Article

Large-scale analyses of heat shock transcription factors and database construction based on whole-genome genes in horticultural and representative plants

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Abstract

Heat shock transcription factor (Hsf) plays a critical role in regulating heat resistance. Here, 2950 Hsf family genes were identified from 111 horticultural and representative plants. More Hsf genes were detected in higher plants than in lower plants. Based on all Hsf genes, we constructed a phylogenetic tree, which indicated that Hsf genes of each branch evolved independently after species differentiation. Furthermore, we uncovered the evolutionary trajectories of Hsf genes by motif analysis. There were only six motifs (M1–M6) in lower plants, and then four novel motifs (M7–M10) appeared in higher plants. However, the motifs of some Hsf genes were lost in higher plants, indicating that Hsf genes have undergone sequence variation during their evolution. The number of Hsf genes lost was greater than the number of genes that were duplicated after whole-genome duplication in higher plants. The heat response network was constructed using 24 Hsf genes and 2421 downstream and 222 upstream genes of Arabidopsis. Further enrichment analysis revealed that Hsf genes and other transcription factors interacted with each other in the response to heat stress. Global expression maps were illustrated for Hsf genes under various abiotic and biotic stresses and several developmental stages in Arabidopsis. Syntenic and phylogenetic analyses were conducted using Hsf genes of Arabidopsis and the pan-genome of 18 Brassica rapa accessions. We also performed expression pattern analysis of Hsf and six Hsp family genes using expression values from different tissues and heat treatments in B. rapa. The interaction network between the Hsf and Hsp gene families was constructed in B. rapa, and several core genes were detected in the network. Finally, we constructed an Hsf database (http://hsfdb.bio2db.com) for researchers to retrieve Hsf gene family information. Therefore, our study will provide rich resources for the study of the evolution and function of Hsf genes.

Introduction

Nowadays, nearly all plants often suffer from several abiotic stresses under a vast range of different environments [1, 2]. Generally, the abiotic stresses mainly include drought, high salinity, water deficiency, radiation, acidity, and low or high temperature [1–3]. Among these extreme stresses, high temperature is one of the most important factors as it affects plant distribution [4]. Heat stress also affects plant photosynthesis, induces cellular death, and destroys cell membranes [5, 6]. In response to heat stress, plants need to alleviate its adverse effects to maintain their yield or production quality [7, 8]. Therefore, plant growth and development need to adapt to heat stress [9, 10]. It is of critical importance to control the expression of stress-responsive genes to adapt to heat stress [11–13]. Hsf is a key transcription factor gene family, which responds to heat stress and plays an important role in heat resistance [14–16].

Previous reports showed that Hsf family genes contained several conserved domains [17]. At the N-terminus, there was a DNA-binding domain (DBD), which could recognize the elements of promoters in heat-response genes. An adjacent oligomerization domain (OD or HR-A/B) was found in all Hsf family genes, and this domain mainly comprised hydrophobic heptad repeats [18, 19].

An Hsf family gene was first cloned and characterized in Saccharomyces cerevisiae (yeast) [20]. Until now, the Hsf gene family has been detected in the whole genome in most species, such as in Arabidopsis thaliana (21 genes) [9], Oryza sativa (25) [9], Solanum lycopersicum (26) [21], Brassica oleracea (35) [22], Brassica rapa (36) [17, 23], Brassica juncea...
Currently, with the release of more and more species genomes, it is possible to identify a large number of Hsf family genes at the whole-genome level in different species. Several databases have been constructed to collect the transcription factors, such as the plant transcription factor database (PlantTFDB, http://planttfdb.gao-lab.org/) [37]. However, this database was not specifically made for the Hsf gene family, and much detailed information and comparative analysis is absent. In addition, a heat shock protein information resource (HSPIR, http://pdslab.biochem.iisc.ernet.in/hspir/) was constructed to collect information on heat shock proteins [38]. This database contains six major heat shock proteins, but does not contain the Hsf gene family. Most existing databases have not been updated with the newest information on genome sequences. Compared with existing databases, all the selected species used in our database have complete genome sequencing.

More importantly, we not only provide more comprehensive Hsf gene family information through constructing a database but also have carried out large-scale systematic and comprehensive comparative analyses of the Hsf gene family in 111 plants to explore their expression pattern and evolutionary mechanism.

Results

Identification of the Hsf gene family in 111 plants

Here, we identified a total of 2950 Hsf family genes from 3911383 gene sequences of 111 species (Fig. 1; Table S1). All of these species have complete whole-genome sequencing, and could well represent the main taxa of plants. The examined species contained 8 lower plants (Fig. S1b). The length of Hsf family genes was 26.58, and most species (97, 87.39%) had >10 Hsf genes (Fig. S1a). We further compared the average length of Hsf family genes and all genes of the whole genome in each species (Fig. S1b). The length of Hsf family genes was 1.2 times greater than the length of all genes in only 7.21% of plants.

Comparative analysis of the Hsf gene family in plants

More Hsf family genes were detected in higher plants than in lower plants (Figs 1 and 2a). Among the top 10 species with a higher percentage of Hsf family genes, all species belonged to the higher plants, including 9 eudicots and 1 monocot (Fig. 2b; Table S1). The only monocot species was M. acuminata (banana), which is a tropical fruit and might need more heat-resistant genes to adapt to a high-temperature environment. Interestingly, all three species with the highest percentage of Hsf family genes belonged to the Brassicaceae, including B. juncea, C. grandiflora, and B. rapa (Fig. 2b). This phenomenon suggested that the Brassicaceae family might contain a higher proportion of heat-resistant genes than other families.

Among the top 10 species with a lower percentage of Hsf family genes, most species (6) belonged to the lower plants (Fig. 2b; Table S1). No Hsf family genes were detected in two species, Coccomyxa subellipsoida (lower plant) and Oropetium thomaeum (monocot). Besides, the three species with the lowest percentage of Hsf family genes were all lower plants: Volvox carteri, Chlorella variabilis, and Chlamydomonas reinhardtii. In V. carteri, only 1 Hsf family gene was detected from the 15285 genes of the whole genome, only accounting for 0.0065% of all genes. This phenomenon indicated that these species might be using ways to regulate heat stress other than using Hsf family genes.

To explore the evolution of the Hsf gene family in plants, we constructed a phylogenetic tree using all 2950 protein sequences of the Hsf gene family from 111 plants (Fig. 2c). The results showed that most branches contained Hsf genes from various plant taxa, and Hsf genes of the lower-level species were located at the base of the branches in the phylogenetic tree (Fig. 2c). This phenomenon indicated that the Hsf genes of each branch evolved independently after species differentiation. However, there were also some branches that only contained the Hsf family genes of monocots and dicots, and they could be clearly separated (Fig. 2c). These results
indicated that the Hsf genes might have expanded or produced sequence variation in angiosperms.

**Gene duplication and loss inference in the Hsf gene family in plants**

Gene family duplication or loss has often occurred in plants, led by whole-genome duplication (WGD) or whole-genome triplication (WGT) events [39–41]. To clarify the evolution of the Hsf gene family, we performed gene duplication and loss analysis in 16 representative plants, including three Chlorophyta, one Marchantiophyta, two Bryophyta, one Lycopodiophyta, one Gymnospermae, one basal Angiospermae, one eudicot, and six monocots (Fig. 3a). The duplication and loss analysis was conducted using Notung software through reconciliation between gene and species phylogenetic trees. Here, we obtained the number of variations of Hsf family genes at different stages of evolution according to the reconstructed phylogenies.

In the lineage leading to the common ancestor of all 16 plants, 16 ancestral genes were duplicated, while no gene was lost (Fig. 3a). However, there were 15 gene losses, while no gene was duplicated in the lineage of the common ancestor of three Chlorophyta. Similarly, there were 23 gene losses, while no gene was duplicated in the lineage of the common ancestor of three Bryophyta. In the lineage of the common ancestor of bryophytes and other higher plants, there were 11 gene duplications but only one gene loss. Although most WGD and WGT events occurred in most higher plants, losses of Hsf family genes were more than duplications in all the representative Lycopodiophyta, Gymnospermae, and Angiospermae species (Fig. 3a and c). This phenomenon indicated that the losses of Hsf family genes were accompanied by WGD or WGT events during plant evolution.

Furthermore, we also conducted Hsf family gene duplication and loss analysis in 20 representative eudicot species (Fig. 3b). In the lineage leading to the common ancestor of all these 20 plants, 94 genes were duplicated, while 23 gene losses, while no gene was duplicated in the lineage of the common ancestor of three Chlorophyta. Similarly, there were 23 gene losses, while no gene was duplicated in the lineage of the common ancestor of three Bryophyta. In the lineage of the common ancestor of bryophytes and other higher plants, there were 11 gene duplications but only one gene loss. Although most WGD and WGT events occurred in most higher plants, losses of Hsf family genes were more than duplications in all the representative Lycopodiophyta, Gymnospermae, and Angiospermae species (Fig. 3a and c). This phenomenon indicated that the losses of Hsf family genes were accompanied by WGD or WGT events during plant evolution.
Figure 2. Comparative and phylogenetic analysis of Hsf family genes in plants. a Boxplot of Hsf gene number for different categories of plants. b The −log10 number ratio of Hsf family genes compared with whole-genome genes in representative plants. Orange color indicates the 10 plants with the highest number ratio of Hsf family genes, and green indicates the 10 plants with the lowest number ratio of Hsf family genes. c Phylogenetic tree using the protein sequences of all the Hsf family genes (2950) from 111 plants.

representative eudicots species except *Populus trichocarpa*. This phenomenon further indicated that Hsf family genes were lost after WGD or WGT events in most eudicot plants.

**Phylogenetic, conserved motif, and evolutionary trajectory analyses of Hsf family genes in seven representative species**

Conserved motif analysis can uncover the conservative patterns of a gene family. Here, we explore the conserved motif of Hsf family genes in seven representative plants, ranging from a lower plant (*C. reinhardtii*) to higher plants, including the eudicot model species *A. thaliana*, the monocot model species *O. sativa*, the basal angiosperm species *Amborella trichopoda*, the Gymnospermae species *Picea abies*, the Lycoperidiophyta species *Selaginella moellendorffii*, and the Bryophyta species *Physcomitrella patens* (Fig. 4).

In total, 94 Hsf family genes were identified from the whole genomes of seven species. The largest number of genes was detected in *O. sativa* (25), followed by *A. thaliana* (24) and *P. abies* (16) (Fig. 4a; Fig. S2a). However, there were only two Hsf family genes in the lower plant *C. reinhardtii*. To explore the phylogenetic relationship and classification of Hsf family genes, we constructed a tree using amino acid sequences. Our analysis revealed that all the Hsf family genes could be divided into seven groups, and we defined them as groups A–F (Fig. 4a). There were 36 and 39 Hsf family genes in groups B and C, respectively, i.e. more than in the other four groups (Fig. S2b).

Ten motifs [motifs (M) 1–10] were detected in the Hsf family genes using the MEME program (Fig. 4a; Table S2). M6 was the longest, followed by M1 and M4 (Fig. S2c, Fig. S3). Four motifs, M3, M5, M2, and M1, were present in almost all Hsf family genes. However, some of these motifs were absent in genes *Ath-Hsf008*, *Ath-Hsf012*, and *Ath-Hsf018*. Furthermore, the Pfam analysis showed that the domain of these three genes was also partial in Arabidopsis (Fig. 4). Similar, another four genes also had the partial domain, including *Atr-Hsf012*, *Smo-Hsf002*, *Smo-Hsf003*, and *Osa-Hsf009* (Fig. S2d). In addition, we also found some motifs specific to the related groups. For example, 7M8 was detected in almost all Hsf family genes of group B, while it was nearly absent in genes of other groups. Similarly, M7 and M10 were only detected in the Hsf genes of groups A and C. These motifs might be associated with the functional specificity of different groups.
of Hsf family genes. In conclusion, these results indicate that the motifs in the same group were highly similar, which is consistent with the phylogenetic relationship of these genes.

Furthermore, we tried to explore the evolutionary trajectories of the Hsf family gene domains (Fig. 4b). In the lower plant C. reinhardtii, only six motifs (M1–M6) were detected, while four motifs (M7–M10) were completely lost. In the bryophyte P. patens, all 10 motifs were present, but M6–M10 were partially lost in some Hsf genes. Then, we found that M6, M7, and M10 were completely lost in the Lycopodidiophyta plant S. moellendorffii. In Gymnospermae and angiosperms, all the motifs were present, while some motifs were partially lost in some genes. In A. thaliana, only M3 was completely retained in all Hsf genes. Similarly, only M1 and M2 were completely retained in all Hsf genes in O. sativa. This phenomenon indicated that Hsf family genes generated sequence divergence during the evolution of the species.

**Duplication type of Hsf gene family in seven representative species**

We explored the gene family expansion mechanism through gene duplication type analysis. Five types of gene duplication were identified for whole-genome genes and Hsf family genes, including singleton, dispersed, proximal, tandem, and WGD/segmental (Figs 4 and 5; Table S3). No Hsf family gene belonged to the singleton, proximal, and tandem duplication type in most species except C. reinhardtii. The two Hsf genes of C. reinhardtii were of the singleton type, which was different from the other six higher plants. However, all the Hsf genes from A. trichopoda, P. abies, and P. patens were of the dispersed type.

Several Hsf genes of A. thaliana, O. sativa, and S. moellendorffii belonged to the WGD/segmental duplication type (Figs 4 and 5; Table S3). The percentage of Hsf genes of the WGD/segmental type was 56.00 in O. sativa (Table S4). This ratio was significantly higher than the average for whole-genome genes (10.27%) belonging to the WGD/segmental type ($p < .01$). Therefore, the WGD/segmental type played critical roles in the Hsf gene family expansion of O. sativa. However, we did not find other duplication types for Hsf gene family expansion in other species.

**Interaction network construction using target genes of the Hsf gene family**

Here, we constructed an interaction network for Hsf family genes and their target genes in Arabidopsis (Fig. 6a; Table S5). A total of 4788 gene pairs made up the regulatory network, which contained 2421 downstream genes (regulated by Hsf family genes) and 222 upstream genes (regulated Hsf family genes). We found that the number of target genes regulated by different Hsf genes varied greatly. Gene AT4G36900 had the...
largest number of downstream genes (845), followed by AT4G18880 (593) and AT1G46264 (379) (Fig. 6b; Tables S5–S7). However, there was no downstream gene for three Hsf family genes, including AT1G77570, AT4G18870, and AT4G19630. Similarly, gene AT4G11660 had the largest number of upstream genes (71), while no upstream gene was found for AT4G13980 and AT5G54070.

Interestingly, we found that 60 genes belonged to the downstream and upstream genes, indicating that they were likely to be feedback-regulated genes (Fig. 6c; Table S6). Furthermore, we explored the gene expression patterns of these target genes under heat treatment for different lengths of time (Table S8). The cluster analysis results showed that these genes had obvious tissue-specific expression patterns after heat treatment, i.e. they were clearly divided into two groups according to root and shoot (Fig. 6d). For example, in each heat treatment, the expression level of the AT1G13300 gene in root was higher than that in shoot. The expression pattern of the AT2G18300 gene was exactly the opposite of that of AT1G13300. The expression level of most genes changed obviously after heat treatment. For example, the expression level of the AT5G05410 gene increased significantly in root and shoot after treatment for 1 and 3 hours, and then

Figure 4. Phylogenetic, conserved motif, duplication type, and evolutionary trajectory analyses of Hsf family genes from seven representative species. 

a Maximum-likelihood trees were generated based on the amino acid sequences of the Hsf gene family. The tree was constructed using FastTree software. The motifs were identified using the MEME program. The duplication type was detected by the MCScanX program. b Evolutionary trajectories of the Hsf gene family. The white X indicates that the motif was lost in some genes. The dashed box indicates that the motif was completely lost or did not exist.
decreased with the extension of the treatment time. This expression map will provide a good reference for further research on the functions of these target genes in heat resistance.

**Functional enrichment analysis of the target genes in the network**

To explore the function of the target genes involved in the network in *Arabidopsis* constructed above, we conducted enrichment analysis of all the target genes of the *Hsf* gene family. We identified 16 significantly enriched terms (q-value < 0.05 and fold-change > 2) for upstream genes (Table S9). The fold change indicated that the percentage of terms enriched for annotated target genes was comparable to that for all annotated genes. The most significantly enriched term was *Apetala 2* (AP2) (q-value = 2.30 × 10^{-41} and fold change = 27.29), followed by *Myb_DNA-binding* and the *TCP* gene family (Table S9). However, only three terms were significantly enriched for the downstream genes. As expected, we found that HSP20 was the most significantly enriched term (q-value = 9.82 × 10^{-4} and fold change = 5.22). This phenomenon indicated that the *Hsf* gene family might play more important roles in regulating HSP20 than other heat-shock proteins in *Arabidopsis*. Furthermore, we analyzed specific and shared enriched functional terms among 16 upstream, 3 downstream, and 11 common gene terms. As shown by the Venn diagram, three, and one enriched functional terms were specific to upstream, downstream, and common genes, respectively (Fig. 6e). By contrast, no significantly enriched terms were detected among these three gene datasets. The results also showed that 10 terms were significantly enriched in upstream and common genes, accounting for 50.0% of all significantly enriched functional terms. Most of these enriched functional terms were related to various stresses, such as AP2, WRKY, zf_C2H2, bZIP, and Myb_DNA-binding. This phenomenon indicated that *Hsf* gene family and other transcription factors interact with each other in response to heat resistance and other stresses in plants.

**Exploring the expression pattern of the *Hsf* gene family under various conditions**

Here, we performed expression pattern analysis of the *Hsf* gene family using large-scale expression datasets under various stresses and developmental stages. We collected 154 samples from 18 groups under various abiotic stresses, including cold, osmotic, salt, drought, genotoxic, oxidative, UV-B, wounding, and heat stress treatment after 0, 0.25, 0.5, 1.0, 3.0, 4.0, 6.0, 12.0, and 24.0 hours in shoot and root of *Arabidopsis* (Fig. 7a; Table S10). The cluster analysis showed that all the genes could be divided into two groups. On the whole, the genes in cluster I had higher expression values than those in cluster II under most stress treatments. However, we found that gene AT2G26150 from cluster II had a
Figure 6. The interaction network among Hsf family genes, and their upstream and downstream-regulated genes in Arabidopsis. a Construction of the network using Gephi software. b Number of upstream and downstream genes for each Hsf family gene in the network. c Specific and shared genes between downstream and upstream genes in the network. d Absolute expression values of common target genes up- and downstream from Hsf under heat treatment for different lengths of time. e Specific and shared terms among upstream, downstream, and common gene enriched terms.

relatively high expression level under heat treatment after 0.5, 1.0, and 3.0 hours in shoot and root. Similarly, genes AT2G26150 and AT3G22830 had a relatively high expression level under salt treatment after 3.0, 6.0, 12.0, and 24.0 hours in the root. In addition, we found that gene AT4G36990 had higher expression values in root than shoot in most stress treatments.

For biotic stresses, we collected 70 samples from 27 groups under various biotic stresses (Fig. 7b; Table S11). Three groups were obtained by cluster analysis according to gene expression in different biotic stresses. All four genes (AT4G18880, AT1G67970, AT5G62020, and AT4G36990) in cluster II had the highest gene expression among the three clusters, followed by cluster III and cluster I.

In addition, we examined the gene expression of 47 samples from different developmental stages of several tissues, including seed, leaf, root, flower, and siliques (Fig. 7c; Table S12). Similarly, two groups were obtained by the cluster analysis. Most genes in cluster I had a higher expression level than those in cluster II. However, we found that gene AT2G26150 from cluster II had a relatively high expression level in the carpels, petals, and sepals of flower stages 12 and 15 in Arabidopsis. We also found that gene AT5G54070 from cluster II had a relatively high expression level in the dry seed and siliques of seed stages 8–10. All of these gene expression patterns in the model species Arabidopsis will provide a guide for studying the Hsf gene family in other species in the future.
Pan-genome analysis of Hsf gene family in B. rapa

To study Hsf gene family variation among different varieties of the same species, we performed Hsf gene family analysis in the pan-genome of 18 B. rapa accessions (Fig. 8a). A total of 952 Hsf family genes were identified in 18 accessions, the number ranging from 47 (CXA accession) to 56 (OIA, TCA, and TUE accessions) in each B. rapa accession (Table S13). The syntenic analysis showed that >57.14% of Hsf genes were located in the syntenic regions of the genome. The number of Hsf family genes in the collinear region varied little among accessions, ranging from 32 to 35, suggesting that the genes in the collinear region were relatively stable (Fig. 8b; Table S13).

Based on gene loss and retention, the B. rapa genome was further divided into three subgenomes, the less fractioned subgenome (LF) and the more fractioned subgenomes (MF1 and MF2). The Hsf gene number was from 12 to 14 in the LF subgenome, from 9 to 12 in the MF1 subgenome, and from 8 to 9 in the MF2 subgenome (Fig. 8a and c; Tables S13 and S14). Cluster analysis was conducted according to the number of Hsf family genes in each subgenome, and the results could be divided into three groups, corresponding to the subgenomes of each genome of 18 accessions (Fig. 8a).

The B. rapa genome has experienced an additional WGT event compared with Arabidopsis. Among all the Hsf family genes, we found that genes AT3G22830 and AT4G18870 were well preserved with three copies after duplication in most accessions. However, most Arabidopsis Hsf family genes only retained one or two copies in B. rapa, which means that some gene loss occurred after the WGT event. Interestingly, we found that the syntenic gene of AT5G45710 was completely lost in CXA and CXB accessions (Fig. 8d; Tables S14 and S15). The syntenic genes of four Arabidopsis genes (AT1G77570, AT3G02990, AT4G18880, and AT5G54070) were also completely lost in all 18 B. rapa accessions.

Furthermore, we constructed a phylogenetic tree using the protein sequences of these genes. The tree (Fig. 8e) shows that most branches contained Hsf family genes from the B. rapa species and Arabidopsis. Therefore, we could further explore the evolution and function of homologous genes located on the same branch in B. rapa based on Arabidopsis genes.

Phylogenetic and expression analysis of Hsf and Hsp gene family in B. rapa

In addition to identifying 54 Hsf family genes in B. rapa, we also detected 393 Hsp family genes, which were generally regulated by the Hsf family genes (Table S16). All of these Hsp family genes were further divided into Hsp20 (38), Hsp40 (155), Hsp60 (53), Hsp70 (34), Hsp90 (16), and Hsp100 (97) gene families.

To further explore their relationship, we constructed a phylogenetic tree using the protein sequences of these
Figure 8. Syntenic and phylogenetic analysis of the Hsf gene family in A. thaliana and genomes of 18 B. rapa accessions. a Syntenic genes between Arabidopsis and each subgenome (LF, MF1, MF2) of 18 B. rapa genomes. b Numbers of all Hsf family genes and syntenic Hsf genes and their ratio in 18 B. rapa genomes. c Syntenic gene number in each subgenome (LF, MF1, MF2) of 18 B. rapa genomes. d Heat map of syntenic gene number for each Arabidopsis Hsf gene in each of 18 B. rapa genomes. e Phylogenetic tree using the Hsf family genes of Arabidopsis and 18 B. rapa accessions.

gene families (Fig. 9). The results showed that the Hsf gene family had a closer relationship with the Hsp20 gene family than other families. Hsp90 had a close relationship with the Hsp40 gene family, and Hsp60 had a closer relationship with the Hsp70 gene family. The Hsp100 gene family was divided into two branches in the phylogenetic tree, and it had a close relationship with Hsp60 and Hsp70. Therefore, our analysis first clearly revealed their relationship in B. rapa, and could be further used to explore their evolution and comparative analysis in B. rapa and other plants.

Furthermore, we conducted expression pattern analysis of the Hsf and Hsp gene families using expression datasets under heat stress in B. rapa (Fig. 9; Table S16). For the heat treatment, the samples were treated at 38°C for 1, 4, 8, and 12 hours. The expression heat map shows that several genes from Hsp20, Hsp60, Hsp70, Hsp90, and Hsp40 had highest expression at 1 hour of treatment, and expression then gradually decreased, such as for Hsp20 family genes BraA01g018560, BraA02g010780, BraA03g011180, and BraA10g017550 (Fig. 9). We also found that some genes had higher expression under all heat treatments than control, such as BraA03g027160 (Hsp20) and BraA06g020450 (Hsp20). The genes in other families showed trends similar to those of the Hsp20 gene family.

We also performed expression pattern analysis of the Hsf and Hsp gene families using the expression datasets from different tissues, including root, stem, leaf, flower, silique, and callus of B. rapa (Fig. 9; Table S17). By integrating the expression pattern heat map of heat treatment and various tissues, we found that several genes showed high expression in both expression maps, such as genes in the Hsp60 family (BraA07g018460 and BraA08g000690), Hsp70 (BraA03g016490 and BraA03g019440), Hsp90 (BraA01g014950, BraA02g013110, and BraA06g038950) and Hsp100 (BraA03g015750 and BraA07g032970) (Fig. 9).

Overall, this expression pattern map will provide a guide for the study of their function and regulatory network in B. rapa.

**Interactive network between Hsf and Hsp gene families in B. rapa**

Based on the expression of the Hsf and Hsp family genes mentioned above, we calculated the Pearson correlation coefficients (PCCs) between any two genes of these gene families, and then constructed regulatory networks. After filtering, a total of 1275 connections...
Figure 9. Phylogenetic and expression analyses of the Hsf and various Hsp gene families in B. rapa. (a) Each gene family is marked with a different color in the branches of the phylogenetic tree. The bootstrap was set to 1000 replicates, and values >40% are indicated with a circle. The heat map represents gene expression under different heat treatments (b, control; c, T1; d, T4; e, T8; f, T12) and in different tissues (g, root; h, stem; i, leaf; j, flower; k, silique; l, callus). Expression values are normalized by FPKM, and are log2-transformed in the heat map.

(edges) between any two genes in the network with PCC > .95 (Fig. 10a; Table S18). Among these connections, only 65 (5.10%) were negative regulatory with PCC < −.95, while all of the other 1210 connections had a positive relationship (Table S18). This network included 22 Hsf and 216 Hsp family genes, accounting for 40.74 and 54.96% of all Hsf and Hsp gene family genes, respectively (Fig. 10a; Table S19). Furthermore, we found that Hsf gene BraA02g044030 had the most negative connections (22) with other Hsp family genes, accounting for 33.85% of all 65 negative edges (Fig. 10b; Table S18).

Among Hsf family genes, BraA01g010910 had the most connections (30) with other genes, followed by BraA02g044030 (28) and BraA01g022930 (16) (Fig. 10c; Table S19). Among Hsp family genes, BraA01g027780 (Hsp60) had the most connections (49), followed by BraA07g018250 (Hsp100) and BraA03g051700 (Hsp90). Interestingly, most genes (19, 95%) belonged to Hsp40, Hsp60, and Hsp100 among the top 20 connections of Hsp family genes (Fig. 10c; Table S19). These results indicate that genes with more connections might play a core role in the regulatory network of heat resistance in B. rapa.

Furthermore, we predicted the cis-acting elements from the promoter sequences of Hsp family genes contained in the network. The results showed that the identified cis elements were involved in light response, abiotic stress, circadian control, and hormone signaling (Fig. S4, Table S20). This indicated that Hsp family genes also participated in a large number of other physiological functions in addition to participating in the regulation of plant heat resistance. This phenomenon is also consistent with previous reports [42–45].

Moreover, we identified the heat shock element (HSE) from the promoter sequences of Hsp family genes in the network. Finally, the HSE was detected in 28 Hsp family genes, which might play a direct and important role in the regulation of heat resistance. Then, using these genes and the interacting Hsf and Hsp genes, the heat-resistant regulatory network was reconstructed (Fig. 10d; Table S21). In the network, BraA07g025530 (Hsp40) had the most connections (28) with other genes, followed by BraA05g039740 (Hsp100, 27) and BraA03g026430 (Hsp40, 17) (Fig. 10e; Table S22).
In conclusion, these networks well reveal the regulatory relationship between Hsf and various Hsp family genes in *B. rapa*, and also lay the foundation for future experimental studies on the functional interaction of these genes.

**Database construction for the Hsf family genes**

Using these available datasets, we constructed an Hsf database, which should help users to easily query, compare, and download Hsf family genes of all species simultaneously. All species were taxonomically classified to facilitate checking of the Hsf family genes according to their evolutionary relationship. All Hsf family gene information was stored in backend tables using MySQL, which could be accessed using the front-end web application of the Hsf database. Here, we provide a detailed description of the interfaces of this database, including the home, browse, download, resource, help, and contact interfaces (Fig. 11).
Figure 11. Architecture of the Hsf Database, including the home, browse, source, download, and help modules. a Home page. b Hsf gene and sequence information of each species in the browse module. c Hsf gene number of each species in the chart function of the browse module. d Download function of Hsf family genes for each species.

**Browse**

To make it easier for users to check, we further sorted the examined species according to the first letter of their Latin name. The multi-select dropdown allows users to select each species for browsing. We provide detailed information for each species, including the total number, Hsf name, gene ID, and protein sequences. Furthermore, we also integrated the search function in the browser interface, which makes it easier for users to find the assigned Hsf family gene information according to gene ID.

Besides, the browse home page provides interactive plots and line and bar graphs to view the number of species. Then, bar plots, and line charts are also used to show the Hsf family gene number of each species, which makes it easier and faster for users to compare these genes in different species. Finally, all the related information can be downloaded in Excel format. These documents will allow researchers to conduct a local batch comparative analysis of the Hsf gene family.

**Download, resource, and help**

The information (Excel format) and sequences (Fasta format) of Hsf family genes for each species can be obtained from the download interface. In the resource interface we have collected most Hsf research-related databases and provided relevant links for users to easily query and compare studies. In the help interface, we provide users with a detailed Hsf database manual. We also provide the e-mail, mobile number, and address to allow users to contact us conveniently.

With the novel genome sequences released in the future, we will continuously identify Hsf family genes from these datasets and add them to our Hsf database. We also encourage users to submit new Hsf family genes to us to further enrich the database. Finally, we welcome feedback from all users for further improvement of our database. We believe that this database will be useful and friendly for all researchers.

**Discussion**

In this study, we comprehensively identified 2950 Hsf family genes from the whole genome of 111 representative plants, most of which are horticultural plants. Compared with the PlantTFDB, we have also performed Hsf gene family analysis on some species recently sequenced. Most importantly, we have conducted systematic comparative analysis to reveal their basic characteristics, conserved motif, duplications and losses, evolutionary, expression patterns, and phylogenetic relationships.

Our analysis showed that more Hsf family genes were detected in most higher plants than in lower plants. This might be due to the multiple WGD or WGT events in plants, especially higher plants, leading to the production
of more heat-resistant genes to adapt to high temperatures [46–48]. Interestingly, the three species with the highest percentage of Hsf family genes were all Brassicaceae species. This phenomenon might also relate to the WGD and WGT events of the Brassicaceae species [49–51]. The number of Hsf gene losses was more than gene duplication in most Lycoperidophyta, Gymnospermae, and Angiospermae species, which indicated that the Hsf family genes were lost after the WGD or WGT events in higher plants.

A. thaliana is the typical model plant that has provided the reference for studying gene function and evolution in other plants [52]. In this study, we constructed the interaction network for Hsf family genes and their target genes in Arabidopsis. Then, enrichment analysis of target genes indicated that the Hsf gene family interacted with other transcription factors, such as AP2, WRKY, and Myb_DNA-binding, in response to heat stress. Furthermore, the global expression pattern of the Hsf gene family in Arabidopsis was explored. All of these results in the model species Arabidopsis will provide a guide for studying the Hsf gene family in other species.

B. rapa is one of the most economically important Brassica species and is the main vegetable crop worldwide [53, 54]. B. rapa (Chinese cabbage), is a typical horticultural plant. It experienced a WGT event after it diverged from Arabidopsis [50, 55]. Here, we conducted Hsf gene family analysis using the pan-genome of 18 B. rapa accessions reported recently [53]. The syntenic analysis showed that 0–3 copies of Arabidopsis Hsf family genes were detected in the three subgenomes of 18 B. rapa accessions. The results lay the foundation for better research on the function of duplicated genes.

Furthermore, we performed expression pattern analysis of Hsf family genes and several Hsp gene families using expression datasets from different tissues and under different heat treatments in B. rapa. Then, the interaction network between Hsf gene family and Hsp gene families was constructed in B. rapa, and several core genes were detected in the network. This network well reveals the regulatory relationship of Hsf and various Hsp gene families, and also provides a guide for experimental studies on the functional interaction of these family genes in future.

Finally, we constructed a database (Hsfdb) for all identified Hsf family genes in 111 representative plants. Compared with the PlantTFDB, we have also performed Hsf gene family analysis on some species recently sequenced.

Second/next-generation sequencing (NGS) has tremendously improved sequencing output, and has made genome sequencing much faster and cheaper than Sanger sequencing. However, NGS technology has several drawbacks, most obviously the short reads [56]. Third-generation long-read sequencing overcomes the limitation of short-read sequencing, and can produce high-quality genome assemblies. Therefore, third-generation sequencing has the ability to resolve repeat sequences and large chromosomal rearrangements [56, 57]. Second-generation sequencing leads to incomplete genomes of species, and may lead to the loss of some genes. However, the most influential region should be the repetitive sequence, and the repetitive region contains a relatively small number of genes. Moreover, we use the latest version or the version with higher assembly quality when the species has multiple versions of the genome. Of course, with the development of three-generation sequencing and the reduction of costs, we believe that nearly complete genomes will be obtained for more and more species, just like the human genome and the Arabidopsis genome recently reported [58, 59]. Our database will be continuously updated with updates of the species genome.

In conclusion, we performed comprehensive analyses of the Hsf gene family in 111 horticultural and other representative plants, especially for the study of the horticultural plant B. rapa. We also constructed a database for all identified Hsf family genes. This study will serve as a useful resource for future studies on the biological function and evolutionary history of the Hsf gene family.

Materials and methods

Sequence collection

The protein sequences of examined plants in Fasta format were downloaded from the Ensembl database (http://useast.ensembl.org/index.html). The 18 B. rapa pan-genome sequences were downloaded from the BRAD database (http://brassicadb.cn) [60]. Alternative splice sequences were deleted using a custom Perl script to ensure the non-redundancy of the sequences that were used. The phylogenetic trees of species were made using the iTOL website according to the relationship of species in NCBI taxonomy [61].

Hsf gene family identification and statistics

The Hsf family genes were identified using the Pfam database (PF00447) with an e-value <1e-4 [62]. Furthermore, the SMART and CDD databases were used to conduct domain validation to ensure the results’ accuracy [63, 64]. The SMART and MARCOIL databases were also used for DBD and HR-A/B domain detection, respectively [63, 65]. Violin plots with a boxplot of the Hsf family gene number of each kingdom were drawn using the ggplot function in the ggpubr package of the R program (https://cran.r-project.org/web/packages/ggpubr/index.html).

Phylogenetic relationship construction

Firstly, the protein sequences of the Hsf gene family were aligned by Mafft v7.471 software with maxiterate at 1000 [66]. The maximum likelihood tree was constructed using FastTree (v2.1.11) software [67]. The JTT (Jones–Taylor–Thornton) model was used, and bootstrap replications were set at 1000.
Conserved motif identification and gene duplication and loss inference

The amino acid sequences of the Hsf family genes from seven representative plants were used for motif analysis using Multiple Expectation Maximization for Motif Elicitation (MEME) with the default parameters [68]. Hsf family gene duplication and loss were identified by Notung2.9 software [69]. All of this information on the phylogenetic trees was illustrated by the iTOL program [61].

Duplication type detection for Hsf family genes

The collinearity of Hsf family genes among seven representative plants was detected using MCScanX software [70]. Firstly, the protein sequences of these species were aligned using the Blastp program with an e-value of $1 \times 10^{-5}$. Then, the collinear blocks were detected using MCScanX with the default parameters. Finally, the gene duplication types were identified using a duplicate_gene_classifier program from MCScanX software. Significance analysis of the duplication type for Hsf family genes compared with whole-genome genes was conducted using the $\chi^2$ test ($P < 0.01$).

Target gene identification and interaction network construction

The target genes of the Hsf gene family in Arabidopsis were identified using the integrated gene regulatory network (iGRN) database (http://bioinformatics.psb.ugent.be/webtools/iGRN/) with score $\geq 0.60$ [71]. In this study, we divided the target genes into downstream and upstream genes. The downstream genes were regulated by Hsf family genes, while the upstream genes regulated the Hsf family genes. The interaction network between Hsf family genes and target genes was constructed using Gephi software (v0.9.2) with a continuous graph layout algorithm ForceAtlas2 (https://gephi.org) [72].

Functional annotation and enrichment analysis of target genes

Functional annotation of the above-identified target genes and all genes of Arabidopsis was performed using the Pfam database (http://xfam.org) [73]. Then, enrichment analysis was conducted by comparing four groups: target genes with a related functional term; target genes with annotation; all genes with a related functional term; and all genes with annotation in Arabidopsis. Finally, the scipy package of Python was used to perform enrichment analysis [74]. The P-values obtained by significance analysis were further corrected using the Bonferroni method in the R program. The corrected P-value ($q$-value) $< 0.05$ and fold change $> 2$ were used to define significant enrichment terms. The TBtools program was used to generate the Venn diagram, which indicated the specific or shared enriched terms for the downstream and upstream target genes [75].

Gene expression data retrieval and analysis

The large-scale expression datasets under various stresses and developmental stages were collected from the website of the Arabidopsis eFP browser (http://www.bar.utoronto.ca) [76]. We collected 154 samples from 18 groups under various abiotic stresses, 70 samples from 27 groups under various biotic stresses, and 47 samples from different developmental stages of Arabidopsis. Then, we explored the expression of the Hsf gene family using these large-scale biological datasets. The TBtools program was used to generate a heat map according to cluster analysis of the expression values [75].

For B. rapa, the expression dataset under heat stress was obtained from the Genome Sequence Archive (GSA) in the Big Data Center (accession number CRA002707) according to our previous report [54]. Heat treatment was conducted at $38^\circ$C for 1 hour (T1), 4 hours (T4), 8 hours (T8), and 12 hours (T12); the control was without heat treatment. The expression dataset in different tissues was obtained from Gene Expression Omnibus (GEO) in NCBI (accession number GSE43245) according to a previous report [77]. The expression values were normalized as fragments per kilobase of transcript per million mapped reads (FPKM) [78]. Integration of the phylogenetic tree and creation of the expression heat map were performed using the iTOL program [61].

Hsp gene family identification and interaction network construction

The six main Hsp family genes were identified in this study. Among these, Hsp20, Hsp70, and Hsp90 family genes were directly extracted from the Pfam database using the identifier numbers PF00011, PF00012, and PF00183, respectively (e-value $< 1e^{-4}$) [62]. The Hsp40, Hsp60, and Hsp100 family genes were identified using the hmmsearch program because there was no Pfam identifier number [79]. PCCs between Hsf and Hsp were calculated using in-house Perl scripts according to the gene expression value under heat treatment. The positive and negative regulatory relationships were defined as PCC $> 0.95$ and PCC $< -0.95$, respectively [80, 81]. The interaction network between Hsf and Hsp was constructed using Gephi software [72].

Identification and visualization of cis-acting elements

The promoter sequences were extracted from the 2 kb upstream of the translation initiation site of each Hsp family gene. Then, the cis-acting elements were estimated in these promoter sequences using PlantCARE [82]. The visualization of cis-acting elements in each Hsp promoter region was realized by TBtools software [75]. HSEs were detected from the promoter sequence of each Hsp family gene using the in-house Perl script. The consensus sequence of GAAnTTmnnGAA was used to predict HSEs according to previous reports [83–85].
Pan-genome analysis of Hsf gene family in B. rapa

We performed Hsf gene family identification and analysis using the pan-genome of 18 B. rapa accessions reported recently [53]. The B. rapa genome was further divided into three subgenomes, comprising the less fractioned subgenome (LF) and the more fractioned divided into three subgenomes, comprising the less fractioned subgenome (LF) and the more fractioned subgenome (MF) according to a previous report [86]. Thus, we explored the absence or presence of Hsf family genes in three subgenomes of B. rapa.

Database construction

The Hsf database was built using several software packages, including Microsoft .NET Framework 4.5, MySQL database management, PHP, HTML, CSS, and JavaScript. The collected data were processed using a Perl program for conducting biological data analysis. The interactive web interface was constructed to enable users to conveniently access our database. HTML, PHP, and JavaScript were used to transmit query requirements and extract Hsf gene family-related data from the MySQL database to show in the report pages. The interactive plotting system was made using d3.js and the nvd3 helper library [87].

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Author contributions

X.S. conceived the project and was responsible for project initiation. X.S. and T.Y. supervised and managed the project and research. Data generation and collection were performed by X.S., T.Y., Y.B., S.F., and T.W. Bioinformatics analysis and database construction were led by X.S., T.Y., Z.L., Z.W., Q.Y., S.S., and Y.Z. The manuscript was organized, written, and revised by X.S., T.Y., Y.B., Q.L., and L.G. All authors read and revised the manuscript.

Data availability

All materials and related datasets in this study are available in our Hsf database (http://hsfdb.bio2db.com).

Conflict of interest

The authors declare no competing interests.

Supplementary data

Supplementary data is available at Horticulture Research online.

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