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Genome-wide identification and characterization of WRKY gene family in *Salix suchowensis*

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WRKY proteins are the plant-specific zinc finger transcription factors. They can specifically interact with the W-box ([C/T]TGAC[T/C]), which can be found in the promoter region of a large number of plant target genes, to regulate the expressions of downstream target genes. They also participate in diverse physiological and growing processes in plants. Prior to the present studies, plentiful WRKY genes have been identified and characterized in herbaceous species, but there is no large-scale study of WRKY genes in willow. With the whole genome sequencing in *Salix suchowensis*, we have the opportunity to conduct the genome-wide research for willow WRKY gene family. In this study, we identified 85 WRKY genes in the willow genome and renamed them from SsWRKY1 to SsWRKY85 on the basis of their specific distributions on chromosomes. Due to their diverse structural features, the 85 willow WRKY genes could be further classified into three main groups (group I - III), with five subgroups (IIa - IIe) in group II. With the multiple sequence alignment and the manual search, we found three variations of the WRKYGQK heptapeptide: WRKYGRK, WKKYGQK and WRKYGKK, and four variations of the normal zinc finger motif, which might execute some new biological functions. In addition, the SsWRKY genes from the same subgroup share the similar exon-intron structures and conserved motif domains. Further studies of SsWRKY genes revealed that segmental duplication events played the prominent roles in the expansion of SsWRKY genes. Distinct expression profiles of SsWRKY genes with RNA sequencing data revealed that diverse expression patterns among five tissues, including tender roots, young leaves, vegetative buds, non-lignified stems and barks. With the analyses of WRKY gene family in willow, it is not only beneficial to complete the functional and annotation information of WRKY genes family in woody plants, but also provide important references to investigate the expansion and evolution of this gene family in flowering plants.

1 **Genome-wide identification and characterization of** 2 **WRKY gene family in *Salix suchowensis***

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11 **Abstract**

12 WRKY proteins are the plant-specific zinc finger transcription factors. They can specifically
13 interact with the W-box ([C/T]TGAC[T/C]), which can be found in the promoter region of a
14 large number of plant target genes, to regulate the expressions of downstream target genes.
15 They also participate in diverse physiological and growing processes in plants. Prior to the
16 present studies, plentiful WRKY genes have been identified and characterized in herbaceous
17 species, but there is no large-scale study of WRKY genes in willow. With the whole genome
18 sequencing in *Salix suchowensis*, we have the opportunity to conduct the genome-wide
19 research for willow WRKY gene family. In this study, we identified 85 WRKY genes in the
20 willow genome and renamed them from SsWRKY1 to SsWRKY85 on the basis of their
21 specific distributions on chromosomes. Due to their diverse structural features, the 85 willow
22 WRKY genes could be further classified into three main groups (group I - III), with five
23 subgroups (IIa - IIe) in group II. With the multiple sequence alignment and the manual search,
24 we found three variations of the WRKYGQK heptapeptide: WRKYGRK, WKKYGQK and

1 WRKYGKK, and four variations of the normal zinc finger motif, which might execute some
2 new biological functions. In addition, the SsWRKY genes from the same subgroup share the
3 similar exon–intron structures and conserved motif domains. Further studies of SsWRKY
4 genes revealed that segmental duplication events played the prominent roles in the expansion
5 of SsWRKY genes. Distinct expression profiles of SsWRKY genes with RNA sequencing
6 data revealed that diverse expression patterns among five tissues, including tender roots,
7 young leaves, vegetative buds, non-lignified stems and barks. With the analyses of WRKY
8 gene family in willow, it is not only beneficial to complete the functional and annotation
9 information of WRKY genes family in woody plants, but also provide important references to
10 investigate the expansion and evolution of this gene family in flowering plants.

11 **Keywords:** WRKY, Phylogenetic analysis, Evolution, Duplication, Expression, Willow

12 Introduction

13 Plants form a series of adjustment mechanisms to adapt diverse environment stress in their
14 long evolutionary processes. Among the numerous adjustment mechanisms, transcription
15 factors play important roles [1]. In plants, WRKY proteins constitute a large family of
16 transcription factors, involving in various physiological and developmental processes [2, 3].
17 Since the first WRKY gene was cloned and characterized from sweet potato [4], many
18 corresponding studies have been conducted rapidly, such as *Arabidopsis thaliana*, desert
19 legume (*Retama raetam*), cotton (*Gossypium arboreum*), rice (*Oryza sativa*), *Pinus monticola*,
20 barley (*Hordeum vulgare*), sunflower, cucumber (*Cucumis sativus*), poplar (*Populus*
21 *trichocarpa*), tomato (*Solanum lycopersicum*) and grapevine (*Vitis vinifera*) [2, 5-14].

22 The existence of either one or two highly conserved WRKY domains is the most vital
23 structural characteristic of WRKY gene. WRKY gene consists of about 60 amino acid
24 residues with a conserved WRKYGQK heptapeptide at its N-termini, and a zinc finger motif
25 (C-X₄₋₅-C-X₂₂₋₂₃-H-X₁-H or C-X₇-C-X₂₃-H-X₁-C) at the C-terminal region. Previous
26 functional studies indicated that WRKY genes could specifically interact with the W-box, the
27 promoter region of plant target genes, to adjust the expressions of downstream target genes

1 [15]. What's more, SURE (sugar responsive elements), another prominent cis-element that
2 can promote transcription processes, was also found to bind to the WRKY transcription
3 factors under a convincing research [16]. The proper DNA-binding ability of WRKY genes
4 could be influenced by the variation of the conserved WRKYGQK heptapeptide [17, 18].

5 The WRKY proteins can be classified into three main groups (I, II and III) on the basis of
6 the number of their WRKY domains and the pattern of the zinc finger motif. Proteins from
7 group I contain two WRKY domains followed by a C₂H₂ zinc finger motif, while the other
8 WRKY proteins from group II and III only contain one WRKY domain followed by a C₂H₂ or
9 C₂HC correspondingly [19]. Group II can be further divided into five subgroups from IIa to
10 IIe based on additional amino acid motifs present outside the WRKY domain. Apart from the
11 conserved WRKY domains and the zinc finger motif, there are also some WRKY proteins
12 appearing to have basic nuclear localization signal, LZs (leucine zipper) [20],
13 serine-threonine-rich region, glutamine-rich region and proline-rich region [21]. Throughout
14 the studies of WRKY gene family in many higher plants [3, 10, 13], WRKY genes have been
15 identified to be involved in various regulatory processes mediated by different biotic and
16 abiotic stresses [22]. In plant defense against various biotic stresses, such as bacterial, fungal
17 and viral pathogens, it has been well documented that the WRKY genes play vital roles [14,
18 23, 24]. They are also involved in abiotic stress-induced gene expression. In *Arabidopsis*,
19 with the either heat or salt treatments, the expressions of AtWRKY25 and AtWRKY33 are
20 transformed apparently [25]. Furthermore, the expression of TcWRKY53 that belonged to
21 alpine penny grass (*Thlaspi caerulescens*) is affected by salt, cold, and polyethylene glycol
22 treatments [3]. In rice, a total of 54 OsWRKY genes showed noticeable differences in their
23 transcript abundance under the abiotic stress such as cold, drought, and salinity [22]. There is
24 also accumulating evidence that WRKY genes are involved in regulating developmental
25 processes, such as embryo morphogenesis [26], senescence [27], trichome initiation [28], and
26 some signal transduction processes mediated by plant hormones including gibberellic acid
27 [29], abscisic acid [30], or salicylic acid [31].

1 The number of WRKY genes in different species varies tremendously. For instance, there
2 are 72 members in *Arabidopsis thaliana*, at least 45 in barley, 57 in cucumber, 58 in physic
3 nut (*Jatropha curcas*), 59 in grapevine, 104 in poplar, 105 in foxtail millet (*Setaria italica*),
4 112 in *Gossypium raimondii* and more than 109 in rice [2, 6, 7, 9, 11, 13, 32-34]. Zhang et al.
5 also identified the most basal WRKY genes in the lineage of non-plant eukaryotes and green
6 alga [35]. The study in bryophyte (*Physcomitrella patens*) found at least 12 WRKY genes
7 [21], and the study in gymnosperm (*Cycas revolute*) identified at least 21 WRKY genes [36].
8 Interestingly, the WRKY genes in eukaryotic unicellular chlamydomonas, protoctist (*Giardia*
9 *lamblia*), bryophyte (*Physcomitrella patens*) and fern (*Ceratopteris richardii*) all belonged
10 to group I [2, 37]. The WRKY genes in *Cycas revolute* were divided into two groups, 15
11 WRKY genes therein belonged to group I and the other 6 WRKY genes belonged to group II.
12 Further study suggested that the core WRKY domains of group II and III were similar to the
13 C-terminal domain of group I, and the group II WRKY genes might emerge from the
14 breakage of the C-terminal domain in group I and the group III probably evolve from group
15 III [21]. Above of all indicated that the group I WRKY genes might be the oldest type, which
16 evolved from the origin of eucaryon, and group II and III might generate after the origin of
17 bryophyte [35, 38]. In the evolution of WRKY genes, gene duplication events played
18 prominent roles. As we all know, gene duplication events can lead to the generation of new
19 genes. Take this an example, there are approximately 80% of OsWRKY (rice) genes located
20 in duplicated regions [13], as well as 83% of PtWRKY (poplar) genes [7]. However, no gene
21 duplication events have occurred in cucumber [9].

22 Willow, an important broad-leaf plant, grows quickly and reproduces simply. It can survive
23 under a variety of different ecological environment and grow well. With its broad leaf, willow
24 becomes a prominent part of the protection forest, soil and water conservation forest specie.
25 Therefore, willow has higher ecological and economic value. With these various factors and
26 the draft of the *Salix suchowensis* genome sequence was finished recently [39], we had the
27 opportunity to analyze the willow WRKY gene family. In this study, we identified 85
28 members of the WRKY genes in the willow genome. Subsequently, the distribution of

1 WRKY genes on chromosomes, phylogenetic analysis, classification of WRKY genes,
2 exon-intron organization, conserved motif analysis, and expression analyses were also
3 conducted, which provide a solid foundation for further studies of SsWRKY gene family
4 function and evolution.

5 **Materials and methods**

6 **Datasets and sequence retrieval**

7 The sequence of a shrub willow *Salix suchowensis* (*S. suchowensis*), which flowers within
8 two years, was conducted with a combined approach using Roche/454 and
9 Illumina/HiSeq-2000 sequencing technologies [39]. The latest v5.2 *S. suchowensis* genome
10 annotation information (version5_2.gff3) and protein sequences (Willow.gene.pep) were
11 downloaded from our laboratory website (http://bio.njfu.edu.cn/ss_wrky/). Sequences of 72
12 *Arabidopsis* WRKY proteins were obtained from TAIR (release 10,
13 <http://www.arabidopsis.org/>) [2], and 104 poplar WRKY proteins were obtained from the
14 Supplementary material 3 of poplar [7].

15 **Identification and distribution of WRKY genes in willow**

16 The procedure performed to identify putative WRKY proteins in willow was similar to the
17 method described in other species [6, 7, 13]. The Hidden Markov Model (HMM) profile for
18 the WRKY transcription factor was downloaded from the Pfam database
19 (<http://pfam.sanger.ac.uk/>) with the keyword 'PF03106' [40]. The HMM profile was applied
20 as a query to search against the all willow protein sequences (Willow.gene.pep) using
21 BLASTP program (E-value = $1e^{-3}$) [41]. Another procedure was performed to validate the
22 putative accuracy. An alignment of WRKY seed sequences in Stockholm format from Pfam
23 database was used by HMMER program (hmmbuild) to build a HMM model, and then the
24 model was used to search the willow protein sequences by another HMMER program
25 (hmmsearch) with default parameters [42]. Finally, we employed the SMART program

1 (<http://smart.embl-heidelberg.de/>) to confirm the candidates from the two procedures
2 correlated with the WRKY structure features [43].

3 Additionally, we calculated the length, MW (molecular weight), PI (isoelectric point) of
4 these putative WRKY proteins by ExPasy site (http://au.expasy.org/tools/pi_tool.html). Every
5 WRKY genes were mapped onto chromosomes assembled ourselves
6 (http://bio.njfu.edu.cn/ss_wrky/version5_2.fa) with an in-house Perl script
7 (http://bio.njfu.edu.cn/willow_chromosome/BuildGff3_Chrom.pl), and then rename based on
8 their orderly given chromosomal distribution. The distribution graph of every WRKY gene
9 was drawn by MapInspect software (<http://mapinspect.software.informer.com/>).

10 **Sequence alignments, phylogenetic analysis and classification of** 11 **willow WRKY genes**

12 Using the online tool SMART, we obtained the conserved WRKY core domains of predicted
13 SsWRKY genes, and then multiple sequence alignment based on these domains was
14 performed using ClustalX (version 2.1) [44]. After alignment, we used Boxshade
15 (http://www.ch.embnet.org/software/BOX_form.html) to color the alignment result online. To
16 gain better classification of these SsWRKY genes, a further multiple sequence alignment
17 including 103 SsWRKY domains and 82 WRKY domains from *Arabidopsis* (AtWRKY) was
18 performed using ClustalW [44], and a phylogenetic tree based on this alignment was built by
19 MEGA 6.0 with the Neighbor-joining (NJ) method [45]. Bootstrap values have been
20 calculated from 1000 iterations in the pairwise gap deletion mode, which is conducive to the
21 topology of the NJ tree by divergent sequences. Based on the phylogenetic tree constructed by
22 SsWRKY and AtWRKY domains, these SsWRKY genes were classified into different groups
23 and subgroups. In order to get a better comparison of WRKY family in Salicaceae, a
24 phylogenetic tree including all SsWRKY domains and 126 WRKY domains from poplar
25 (PtWRKY) was constructed with the similar method to *Arabidopsis*. Additionally, a
26 phylogenetic tree based on full-length SsWRKY genes was also constructed to get a better
27 classification. The ortholog of each SsWRKY gene in *Arabidopsis* and poplar was based on

1 the phylogenetic trees of their respective WRKY domains, and the members of group I
2 WRKY genes were considered as orthologs unless the same phylogenetic relationship can be
3 detected between N-termini and C-termini in the tree. Another method, BLAST-based method
4 (Bi-direction best hit) [46], was used to verify the putative orthologous genes (e-value =
5 1e-20).

6 **Evolutionary analysis of WRKY III genes in willow**

7 The group of WRKY III genes, only found in flowering plants, was considered as the
8 evolutionary youngest groups, and played crucial roles in process of plant growth and
9 resistance [7, 13]. Previous study of Zhang et al. held the opinion that duplications and
10 diversifications were plentiful in WRKY III genes, and they appeared to have confronted
11 different selection challenges [35]. Phylogenetic analysis of WRKY III genes was performed
12 using MEGA6.0 with 65 WRKY III genes from *Arabidopsis* (AtWRKY), *Populus*
13 (PtWRKY), grape (VvWRKY), willow (SsWRKY) and rice (OsWRKY). A NJ tree was
14 constructed with the same method described before. Additionally, we estimated the
15 non-synonymous (Ka) and synonymous (Ks) substitution ratio of SsWRKY III genes to verify
16 whether selection pressure participated in the expansion of SsWRKY III genes. Each pair of
17 these WRKY III protein sequences was first aligned using ClustalW. The alignments
18 generated by ClustalW and the corresponding cDNA sequences were submitted to the online
19 program PAL2NAL (<http://www.bork.embl.de/pal2nal/>) [47], which automatically calculates
20 Ks and Ka by the codeml program in PAML [48].

21 **Analysis of exon-intron structure, gene duplication events and** 22 **conserved motif distribution of willow WRKY genes**

23 The exon-intron structures of the willow WRKY genes were obtained based on the protein
24 annotation files which we assembled ourselves
25 (http://bio.njfu.edu.cn/ss_wrky/version5_2.gff3), and the diagrams were obtained from the
26 online website Gene Structure Display Server (GSDS: <http://gsds.cbi.pku.edu.cn/>) [49].

1 Gene duplication events were always considered as the vital sources of biological evolution.
2 Blastp (e-value, $1e-20$) was performed to identify the gene duplication events in SsWRKY
3 genes with the following definition [7, 50]: (1) the coverage of the aligned sequence $\geq 80\%$ of
4 the longer gene; and (2) the similarity of the aligned regions $\geq 70\%$.

5 To better exhibit the structural features of SsWRKY proteins, the online tool MEME
6 (Multiple Expectation Maximization for Motif Elicitation) was used to identify the conserved
7 motifs in the encoded SsWRKY proteins [51]. The optimized parameters were employed as
8 the following: any number of repetitions, maximum number of motifs = 20, and the optimum
9 width of each motif was constrained to between 6 to 50 residues. The online program 2ZIP
10 (<http://2zip.molgen.mpg.de/>) was used to verify the existence of the conserved Leu zipper
11 motif [52], whereas some other important conserved motifs, HARF, LXXLL (X, any amino
12 acid) and LXLXLX, were identified manually.

13 **Expression analyses of willow WRKY genes**

14 The sequenced *S. suchowensis* RNA-HiSeq reads from five tissues including tender roots,
15 young leaves, vegetative buds, non-lignified stems and barks were separately mapped back
16 onto the SsWRKY gene sequences using BWA (mismatch ≤ 2 bp, other parameters as
17 default) [53], and the number of mapped reads for each WRKY gene was counted.
18 Normalization of the mapped reads was done using RPKM (reads per kilo base per million
19 reads) method [54]. The heat map for tissue-specific expression profiling was generated based
20 on the \log_2 RPKM values for each gene in all the tissue samples using R package [55].

21 **Results**

22 **Identification and characterization of 85 WRKY genes in willow** 23 **(*Salix suchowensis*)**

24 In this study, we obtained 92 putative WRKY genes by using HMMER to search the Hidden
25 Markov Model profile of WRKY DNA-binding domain against willow protein sequences,

1 and validated the accuracy of the consequence by BlastP. After submitting the 92 putative
2 WRKY genes to the online program SMART, seven genes without a complete WRKY
3 domain were removed (willow_GLEAN_10004672, willow_GLEAN_10009126,
4 willow_GLEAN_10011436, willow_GLEAN_10011470, willow_GLEAN_10018393,
5 willow_GLEAN_10019671 and willow_GLEAN_10024347), and the other 85 WRKY genes
6 were selected as possible members of the WRKY superfamily.

7 WRKY genes contain one or two WRKY domains, comprising a conserved WRKYGQK
8 heptapeptide at the N-termini and a novel zinc finger motif (C-X_{4,7}-C-X_{22,23}H-X-H/C) at the
9 C-termini [2]. The variations of WRKY core domain or zinc finger motif may lead to the
10 binding specificities of WRKY genes, but this remains to be largely demonstrated [19, 56, 57].
11 In order to identify the variations in WRKY core domains, a multiple sequence alignment of
12 85 SsWRKY core domains was conducted, and the result was shown in Fig. 1. Among the
13 selected 85 WRKY genes, 81 (95.3%) were identified to have highly conserved sequence
14 WRKYGQK, whereas the other four WRKY genes (SsWRKY14, SsWRKY23, SsWRKY38
15 and SsWRKY78) had a single mismatched amino acid in their core WRKY domains (Fig. 1).
16 In SsWRKY14 and SsWRKY38, the WRKY domain has the sequence WRKYGKK, while
17 SsWRKY23 contains a WKKYGQK sequence, and SsWRKY78 contains WRKYGRK
18 sequence. Eulgem et al. previously described that the zinc finger motif (C-X_{4,5}-X_{22,23}-H-X₁-H
19 or C-X₇-C-X₂₃-H-X₁-C) is another vital features of the WRKY family [2]. As illustrated in
20 Fig. 1, four WRKY domains (SsWRKY76C, SsWRKY64, SsWRKY12 and SsWRKY28) do
21 not contain any distinct zinc finger motif, but they were still reserved in the succeeding
22 analyses, as performed in barley and poplar [7, 11]. Additionally, some zinc-finger-like motifs,
23 including C-X₄-C-X₂₁-H-X₁-H in SsWRKY23 and C-X₅-C-X₁₉-H-X₁-H in SsWRKY73 and
24 SsWRKY17, were identified in willow WRKY genes. Both the two zinc-finger-like motifs
25 were also found in poplar (PtWRKY39, 57, 42 and 53).

26 Detailed characteristics of SsWRKY genes are list in Table 1, including the WRKY gene
27 specific group numbers, chromosomal distribution, *Arabidopsis* and poplar orthologs. The
28 molecular weight (MW), isoelectric point (PI) and the length of each WRKY protein

1 sequence are also shown in Table 1. According to the particularization (Table 1), the average
2 length of these protein sequences is 407 residues, and the lengths ranged from 109 residues
3 (SsWRKY23) to 1,593 residues (SsWRKY78). Additionally, the isoelectric point (PI) ranged
4 from 5.03 (SsWRKY38, SsWRKY60) to 10.27 (SsWRKY28), and the molecular weight
5 (MW) ranged from 12.9 (SsWRKY23) to 179.0 kDa (SsWRKY78).

6 **Locations and gene clusters of willow WRKY genes**

7 84 of the 85 putative SsWRKY genes could be mapped onto 19 willow chromosomes and
8 then renamed from SsWRKY1 to SsWRKY84 based on their specific distributions on the
9 chromosomes. Only one SsWRKY gene (willow_GLEAN_10002834), renamed as
10 SsWRKY85, could not be conclusively mapped onto any chromosome. As shown in Fig. 2,
11 Chromosome (Chr) 2 possessed the largest number of SsWRKY genes (11 genes), followed
12 by Chr14 (10 genes). Eight SsWRKY genes were found on Chr6, six on Chr1 and Chr16, and
13 five on Chr5. Additionally, four chromosomes (Chr4, Chr11, Chr17, Chr18) had four
14 SsWRKY genes, as well as three SsWRKY genes were found on Chr8, Chr13 and Chr19.
15 Chr10 and Chr15 had two SsWRKY genes, and only one SsWRKY gene was identified on
16 Chr7, Chr9 and Chr12. The distribution of each SsWRKY genes was extremely irregular,
17 indicating the reduction of the tandem duplication events in willow WRKY genes.

18 Gene clusters, defined as a single chromosome containing two or more genes [58], are very
19 important for predicting co-expression genes or potential function of clustered genes in
20 angiosperms [59]. According to this description, a total of 23 SsWRKY genes were clustered
21 into 11 clusters in willow (Fig. 2). The chromosomal distribution of gene cluster was irregular,
22 and only seven chromosomes were identified to have gene clusters. Three clusters, including
23 seven SsWRKY genes, were found on Chr2, and two clusters were found on both Chr6 and
24 Chr14. Only one cluster was distributed on each of Chr3, Chr8, Chr10 and Chr18, whereas
25 none was identified on other eleven chromosomes. Further analysis of SsWRKY
26 chromosomal distribution showed that a high WRKY gene density region in only 2.23 Mb
27 regions on Chr2, which had also been observed in rice and poplar [7, 13].

1 **Phylogenetic analysis and classification of WRKY genes in willow**

2 In order to get a better separation of different groups and subgroups in SsWRKY genes, a
3 total of 185 WRKY domains, including 82 AtWRKY domains and 103 SsWRKY domains,
4 were used to construct the NJ phylogenetic tree. On the basis of the phylogenetic tree and
5 structural features of WRKY domains, all 85 SsWRKY genes were clustered into three main
6 groups (Fig. 3). Nineteen members containing two WRKY domains and C₂H₂-type zinc finger
7 motifs were categorized into group I, except SsWRKY78, which contains only one WRKY
8 domain and two zinc finger motifs. Domain acquisition and loss events appear to have shaped
9 the WRKY family [60, 61]. Thus, SsWRKY78 may have evolved from a two-domain WRKY
10 gene but lost one WRKY domain during evolution. Additionally, as shown in Fig. 3,
11 SsWRKY78 shows high similarities to SsWRKY40N, implying a common origin of their
12 domains. The similar phenomenon was also found in PtWRKY90 of poplar [7].

13 The largest number of SsWRKY genes, comprising a single WRKY domain and C₂H₂ zinc
14 finger motif, were categorized into group II. SsWRKY genes of group II could be further
15 divided into five subgroups: IIa, IIb, IIc, IId and IIe. As shown in Fig. 3, subgroup IIa (4
16 members) and IIb (8 members) were clustered into one clade, as well as subgroup IId (13
17 members) and IIe (11 members). Strikingly, SsWRKY genes in subgroup IIc (21 members)
18 and group IC are classified into one clade, suggesting that group II genes are not
19 monophyletic and the group IIc WRKY genes may evolve from the group I genes by the loss
20 of the WRKY domain in N-terminal. As shown in Fig. 3 and Fig. 4, SsWRKY23,
21 SsWRKY34 and their orthologous genes, AtWRKY49, PtWRKY39, PtWRKY57,
22 PtWRKY34 and PtWRKY32, seem to form a new subgroup, and shown to be closer to the
23 group III according to the phylogenetic analysis. However, SsWRKY23 and SsWRKY34
24 exhibit the zinc finger motif C-X₄-C-X₂₁-H-X-H and C-X₄-C-X₂₃-H-X-H as observed in the
25 subgroup IIc and group IC. Thereby, they were classified into subgroup IIc in this study.

26 Different from the C₂H₂ zinc finger pattern in group I and II, group III WRKY genes (7
27 members), broadly considered as playing vital roles in plant evolution process and
28 adaptability, contained one WRKY domain and a C-X₇-C-X₂₃-H-X-C zinc finger motif.

1 Intriguingly, a subgroup IIIb containing a $CX_7CX_nHX_1C$ ($n \geq 24$) zinc finger motif was
2 identified in rice and barley [11, 13]. However, this $C-X_7-C-X_n-H-X-C$ ($n \geq 24$) zinc finger
3 motif was never found in poplar, grape, *Arabidopsis* and willow, suggesting that this feature
4 perhaps only belong to monocotyledonous species.

5 In order to obtain a better study in woody plant species, a phylogenetic tree based on the
6 WRKY domains between willow and poplar was constructed (Fig. 4). The tree showed that
7 most of the WRKY domains from willow and poplar were clustered into sister pairs,
8 suggesting that gene duplication events played prominent roles in the evolution and expansion
9 of WRKY gene family. Furthermore, a total of twenty SsWRKY domains show extremely the
10 same domains (similarity: 100%) to poplar, i.e., SsWRKY39 and PtWRKY9, SsWRKY39
11 and PtWRKY9, SsWRKY39 and PtWRKY9, SsWRKY39 and PtWRKY9, and so on. Further
12 functional analyses of these genes in willow or poplar will provide a useful reference for
13 another one.

14 **The ortholog of SsWRKY genes in *Arabidopsis* and poplar**

15 The clustering of orthologous genes emphasizes the conservation and divergence of gene
16 families, and they may contain the same functions [9]. In this study, a phylogeny-based
17 method was used to identify the putative orthologous SsWRKY genes in *Arabidopsis* and
18 poplar (Fig. 3 and Fig. 4), and BLAST-based method (Bi-direction best hit) was used to
19 confirm the true orthologs. The WRKY genes of group I contained two WRKY domains, and
20 both of them were used to construct the phylogenetic trees. To avoid the mistakes of
21 orthologous genes in group I, the members of group I WRKY genes were considered as
22 orthologous genes unless the same phylogenetic relationship can be detected between
23 N-termini and C-termini in the phylogenetic tree. For example, SsWRKY37 and AtWRKY44
24 were considered as an orthologous gene pair because they clustered into a clade of their
25 N-termini and C-termini (Fig. 3), while SsWRKY80 and PtWRKY30 were excluded from
26 orthologous gene pairs due to their different clusters of N-termini and C-termini (Fig. 4).
27 Totally, 75 orthologous gene pairs were found between willow and *Arabidopsis*, less than 82

1 orthologous gene pairs between willow and poplar (Table 1), which was congruent with the
2 evolutionary relationship among the three plant species.

3 **Evolutionary analysis of WRKY III genes in willow**

4 The WRKY III genes were considered as the evolutionary youngest groups, and played
5 crucial roles in the process of plant growth and resistance. In order to further probe the
6 duplication and diversification of WRKY III genes after the divergence of the monocots and
7 dicots, a phylogenetic tree was constructed using 65 WRKY III genes from *Arabidopsis* (13),
8 rice (29), poplar (10), willow (7) and grape (6). As shown in Fig. 5, willow SsWRKY III
9 genes were closer to the eurosids I group (poplar and grape) than eurosids II group
10 (*Arabidopsis*) and monocots (rice). Meanwhile, most *Arabidopsis* and rice WRKY III genes
11 formed the relatively independent clades, suggesting that two gene duplication events,
12 including tandem and segmental duplication, perhaps were the main factors in the expansion
13 of WRKY III genes in *Arabidopsis* and rice. What's more, the results also indicated that
14 WRKY III might arise after the divergence of the *Arabidopsis* (eurosids I) and eurosids II
15 (poplar, willow and grape). The study by Ling et al. in cucumber [9] showed the similar
16 results and hence proved the validity. Interestingly, seven rice WRKY III genes (OsWRKY55,
17 84, 18, 52, 46, 114 and 97) contained the variant domain WRKYGEK, but the variant was not
18 found in other four dicots, implying that this may be a feature of WRKY III genes in
19 monocots and these OsWRKY genes may respond to different environmental signals.

20 According to the comparison of the number of WRKY III genes in the five observed plants,
21 the number is smaller in eurosids I (poplar, grape and willow) than *Arabidopsis* (eurosids II)
22 and rice (monocots), which may be caused by different patterns of duplication events. Genes
23 generated by duplication events are not stable, and can be retained or lost due to different
24 selection pressure and evolution [62]. In order to determine which selection pressure played
25 prominent roles in the expansion of willow WRKY III genes, we estimated the Ka/Ks ratios
26 for all pairs (21 pairs) of willow WRKY III genes. As shown in Fig. 6, all the Ka/Ks ratios

1 were less than 0.5, suggesting willow WRKY III genes had mainly been subjected to strong
2 purifying selection and they were slowly evolving at the protein level.

3 **Exon–intron structures of SsWRKY genes**

4 The exon-intron structures of multiple gene families play crucial roles during plant evolution.
5 As shown in Fig. 7, the SsWRKY gene phylogenetic tree and the corresponding exon-intron
6 structures are shown in A and B, respectively. Exon-intron structures of each group were
7 shown in Fig. 7B, a large number of WRKY genes had two to five introns (94%, 80 of 85),
8 including 8 WRKY genes contained one intron; 39 contained two introns; 13 contained three
9 introns; 15 contained four introns and 5 contained five introns. The number of exons in
10 remaining WRKY genes was quite different: SsWRKY49, SsWRKY76 and SsWRKY78 had
11 six, eleven and ten introns, respectively; SsWRKY17 had the largest number of introns
12 (seventeen introns), while no intron was found in SsWRKY12. The intron acquisition or loss
13 occurred during the evolution of WRKY gene family, while WRKY genes in the same group
14 shared the similar number of introns [6]. In our study, most of WRKY genes in group I had
15 three to six introns, except SsWRKY76 and SsWRKY78, which might acquire some introns
16 during evolution. The number of introns of WRKY genes in group II was extremely different,
17 ranging from one to five introns, except SsWRKY17 with 17 introns and SsWRKY12 with
18 zero intron might obtain or lose some introns during evolution. Strikingly, WRKY genes in
19 group III had the most stable number of introns with all of seven WRKY III genes had two
20 introns, suggesting that WRKY III genes may be the most stable genes in the environmental
21 stress. The stable number of introns in SsWRKY III genes was consistent with the results of
22 Ka/Ks analysis, which reflected that purifying selection pressure played vital roles in willow
23 WRKY III genes.

24 A great deal of research in WRKY genes proved that nearly all of the WRKY genes
25 contained an intron in their WRKY core domains [2, 6-9, 30]. According to the further
26 analysis of SsWRKY genes, two major types of splicing introns, R-type and V-type, introns
27 were observed in numerous SsWRKY domains. The R-type intron was spliced exactly at the

1 R residue, about five amino acids before the first Cys residue in the C₂H₂ zinc finger motif.
2 The V-type intron was localized before the V residue, six amino acids after the second Cys
3 residue in the C₂H₂ zinc finger motif. As shown in Fig. 7B, the R-type introns could be
4 observed in more groups, including group IC, subgroup IIc, IId, IIe and group III, while
5 V-type introns were only observed in subgroup IIa and IIb. Moreover, there was no intron
6 found in group IN. The similar results were also observed in *Arabidopsis*, poplar and rice,
7 suggesting that the special distribution of introns in WRKY domains was a feature of WRKY
8 family.

9 **Identification of gene duplication events and conserved motifs in** 10 **willow**

11 Gene duplication events were always considered as the vital sources of biological evolution
12 [63, 64]. Two or more adjacent homologous genes located on a single chromosome were
13 considered as tandem duplication events (TDs), while homologous gene pairs between
14 different chromosomes were defined as segmental duplication events (SDs) [10]. In our study,
15 a total of 33 homologous gene pairs, including 66 SsWRKY genes, were identified to
16 participate in gene duplication events. The composition of gene duplication events in each
17 group in ascending order was group I: 73.7% (14 of 19), group II: 78% (46 of 59) and group
18 III: 85.7% (6 of 7). Among the 33 homologous gene pairs, none of them appeared to have
19 undergone TDs, on the contrary, all of the 66 genes (77.6% of all SsWRKY genes)
20 participated in SDs, implying that segmental duplication events played major roles in the
21 expansion of willow WRKY genes.

22 WRKY genes shared more functional and homologies in their conserved WRKY core
23 domains (about 60 residues), while the rest sequences of WRKY genes shared a little [2]. In
24 order to get a more comprehensive understanding of the structural feature in WRKY domains,
25 the conserved motifs of SsWRKY genes were predicted using the online program MEME
26 (Fig. 8 and Table 2). Among the 20 putative motifs, motifs 1, 2, 3 and 5, broadly distributed
27 across SsWRKY genes, were characterized as the WRKY conserved domains. The motif 6

1 was characterized as nuclear localization signals (NLS), which mainly distributed in subgroup
2 II d and IIe and group III. Some other motifs with poorly defined recently were also predicted
3 by MEME: the motif 4 was only found in group IC and subgroup IIc; motifs 7 and 9 were
4 limited to subgroup IIa and IIb; the motif 8 was found in group I and a few genes of subgroup
5 IIc; motifs 10, 13, 15 and 17 were unique in subgroup IIId; the motif 12 was only observed in
6 subgroup IIb; the motif 16 was mainly found in group II; the motif 18 was found in subgroup
7 IIc; motifs 19 and 20 were only observed in subgroup I. The distinct conserved motifs of
8 different groups could be an important foundation for future structural and functional study in
9 WRKY gene family.

10 Some other important motifs, including Leu zipper motif, HARF, LXXLL and LXLXLX,
11 could be also identified in WRKY genes. Using the online program 2ZIP, the conserved Leu
12 zipper motif, described as a common hypothetical structure to DNA binding proteins [65],
13 was identified in only two SsWRKY genes (SsWRKY61 and SsWRKY39). With manual
14 inspection, the conserved HARF (RTGHARFRR[A/G]P) motifs, whose putative functions
15 were not distinguished clearly, were only observed in seven WRKY genes of subgroup IIId,
16 including SsWRKY82, 33, 45, 81, 9, 30 and 56. In the meantime, the conserved LXXLL and
17 LXLXLX (L: Leucine; X: any amino acid) motifs, which respectively defined as the
18 co-activator and active repressor motifs, were also found in SsWRKY genes. A total of seven
19 SsWRKY genes (SsWRKY19, 45, 72, 61, 76, 30 and 59) contained the helical motif LXXLL,
20 whereas eight genes (SsWRKY66, 26, 35, 81, 83, 75, 73 and 3) shared the LXLXLX motif.
21 The plenty of conserved motifs in WRKY genes with different lengths and variant functions,
22 suggesting that the WRKY genes might play more vital roles in gene regulatory network.

23 **Distinct expression profiles of SsWRKY genes in various tissues**

24 In order to gain more information about the roles of WRKY genes in willow, RNA-seq data
25 from the sequenced genotype were used to quantify the expression level of WRKY genes in
26 five tissues of *Salix suchowensis*. As illustrated in Fig. 9, the expression of all 85 SsWRKY
27 genes were detected in at least one of the five examined tissues, such as 84 genes in roots, 80

1 in stems, 84 in barks, all in buds and 73 in leaves. Meanwhile, the cluster analysis of the
2 expression pattern in five tissues showed that SsWRKY genes shared more similarities
3 between stem and leaf, as well as bark and bud, and root was more similar to the clade formed
4 by bark and bud. The results detected here were consistent with their biological characteristics.
5 SsWRKY38, not detected in roots and leaves, was also lowly expressed in other tissues.
6 Similarly, SsWRKY74, not detected in stems, barks and leaves, was only expressed in roots
7 and buds with extremely low levels. Among the five genes not expressed in stems,
8 SsWRKY66, 74 and 79 were also not detected in leaves. The largest number of expressed or
9 unexpressed SsWRKY genes (12 genes) was found in buds or leaves, respectively, suggesting
10 that WRKY genes might play more roles in buds than leaves.

11 According to the expression annotation of 85 SsWRKY genes by RPKM method in Fig. 9
12 and Table S1, the total transcript abundance of SsWRKY genes in tender root (RPKM =
13 1181.21), bark (RPKM = 1363.01) and vegetative bud (RPKM = 928.58) was relatively larger
14 than that in other two tissues, including non-lignified stem (RPKM = 537.88) and young leaf
15 (RPKM = 349.84). As shown in Table S1, SsWRKY81 (RPKM = 97.75), the most expressed
16 SsWRKY genes in roots, was also expressed in other four tissues, though the expression
17 levels were relatively low; SsWRKY56 (RPKM = 32.54), the most expressed SsWRKY genes
18 in stem, was also highly expressed in other examined tissues. Similarly, SsWRKY67, the
19 most expressed SsWRKY genes in barks (RPKM = 188.16), was also detected in vegetative
20 buds (RPKM = 82.07) and young leaves (RPKM = 26.11) with high expression levels.
21 Similarly, SsWRKY6 (RPKM = 26.31), the most expressed genes in leaves, was also highly
22 detected in other tissues. A few genes, i.e., SsWRKY52, SsWRKY2 and SsWRKY35, were
23 expressed highly in barks, but lowly in other four tissues. The results mentioned above may
24 be an important foundation for the specific expression analysis of each WRKY gene in
25 willow.

26

1 Discussion

2 WRKY genes are the induced plant TFs, which can specifically interact with the W-box to
3 regulate the expressions of downstream target genes. They also play prominent roles in
4 diverse physiological and growing processes, especially in various abiotic and biotic stress
5 responses in plants. Previous studies about the features and functions of WRKY family have
6 been conducted in many model plants, including *Arabidopsis* for annual herbaceous dicots [2],
7 grape for perennial dicots [6], poplar for woody plants and rice for monocots [7, 13]. Here,
8 the comprehensive analyses of WRKY family in willow (*Salix suchowensis*) would not only
9 provide valuable information for future functional analysis of WRKY genes in woody plants,
10 but also provide an important reference to investigate the complex structures, evolution and
11 gene expansion in this gene superfamily. In this study, a total of 85 SsWRKY genes were
12 identified from willow, accompanying with analyses of their complex structures,
13 classification, gene expansion patterns, conserved motifs and distinct expression profiles.

14 Comparing the two phylogenetic trees based on the SsWRKY domains (Fig. 3) and
15 proteins (Fig. 7 A), we obtained the nearly same classification of all SsWRKY genes,
16 suggesting that the conserved WRKY domain is an indispensable unit in WRKY genes. The
17 variation of the WRKYGQK heptapeptide may influence the proper DNA-binding ability of
18 WRKY genes [17, 18]. A recent binding study by Brand et al. disclosed that a reciprocal Q/K
19 change of the WRKYGQK heptapeptide might result in different DNA-binding specificities
20 of the respective WRKY genes [56]. For instance, the soybean WRKY genes, GmWRKY6
21 and GmWRKY21, which contains the WRKYGKK variant, can't bind normally to the W-box
22 ([C/T]TGAC[T/C]) [66]. Another NtWRKY12 gene in tobacco with the WRKYGKK variant
23 recognizes another binding sequence 'TTTTCCAC' instead of normal W-box [67]. Strikingly,
24 many WRKY genes with WRKYGKK variant recognize a much more degenerate consensus
25 with only a central GAC-core motif, i.e., AtWRKY50 in *Arabidopsis* [56]. Therefore, further
26 investigation of the functions and binding specificities of the variants of WRKYGQK
27 heptapeptide in plants would be very interesting. In our study, four WRKY genes

1 (SsWRKY14, SsWRKY23, SsWRKY38 and SsWRKY78) had a single mismatched amino
2 acid in their conserved WRKYGQK heptapeptide (Fig. 1), including WRKYGKK,
3 WKKYGQK and WRKYGRK. The variants detected in willow were extremely congruent
4 with that in another salicaceous plant, poplar, which also contains the three variants in seven
5 PtWRKY genes [7]. Additionally, two variants, WRKYGKK and WRRKGQK, were found in
6 grape and tomato [6, 8]; WRKYGKK, the most common variant in plants, was the only one
7 found in castor bean and cucumber [9, 68]. The variants may be different between dicots and
8 monocots. Four variants, including WQKYGQK, WRKYGKK, WSKYGQM and
9 WRKYGEK, were found in barley [11]. Meanwhile, the largest number of variants was found
10 in rice [13], including WQKYGQK, WRKYGEK, WIKYGQK, WRKYSEK, WKKYGQK,
11 WKRYGQK, WSKYEQK and WRKYGKK, perhaps due to its various habitats. Strikingly,
12 WRKYGEK, a prevalent variant in plants, was only found in WRKY III genes of rice and
13 barley among the above plants examined, implying that this variant may be a feature of
14 WRKY III genes in monocots and they may respond to different environmental signals.
15 Moreover, many previous studies have disclosed that the binding specificities of variable
16 WRKYGQK heptapeptide vary tremendously [56], however, few studies were shown about
17 the effect of variable zinc finger motif. In this study, four WRKY domains (SsWRKY76C,
18 SsWRKY64, SsWRKY12 and SsWRKY28) without complete zinc finger motif may lack the
19 ability of interacting with W-box, as well as PtWRKY83, 40, 95 and 10 in poplar [7]. It is still
20 indispensable to further investigate the function or the expression patterns of the regulated
21 gene targets in the variant sequences of the WRKY conserved domains.

22 Different classification methods may lead to different numbers of WRKY genes in each
23 group. The classification method in our study was categorized as described in *Arabidopsis*,
24 grape, cucumber, castor bean and many other plant species [2, 6, 9, 68]. According to this
25 method, the WRKY genes were classified into three main groups (I, II and III), with five
26 subgroups in group II (IIa, IIb, IIc, IId and IIe) based on the number of WRKY domains and
27 the features of diverse zinc finger motifs. However, the strategy described in rice and poplar
28 was a little different [7, 13], and they classified the subgroup IIc categorized above into a new

1 subgroup Ib based on the fact that the C-termini of group I and the domains of the above
2 subgroup IIc shared more similar consensus structures. At the meantime, subgroup IId and IIe
3 categorized above were reclassified into subgroup IIc and IId, respectively. Thus, the number
4 of WRKY genes in poplar and rice was different from other plant species (Table 3). With the
5 same classification method as described in *Arabidopsis* and many other plants, the number of
6 different groups in poplar was as follows: group I: 23, subgroup IIa: 5, IIb: 9, IIc: 31, IId: 13,
7 IIe: 13 and group III: 10, and the number of OsWRKY genes in rice: group I: 14, subgroup IIa:
8 4, IIb: 8, IIc: 20, IId: 7, IIe: 11 and group III: 36. WRKY genes of subgroup IIa, the smallest
9 number of members, appear to play crucial roles in regulating stress responses (both biotic
10 and abiotic) [3]. As illustrated in Table 3, the WRKY genes of subgroup IIa and IIb in willow
11 are extremely similar to that of other plant species, suggesting that all SsWRKY genes of
12 these subgroups have been identified. Subgroup IIa genes, the smallest number of members,
13 appear to play many important roles in regulating biotic and abiotic stress responses [3].
14 Nevertheless, the number of WRKY III in eurosids I group, such as cucumber (6), poplar (10),
15 grape (6) and willow (7) is less than that of eurosids II (*Arabidopsis*: 14) and monocots (rice:
16 36), suggesting that different duplication events or selection pressures occurred in WRKY III
17 genes after the divergence of eurosids I and eurosids II group. Interestingly, the previous
18 study in *Arabidopsis* showed that nearly all WRKY III members respond to diverse biotic
19 stresses, suggesting that this group probably evolved with the increasing biological
20 requirements and the larger number of WRKY III genes in *Arabidopsis* and rice probably due
21 to their various biotic stresses during evolution.

22 WRKY transcription factors play important roles in the regulation of developmental
23 processes and response to biotic and abiotic stress [56]. The evolutionary relationship of
24 WRKY gene family promises to obtain significant insights into how biotic and abiotic stress
25 responses from single cellular aquatic algae to multicellular flowering plants [57]. The first
26 work by Eulgem et al. defined the seven major groups of WRKY genes observed in flowering
27 plants, which has proven over time to be an accurate representation of groups of WRKY
28 genes [2, 3]. Previous studies hypothesized that group I WRKY genes were generated by

1 domain duplication of a proto-WRKY gene with a single WRKY domain, group II WRKY
2 genes evolved through the subsequent loss of N-terminal WRKY domain, and group III genes
3 evolved from the replacement of conserved His residue with a Cys residue in zinc motif [13].
4 However, recent study proposed two alternative hypotheses of WRKY gene evolution [57]:
5 the "Group I Hypothesis" suggests that all WRKY genes in higher plants evolved from group
6 I genes, while the "IIa + b Separate Hypothesis" considers that subgroup IIa and IIb with their
7 hallmark V-type intron are evolved from a single domain of ancestral algal WRKY gene
8 instead of evolving from group I genes. Additionally, another recent study by Brand et al.
9 concluded that subgroup IIc WRKY genes evolved directly from IIc-like ancestral WRKY
10 domains, and group I genes evolved independently due to a duplication of the IIc-like
11 ancestral WRKY domains [56]. In his study, subgroup IIa genes evolved from group I genes
12 through loss of their N-terminal domains; subgroup IIb genes were descendants from IIa
13 genes, because IIb representatives can only be found in monocots and dicots; subgroup IIc
14 genes evolved most probably from IIa, and IIe are most likely the descendants from IIc
15 WRKY genes; and group III WRKY genes are considered as the evolutionary youngest genes.
16 Phylogenetic analysis in our study shows that subgroup IIc and group IC are evolutionarily
17 close, as well as subgroups IIa and IIb, subgroups IIc and IIe, and this result is consistent with
18 the conclusion drew by Brand et al [56]. Additionally, the V-type introns of SsWRKY genes
19 are only found in subgroup IIa and IIb, while R-type introns are found in other groups except
20 group IN. The results are congruent with the "IIa + b Separate Hypothesis". Therefore, further
21 information is still required to determine the accurate evolutionary relationship of WRKY
22 gene family.

23 Gene duplication events played prominent roles in a succession of genomic rearrangements
24 and expansions, and it is also the main motivation of plants evolution [69]. The gene family
25 expansion occurs via three mechanisms: tandem duplication events (TDs), segmental
26 duplication events (SDs) and transposition events [70], and we only focused on the tandem
27 and segmental duplication events in this study. In willow, a total of 66 SsWRKY genes were
28 identified to participate in gene duplication events, and all of these genes appeared to have

1 undergone SDs. In poplar, only one homologous gene pair participated in TDs, while 29 of 42
2 (69%) homologous gene pairs were determined to participate in SDs. The WRKY gene
3 expansion patterns in willow and poplar perhaps showed that SDs were the main factors in the
4 expansion of WRKY genes in woody plants. However, in cucumber, no gene duplication
5 events have occurred in CsWRKY gene evolution, probably because there were no recent
6 whole-genome duplication and tandem duplication in cucumber genome [71]. In rice and
7 *Arabidopsis*, many WRKY genes were generated by TDs, which was incongruent with the
8 duplication events in willow, poplar and cucumber. The different WRKY gene expansion
9 patterns of the above plant species could be due to their different life habits and selection
10 pressures in a large scale.

11 The WRKY gene family plays crucial roles in response to biotic and abiotic stresses, as
12 well as diverse physiological and developmental processes in plant species. Because of the
13 lack of researches on the function of willow WRKY genes, our study provided putative
14 functions of SsWRKY genes by comparing the orthologous genes between willow and
15 *Arabidopsis*. The details of the functions or regulations of AtWRKY genes can be obtained
16 from TAIR (<http://www.arabidopsis.org/>). For example, AtWRKY2, the ortholog to
17 SsWRKY6, which highly expressed in the five examined tissues, plays important roles in
18 seed germination and post germination growth [72]. AtWRKY33, the ortholog to SsWRKY1,
19 35, 55 and 84, influences the tolerance to NaCl, inc sensitivity to oxidative stress and abscisic
20 acid [25]. A large number of AtWRKY genes, i.e. AtWRKY3, 4, 18, 53, 41, work in the
21 resistance to *Pseudomonas syringae* [73-76], so do their orthologs in willow (SsWRKY42, 47,
22 39, 79, 20 and 70). Based on the comparison of willow WRKY genes with their *Arabidopsis*
23 orthologs, we could speculate that the functional divergence of SsWRKY genes has played
24 prominent roles in the responses to various stresses.

25

1 **Conflict of Interests**

2 The authors declare that there is no conflict of interests regarding the publication of this
3 paper.

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9

Figure 1 (on next page)**Comparison of the WRKY domain sequences from 85 SsWRKY genes.**

The WRKY gene with the suffix -N and -C indicates the N-terminal and C-terminal WRKY domain of group I members, respectively. "-" has been inserted for the optimal alignment. Red indicates the highly conserved WRKYGQK heptapeptide, and the zinc finger motifs are highlighted in green. The position of a conserved intron is indicated by an arrowhead.

Figure 2 (on next page)**Chromosomal location of SsWRKY genes.**

Red triangle indicates group I, red star indicates group II and red diamond indicates group III. The chromosome numbers are given at the top of each chromosome and the left side of each chromosome is related to the approximate physical location of each WRKY gene. Only one unmapped SsWRKY gene is shown on SsChrN.

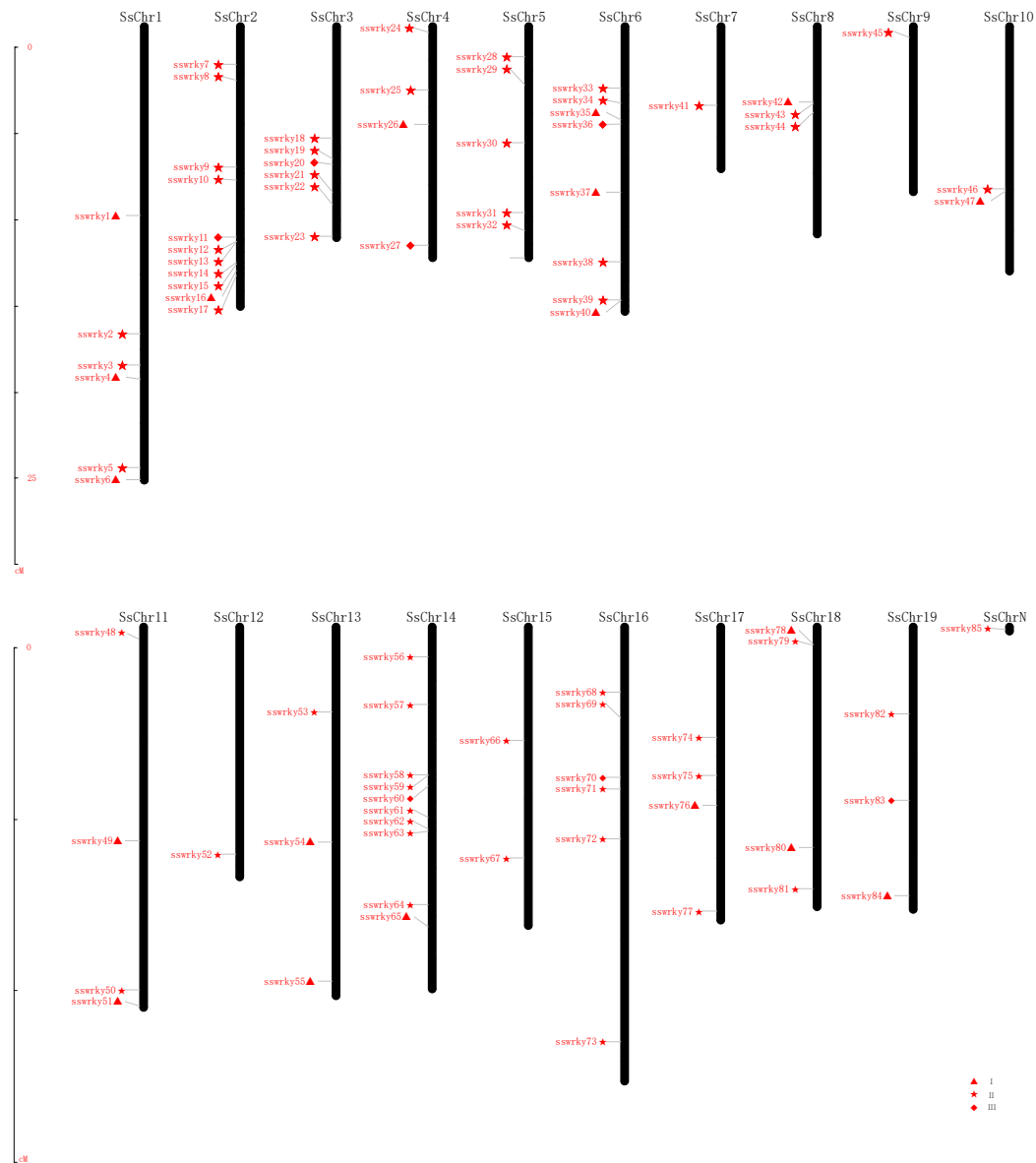


Figure 3 (on next page)**Phylogenetic tree of WRKY domains from willow and *Arabidopsis*.**

The phylogenetic tree was constructed using the neighbor-joining method in MEGA 6.0. The WRKY genes with the suffix 'N' and 'C' indicate the N-terminal and the C-terminal WRKY domains of group I, respectively. The different colors indicate different groups (I, II and III) or subgroups (IIa, b, c, d and e) of WRKY domains. Circles indicate WRKY genes from willow, and diamonds represent genes from *Arabidopsis*.

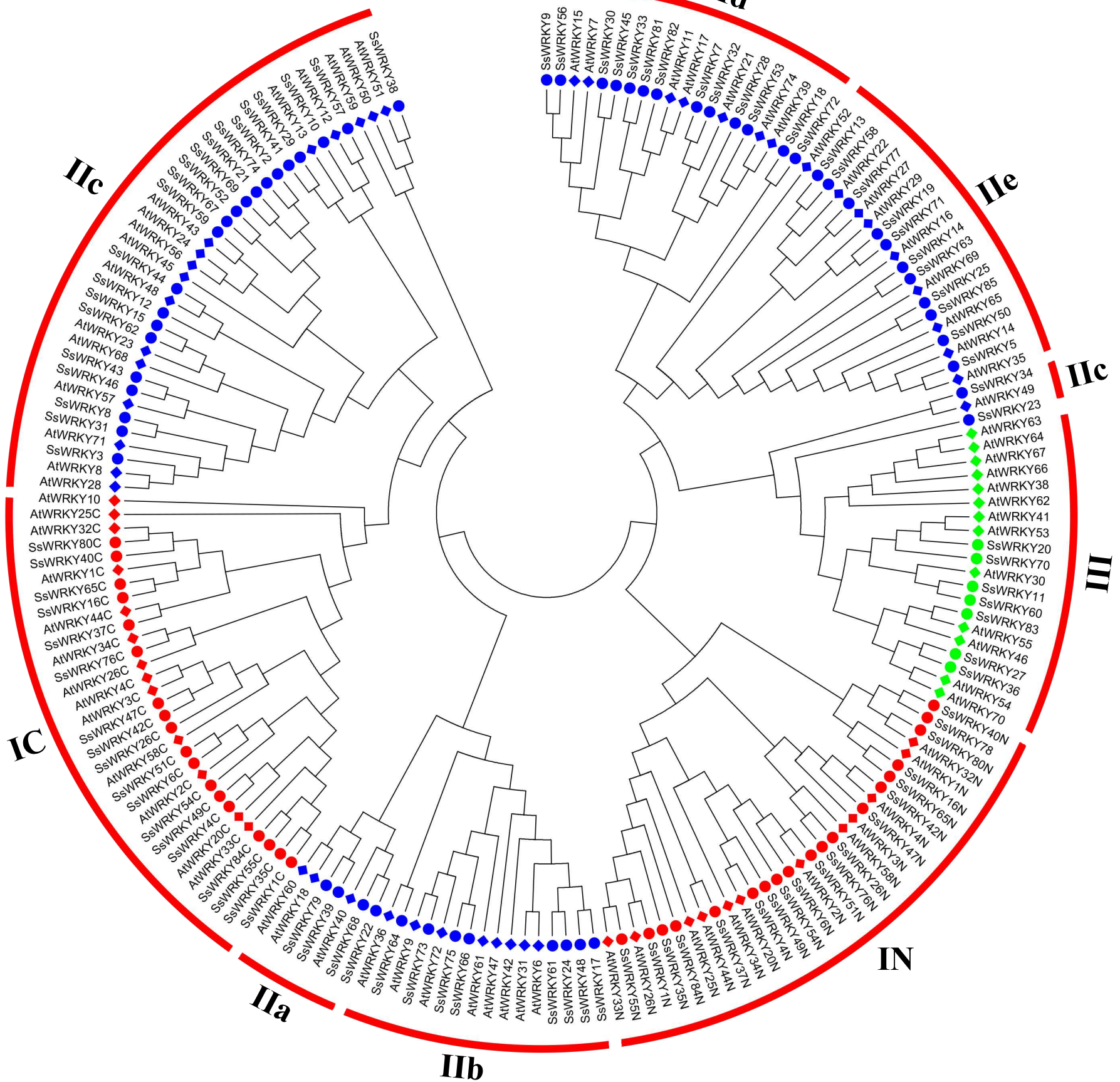


Figure 4(on next page)**Phylogenetic tree of WRKY domains from willow and poplar.**

The phylogenetic tree was constructed using the neighbor-joining method in MEGA 6.0. The WRKY genes with the suffix 'N' and 'C' indicate the N-terminal and the C-terminal WRKY domains of group I, respectively. The different colors indicate different groups (I, II and III) or subgroups (IIa, b, c, d and e) of WRKY domains. Circles indicate WRKY genes from willow, and triangles represent genes from poplar.

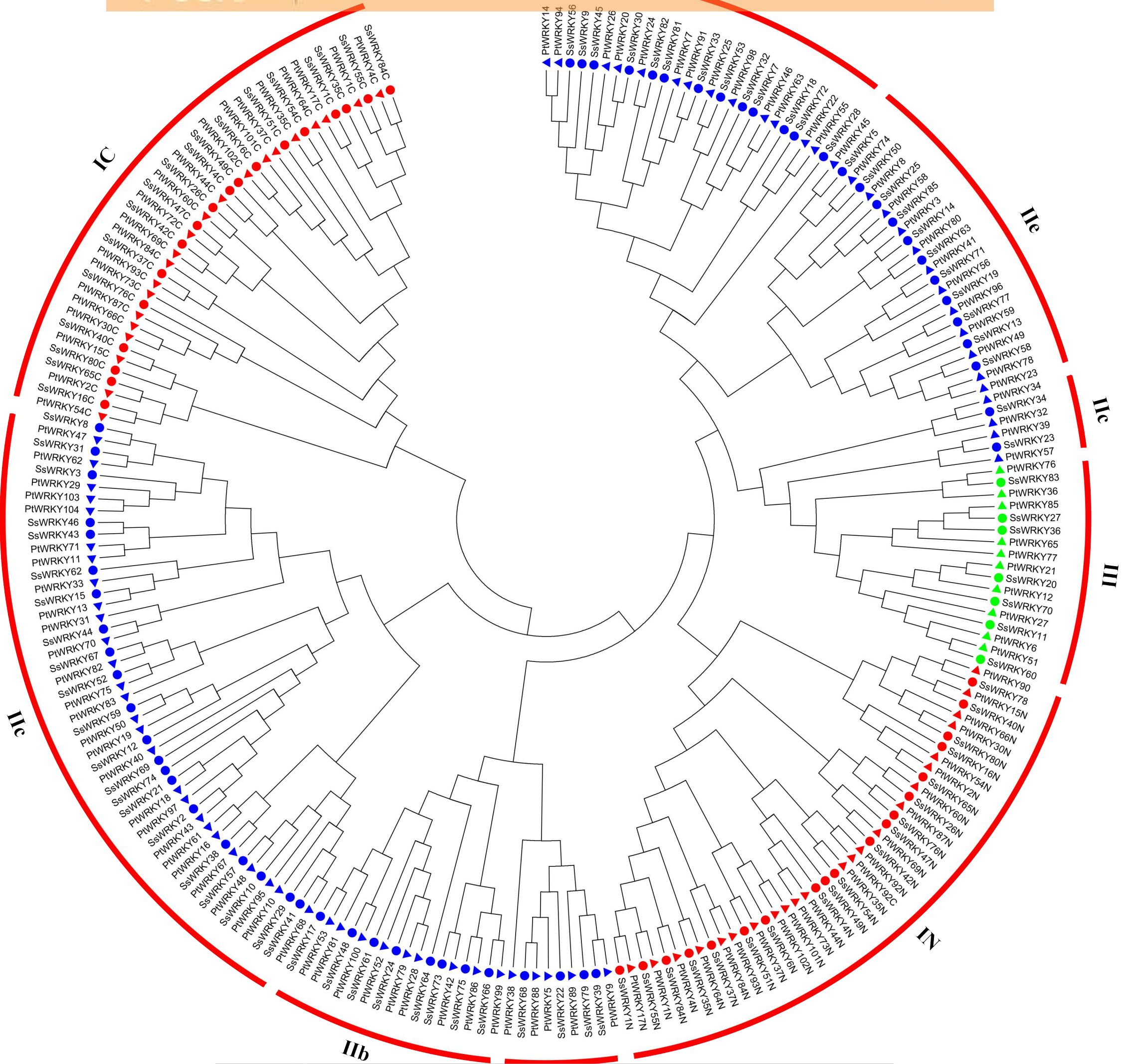


Figure 5 (on next page)

Phylogenetic tree of full-length group III WRKY genes from *Arabidopsis* (AtWRKY), rice (OsWRKY), grape (VvWRKY), poplar (PtWRKY) and willow (SsWRKY).

The phylogenetic tree was constructed using the neighbor-joining method in MEGA 6.0. Dicotyledonous (*Arabidopsis*, grape, poplar and willow) and monocotyledonous (rice) WRKY III genes are marked with colored dots.

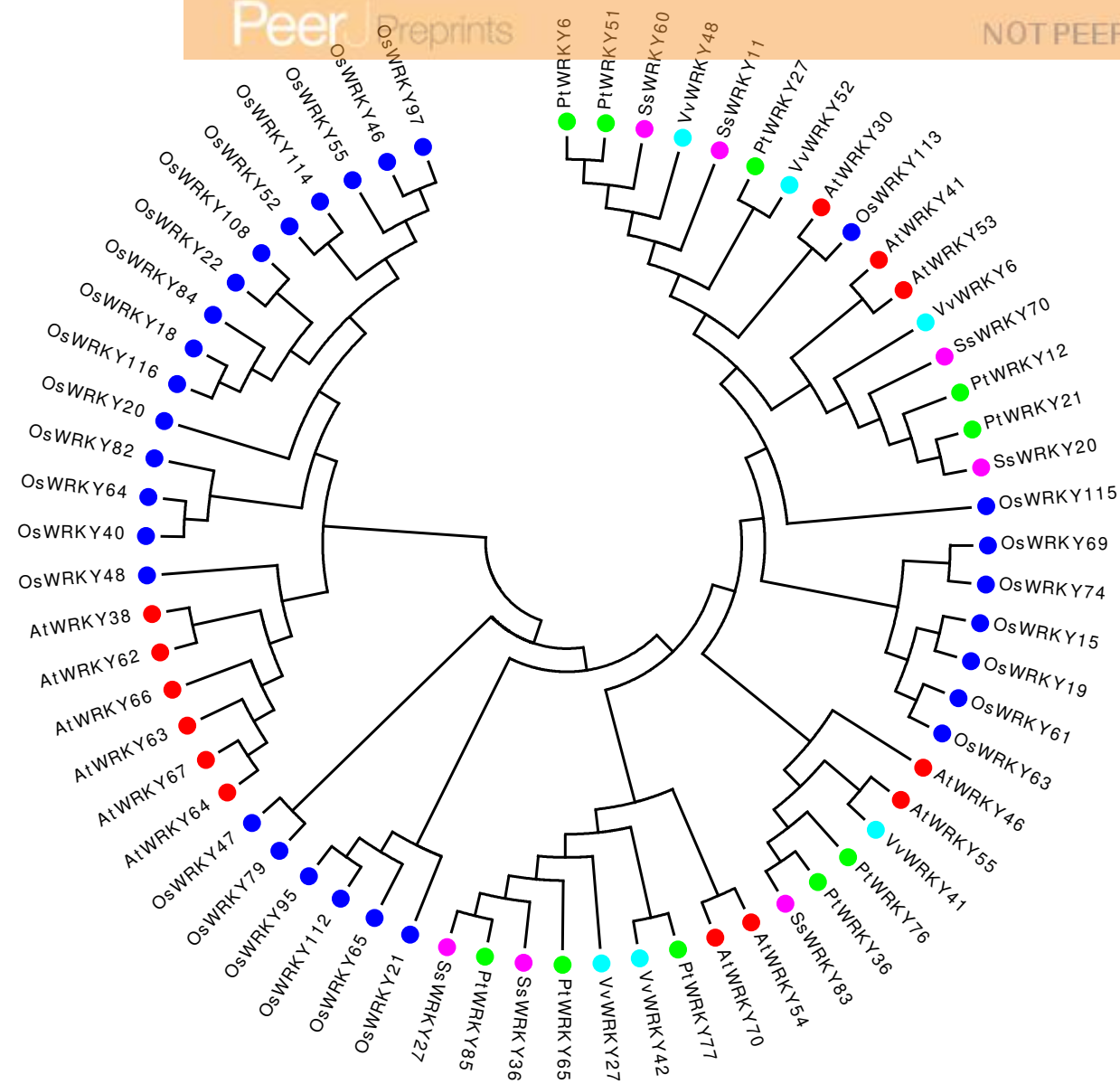


Figure 6 (on next page)**Scatter plots of the Ka/Ks ratios of WRKY III genes in willow.**

The Y- and X-axes denote the Ka/Ks ratio and Ka for each pair, respectively.

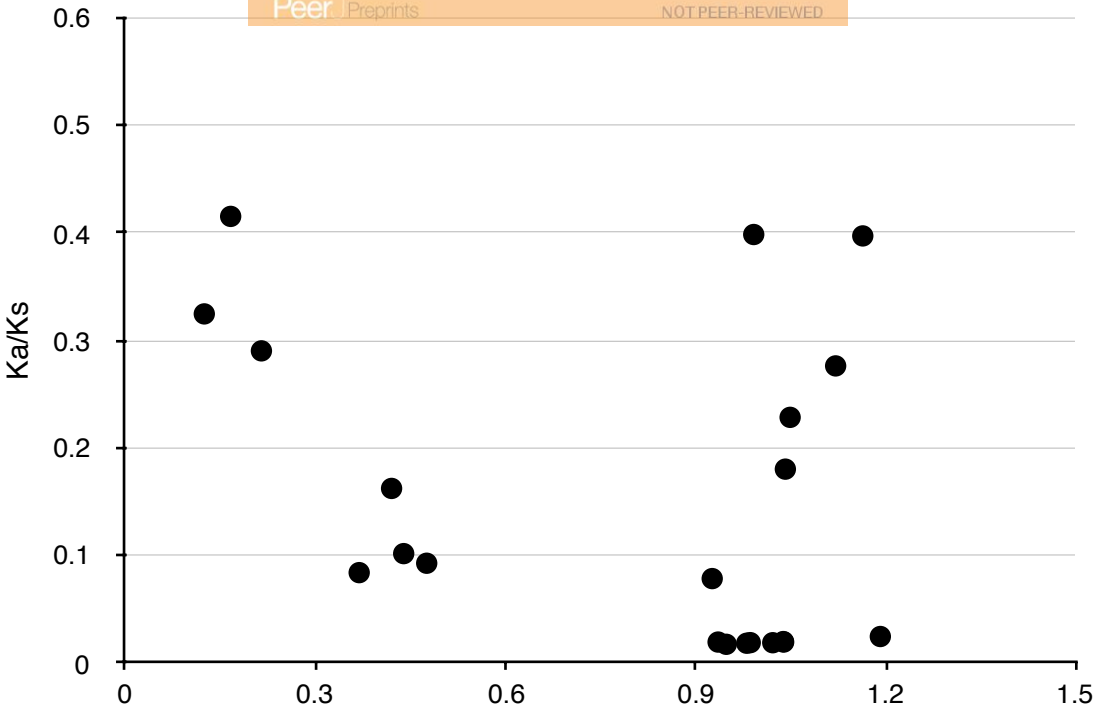


Figure 7 (on next page)**Genomic organization of SsWRKY genes.**

(A) The phylogenetic tree built on the basis of full-length SsWRKY genes was depicted using the neighbor-joining method in MEGA 6.0. The short black lines indicate the existence of duplicated gene pairs; (B) The graphic exon-intron structure of SsWRKY genes is displayed using GSDS. Green indicates exons, and gray indicates introns. The introns phases 0, 1 and 2 are indicated by numbers 0, 1 and 2, respectively.

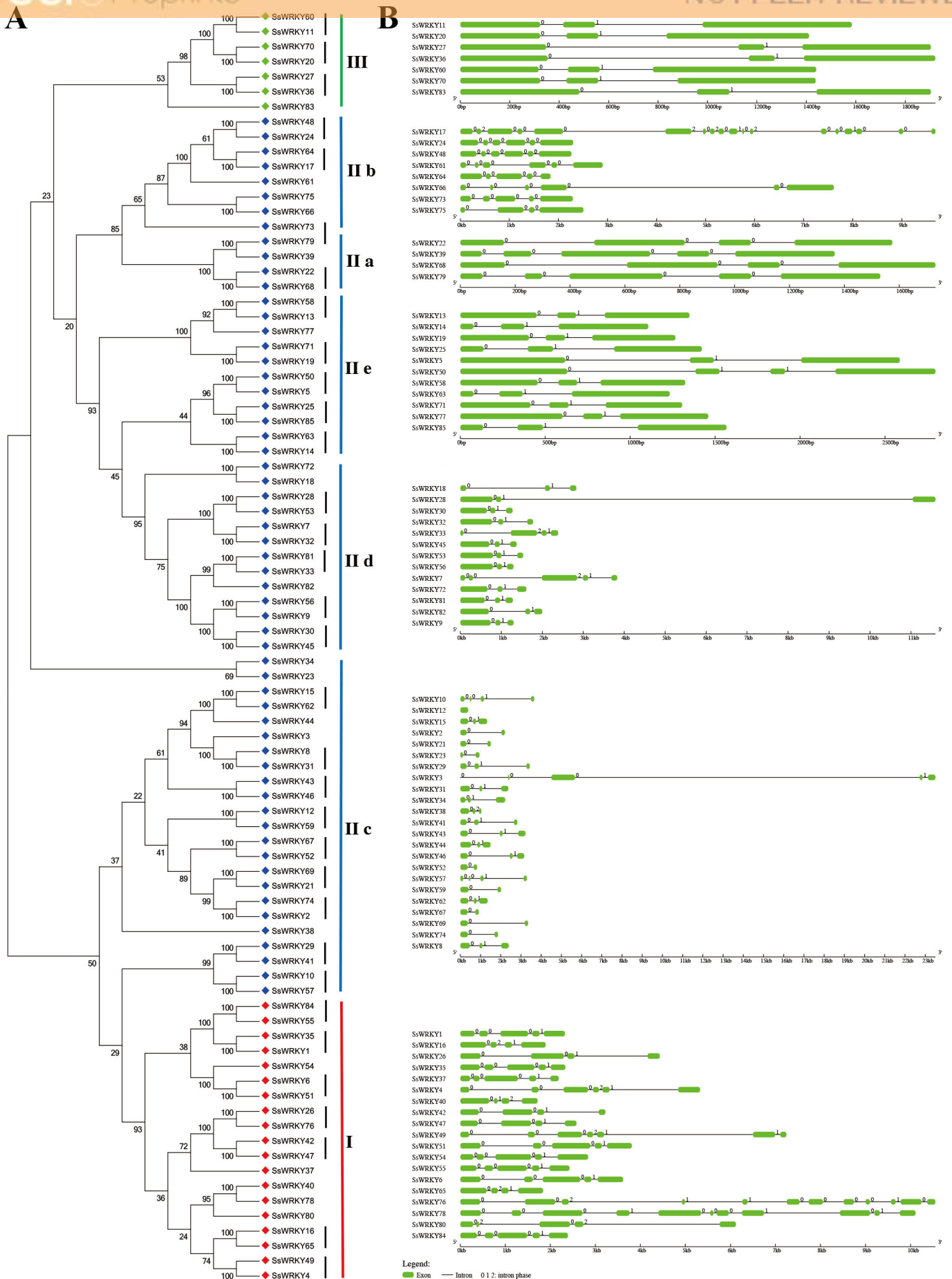


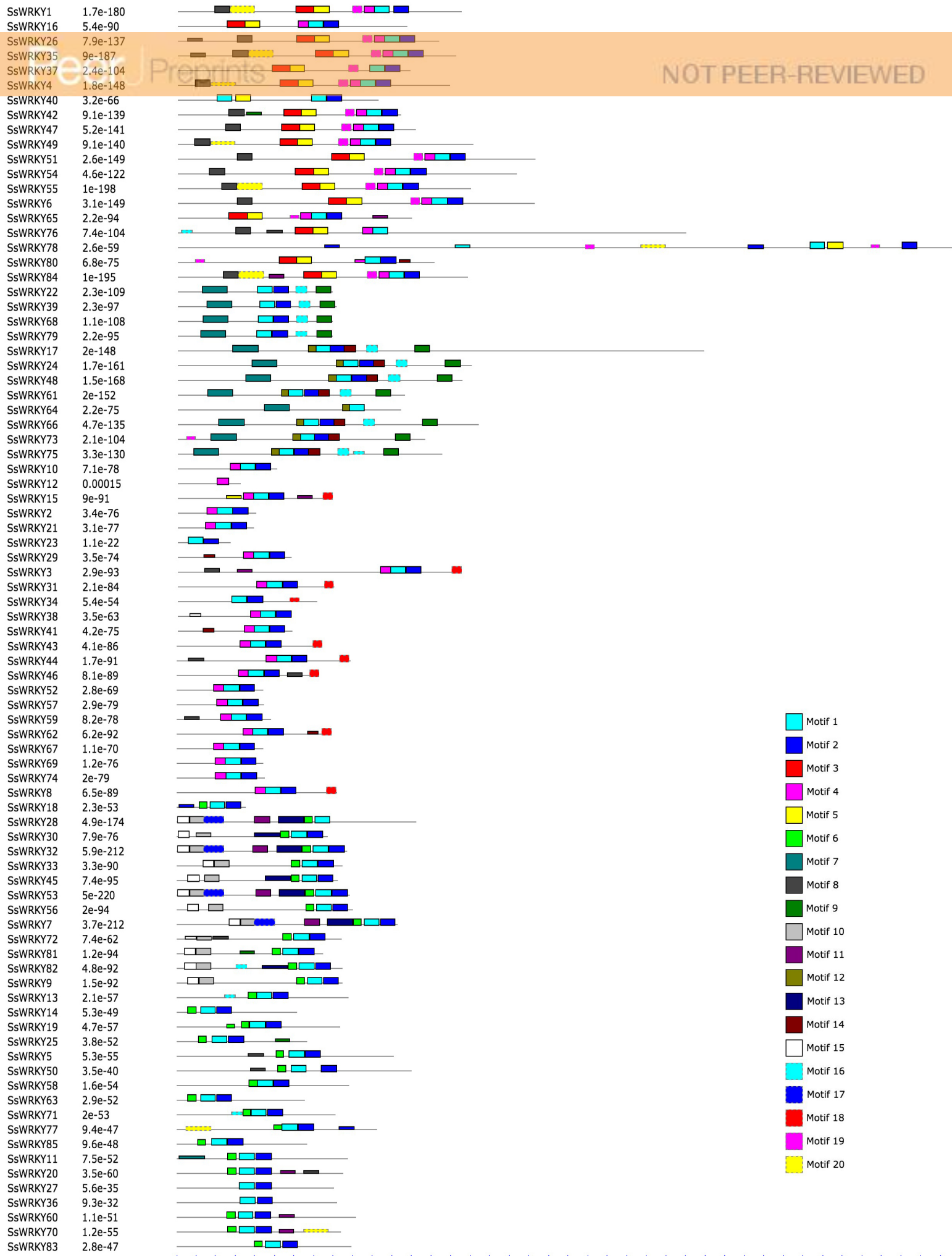
Figure 8(on next page)

The distribution of twenty conserved motifs of SsWRKY genes was identified by the online program MEME.

The names of all members are displayed on the left side of the figure. Different motifs are displayed in different colored boxes as indicated on the right side. The conserved motifs 1, 2, 3, and 5, broadly distributed across SsWRKY genes, were definitely characterized as the WRKY conserved domains.

Name Expect

Motif Location



- Motif 1
- Motif 2
- Motif 3
- Motif 4
- Motif 5
- Motif 6
- Motif 7
- Motif 8
- Motif 9
- Motif 10
- Motif 11
- Motif 12
- Motif 13
- Motif 14
- Motif 15
- Motif 16
- Motif 17
- Motif 18
- Motif 19
- Motif 20

Figure 9 (on next page)**Expression profiles of the 85 SsWRKY genes in root, stem, bark, bud and leaf.**

Color scale represents RPKM normalized log₂ transformed counts and red indicates high expression, blue indicates low expression and white indicates the gene is not expressed in this tissue.

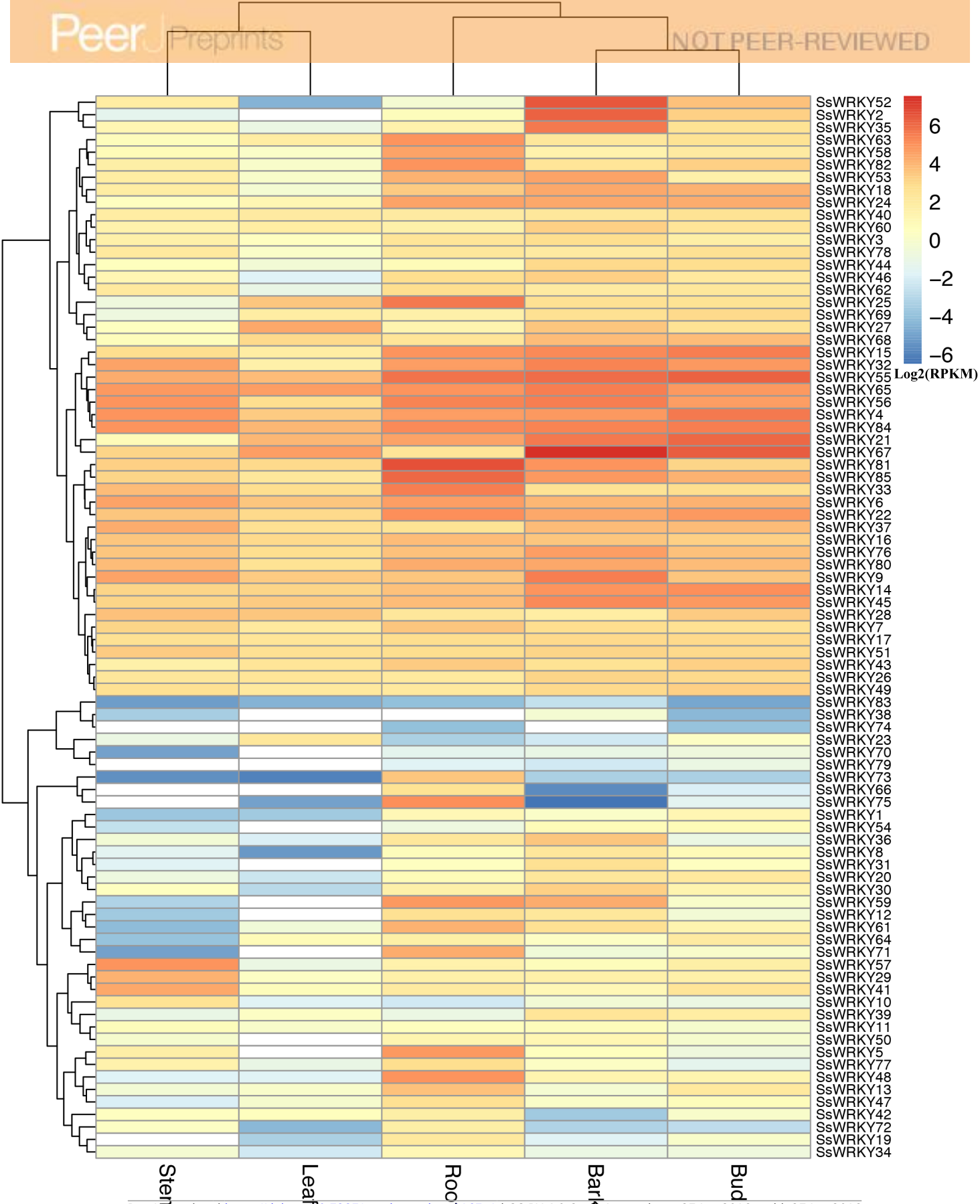


Table 1 (on next page)

The detailed characteristics of WRKY genes identified in willow.

Gene	SequenceID	Chr	Group	Ortholog		Deduced polypeptide			Introns
				AtWRKY	PtWRKY	Length(aa)	PI	MW(kDa)	
SsWRKY1	willow_GLEAN_10011238	1	I	33	17	583	7.14	64.7	4
SsWRKY2	willow_GLEAN_10019192	1	II c	45	43	162	9.47	18.6	1
SsWRKY3	willow_GLEAN_10017208	1	II c	28,71	29	584	9.42	65.6	4
SsWRKY4	willow_GLEAN_10017139	1	I	20	44	560	6.99	60.9	5
SsWRKY5	willow_GLEAN_10007860	1	II e	35	45	445	5.92	48.4	2
SsWRKY6	willow_GLEAN_10003806	1	I	2	37,101,102	733	5.69	78.8	4
SsWRKY7	willow_GLEAN_10022392	2	II d	21	46,63	453	9.53	49.9	4
SsWRKY8	willow_GLEAN_10022273	2	II c	71	47	328	6.89	37.0	2
SsWRKY9	willow_GLEAN_10009329	2	II d	15	14,94	339	9.77	37.5	2
SsWRKY10	willow_GLEAN_10009231	2	II c	12	48	204	7.64	23.6	3
SsWRKY11	willow_GLEAN_10016913	2	III	30	6,51	351	6.27	39.2	2
SsWRKY12	willow_GLEAN_10016886	2	II c	-	19,50	129	6.75	14.6	0
SsWRKY13	willow_GLEAN_10016883	2	II e	22	23,49,78	352	5.81	38.3	2
SsWRKY14	willow_GLEAN_10019911	2	II e	-	3	247	5.58	28.1	2
SsWRKY15	willow_GLEAN_10019925	2	II c	23	13,33	319	6.46	35.6	2
SsWRKY16	willow_GLEAN_10019982	2	I	1	54	472	6.88	52.2	3
SsWRKY17	willow_GLEAN_10020022	2	II b	47	53	1081	5.25	116.8	17
SsWRKY18	willow_GLEAN_10025583	3	II d	-	55	142	9.60	16.5	2
SsWRKY19	willow_GLEAN_10025423	3	II e	29	41	335	5.54	37.9	2
SsWRKY20	willow_GLEAN_10025378	3	III	41/53	21	342	5.25	38.4	2
SsWRKY21	willow_GLEAN_10008020	3	II c	45	18	157	9.41	17.8	1
SsWRKY22	willow_GLEAN_10006448	3	II a	40	88	320	8.38	35.4	3
SsWRKY23	willow_GLEAN_10013342	3	II c	-	39	109	8.03	12.9	1
SsWRKY24	willow_GLEAN_10009960	4	II b	42	28,79	604	6.93	65.3	5
SsWRKY25	willow_GLEAN_10017267	4	II e	65	8,58	267	5.43	29.7	2
SsWRKY26	willow_GLEAN_10018559	4	I	58	60	537	8.72	58.9	3
SsWRKY27	willow_GLEAN_10004854	4	III	54	85	323	5.70	36.3	2
SsWRKY28	willow_GLEAN_10008312	5	II d	-	-	490	10.27	54.0	2
SsWRKY29	willow_GLEAN_10009112	5	II c	13	68	235	8.70	26.7	2
SsWRKY30	willow_GLEAN_10003565	5	II d	15	20	310	9.48	34.3	2
SsWRKY31	willow_GLEAN_10016009	5	II c	28,71	62	322	6.67	36.2	2
SsWRKY32	willow_GLEAN_10018195	5	II d	21	46,63	349	9.69	38.8	2
SsWRKY33	willow_GLEAN_10026833	6	II d	7	91	339	9.89	36.8	3
SsWRKY34	willow_GLEAN_10026721	6	II c	49	34	287	5.25	32.1	2
SsWRKY35	willow_GLEAN_10026591	6	I	33	64	572	6.41	62.7	4
SsWRKY36	willow_GLEAN_10026566	6	III	54	85	329	6.13	36.7	2
SsWRKY37	willow_GLEAN_10020588	6	I	44	93	478	9.25	52.5	4
SsWRKY38	willow_GLEAN_10026166	6	II c	51	67	233	5.03	26.1	2

SsWRKY39	willow_GLEAN_10026455	6	II a	18/60	9	327	9.02	36.2	4
SsWRKY40	willow_GLEAN_10026458	6	I	32	15	413	8.26	44.9	3
SsWRKY41	willow_GLEAN_10008192	7	II c	13	68	236	9.21	26.6	2
SsWRKY42	willow_GLEAN_10025108	8	I	3/4	69	460	8.80	50.6	3
SsWRKY43	willow_GLEAN_10025123	8	II c	57	71	295	6.32	32.3	2
SsWRKY44	willow_GLEAN_10015641	8	II c	48	70	357	6.11	39.9	2
SsWRKY45	willow_GLEAN_10008155	9	II d	15	20,26	331	9.57	36.4	2
SsWRKY46	willow_GLEAN_10013562	10	II c	57	71	289	6.26	31.9	2
SsWRKY47	willow_GLEAN_10013586	10	I	3/4	72	490	8.60	53.7	3
SsWRKY48	willow_GLEAN_10004012	11	II b	42	100	585	6.48	63.3	5
SsWRKY49	willow_GLEAN_10006060	11	I	20	44	607	7.09	6.6	6
SsWRKY50	willow_GLEAN_10007614	11	II e	35	74	481	5.39	51.6	3
SsWRKY51	willow_GLEAN_10007542	11	I	2	37	734	6.10	79.7	4
SsWRKY52	willow_GLEAN_10013801	12	II c	-	75	178	9.08	20.5	1
SsWRKY53	willow_GLEAN_10012158	13	II d	74	25	356	9.66	40.0	2
SsWRKY54	willow_GLEAN_10004417	13	I	2	35	697	6.52	76.1	4
SsWRKY55	willow_GLEAN_10007732	13	I	33	1	602	7.65	66.0	4
SsWRKY56	willow_GLEAN_10009039	14	II d	15	14,94	362	9.39	40.0	2
SsWRKY57	willow_GLEAN_10016668	14	II c	12	48	180	8.47	20.7	3
SsWRKY58	willow_GLEAN_10016177	14	II e	22	23,49,78	354	6.35	38.8	2
SsWRKY59	willow_GLEAN_10016180	14	II c	43	19,50	193	9.47	21.7	1
SsWRKY60	willow_GLEAN_10016220	14	III	30	6	368	5.03	41.3	2
SsWRKY61	willow_GLEAN_10018940	14	II b	42	28,79	467	8.78	50.0	5
SsWRKY62	willow_GLEAN_10018891	14	II c	23	13,33	318	5.71	35.6	2
SsWRKY63	willow_GLEAN_10018881	14	II e	-	80	263	5.05	29.7	2
SsWRKY64	willow_GLEAN_10020302	14	II b	36	-	460	6.28	50.0	4
SsWRKY65	willow_GLEAN_10020380	14	I	1	2	481	5.98	52.8	3
SsWRKY66	willow_GLEAN_10011119	15	II b	9	99	618	6.55	66.2	5
SsWRKY67	willow_GLEAN_10016438	15	II c	-	82	178	9.35	20.5	1
SsWRKY68	willow_GLEAN_10023347	16	II a	40	88	320	8.82	35.3	3
SsWRKY69	willow_GLEAN_10023447	16	II c	45	18	178	9.17	20.1	1
SsWRKY70	willow_GLEAN_10023687	16	III	41/53	21	336	5.17	37.2	2
SsWRKY71	willow_GLEAN_10023735	16	II e	29	41	325	5.54	36.6	2
SsWRKY72	willow_GLEAN_10014752	16	II d	-	55	338	9.24	37.9	2
SsWRKY73	willow_GLEAN_10009602	16	II b	9	42	509	5.51	55.3	4
SsWRKY74	willow_GLEAN_10010473	17	II c	45	43	182	9.92	20.9	1
SsWRKY75	willow_GLEAN_10015128	17	II b	9	86	544	6.01	59.0	3
SsWRKY76	willow_GLEAN_10015184	17	I	58	87	1044	8.94	116.1	11
SsWRKY77	willow_GLEAN_10005468	17	II e	27	96	411	5.96	45.7	2
SsWRKY78	willow_GLEAN_10006860	18	I	-	90	1593	8.67	179.0	10

SsWRKY79	willow_GLEAN_10006862	18	II a	18/60	9	320	8.57	35.6	4
SsWRKY80	willow_GLEAN_10011608	18	I	32	-	528	5.74	57.8	4
SsWRKY81	willow_GLEAN_10004546	18	II d	7	7,91	300	9.80	32.8	2
SsWRKY82	willow_GLEAN_10003422	19	II d	11/17	24	339	9.58	37.1	2
SsWRKY83	willow_GLEAN_10011321	19	III	55	36,76	358	5.63	38.7	2
SsWRKY84	willow_GLEAN_10005288	19	I	33	4	597	6.69	65.6	4
SsWRKY85	willow_GLEAN_10002834	N/A	II e	65	58	268	5.83	30.2	2

Chr, chromosome numbers.

N/A, not available.

"-", not detected.

Table 2 (on next page)

The details of twenty conserved motif sequences identified in SsWRKY genes.

Motif	Width	Best possible match
1	29	ILDDGYRWRKYGQKVIKGNPYPRSYRCT
2	29	CPVRKHVERCWEDPTMVITTYEGEHNHPW
3	37	PSDDGYNWRKYGQKQVKGSEYPRSYKCTHPNCPVKK
4	21	KKGHKKIREPRFAFQTRSEVD
5	29	KVECSHDGHITEIYKGTNHPKQPNCR
6	15	KRRKNRVKVVVRVPA
7	50	KEELAVLQEELNRMKEENKRLKEMLDQICENYNALQMFMQLMQQNE
8	29	PVIRSPYFTIPPGLSPTTELLDSPVFFSNS
9	29	LVEQMTAAITADPNFTAALAAAISGIMGQ
10	28	QVQYRNCMVITDETVEFKFKVISLLNRT
11	29	LQQQQQQMKYQADMMYRKSNSGINLNF
12	15	MRKARVSVRARCEAP
13	50	MDGTVANLDGDAFHLMGMPHSSDHISQQHKRKCGRGEDGNVCKGSSGI
14	21	PPAAMAMASTTSAAASMLLSG
15	21	VEEAARAGIESCEHVIRLLCQ
16	21	MATISASAPFPTITLDTQNP
17	40	LGHGRVRKLKLP SHLPQNIFLDNPHCKTIHAPKPPQMPV
18	17	LLPDYGLLQDIVPSMH
19	17	GGEDDEDEPEPKRWKIE
20	49	PSPTTGTFPGQAFNWKSNSGDNQQGVKGEDKDFSDFSFQTPARPPATSS

Table 3 (on next page)

The number of WRKY genes identified in *Arabidopsis thaliana*, *Cucumis sativus*, *Populus trichocarpa*, *Vitis vinifera*, *Salix suchowensis* and *Oryza sativa*.

Species	Group						
	I	IIa	IIb	IIc	IId	IIe	III
<i>Arabidopsis thaliana</i>	13	4	7	18	7	9	14
<i>Cucumis sativus</i>	10	4	4	16	8	7	6
<i>Populus trichocarpa</i>	50	5	9	13	13	4	10
<i>Vitis vinifera</i>	12	4	8	16	7	6	6
<i>Salix suchowensis</i>	19	4	8	23	13	11	7
<i>Oryza sativa</i>	34	4	8	7	11	0	36