

**Supplemental Table S1.** Polymerase chain reaction (PCR) primers used for construction of vectors, gene structure analysis, and chromatin immunoprecipitation (ChIP) analysis

Gene (GenBank accession number or locus number)	Primer name	Forward primer (5'-3')	Reverse primer (5'-3')	Product size (nt)	Use
WRKY42 (AK110587)	153U3F/153 U3R	AAG <u>G</u> TAC <u>C</u> TTCAATT <u>T</u> GCTTGCTT <u>G</u> <sup>a</sup>	AAG <u>G</u> GTAC <u>C</u> CGAGTGAGG GAGGTAGCAGAT <sup>a</sup>	1187	Amplifying genomic DNA for constructing WRKY42-overexpressing vector
	153R4F/153 R4R	ACTAG <u>T</u> GGTAC <u>C</u> CAACGG AAGGCGTC <u>G</u> TGGAG <sup>b</sup>	GAG <u>C</u> TC <u>G</u> GAT <u>C</u> CTGGTC AGTC <u>G</u> TCC <u>T</u> GTGCTCC <sup>c</sup>	313	Amplifying cDNA fragment for constructing WRKY42 RNAi vector
	1536F/1536 R	T <u>C</u> TA <u>G</u> AGGCCATGGCGGA TCC <u>G</u> TTC <sup>d</sup>	T <u>C</u> TA <u>G</u> ACAAT <u>T</u> CCAGT AGTT <u>G</u> GTC <sup>d</sup>	759	Amplifying the complete coding sequence for subcellular localization analysis and transcriptional activity vector
	N1F/N1R	AGGAACC <u>G</u> TAAACATCA GCC	GAGAAG <u>C</u> AAGCAA <u>G</u> C AAAAT	336	Analyzing the ChIP product and constructing reporter vector
	N2F/N2R	TTAT <u>G</u> TTTG <u>A</u> ATCGCTG GTA	CGGG <u>C</u> TGAT <u>G</u> TTACGG T	215	Analyzing the ChIP product and constructing reporter vector
	N3F/N3R	ATT <u>C</u> TG <u>T</u> AGTCAAT <u>C</u> TG AGT	ATAT <u>C</u> TG <u>T</u> CT <u>C</u> TCAA TCT	357	Analyzing the ChIP product and constructing reporter vector
	1532F/R	AT <u>G</u> A <u>T</u> CGAT <u>C</u> CGTT <u>C</u> CGGC <u>GG</u> C <sup>e</sup>	AT <u>G</u> AG <u>G</u> AT <u>C</u> CCAGCAAT <u>T</u> TT CCAGTAG <u>T</u> TTGGTCA <sup>c</sup>	750	Amplifying cDNA fragment for constructing WRKY42 pGAL4-BD vector and effector

					vector
	153Y1F/153 Y1R	ATGGCGGATCCGTTCCC G	TCCGCCGGGATGTCGGC G	507	Amplifying cDNA fragment for constructing <i>WRKY42</i> pGAL4-BD vector
	153Y2F/153 Y2R	CCGGCGGACGACTATTG G	GAGCAATCTTCCAGTAG TTT	261	Amplifying cDNA fragment for constructing <i>WRKY42</i> pGAL4-BD vector
	153GAL4F/ 153GAL4R	A <u>AGGTACCGCCATGGCG</u> GATCCGTT <sup>c</sup>	A <u>AGGTACCCTAGAGCAA</u> TCTTCCAGTAG <sup>c</sup>	759	Amplifying the complete coding region for constructing transcriptional activation vector
	153JGF/153 JGR	GAATT <u>CATGGCGGATCC</u> GTTCCCGG <sup>e</sup>	CTCGAGCTAGAGCAATC TTCCAGTAG <sup>f</sup>	759	Amplifying the complete coding region to construct yeast one hybrid vector
<i>WRKY13</i> (EF143611)	WRKY1322 F/WRKY13 22R	ACTAG <u>TTGGTACCA</u> GAAAG CGGGTGGTGTCCG <sup>b</sup>	<u>GAGCTCGGATCCATT</u> CG TTCTCCGCTGGCTC <sup>c</sup>	366	Amplifying cDNA fragment for constructing <i>WRKY13</i> RNAi construct
	WRKY1323 F/WRKY13 23R	GCGTATCCTCCGCCGTT TGA	CACCACCACCACTCCCG GCG	1001	Amplifying the promoter sequence for constructing reporter vector
	WRKY1320 F/WRKY13 20R	<u>TCTAGAGGAGTGGTGGT</u> GGTGATG <sup>d</sup>	<u>TCTAGAGGAGCACGGCG</u> CGGTGGC <sup>d</sup>	964	Amplifying the complete coding region for constructing transcriptional activity vector
	WRKY1316 F/WRKY13 13R	AT <u>GAATT</u> CGGAGTGGTG GTGGTGATG <sup>e</sup>	AT <u>AGGATCC</u> AGGAGCAC GGCGCGGTGGC <sup>c</sup>	963	Amplifying the complete coding region for constructing effector vector
<i>WRKY45-2</i> (GQ331927)	WRKY45F 5/WRKY45	AT <u>GAATT</u> CTTGCTTGGA TTGAGGATG <sup>e</sup>	AT <u>GGATC</u> TCACCGAAG AATCATGGATG <sup>c</sup>	978	Amplifying the complete coding region for constructing

	R5				effector vector
WRKY45T F/WRKY45 TR	A <u>AGGTACCATGACGTCA</u> TCGATGTCGCCG <sup>a</sup>	A <u>AGGTACCAAAGCTCAA</u> ACCCATAATG <sup>a</sup>		978	Amplifying the complete coding region for constructing transcriptional activation vector
WRKY45F 4/WRKY45 R3	GACATCAAGGAGGCAA AGGC	GGGTGGAGGCAGGTG GTAT		959	Amplifying the promoter sequence for constructing reporter vector
WRKY45J GF/WRKY 45JGR	<u>GAATT</u> CATGACGTAC GATGTCGC <sup>e</sup>	<u>CTCGAG</u> TCAAAAGCTCA AACCCATAA <sup>f</sup>		959	Amplifying the complete coding region to construct yeast one hybrid vector
<i>bZIP23</i> (AK072062)	<i>bZIP23F/bZ</i> IP23R	<u>AAGGTACCATGGATT</u> CCGGGAGGG <sup>a</sup>	<u>AAGGTACCCC</u> CATGGACC CGTCAGAGT <sup>a</sup>	1071	Amplifying the complete coding region for constructing transcriptional activation vector
<i>Pot</i> (Z33638)	PotF/PotR	ACGACCCGTCTTACTT ATTGGA	AAGTAGCGTTGGTTTG TTGGAT	98	Quantitatively amplifying the <i>M. oryzae</i> genomic DNA for fungal growth analysis
<i>Ubiquitin</i> (Os03g13170)	UbiqF/Ubiq R	AACCAGCTGAGGCCAA GA	ACGATTGATTAAACCAG TCCATGA	76	Quantitatively amplifying the rice genomic DNA to standardize DNA samples for fungal growth analysis

<sup>a</sup>The underlined nucleotides are the digestion site of *Kpn*I.

<sup>b</sup>The underlined nucleotides are the digestion site of *Spe*I and the double underlined nucleotides are the digestion site of *Kpn*I.

<sup>c</sup>The underlined nucleotides are the digestion site of *Sac*I and the double underlined nucleotides are the digestion site of *Bam*HI.

<sup>d</sup>The underlined nucleotides are the digestion site of *Xba*I.

<sup>e</sup>The underlined nucleotides are the digestion site of *Eco*RI.

<sup>f</sup>The underlined nucleotides are the digestion site of *Xho*I.

**Supplemental Table S2.** Primers used for quantitative polymerase chain reaction (qPCR) in gene expression analysis

Gene (GenBank accession number)	Primer name	Forward primer (5'-3')	Reverse primer (5'-3')	Product size (nt)
<i>WRKY13</i> (EF143611)	WRKY13-F/R	TCAGTGGAGAACGCGGTGGTG	GGGTGGTTGTGCTCGAAGGAG	253
<i>WRKY42</i> (AK110587)	WRKY42-F/R	ATGCAGTCTGCTTCAGATTATGCT	GACGCCTCCGTTTTCTTG	100
<i>AOS2</i> (AY062258)	AOS2-F/R	CAATACGTGTACTGGTCGAATGG	AAGGTGTCGTACCGGAGGAA	120
<i>JAZ8</i> (AK108738)	JAZ8-F/R	GAAGGGCTAACAGCTGACCAT	TTGGTGGACGGGAAGTTCTC	120
<i>Actin</i> (X15865)	Actin-F/R	TGTATGCCAGTGGTCGTACCA	CCAGCAAGGTGAGACGAA	121