration stress, soybean seedlings were removed from cultural solution and dried in air. Roots were harvested after 0 h, 1 h, 3 h, 6 h, 12 h, and 24 h. Three individual plants per point were used for each stress level.

Total RNA was extracted by using the TRIZOL reagent (Invitrogen, Waltham, MA, USA). Then, the cDNA was synthesized through the HiScript III RT SuperMix for qPCR (Vazyme, Nanjing, China). TransStart® Top Green qPCR SuperMix (TransGen Biotech, Beijing, China) was used to perform quantitative real-time PCR (qRT-PCR). The relative expression levels were calculated according to the formula $2^{-\Delta\Delta C_T}$ [44]. The expression levels were normalized to 1 at 0 h. Then, the relative fold changes of other points were calculated compared with 0 h. Gene specific primers of *GmCHYRs* and the internal reference gene (*soybean ubiquitin 3, Glyma.20g141600*) [45] were listed in Table S4.

All of the above numerical data were subjected to statistical analyses using EXCEL 2010 and Prism 9 statistical software by Student’s *t*-test.

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