Genome-wide analysis of basic helix-loop-helix transcription factors in papaya (Carica papaya L.)

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Abstract

The basic helix-loop-helix (bHLH) transcription factors (TFs) have been identified and functionally characterized in many plants. However, no comprehensive analysis of the bHLH family in papaya (Carica papaya L.) has been reported previously. Here, a total of 73 CpbHLHs were identified in papaya, and these genes were classified into 18 subfamilies based on phylogenetic analysis. Almost all of the CpbHLHs in the same subfamily shared similar gene structures and protein motifs according to analysis of exon/intron organizations and motif compositions. The number of exons in CpbHLHs varied from 1 to 10 with an average of 5. The amino acid sequences of the bHLH domains were quite conservative, especially Leu-27 and Leu-63. Promoter cis-element analysis revealed that most of the CpbHLHs contained cis-elements that can respond to various biotic/abiotic stress-related events. Gene ontology (GO) analysis revealed that CpbHLHs mainly functions in protein dimerization activity and DNA-binding, and most CpbHLHs were predicted to localize in the nucleus. Abiotic stress treatment and quantitative real-time PCR (qRT-PCR) revealed some important candidate CpbHLHs that might be responsible for abiotic stress responses in papaya. These findings would lay a foundation for further investigate of the molecular functions of CpbHLHs.

Introduction

Since plants are unable to move, plant growth and development are regularly affected by abiotic and biotic stresses, which impair yields and result in losses to farmers. For better growth and development, plants have to make use of a series of physiological and biochemical processes in their responses to multiple abiotic stresses by regulating gene expression (Agarwal et al. 2006; Feller et al. 2011; Pires & Dolan 2010). And previous studies have demonstrated that these physiological and biochemical mechanisms are more likely to be a polygenic cooperative defense response induced by various stresses, rather than the single response of a single gene (Zhang et al. 2004). Therefore, the traditional method of obtaining the stress tolerance of plants by modification a single resistance/sensitive gene is limited. Comprehensive analysis of important gene families are very important for molecular breeding.

As an important and popular fruit, papaya is famous for its high nutritional and medical values. Papaya is widely grown in southern China, the tropics and subtropics areas, and its demand is increasing every year. However, the production and quality of papaya were often threatened by various abiotic stresses, such as salt, drought, and cold. These stresses often cause severe economic losses in papaya production in China. So it is very important to study the functions of gene families that involved in abiotic stresses response in papaya. Since obtaining the whole genome sequences of papaya (Ming et al. 2008), several important gene families have been identified by the tool of genome-wide analysis in papaya, including Aux/IAA gene family, ARF family, SQUAMOSA promoter binding protein-like (SPL) gene family, NBS resistance gene family and NPR1 family. These families were essential for papaya fruit ripening, flower and fruit development, fitness and disease resistance (Liu et al. 2017a; Liu et al. 2015; Peraza-Echeverria et al. 2012; Porter et al. 2009; Xu et al. 2020).
In various stresses regulation network and signaling pathways, transcription factors (TFs) are a kind of important proteins that regulate gene expression by activating and repressing related downstream genes. Among them, WRKY and bHLH families are the most common TF families in higher plants (Kosugi & Ohashi 2002). And the WRKY transcription factors has been reported to be related to abiotic and biotic stresses responses in papaya (Pan & Jiang 2014). Basic/helix-loop-helix (bHLH) TFs are widely found in almost all eukaryotes and are the second largest TFs family in plants (Carretero-Paulet et al. 2010; Feller et al. 2011; Jones 2004; Pires & Dolan 2010). The bHLH superfamily proteins are defined by one highly conserved bHLH domain, which comprises approximately 60 amino acids in length and contains two different functional regions: the basic region and the HLH region (Li et al. 2006; Toledo-Ortiz et al. 2003). The basic region is located at the N-terminal end of the bHLH domain and consists approximately 15 amino acids. It is a DNA-binding region that enables bHLH TFs to bind to a specific E-boxes (CANNTG) (Atchley & Fitch 1997; Atchley et al. 1999). The HLH region, at the C-terminal end, is mainly composed of hydrophobic residues, containing two amphipathic α-helices linked by a loop region that has variable sequences and acts as a dimerization domain (Heim et al. 2003; Li et al. 2006). Outside of the two conserved regions, the rest of the bHLH protein sequences are usually very different (Morgenstern & Atchley 1999).

In animals, the bHLH TFs can be divided into six main groups (designated A to F) based on phylogenetic analysis, functional properties and DNA-binding specificity (Atchley & Fitch 1997). These bHLH groups can be divided into several small subfamilies (Ledent & Vervoort 2001; Simionato et al. 2007). The bHLHs mainly function in sensing the external environment, cell cycle regulation and tissue differentiation (Amoutzias et al. 2004; Atchley & Fitch 1997; Stevens et al. 2008; Vervoort & Ledent 2001). Compared to animals, the research on bHLH proteins in plants is limited, even the exact number subfamilies of bHLH TFs has not been determined. Generally, the bHLH proteins is thought to cover 15–25 subfamilies (Buck & Atchley 2003; Pires & Dolan 2010), but some atypical bHLHs have extended the number to 32 based on phylogenetic analysis in plants (Carretero-Paulet et al. 2010). With the availability of genome sequence data and the rapid development of molecular biology, increasing numbers of bHLH subfamily genes have been identified and characterized in a wide range of plant species, including Arabidopsis (Toledo-Ortiz et al. 2003), peanut (Gao et al. 2017), apple (Mao et al. 2017), tomato (Sun et al. 2015), potato (Wang et al. 2018b), peach (Zhang et al. 2018), grapes (Wang et al. 2018a), sweet orange (Geng & Liu 2018), and bamboo (Cheng et al. 2018). The results from these research have shown that bHLH TFs have versatile biological functions, such as regulating light morphogenesis (Leivar et al. 2008; Roig-Villanova et al. 2007), hormone signals (Friedrichsen et al. 2002; Lee et al. 2006), the developmental of root (Feng et al. 2017) and anther (Farquharson 2016), regulating epidermal cell fate determination (Bernhardt et al. 2003), participating in various biotic and abiotic stress responses (Jiang et al. 2009; Liu et al. 2014; Wang et al. 2018b), etc.

In recent years, some studies demonstrated that bHLH transcription factors play important roles in the stress-related regulation network and signaling pathways in many species. However, no
systematic analysis of the bHLH TFs have previously been performed in papaya. In this study, a total of 73 CpbHLH genes were identified in papaya, and phylogenetic analyses were carried out to analyze the relationships among these genes. Meanwhile, gene structure, protein physicochemical properties and conserved motifs, the cis-element of the promoter region, and gene ontology (GO) analysis were investigated. Furthermore, to analyze the functions of CpbHLHs responsible for responding to abiotic stresses, the expression profiles of 22 selected genes under salt, drought, ABA and cold stresses were investigated by using quantitative real-time PCR (qRT-PCR). We identified several important candidate genes that might be responsible for abiotic stress responses. We completed the first comprehensive genome-wide analysis of the bHLH gene family in papaya, and our results provide information necessary for further functional research of the bHLH family in papaya.

### Materials & Methods

#### Identification of CpbHLH genes, gene structure and physicochemical analysis

Papaya (*Carica papaya L.*) bHLH protein sequences were downloaded from the Plant TFDB V4.0 database (Jin et al. 2017). Furthermore, we used the SMART online software (http://smart.embl-heidelberg.de/) and the InterProScan tool (http://www.ebi.ac.uk/Tools/pfa/iprscan/) to identify integrated bHLH domains in putative papaya bHLH proteins. The physicochemical properties of CpbHLH proteins were predicted by ProPAS (Wu & Zhu 2012). The genomic sequences, ID numbers and coding sequences (CDS) corresponding to each predicted CpbHLH gene were obtained from the Phytozome database (https://phytozome.jgi.doe.gov/pz/portal.html). The intron numbers, exon–intron organizations and locations of the CpbHLH genes were analyzed by Gene Structure Display Server (GSDS) v2.0 (Hu et al. 2015).

#### Phylogenetic tree building, motif identification and multiple sequence alignment

To research the phylogenetic relationship of CpbHLH proteins, protein sequences of papaya were pre-aligned using HMM align (Eddy 1998) and the pHMM HLH ls.hmm from PFAM (https://pfam.xfam.org/family/PF00010) to identify the domains of bHLH TFs. Based on the manually aligned bHLH region of 158 bHLH proteins from *Arabidopsis* and 173 from rice (Pires & Dolan 2010), the identified bHLH domains were later aligned using MAFFT v7.305b (Katoh et al. 2002) with default settings. Phylogenetic tree was constructed based on the neighbor-joining method using FastTree v2.1.11 (Price et al. 2009) with default settings. Bootstrapping with 1000 replicates was used to assess the statistical reliability of nodes in the tree. Multiple sequence alignment based on protein sequences of these 73 CpbHLH TFs was generated by MAFFT v7.305b (Katoh et al. 2002) with default settings.

To identify the conserved motifs among the CpbHLH proteins, we uploaded the 73 amino acid sequences of the CpbHLH family to the Multiple EM for Motif Elicitation (MEME, version 5.02) (http://meme-suite.org/tools/meme). The parameter settings were as follows: zero or one, occurrence of a single motif per sequence; 3, maximum number of motifs found. All other parameters were set to the default values.

#### Promoter cis-acting Regulatory Element Analysis and Gene Ontology (GO) Annotation
To predict and compare the putative promoter *cis*-elements of *bHLHs* in papaya and *Arabidopsis*, the upstream 2000 bp genomic DNA sequences of 73 *CpbHLH* genes in papaya, and 47 *AtbHLH* genes in *Arabidopsis* (the putative orthologous genes corresponding to *CpbHLHs*) were downloaded and then submitted to the PlantCARE (Magali et al. 2002). The full-length protein sequences of papaya *bHLH* were blasted against *Arabidopsis* proteins with default parameters. The best hits were submitted to AgriGOv2.0 (http://systemsbiology.cau.edu.cn/agriGOv2/) for GO annotation (Tian et al. 2017). GO terms include three aspects: biological process, cellular component and molecular function.

**Plant materials, growth conditions and stress treatments**

In this experiment, stems with axillary buds were selected as explants from two-year-old ‘Yi Chi Gua’ papaya trees grown under standard field conditions in the Institute of Fruit Tree Research, Guangdong Academy of Agriculture Science, Guangzhou, China, and cultured in vitro to obtain the complete papaya seedlings with normal leaves and roots using tissue culture techniques. Healthy and uniform papaya seedlings were used for different treatments. For the selection of stress conditions for papaya, we designed different gradients of stress conditions for pre-experiments: the concentration gradients of salt stress are 100 mM Nacl, 200 mM Nacl, and 300 mM Nacl; the concentration gradients of PEG6000 (to mimic drought stress) are 15% PEG6000, 20% PEG6000, 25% PEG6000 and 30% PEG6000; the concentration gradients of ABA are 50μM ABA, 100μM ABA and 150μM ABA; the temperature gradients are 0 ℃, 4 ℃ and 10 ℃, and finally determined the suitable stress conditions used in this manuscript. For salt, drought and ABA stresses, seedlings were treated with MS liquid medium containing 200 mM Nacl, 25% PEG6000 and 100 μM ABA for 2 hours respectively, and then the roots were collected. For cold treatment, seedlings were subjected to 4℃ for 2 hours and the leaves were collected. All of the collected materials were immediately frozen in liquid nitrogen and stored at -80℃ for RNA isolation. Untreated seedlings were used as the control groups. Three biological replications were carried out for each treatment.

**RNA extraction and quantitative real-time PCR (qRT-PCR) analysis**

Total RNA from papaya after different treatments was isolated using TRIzol reagent (Invitrogen). The extracted RNA was treatment with DNase (TaKaRa), and then reverse transcribed into cDNA using the PrimeScriptTM RT Reagent Kit (TaKaRa). The qRT-PCR was conducted on the ABI StepOne Real Time PCR system using 2X SG Fast qPCR Master Mix (High Rox) (TaKaRa) according to the manufacturer’s instructions. TATA binding protein 2 (*CpTBP2*) amplification was used as an internal control (Zhu et al. 2012). The qRT-PCR reactions used three biological replicates, and each biological repeat had three technical replicates. Gene-specific primers for qRT-PCR of the 22 *CpbHLH* genes were designed based on the CDSs of the *CpbHLH* genes using Primer Premier 5.0 (Data S1). The relative expression levels of each gene were calculated using the 2-ΔΔCT method, the raw data was showed in supplemental file 1.

**Results**
Identification and characterization of CpbHLHs

A total of 105 putative bHLH transcription factors of papaya (C. papaya) were downloaded from the PlantTFDBv2.0 (http://planttfdb.cbi.pku.edu.cn/). To verify the reliability of these results, the 105 CpbHLH proteins sequences were filtered by Interproscan and SMART domain annotation, and a total of 73 predicted CpbHLH proteins were identified. They were named CpbHLH001 to CpbHLH073 at random except for 32 proteins that were explicitly excluded by Interproscan and SMART (Table S1). The detailed information on these predicted CpbHLHs, including protein ID, locus ID, opening reading frame (ORF) lengths, amino acid sequences/lengths, molecular weight, isoelectric point and exon/intron numbers, are listed in Data S2. In previous studies, 129/132, 188, 159, 124, 95, 94 and 56 bHLH genes were identified in peanut (Gao et al. 2017), apple (Mao et al. 2017), tomato (Sun et al. 2015), Arabidopsis (Toledo-Ortiz et al. 2003), potato (Wang et al. 2018b), peach (Zhang et al. 2018), grapes (Wang et al. 2018a) and sweet orange (Geng & Liu 2018), respectively. Compared with the above dicotyledonous plants, the density of bHLHs genes in papaya genome was about 0.26%, which is lower than the density of peanut (Gao et al. 2017), apple (Mao et al. 2017), tomato (Sun et al. 2015), Arabidopsis (Toledo-Ortiz et al. 2003), potato (Wang et al. 2018b) and sweet orange (Geng & Liu 2018), and similar to peach (Zhang et al. 2018) and wine grapes (Wang et al. 2018a) (Table 1). This is probably associated with the whole-genome duplications during evolution. Among the above plants, some plants with recent whole-gene duplication like peanut, apple, tomato, Arabidopsis, potato and sweet orange while the plants without whole-gene duplication like papaya, peach and wine grapes.

To further characterize the bHLHs in papaya, the physicochemical properties of these putative proteins were analyzed and are shown in Data S2. The size of deduced CpbHLHs ranged from 100 (CpbHLH053) to 679 (CpbHLH068) amino acids, the corresponding molecular weights from 11.525 KDa to 75.899 KDa. The predicted theoretical isoelectric points (PI) values of CpbHLHs were between 4.71(CpbHLH028) and 11.07(CpbHLH003). Similar molecular weights and isoelectric points have been made in potato (Wang et al. 2018b). And all of predicted CpbHLH proteins were hydrophilic characteristic proteins, the grand average of hydropathy values were negative, ranging from -0.2098(CpbHLH033) to -1.0125(CpbHLH006). Similar result has been made in Brachypodium distachyon (Niu et al. 2017). That is, the predicted CpbHLH proteins showed diversities in their length, molecular weight, PI and the grand average of hydropathy values.

Phylogenetic analysis, gene structure, conserved motifs analysis and multiple sequence alignment of CpbHLHs

To evaluate the evolutionary relationships of the CpbHLH proteins, a neighbor-joining phylogenetic tree was generated using conserved bHLH domains from papaya, Arabidopsis and rice. The phylogenetic tree showed that the 73 CpbHLH members were clustered into 18 subfamilies with one orphan (Fig.1A and Data S3), consistent with the earlier results showing that the bHLH subfamily in plants can be divided into 15-25 subfamilies (Pires & Dolan 2010). Previous research have named the bHLH subfamilies using English letters (Li et al. 2006; Mao et
al. 2017), Roman numerals (Song et al. 2017; Sun et al. 2015), or Arabic numerals (Chen et al. 2015; Toledo-Ortiz et al. 2003). In this study, we named CpbHLH subfamilies using Roman numerals. As shown in figure 1, the subfamily XII was the largest subfamily among all three species, and all of subfamilies include at least two species. In papaya, none of the bHLHs were grouped into IVd, II, X V, X, XI V and X III subfamilies compared to rice and Arabidopsis, which may be due to these bHLHs were lost during the process of evolution.

Exon/intron organization, as a type of structural divergence, plays an important role in the evolution of multiple gene families (Xu et al. 2012). The annotation features of the CpbHLH genes were submitted to Gene Structure Display Server (GSDS) together to show their gene structures. As described in Data S2 and Figure 2A, the number of introns varied from zero to ten, representing a complex distribution pattern. Most (63 (86.3%)) of the CpbHLHs were found to possess introns among the 73 CpbHLH genes, while 10 (13.7%) of the genes were intron-less, 8 (11.0%) genes contained one intron, and the remaining genes had two or more introns. In addition, members of the same subfamily also displayed similar intron distribution patterns in view of the full-length genome sequences. For instance, all of the CpbHLHs in subfamily Vb had one intron and two exons, the whole members of subfamily IIIf had six introns and seven exons, the IVa subfamily members showed three introns and four exons, and all members of VIIIb subfamily consisted only one exon.

Most importantly, members of the same bHLHs subfamily are usually participated in the same signaling pathway or biological process, and the functions of these members are often partially or totally redundant (Pires & Dolan 2010). For example, AtbHLH10, AtbHLH89 and AtbHLH91, corresponding rice orthologs OsbHLH141, OsbHLH142 are members of subfamily II, they are all involved in the process of pollen development (Li et al. 2006; Liu et al. 2017; Zhu et al. 2015). Especially in Arabidopsis, there is no obvious phenotype in single mutant of AtbHLH10, AtbHLH89 or AtbHLH91, only their various double or triple mutants showed the phenotype of pollen development deficiency (Liu et al. 2017). In subfamily IIIb, OsbHLH001 (OsICE2), OsbHLH002 (OsICE1), CpbHLH027, CpbHLH062, AtbHLH116 (ICE1) and AtbHLH33 (ICE2) were clustered within one clade. In previous studies, AtbHLH116(ICE1) and AtbHLH33(ICE2) and corresponding orthologs in rice (OsbHLH001/OsICE2, OsbHLH002/OsICE1) have been reported to function in the stress of chilling (Chinnusamy et al. 2003; Deng et al. 2017; Fursova et al. 2009; Li et al. 2010; Zhang et al. 2017). And we also found transcripts of CpbHLH027 and CpbHLH062 were increased under chilling stress in this study, implying that CpbHLH027 and CpbHLH062 are involved in the process of chilling stress in papaya.

To further study the sequence characteristics of the predicted bHLH domains at the amino acid level, we carried out a multiple sequence alignment of the 73 predicted CpbHLH protein sequences (Fig.3). The result showed that the 73 putative CpbHLH proteins contained two conserved regions in the bHLH domains: the basic region plus helix 1 and the loop region plus helix 2 (Fig. 3 and Table S2). Additionally, we used the online MEME program to identify the conserved motifs (Bailey & Elkan 1994). The result also showed that most of the sequences
excise Cp\textit{bHLH003} exhibited two highly conserved motifs: one is contains 29 amino acids, and the other consists of 21 amino acids, are shown in red and blue blocks, respectively (Fig.2B). Among the two motifs, motif 1 comprises basic residues and helix 1, and motif 2 comprises a loop and helix 2. And the space between motif 1 and 2 consists of a loop, which is variable in length in some bHLH proteins. The sequence logos of motif 1 (in red) and motif 2 (in blue) are shown in Figure 4A. The backbones of motif 1 and 2 are also conserved in most plant species (Guo & Wang 2017; Heim et al. 2003; Sun et al. 2015), and these highly conserved residues in bHLH domains may be responsible for protein dimerization (Heim et al. 2003).

Besides these two common conserved motifs, some \textit{Cp} bHLHs that are mainly distributed into eight subfamilies (including \textit{V}a, \textit{V}b, \textit{III}f, \textit{IV}a, \textit{III}b, III, \textit{I}b and \textit{I}a subfamilies) harbor another conserved motif (motif 3) with a length of 36 amino acids. The motif 3 is indicated by the green blocks and the sequence logo is visualized as logo3 (Fig. 2B and Fig. 4B). This result is accord with the previous studies that members of a given subfamily exhibited another conserved nonbHLH motif (motif 3) in plant bHLH superfamily (Pires & Dolan 2010). However, in papaya, members of the bHLH proteins have the same motif that is distributed into eight subfamilies, not just one subfamily. In addition, among the 73 \textit{Cp}bHLHs, one atypical bHLH protein (\textit{Cp}bHLH003) exhibited incomplete bHLH domains, whereas the remaining 72 \textit{Cp}bHLH proteins all presented complete bHLH domains. Similar observations have been made in other plant species, such as peach and blueberry (Song et al. 2017; Zhang et al. 2018).

Promoter analysis of \textit{bHLH} genes in papaya
To further understand \textit{Cp}bHLHs functions and regulation patterns, \textit{cis}-elements in \textit{Cp}bHLH genes promoter sequences were investigated. Regions of 2,000 bp upstream from the start codons of each \textit{Cp}bHLH gene were analyzed using PlantCARE. The results showed that the \textit{cis}-elements could be divided into three main categories (Fig. 5A and Data S4). Category one contained a ubiquitous class of plant light responsive elements among which G-Box, G-box, GT1-motif and Box 4 were common in the \textit{Cp}bHLH promoters. Category two contained important elements that were involved in the process of stress-responsiveness, including MYB binding site involved in drought-inducibility (MBS), low temperature response elements (LTR), defense and stress responsive elements (TC-rich) and wound-responsive elements (WUN-motifs). In addition, more than ten kinds of hormone-responsive \textit{cis}-elements were identified (e.g., gibberellin-\textit{GA}, auxin-\textit{IAA}, methyl jasmonate-\textit{MeJA}, salicylic acid-\textit{SA}, and abscisic acid-\textit{ABA}). Among them, the most common response elements were \textit{ABA} (\textit{ABRE}), \textit{MeJA} (\textit{CGTCA-motif} and \textit{TGACG-motif}) and \textit{SA} (\textit{TCA-element} and \textit{SARE}), which included 158 (29.15%), 128 (23.62%) and 53 (9.78%), respectively (Fig. 5B). Category three contained plant growth and development elements, such as anaerobic induction elements (\textit{ARE}), \textit{O}$_2$-site, CAT-box and so on. Additionally, we also analyze the \textit{cis}-elements in the promoter regions of putative orthologous genes that corresponding to \textit{Cp}bHLH in \textit{Arabidopsis}, and the similar result has been obtained in \textit{Arabidopsis} (Fig. S1 and Data S4). There also existed three main categories: plant light, abiotic and biotic stresses and plant growth and development responsive elements. And the percentage of most stress-responsive elements in \textit{Arabidopsis} were similar to papaya, including
ABA responsive elements, drought-responsive elements, wound-responsive elements, low temperature-responsive elements and IAA responsive elements, implying that most of the promoter cis-elements of bHLH family were conserved in Arabidopsis and papaya.

**GO annotation of CpbHLH proteins**

To understand the functions of papaya bHLHs, we performed a GO annotation of CpbHLHs, and the results are shown in Data S5. A total of 70 CpbHLHs were involved in protein dimerization activity (GO: 0046983). The result is consistent with the earlier studies, which show that the HLH domain was necessary for protein dimerization and DNA binding (Murre et al. 1989). Some conserved amino acid residues are important to the function of bHLH proteins, especially the Leu-27 in helix 1 and the Leu-73 in helix 2 (Carretero-Paulet et al. 2010). In this study, we found 72 (out of 73) CpbHLH proteins have Leu-27 (corresponding to Leu-27 in Arabidopsis bHLHs), and all of the CpbHLH proteins have Leu-63 (corresponding to Leu-73 in Arabidopsis bHLHs) (Fig. 3 and Table S2).

Because of a lack of reported experimental data and databases, we used Arabidopsis as the reference species to perform a GO annotation of CpbHLH proteins, and 54 of 73 predicted CpbHLH proteins were obtained with results compared to Arabidopsis. We summarized the results in Figure 6 and Data S6. The majority of predicted CpbHLH proteins were involved in DNA binding. Almost all of the predicted CpbHLH proteins (37, 68.5\%) were predicted to localize in the nucleus, whereas the remaining predicted CpbHLH proteins were located in other organelles, including plastids, the cytoplasm, and chloroplasts. Additionally, some predicted CpbHLH proteins existed in multiple cellular components. For example, CpbHLH013 was located in three cellular components: chloroplasts, part of the cytoplasm, and the nucleus, which may reflect its multiple functions in various biological processes. The metabolic processes involved the greatest number of putative CpbHLH proteins (47, 87.0\%). Biosynthetic processes and gene expression involved the second greatest number of putative CpbHLH proteins (46, 85.2\%). In addition, CpbHLH proteins could respond to stimulus, morphogenesis, cell differentiation, and developmental process.

**Expression analysis of bHLH superfamily genes under different abiotic stresses**

The bHLH proteins have been characterized functionally in many plants with a vital role in the regulation of diverse biological processes, but little is known about their role in papaya. To analyze the functions of CpbHLHs responding to abiotic stresses, the expression profiles of 22 selected genes under salt, drought, ABA and cold stresses were investigated by using qRT-PCR (Table 2 and Fig. 7). The results showed that 4 (CpbHLH011, CpbHLH022, CpbHLH027 and CpbHLH056) of 22 CpbHLH mRNAs were increased, and 3 CpbHLH (CpbHLH020, CpbHLH053 and CpbHLH062) mRNAs were reduced more than 2-fold in salt (200 mM NaCl) treated papaya seedlings. Under drought stress (25\% PEG), 8 (CpbHLH011, CpbHLH022, CpbHLH027, CpbHLH046, CpbHLH050, CpbHLH052, CpbHLH056 and CpbHLH068) of 22 CpbHLH mRNAs were upregulated, and 3 CpbHLH (CpbHLH020, CpbHLH042 and CpbHLH053) mRNAs were downregulated more than 2-fold. Under ABA treatment (100 \(\mu\)M), 3 (CpbHLH027, CpbHLH052 and CpbHLH056) of 22 CpbHLH mRNAs were upregulated, and 5
CpbHLH (CpbHLH019, CpbHLH020, CpbHLH042, CpbHLH053 and CpbHLH062) mRNAs were downregulated. Under cold stress (4°C), there were 4 CpbHLH genes (CpbHLH027, CpbHLH035, CpbHLH056 and CpbHLH062) whose expression increased more than 1.5-fold, and 4 CpbHLH (CpbHLH046, CpbHLH050, CpbHLH052 and CpbHLH068) mRNAs were reduced more than 2-fold. Interestingly, a few transcripts of CpbHLH responded to all or multiple stresses. For instance, CpbHLH056 was sensitive to all four stresses and was upregulated distinctly under the four stresses. The orthologue of CpbHLH056 in Arabidopsis is BEE1 (AtbHLH044) (Fig. 1), which has been functionally characterized in previous reports. At low temperatures, BEE1 is a positive regulator of flavonoid accumulation (Petridis et al. 2016), which is consistent with our results. In addition, BEE1, BEE2 and BEE3 are functionally redundant positive regulators of BR (brassinosteroid) signaling, but these transcripts are repressed by ABA (Friedrichsen et al. 2002). However, we found the transcription of CpbHLH056 was notably upregulated (>10-fold) under ABA treatment. More interestingly, CpbHLH042, which is an orthologue of BEE2 (Fig.1), was distinctly repressed by ABA (approximately 4-fold). These results suggested that CpbHLH056 and CpbHLH042 may provide different functionalities compared to Arabidopsis. Additionally, CpbHLH027 was also upregulated distinctly under four stresses. In Arabidopsis, the orthologue of CpbHLH027 is AtbHLH116 (ICE1) (Fig. 1), which can be induced by Nacl, ABA and cold stresses, playing an important role in the cold-responsive signaling pathway via an ABA-independent pathway (Chinnusamy et al. 2003). There are two orthologues of CpbHLH027 in rice, one ortholog is OsICE2/OsbHLH001, is induced by salt stress, and its overexpression can enhanced the tolerance to freezing and salt stress (Deng et al. 2017; Li et al. 2010). OsICE1/OsbHLH002 is another ortholog in rice, which is induced by chilling stress. OsbHLH002 can positively regulates cold signaling via targeting OsTPP1, which encodes a keyenzyme for trehalose biosynthesis (Zhang et al. 2017). These results implied CpbHLH027 plays essential roles in abiotic stresses in papaya. In addition, the transcript of CpbHLH062 was also increased under cold treatment, its orthologue is AtbHLH033/ICE2, which involving the cold response and the ABA pathway (Fursova et al. 2009; Kurbidaeva et al. 2014), implying the CpbHLH062 may involved in the cold stress. CpbHLH053 was downregulated under salt, drought and ABA stresses. The orthologue of CpbHLH053 is AtbHLH129 (Fig. 1), which is a transcription repressor that negatively regulates the ABA response in Arabidopsis (Tian et al. 2015), implying CpbHLH053 may have the similar function with the AtbHLH129 in the process of ABA response. We should also noticed a few CpbHLHs that showed distinct increases or decreases in their mRNA levels under different treatments, and these CpbHLHs' orthologues have not been reported in previous studies. For instance, CpbHLH050 is notably upregulated (>10-fold) under PEG treatment, CpbHLH046 is upregulated by PEG treatment, but sharply down regulated under ABA and cold treatments, implying these genes may have additional functions than response to drought by regulating root development. CpbHLH020 and CpbHLH053 were downregulated
 (>2-fold) by Nacl, PEG and ABA stresses distinctly. We should also pay attention to these genes in the following research.

**Discussion**

Transcription factors (TFs) play key roles in the stress regulation network and signal pathways in plants. Basic/helix-loop-helix (bHLH) TFs are the second largest TFs family in plants and have been identified in many species (Cheng et al. 2018; Gao et al. 2017; Geng & Liu 2018; Mao et al. 2017; Sun et al. 2015; Toledo-Ortiz et al. 2003; Wang et al. 2018a; Wang et al. 2018b; Zhang et al. 2018). However, the bHLH TF family has not previously been reported in papaya (*Carica papaya L*.). In this paper, we found 73 *CpbHLH* genes in papaya. This TF family seemed to be one of the moderately sized families compared with other plant species, which might be because of the papaya has a relatively small reference genome, the size is only 372 Mb (Ray et al. 2008).

The gene evolution changes the gene organization. In this study, we found that *CpbHLH* genes were diverse in their number introns, ranging from 0 to 10 (Data S2). This result implied these genes may have undergone numerous of genetic evolution events, and the genes in different subfamilies may have different functions (Cheng et al. 2018). Most *CpbHLHs* in the same subfamily shared similar gene structures and protein motifs according to the analysis of exon/intron organizations and motif compositions (Fig. 2, 3 and 4), indicating that the functions of encoded proteins in each subfamily are probably stable. However, the conserved motif analysis showed that some *CpbHLHs*, which are mainly distributed in eight subfamilies (including V a, V b, IIIf, IV a, IIIb, III, 1 b and 1 a subfamilies) from the phylogenetic tree, harbor another conserved motif (motif 3) with a length of 36 amino acids (Fig. 2 and 4), indicating that these proteins may have additional functions.

Promoter cis-acting regulatory element analysis showed that cis-elements could be divided into three main categories: light responsive, abiotic and biotic stresses and plant growth and development. Especially in abiotic and biotic stresses, the most common response elements were ABA (29.15%), MeJA (23.62%) and SA (9.78%), suggesting that these phytohormones may play important roles in the regulation of papaya growth and development (Fig. 5 and Data S4). In addition, the promoter cis-acting element involved in the abscisic acid responsiveness analysis is consistent with the qRT-PCR results (Data S4 and Table 2), showing four genes (*CpHLH020/-027/-053/-056*) involved in abscisic acid response. Another two genes (*CpHLH020/-062*) were also identified that were involved in abscisic acid response by GO annotation analysis and qRT-PCR (Data S6 and Table 2). We also identified a large number of cis-acting elements in *CpbHLH* genes that may respond to drought (MBS, 8.67%), which is also consistent with the qRT-PCR (including seven genes: *CpHLH027/-050/-056/-011/-068/-042/-053*). Other genes also had important elements, including LTR, TC-rich and WUN-motifs, which indicated plant responses to low temperatures, defense stresses and wound-responsiveness, respectively. These results implied *CpbHLH* genes may have a wide range of functions in papaya growth, disease resistance, and response to environmental conditions. We also analyze the cis-elements in the promoter regions of putative orthologous genes in *Arabidopsis* (Fig. S1 and Data S4). And we...
compared the promoter cis-elements of bHLH genes in Arabidopsis, papaya and previously reported bamboo (Cheng et al. 2018). The result showed that the promoter cis-elements of bHLH genes in these three plants were divided into three categories, and most of elements were the same. These similar results implied that most of the promoter cis-elements of bHLH family were conserved in Arabidopsis, papaya and bamboo. The most notable is, the percentage of MeJA responsive cis-elements in papaya and Arabidopsis (23.62%, 30.61%) were less than bamboo (43.39%). And the percentage of SA responsive cis-elements in papaya and Arabidopsis (9.78%, 5.87%) were also less than bamboo (10.31%). SA is a phytohormone that plays important roles in plant defenses against pathogens (Pokotylo et al. 2019). MeJA also has been identified as a vital cellular regulator that mediates defense processes (Cheong & Choi 2003). So, the stress-responsive elements in papaya and Arabidopsis were corresponding less than bamboo, including drought-responsive elements, wound-responsive elements, low temperature-responsive elements, and defense and stress responsive elements. These results may help explain why papaya is more sensitive to external stresses compared to bamboo.

Many studies have shown that bHLH genes are involved in various abiotic and biotic stresses responses. We randomly selected 22 genes to investigate their expression profiles by using qRT-PCR under salt, drought, ABA and cold stresses (Fig.7 and Table 2). The results revealed some candidate CpbHLH genes that might be responsible for abiotic stress responses in papaya. For example, CpbHLH027, CpbHLH062, AtbHLH116 (ICE1), AtbHLH33 (ICE2), OsbHLH001 (OsICE2) and OsbHLH002 (OsICE1) were clustered within one clade. Among them, AtbHLH116 (ICE1), AtbHLH33 (ICE2), OsbHLH001 (OsICE2) and OsbHLH002 (OsICE1) have been reported function in chilling stress in Arabidopsis and rice (Chinnusamy et al. 2003; Deng et al. 2017; Fursova et al. 2009; Li et al. 2010; Zhang et al. 2017). And the transcripts of CpbHLH027, CpbHLH062 were increased under chilling stress in this study, implying that CpbHLH027 and CpbHLH062 may be also involved in the process of chilling stress. The orthologue of CpbHLH056 in Arabidopsis is BEE1 (At1G18400), which is a positive regulator of flavonoid accumulation (Petridis et al. 2016). BEE1, BEE2 and BEE3 are functionally redundant positive regulators of BR signaling, and their transcription is repressed by ABA in Arabidopsis (Friedrichsen et al. 2002). However, we found that the transcription of CpbHLH056 is notably upregulated (>10-fold) under ABA treatment rather than downregulated in this study. These results imply that CpbHLH056 may involved in the process of ABA stress but has different function compared to Arabidopsis. We have also noticed a few candidate CpbHLHs that showed distinct increases or decreases in their mRNA levels under different treatments, and these CpbHLHs' orthologues have not been reported in other plants. For instance, CpbHLH050, CpbHLH020, CpbHLH046, CpbHLH053, and so on. These findings provide important candidate genes/proteins necessary for further functional research on the bHLH family in papaya.

Conclusions
In conclusion, the study performed a genome-wide analysis of basic helix-loop-helix (bHLH) transcription factors in papaya. As a result, a total of 73 bHLH genes were identified in papaya,
and these CpbHLHs were classified into 18 subfamilies with one orphan, which was consistent with the earlier results showing that the bHLH subfamily in plants can be divided into 15-25 subfamilies. Almost all of the CpbHLHs in the same subfamily shared similar gene structures and protein motifs according to analysis of exon/intron organizations and motif compositions. These results further supported the classification predicted by the phylogenetic tree. Compared to rice and Arabidopsis, the amino acid sequences of the CpbHLH domains were quite conservative, especially Leu-27 and Leu-63. Promoter cis-element and GO annotation analysis revealed that most of the CpbHLHs could respond to various biotic/abiotic stress-related events. Abiotic stress treatment and quantitative real-time PCR (qRT-PCR) assay further supported promoter cis-acting regulatory element and GO annotation analysis, revealed some important candidate CpbHLHs that might be responsible for abiotic stress responses in papaya. We completed the first comprehensive genome-wide analysis of the bHLH gene family in papaya, and our results provide information necessary for further functional research of the bHLH family in papaya.

**Figure Titles and Legends**

**Figure 1:** Phylogenic and family members analysis of bHLHs from papaya, rice and Arabidopsis.

(A) The 73 CpbHLHs are clustered into 18 subfamilies. Phylogenetic tree was constructed based on the neighbor-joining method. Bootstrapping with 1000 replicates was used to assess the statistical reliability of nodes in the tree. (B) Comparison of bHLH family members from papaya, rice and Arabidopsis. Different colors represent the different plants. Green: OsbHLHs, red: AtbHLHs, Blue: CpbHLHs.

**Figure 2:** Gene structure and motif distribution of the papaya bHLH family.

(A) Exon-intron organization of CpbHLH genes. Exons and introns are presented as filled orange sticks and thin black single lines, respectively. The brackets and Roman numerals separate each subfamily and clearly present the member conservation of each subfamily. (B) Arrangements of conserved motifs in 73 CpbHLH proteins. Three predicted motifs are represented by different colored boxes, motif 1 (red block), motif 2 (blue block) and motif 3 (green block).

**Figure 3:** Multiple sequence alignment of the bHLH domains in papaya.

Amino acids with more than 50% identity are labeled with colored boxes.

**Figure 4:** Motif composition and logos of papaya bHLH proteins.

(A) The logos of motif 1 and 2, which together constitute the bHLH domain in papaya. The overall height of the character represents the conservation of an amino acid at the specific position. Each color of the English letters represents a type of amino acid residue. (B) The logo of motif 3, which is another conserved motif.

**Figure 5:** Cis-acting element analysis of the promoter of bHLH genes in papaya.

(A) Percentage of total cis-acting elements in the promoter region of CpbHLH genes. (B) The percentage of each cis-acting element in the abiotic and biotic stresses categories.

**Figure 6:** Gene ontology (GO) annotation of CpbHLH proteins.
The annotation was performed on three categories, (A) molecular function, (B) biological processes and (C) cellular components.

**Figure 7**: Quantitative RT-PCR analysis of 22 selected *CpbHLH* genes under cold stress condition (4°C).

The data are expressed as means ± SD of three independent biological determinations. Untreated seedlings were used as the control groups. *P < 0.05 and **P < 0.01 (Student’s t test) indicate significant differences between treated seedlings and control groups.

**Supplemental Information**

- **Supplemental Data S1**: Primers used for qRT-PCR in this study.
- **Supplemental Data S2**: Detailed information of *CpbHLH* genes and *CpbHLH* proteins.
- **Supplemental Data S3**: bHLH subfamily members of papaya, rice and *Arabidopsis*.
- **Supplemental Data S4**: Promoter analysis of *bHLH* genes in papaya and *Arabidopsis*.
- **Supplemental Data S5 and Supplemental Data S6**: GO annotation of *CpbHLH* proteins.
- **Table S1**: The information of 32 predicted proteins from the PlantTFDBv2.0 that were excluded by Interproscan and SMART.
- **Table S2**: Consensus sequences of bHLH domains in papaya, rice and *Arabidopsis*.
- **Supplemental file 1**: Raw data of qRT-PCR assay.
- **Figure S1**: *Cis*-acting element analysis of the promoter of *bHLH* genes in *Arabidopsis*.
  (A) Percentage of total *cis*-acting elements in the promoter region of *AtbHLH* genes. (B) The percentage of each *cis*-acting element in the abiotic and biotic stresses categories.

**Additional Information and Declarations**

**Authors’ contributions**

Min yang conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, approved the final draft.

Chenping Zhou and Hu Yang performed the experiments, analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, approved the final draft.

Ruibin Kuang and Bingxiong Huang performed the experiments, analyzed the data, authored or reviewed drafts of the paper, approved the final draft.

Yuerong Wei conceived and designed the experiments, authored or reviewed drafts of the paper, approved the final draft.

**Data Availability**

Putative bHLH transcription factors were downloaded from the PlantTFDBv2.0 (http://planttfdb.cbi.pku.edu.cn/). The genomic sequences, ID numbers and coding sequences (CDS) corresponding to each predicted *bHLH* gene were obtained from the Phytozome database (https://phytozome.jgi.doe.gov/pz/portal.html). The online tool Multiple EM for Motif Elicitation (MIME, version 5.02) was used to search for conserved motifs among the bHLH proteins.
by uploading the protein sequences of the papaya bHLH superfamily.

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Figure 1

Phylogenic and family members analysis of bHLHs from papaya, rice and Arabidopsis.

(A) The 73 CpHLHs are clustered into 18 subfamilies. Phylogenetic tree was constructed based on the neighbor-joining method. Bootstrapping with 1000 replicates was used to assess the statistical reliability of nodes in the tree. (B) Comparison of bHLH family members from papaya, rice and Arabidopsis. Different colors represent the different plants. Green: OsbHLHs, red: AtbHLHs, Blue: CpHLHs.
Figure 2

Gene structure and motif distribution of the papaya bHLH family.

(A) Exon-intron organization of CpbHLH genes. Exons and introns are presented as filled orange sticks and thin black single lines, respectively. The brackets and Roman numerals separate each subfamily and clearly present the member conservation of each subfamily. (B) Arrangements of conserved motifs in 73 CpbHLH proteins. Three predicted motifs are represented by different colored boxes, motif 1 (red block), motif 2 (blue block) and motif 3 (green block).
Figure 3

Multiple sequence alignment of the bHLH domains in papaya.

Amino acids with more than 50% identity are labeled with colored boxes.
|    | 1 | 5 | 10 | 15 | 20 | 25 | 30 | 35 | 40 | 45 | 50 | 55 | 60 | 65 |
|----|---|---|----|----|----|----|----|----|----|----|----|----|----|----|
| 1  | Q | E | P | R | M | S | T | V | K | K | N | L | S | P |
| 2  | R | S | T | P | K | K | K | Y | K | R | K | L | S | P |
| 3  | R | G | S | P | T | M | S | L | P | K | Y | V | K | R |
| 4  | E | S | S | K | K | K | L | S | P | K | Y | V | K | R |
| 5  | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 6  | M | N | U | C | R | I | P | T | O | B | E | R | V | E |
| 7  | M | N | U | C | R | I | P | T | O | B | E | R | V | E |
| 8  | C | H | A | N | T | E | R | S | L | P | K | Y | V | K |
| 9  | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 10 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 11 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 12 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 13 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 14 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 15 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 16 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 17 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 18 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 19 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 20 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 21 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 22 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 23 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 24 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 25 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 26 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 27 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 28 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 29 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 30 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 31 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 32 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 33 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 34 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 35 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 36 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 37 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 38 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 39 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 40 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 41 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 42 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 43 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 44 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 45 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 46 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 47 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 48 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 49 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 50 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 51 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 52 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 53 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 54 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 55 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 56 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 57 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 58 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 59 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 60 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 61 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 62 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 63 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 64 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 65 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 66 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 67 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 68 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 69 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 70 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 71 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 72 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |
| 73 | A | S | G | Q | T | P | M | S | L | P | K | Y | V | K |

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Figure 4

Motif composition and logos of papaya bHLH proteins.

(A) The logos of motif 1 and 2, which together constitute the bHLH domain in papaya. The overall height of the character represents the conservation of an amino acid at the specific position. Each color of the English letters represents a type of amino acid residue. (B) The logo of motif 3, which is another conserved motif.
**Figure 5**

*Cis*-acting element analysis of the promoter of *bHLH* genes in papaya.

(A) Percentage of total *cis*-acting elements in the promoter region of *CpbHLH* genes. (B) The percentage of each *cis*-acting element in the abiotic and biotic stresses categories.
Figure 6

Gene ontology (GO) annotation of CpbHLH proteins.

The annotation was performed on three categories, (A) molecular function, (B) biological processes and (C) cellular components.
Figure 7

Quantitative RT-PCR analysis of 22 selected *CpbHLH* genes under cold stress condition (4°C).

The data are expressed as means ± SD of three independent biological determinations. Untreated seedlings were used as the control groups. *P < 0.05 and **P < 0.01 (Student’s t test) indicate significant differences between treated seedlings and control groups.
Table 1 (on next page)

Summary of TFs identified from dicotyledonous plant species with genome sequences
| Plant species            | Common name      | bHLH | Proteins | Ratio (%) |
|-------------------------|------------------|------|----------|-----------|
| C. papaya               | Papaya           | 73   | 27829    | 0.26      |
| A. ipaensis/A. duranensis | Peanut            | 129/132 | 7243    | 1.78/1.82 |
| Malus x domestica       | Apple            | 188  | 15173    | 1.24      |
| S. lycopersicum         | Tomato           | 159  | 15722    | 1.01      |
| A. thaliana             | Arabidopsis      | 147  | 32125    | 0.46      |
| S. tuberosum            | Potato           | 124  | 17445    | 0.71      |
| P. persica              | Peach            | 95   | 28299    | 0.34      |
| V. vinifera             | Wine Grapes      | 94   | 47097    | 0.20      |
| C. sinensis             | Valencia Orange  | 56   | 13522    | 0.41      |
Expression levels of *CpbHLH* genes under salt, drought and ABA stresses

Quantitative RT-PCR was used to investigate the expression levels (shown in fold change) of the *CpbHLHs*. The expression level in the control (CK) was set at 1.0. The means of three replicates of qRT-PCR and standard deviations (SD) values are shown.
Table 2:

Expression levels of *CpbHLH* genes under salt, drought and ABA stresses

| The name of *CpbHLHs* | CK | NaCl (200 mM) | PEG (25%) | ABA (100 μM) |
|-----------------------|----|---------------|------------|--------------|
| *CpbHLH006*           | 1  | 0.74±0.13     | 0.95±0.15  | 0.75±0.07    |
| *CpbHLH011*           | 1  | 2.45±0.04     | 4.06±0.12  | 1.66±0.12    |
| *CpbHLH019*           | 1  | 0.80±0.03     | 0.64±0.02  | 0.18±0.02    |
| *CpbHLH020*           | 1  | 0.20±0.003    | 0.39±0.01  | 0.22±0.01    |
| *CpbHLH022*           | 1  | 4.21±0.03     | 3.05±0.03  | 1.54±0.08    |
| *CpbHLH027*           | 1  | 4.49±0.09     | 2.32±0.06  | 2.27±0.12    |
| *CpbHLH035*           | 1  | 1.62±0.08     | 1.79±0.07  | 0.71±0.02    |
| *CpbHLH037*           | 1  | 0.96±0.02     | 1.04±0.01  | 1.92±0.07    |
| *CpbHLH040*           | 1  | 1.38±0.04     | 0.77±0.03  | 0.51±0.03    |
| *CpbHLH041*           | 1  | 0.55±0.01     | 0.76±0.03  | 0.70±0.03    |
| *CpbHLH042*           | 1  | 0.55±0.03     | 0.33±0.04  | 0.26±0.03    |
| *CpbHLH046*           | 1  | 0.78±0.06     | 5.46±0.10  | 0.53±0.03    |
| *CpbHLH050*           | 1  | 0.53±0.04     | 11.86±0.10 | 1.18±0.02    |
| *CpbHLH052*           | 1  | 1.64±0.06     | 8.71±0.06  | 4.42±0.13    |
| *CpbHLH053*           | 1  | 0.44±0.04     | 0.31±0.02  | 0.19±0.01    |
| *CpbHLH056*           | 1  | 7.11±0.04     | 32.65±0.45 | 12.22±0.21   |
| *CpbHLH060*           | 1  | 0.89±0.07     | 0.76±0.05  | 0.75±0.11    |
| *CpbHLH062*           | 1  | 0.38±0.02     | 0.65±0.02  | 0.30±0.02    |
| *CpbHLH065*           | 1  | 1.64±0.06     | 1.04±0.03  | 0.52±0.05    |
| *CpbHLH068*           | 1  | 1.72±0.05     | 2.37±0.04  | 1.39±0.07    |
| *CpbHLH069*           | 1  | 0.68±0.07     | 0.89±0.18  | 0.52±0.08    |
| *CpbHLH070*           | 1  | 1.11±0.02     | 1.17±0.07  | 1.47±0.04    |

Quantitative RT-PCR was used to investigate the expression levels (shown in fold change) of the *CpbHLHs*. The expression level in the control (CK) was set at 1.0. The means of three replicates of qRT-PCR and standard deviations (SD) values are shown.